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Szellemi Tulajdon Nemzeti Hivatala

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THERAPEUTISCHE VERBINDUNGEN UND DAZUGEHÖRIGE VERWENDUNGSVERFAHREN

COMPOSÉS THÉRAPEUTIQUES ET PROCÉDÉS D'UTILISATION ASSOCIÉS

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- Remarks:
The file contains technical information submitted after the application was filed and not included in this specification

Description

RELATED APPLICATIONS

[0001] This application claims priority to U.S. Provisional Applications Serial Nos. 61/291,544, entitled "Therapeutic Compounds and Related Methods of Use" filed on December 31, 2009; 61/291,550, entitled "Therapeutic Compounds and Related Methods of Use" filed on December 31, 2009; and 61/291,554, entitled "Therapeutic Compounds and Related Methods of Use" filed on December 31, 2009, all of which are herein incorporated by reference in their entireties.

BACKGROUND OF INVENTION

[0002] Tyrosine phosphorylation of synaptic receptors and signaling molecules regulates synaptic activity. A number of protein tyrosine phosphatases specifically expressed within the brain have been identified, including STEP (for STRiatal-Enriched tyrosine Phosphatase, also known as PTPN5). Recent evidence suggests that STEP plays an important role in synaptic plasticity, for review see (Braithwaite SP, et al., (2006), Trends Neurosci, 29 (8): 452; Baum ML, et al., (2010), Commun Integr Biol, 3 (5): 419). STEP is specifically expressed within neurons of the central nervous system. As its name indicates, the highest expression level is within the striatum. However, more recent work has found that it is expressed at lower levels in multiple brain regions including the neocortex, amygdala, hippocampus, and embryonic spinal cord.

[0003] Four groups of proteins that STEP regulates have been identified: the mitogen-activated protein kinases (MAPKs), the tyrosine kinase Fyn, the N-methyl-D-aspartate (NMDA) receptor complex (specifically the NR2B subunit) and AMPA receptors (specifically, GluR2, (Zhang Y, et al., (2008), J Neurosci, 28 (42): 10561)). Three additional new substrates for STEP have also been recently discovered; proline-rich tyrosine kinase 2 (Pyk2; Xu J, et al., (2010), Abstracts of the Society for Neuroscience Meetings), the fragile X mental retardation protein (FMRP) (Goebel-Goody SM, et al., (2010), Abstracts of the Society for Neuroscience Meetings) and the cell-death mediator Bak (Fox JL, et al., (2010), EMBO J, 29 (22): 3853). Tyrosine phosphorylation of one member of the MAPK family, the extracellular signal regulated kinase (ERK), is necessary for the expression and maintenance of synaptic plasticity in many brain regions, and disruption of the ERK pathway leads to a disruption of learning and memory. One of the functions of these src and Pyk2 kinases is to phosphorylate NMDA receptors, thereby modulating their channel conductance properties and facilitating their movement toward the surface of neuronal plasma membranes. Pyk2 and Fyn tyrosine kinases are activated by phosphorylation on tyrosine residues. NR2B phosphorylation on Tyrosine 1452 inhibits the receptor endocytosis. STEP acts as direct or indirect brake of NMDAR mediated signaling by either respectively dephosphorylating NR2B or its associated kinases, Pyk2 and Fyn. Activation of AMPA, NMDA receptors and MAPKs are required for the induction of several forms of long-term potentiation (LTP) and long-term depression (LTD). Hippocampal LTP is increased in transgenic mice model of Alzheimer lacking STEP (Zhang Y, et al., (2010), Proc Natl Acad Sci U S A, 107 (44): 19014). NR2B and AMPA receptor surface expression is increased in STEP KO mice. AMPA receptor endocytosis in group I metabotropic glutamate receptor I (mGluR) mediated LTD is mediated by a tyrosine phosphatase. AMPA receptor endocytosis induced by activation of group I mGluR is blocked in STEP KO mice suggesting that STEP might also control mGluR mediated LTD.

[0004] Compounds that inhibit STEP activity should mimic the effects observed with the STEP KO and may be useful for treating conditions mediated by abnormal NMDA-receptor (NMDA-Rs) and/or MAP kinase pathway signaling. Both may mediate cognition, learning and memory, neurogenesis, and may also affect neuronal plasticity, pain perception, mood and anxiety, and neuroendocrine regulation.

Modulation of NMDA-Rs:

[0005] STEP decreases the tyrosine phosphorylation level of NMDA-Rs. Less phosphorylated NMDA-Rs have lower conductance states and thus will allow less current and fewer ions to pass. The NMDA-Rs will therefore be functionally less active (Alvestad RM, et al., (2003), J Biol Chem, 278 (13): 11020), which can lead to schizophrenic symptoms. Hypofunction of NMDA-Rs has been linked to schizophrenia. For example, phencyclidine, ketamine, and other noncompetitive antagonists at NMDA-type glutamate receptors can exacerbate symptoms in patients (Lahti AC, et al., (1995), Neuropsychopharmacology, 13 (1): 9) and may produce a range of psychotic symptoms in volunteers that are similar to those of schizophrenic patients. NMDA-R hypofunction is also linked to psychosis and drug addiction (Javitt DC and Zukin SR, (1991), Am J Psychiatry, 148 (10): 1301). Chronic treatment of atypical antipsychotic clozapine and risperidone in mice result in significant increase of phosphorylation of ERK, NR2B and Pyk2 on tyrosine residues recognized by STEP (Carty NC, et al., (2010), Abstracts of the Society for Neuroscience Meetings). Treatment of these anti-psychotics also enhances cAMP and STEP phosphorylation. Since PKA mediated phosphorylation of STEP is known to inactivate STEP, these results suggest that STEP inhibition mediates the beneficial effect of antipsychotic drugs. Recent studies

have linked abnormal NMDA-R activity and expression of STEP to the cognitive decline observed in Alzheimer's disease or transgenic mice expressing mutant APP (Tg2576 mice) (Snyder EM, et al., (2005), Nat Neurosci, 8 (8): 1051; Hynd MR, et al., (2004), J Neurochem, 90 (4): 913; Kurup P, et al., (2010), Channels (Austin), 4 (5)). More specifically, STEP KO mice are less susceptible to PCP-induced hyperlocomotion and PCP-induced cognitive deficits in the object recognition tasks (Carty NC, et al., (2010), Abstracts of the Society for Neuroscience Meetings). Compared to the Tg2576 mice expressing STEP, Tg2576 lacking STEP gene showed rescue in their deficits in hippocampal LTP and in different behavioral cognitive tasks. Altogether, these results suggest that STEP inhibitors might represent a novel class of drugs that can treat both positive symptoms and cognitive deficit associated with schizophrenia.

[0006] Medications that modulate glutamatergic neurotransmission via NMDA-Rs may be also effective in treatment for mood and anxiety disorders. Administration of NMDA-R antagonists has anxiolytic effects in rodent models of anxiety (Falls WA, et al., (1992), J Neurosci, 12 (3): 854; Miserendino MJ, et al., (1990), Nature, 345 (6277): 716). NMDA-Rs antagonist like ketamine has been shown to be effective in drug-resistant unipolar depression (Machado-Vieira R, et al., (2009), Pharmacol Ther, 123 (2): 143).

[0007] Abnormal balance between the activity of NMDA receptors at synaptic (prosurvival linked to ERK activation) and extrasynaptic (proapoptotic linked to p38 activation) sites has been proposed in cellular and mouse model of Huntington Disease (HD) (Milnerwood AJ, et al., Neuron, 65 (2): 178). YAC128 mouse model (containing high number of glutamine repeat on huntingtin) of HD showed an increased activity of extrasynaptic NMDA receptors (NR2B subunit) and require p38 and caspase-6 cleavage activation. In YAC128 mice, NR2B synaptic expression is associated with high STEP expression and activity and a reduction in NR2B expression and phosphorylation (Gladding CM, et al., (2010), Abstracts of the Society for Neuroscience Meetings). Extrasynaptic NMDA receptors couple preferentially to excitotoxicity via calpain-mediated cleavage of STEP and activation of p38 (Xu J, et al., (2009), J Neurosci, 29 (29): 9330). Inhibiting STEP activity might therefore shift the balance toward the NMDA receptor/ERK synaptic prosurvival signaling pathway.

Modulation of ERK pathway:

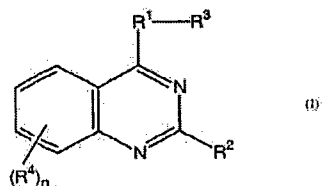
[0008] STEP inhibition may translate into activation of ERK1/2 kinases, for example, in the central nervous system (CNS). Activation of the ERK pathway in the CNS can mediate neurotrophic pathways involved in cellular resilience. ERK signaling directly affects Bak phosphorylation through inhibition of STEP to promote cell survival (Fox JL, et al., (2010), EMBO J, 29 (22): 3853). BDNF and other neurotrophins can block apoptosis and increase cell survival of different type of CNS neurons *in vitro* and *in vivo* via stimulation of the ERK pathway. Mood stabilizers effective in bipolar disorder like valproate and lithium may be potent activators of ERK activity. This effect on ERK activation is believed to be responsible for the neurotrophic effects of mood stabilizers observed *in vitro* or in brains of treated patients with bipolar disorder, for review see (Engel SR, et al., (2009), Mol Psychiatry, 14 (4): 448; Chen G and Manji HK, (2006), Curr Opin Psychiatry, 19 (3): 313; Machado-Vieira R, et al., (2009), Bipolar Disord, 11 Suppl 2 92). *In vivo* disruption of STEP activity was shown to activate MAPK pathway, leading to significant rescue from neuronal cell death after pilocarpine-induced status epilepticus (Choi YS, et al., (2007), J Neurosci, 27 (11): 2999). Increasing cellular resilience could therefore limit or reduce neuronal loss in several neurologic disorders. Recent work has suggested a positive role for STEP inhibition in fragile X syndrome (FXS). This disorder results from the mutation of *fmr1* gene coding for the fragile X mental retardation protein (FMRP). STEP binds to FMRP and its expression is dysregulated in FXS. FMR KO mice model displayed audiogenic seizures. FMR KO mice lacking STEP gene show a significant reduction of these seizures (Goebel-Goody SM, et al., (2010), Abstracts of the Society for Neuroscience Meetings), suggesting that STEP modulators might be therapeutic approach for FXS.

[0009] Various substituted heterocyclic compounds are disclosed in the art. For example, WO 02/062767 discloses quinazoline derivatives; WO 03/000188 discloses quinazolines and uses thereof; WO 2005/042501 discloses norepinephrine reuptake inhibitors for the treatment of central nervous system disorders; WO2006/058201 discloses heterocyclic and bicyclic compounds, compositions and methods; WO 2007/104560 discloses substituted 4-amino-quinazoline derivatives as regulators of metabotropic glutamate receptors and their use for producing drugs; WO 2007/133773 discloses CDKI pathway inhibitors; WO 2008/009078 discloses 4,6-DL- and 2,4,6-trisubstituted quinazoline derivatives useful for treating viral infections; WO 2009/000085 discloses quinoline and quinazoline derivatives useful as modulators of gated ion channels; US 2009/0143399 discloses protein kinase inhibitors; and Japan Publication Number 2007-084494A discloses substituted bicyclic compounds.

[0010] Current Chemical Genomics 3, 2009, 42-49 relates to small molecule modulators of prostatic acid phosphatase (PAP), one of which is 4-Phenylamino-2-(3-Pyridyl)Quinazoline.

[0011] Bioorg. & Med. Chem. Lett. 19(8), 2009, 2277-2281, WO 2005/099711 A1, WO 2004/065392 A1, WO 2004/054582 A1, WO 03/062209 A2, US 2002/025968 A1 relate to substituted quinazolin-4-ylamine analogues which are said to be useful as ALK5 inhibitors, as potassium ion channel modulators, for the treatment or prophylaxis of disorders characterised by overexpression of transforming growth factor beta (TGF-beta), for the treatment of pain, as capsaicin receptors modulators, and for inhibiting neoplastic cells and related conditions.

[0012] WO 2005/042501 A1 relates to compounds of the general formula (I), pharmaceutical compositions containing them and their use in the treatment of central nervous system disorders



[0013] WO 2005/055813 A2 relates to the identification of STEP being as involved in signaling pathways relating to psychotic diseases, including schizophrenia, and other disorders in which NMDA receptor dysfunction is implicated, as well as methods for screening STEP inhibitors that modulate NMDA-R signaling.

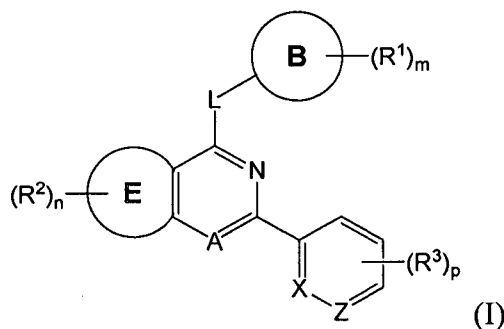
[0014] Molecular Pharmacology Vol. 68, No. 6, 1510-1523 characterizes several partial agonists at the NR1/NR2B receptor and investigates the mechanisms underlying their reduced efficacy by introducing mutations in the glutamate binding site.

SUMMARY OF INVENTION

[0015] Described herein are compounds, pharmaceutical compositions containing the compounds, and methods of using the compounds to treat a disorder, e.g., schizophrenia or cognitive deficit, in a subject. The compounds disclosed herein include quinoline- and quinazoline-containing compounds that modulate (e.g., inhibit) the activity of STEP.

[0016] The present invention provides therapeutic compounds, pharmaceutical composition comprising said compounds, use of said compounds and method for treating or preventing a disorder as described in items 1 to 21 below.

Item 1. A compound of formula (I):



or a salt thereof, wherein:

A is CH or N;

B is phenyl, dihydroindenyl, dihydrobenzoxazinyl, dihydrobenzodioxinyl, chromenyl, tetrahydroisoquinolyl, tetrahydroquinolyl, dihydroquinolyl, quinolyl, tetrahydroquinazolinyl, indolyl, dihydrobenzothiazolyl, dihydrobenzimidazolyl, dihydrobenzoxazolyl, isoindolyl, benzofuryl, benzothienyl, benzodioxolyl, indolyl, indazolyl, benzoimidazolyl, benzotriazolyl, benzisoxazolyl, benzoxazolyl, benzothiazolyl, benzoxadiazolyl, tetrahydrobenzothienyl, thienyl, thiazolyl, imidazolyl, pyrazolyl, thiadiazolyl or pyridyl;

m is 1, 2, 3, or 4;

E is phenyl, thienyl or pyrrolyl;

when E is phenyl, n is 1, or 2; and when E is thienyl or pyrrolyl, n is 0, or 1;

L is NR⁵;

one of X and Z is N and the other is CH;

p is 0, 1, or 2;

R¹ is C₁-C₈ alkyl, halo, haloalkyl, haloalkoxy, alkoxyalkyl, -CN, oxo, -NO₂, -C(Y)NR^bR^{b'}, -NR^cC(Y)R^{c'}, -SO₂NR^bR^{b'}, -OR^d, or -S(O)_qR^f;

R² is C₁-C₈ alkyl, C₃-C₈ cycloalkyl, thiomorpholinylalkyl, pyrrolidinylalkyl, morpholinylalkyl, haloalkyl, haloalkoxy, aminoalkyl, alkylaminoalkyl, dialkylaminoalkyl, hydroxyalkyl, alkoxyalkyl, silyloxyalkyl, -CN, -NO₂, -C(O)OR^a,

-C(Y)NR^bR^{b'}, -NR^bR^{b'}, -OR^d, -C(Y)R^e or -S(O)_qR^f;

R³ is C₁-C₈ alkyl, halo, haloalkyl -NR^bR^{b'} or -OR^d

R⁵ is hydrogen;

Y is O or S;

q is 1 or 2 and

each R^a, R^b, R^{b'}, R^c, R^{c'}, R^d, R^e and R^f is independently hydrogen, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₃-C₈ cycloalkyl, phenyl, dihydroindenyl, morpholinyl, tetrahydropyranyl, piperidyl, pyrrolidinyl, thiomorpholinyl, phenylalkyl, thienylalkyl, pyridylalkyl, tetrahydropyranylalkyl, dihydroindenylalkyl, tetrahydrofurylalkyl, hydroxyalkyl, morpholinylalkyl, pyrrolidinylalkyl, dialkylaminoalkyl, piperidylalkyl, benzodioxolylalkyl, dihydrobenzodioxinylalkyl, C₃-C₈ cycloalkylalkyl, haloalkyl or alkoxyalkyl.

Item 2. The compound according to Item 1 or a salt thereof, wherein

A is N;

B is phenyl or pyridyl;

m is 1, 2 for 3;

E is phenyl;

n is 1, or 2; and

p is 0 or 1

Item 3. The compound according to Item 2 or a salt thereof, wherein

m is 1 or 2 ;

p is 0;

R¹ is C₁-C₈ alkyl, halo, haloalkyl, haloalkoxy, -CN, NO₂, -C(Y)NR^bR^{b'}, -NR^cC(Y)R^{c'}, -SO₂NR^bR^{b'}, -OR^d, or -S(O)_qR^f; and

R² is -OR^d.

Item 4. The compound according to Item 3 or a salt thereof, wherein

R² is -OR^d, in which R^d is C₁-C₈ alkyl; and

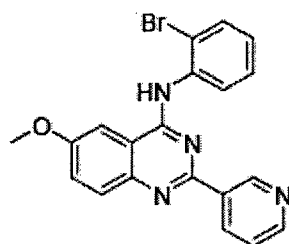
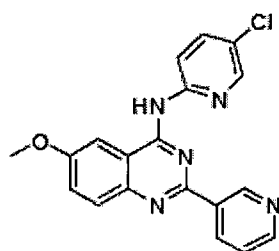
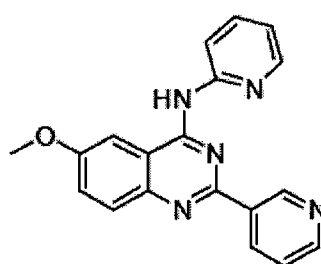
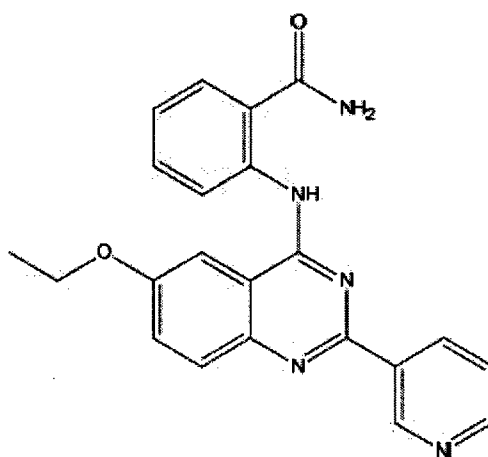
each R^b, R^{b'}, R^c, R^{c'}, R^d and R^f is independently hydrogen, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₃-C₈ cycloalkyl, phenyl, dihydroindenyl, tetrahydropyranyl, phenylalkyl, thienylalkyl, tetrahydropyranylalkyl, dihydroindenylalkyl, or haloalkyl.

Item 5. The compound according to Item 4 or a salt thereof, wherein

R¹ is C₁-C₈ alkyl, halo, haloalkyl, haloalkoxy, -C(Y)NR^bR^{b'} or -OR^d; and

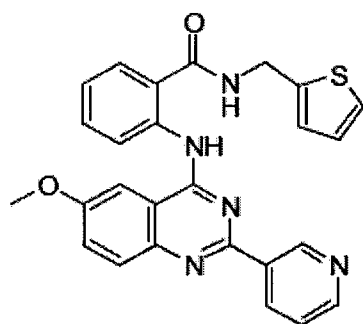
each R^b, R^{b'}, R^d and R^f is independently hydrogen, C₁-C₈ alkyl, C₃-C₈ cycloalkyl or haloalkyl.

Item 6. The compound according to Item 1 or a salt thereof, wherein the compound of formula (I) is a compound selected from the group consisting of



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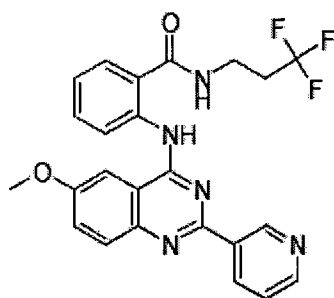
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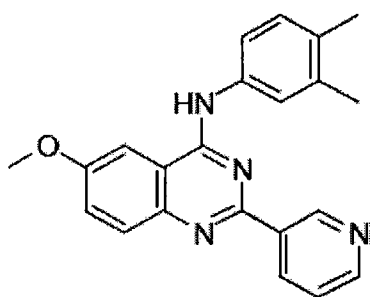
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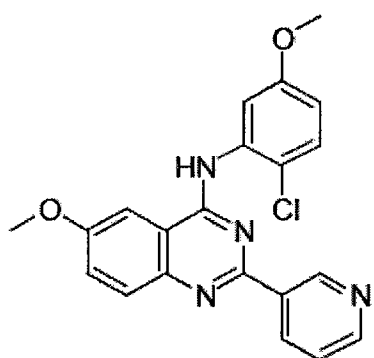


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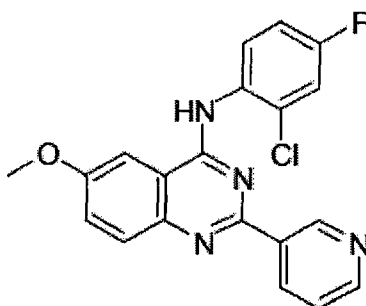
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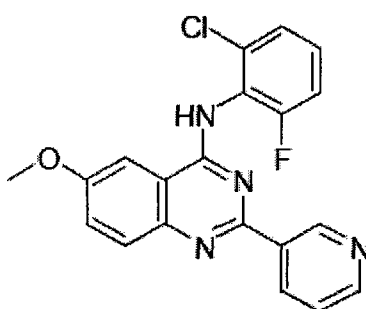


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and



Item 7. A pharmaceutical composition comprising the compound according to any one of Items 1 to 6 or a salt thereof as an active ingredient and a pharmaceutically acceptable carrier.

Item 8. The pharmaceutical composition according to Item 7 for use in preventing or treating central nervous system diseases.

Item 9. The pharmaceutical composition according to Item 8 for use in treating or preventing central nervous system disorders selected from the group consisting of schizophrenia; refractory, intractable or chronic schizophrenia; emotional disturbance; psychotic disorder; mood disorder; bipolar I type disorder; bipolar II type disorder; depression; endogenous depression; major depression; melancholy and refractory depression; dysthymic disorder; cyclothymic disorder; panic attack; panic disorder; agoraphobia; social phobia; obsessive-compulsive disorder; post-traumatic stress disorder; generalized anxiety disorder; acute stress disorder; hysteria; somatization disorder; conversion disorder; pain disorder; hypochondriasis; factitious disorder; dissociative disorder; sexual dysfunction; sexual desire disorder; sexual arousal disorder; erectile dysfunction; anorexia nervosa; bulimia nervosa; sleep disorder; adjustment disorder; alcohol abuse; alcohol intoxication; drug addiction; stimulant intoxication; narcotism; anhedonia; iatrogenic anhedonia; anhedonia of a psychic or mental cause; anhedonia associated with depression; anhedonia associated with schizophrenia; delirium; cognitive impairment; cognitive impairment associated with Alzheimer's disease, Parkinson's disease and other neurodegenerative diseases; cognitive impairment caused by Alzheimer's disease; Parkinson's disease and associated neurodegenerative diseases; cognitive impairment of schizophrenia; cognitive impairment caused by refractory, intractable or chronic schizophrenia; vomiting; motion sickness; obesity; migraine; pain (ache) ; mental retardation; autism disorder (autism); Tourette's disorder; tic disorder; attention-deficit/hyperactivity disorder; conduct disorder; and Down's syndrome.

Item 10. A process for producing a pharmaceutical composition comprising mixing a compound or a salt thereof according to any one of Items 1 to 6 with a pharmaceutically acceptable carrier.

Item 11. A compound or a salt thereof according to any one of Items 1 to 6 for use as a drug.

Item 12. A compound or a salt thereof according to any one of Items 1 to 6 for use in treating schizophrenia or cognitive deficit.

Item 13. A combination of a compound or a salt thereof according to any one of Items 1 to 6 with an additional

therapeutic agent for use in treating a disorder that would benefit by the modulation of STEP.

Item 14. The combination for use according to Item 13, wherein the additional therapeutic agent is an atypical antipsychotic.

Item 15. The combination for use according to Item 13, wherein the additional therapeutic agent is selected from the group consisting of aripiprazole, clozapine, ziprasidone, risperidone, quetiapine, olanzapine, amisulpride, asenapine, iloperidone, melperone, paliperidone, perospirone, sertindole and sulpride.

Item 16. The combination for use according to Item 13, wherein the additional therapeutic agent is a typical antipsychotic.

Item 17. The combination for use according to Item 13, wherein the additional therapeutic agent is selected from the group consisting of haloperidol, molindone, loxapine, thioridazine, molindone, thiothixene, pimozide, fluphenazine, trifluoperazine, mesoridazine, chlorprothixene, chlorpromazine, perphenazine, triflupromazine and zuclopenthixol.

Item 18. A kit comprising a composition comprising a compound or a salt thereof according to any one of Items 1 to 6 and an acceptable carrier.

Item 19. A kit comprising a pharmaceutical composition comprising a compound or a salt thereof according to any one of Items 1 to 6 and a pharmaceutically acceptable carrier.

[0017] In some embodiments, A is N. In some embodiments, A is CH.

[0018] In some embodiments, B is aryl (e.g., phenyl).

[0019] In some embodiments, m is 1. In some embodiments, m is 2. In some embodiments, m is 3.

[0020] In some embodiments, R¹ is in the ortho position. In some embodiments, R¹ is in the meta position. In some embodiments, R¹ is in the para position.

[0021] In some embodiments, R¹ is C₁-C₈ alkyl (e.g., methyl or tert-butyl).

[0022] In some embodiments, m is 1 and R¹ is halo (e.g., fluoro, chloro or bromo). In some embodiments, m is 2 and each R¹ is halo (e.g., fluoro, chloro or bromo). In some embodiments, m is 3 and each R¹ is halo (e.g., fluoro, chloro or bromo). In some embodiments, R¹ is haloalkyl (e.g., trifluoromethyl). In some embodiments, R¹ is haloalkoxy (e.g., difluoromethoxy or trifluoromethoxy).

[0023] In some embodiments, R¹ is -CN.

[0024] In some embodiments, R¹ is -NO₂.

[0025] In some embodiments, R¹ is -NR^cC(Y)R^c. In some embodiments, one of R^c and R^c is hydrogen and the other is C₁-C₈ alkyl (e.g., methyl).

[0026] In some embodiments, R¹ is -OR^d. In some embodiments, R^d is hydrogen. In some embodiments, R^d is C₁-C₈ alkyl (e.g., methyl or ethyl).

[0027] In some embodiments, R¹ is -SO₂NR^bR^b. In some embodiments, R^b and R^b are both hydrogen.

[0028] In some embodiments, R¹ is -C(Y)NR^bR^b. In some embodiments, Y is S. In some embodiments, Y is O. In some embodiments, R^b and R^b are both hydrogen. In some embodiments, R^b is hydrogen. In some embodiments, R^b is hydrogen and R^b is aralkyl. In some embodiments, R^b is hydrogen and R^b is optionally substituted benzyl. In some embodiments, R^b is C₁-C₈ alkyl, e.g., methyl, ethyl, C₃ alkyl (e.g., n-propyl or isopropyl), C₄ alkyl (e.g., n-butyl, sec-butyl or tert-butyl), C₅ alkyl (e.g., n-pentyl, isopentyl or pentan-3-yl), C₆ alkyl (e.g., n-hexyl or 3,3-dimethylbutan-2-yl), or C₇ alkyl (e.g., n-heptyl or 2-heptyl).

[0029] In some embodiments, R^b is bicycyl (e.g., indanyl). In some embodiments, R^b is heterocyclyl, e.g., a 6-membered heterocyclyl. In some embodiments, R^b is a 6-membered oxygen-containing heterocyclyl (e.g., tetrahydropyranyl). In some embodiments, R^b is a 6-membered nitrogen-containing heterocyclyl (e.g., piperidinyl).

[0030] In some embodiments, R^b is aralkyl. In some embodiments, the alkyl is a C₁-C₈ alkyl (e.g., C₁, C₂, C₃ or C₄ alkyl). In some embodiments, the alkyl is a straight-chain alkyl. In some embodiments, the alkyl is a branched alkyl. In some embodiments, the aryl is phenyl. In some embodiments, R^b is benzyl. In some embodiments, R^b is phenylethyl.

[0031] In some embodiments, R^b is heteroaralkyl. In some embodiments, the alkyl is a C₁-C₈ alkyl (e.g., C₁, C₂ or C₃ alkyl). In some embodiments, the alkyl is a straight-chain alkyl. In some embodiments, the alkyl is a branched alkyl. In some embodiments, the heteroaryl is pyridyl. In some embodiments, the heteroaryl is furanyl. In some embodiments, the heteroaryl is thiazolyl. In some embodiments, the heteroaryl is thienyl.

[0032] In some embodiments, R^b is cyclylalkyl. In some embodiments, the alkyl is a C₁-C₈ alkyl (e.g., C₁ alkyl). In some embodiments, the cyclyl group is cyclopropyl. In some embodiments, the cyclyl group is cyclopentyl. In some

embodiments, the cyclyl group is a bicyclic group. In some embodiments, the bicyclic group is indanyl. In some embodiments, $R^{b'}$ is heterocyclalkyl. In some embodiments, the alkyl is a C_1 - C_8 alkyl (e.g., C_1 alkyl). In some embodiments, the heterocycl group is tetrahydropyranyl.

[0033] In some embodiments, $R^{b'}$ is haloalkyl (e.g., fluoroethyl, difluoroethyl, trifluoroethyl or trifluoropropyl).

[0034] In some embodiments, $R^{b'}$ is alkoxyalkyl. In some embodiments, the alkyl is a C_1 - C_8 alkyl (e.g., C_1 , C_2 , C_3 or C_4 alkyl). In some embodiments, the alkyl is a straight-chain alkyl. In some embodiments, the alkyl is a branched alkyl. In some embodiments, the alkoxy is methoxy.

[0035] In some embodiments, R^b and $R^{b'}$ are each independently C_1 - C_8 alkyl (e.g., R^b and $R^{b'}$ are both methyl, both ethyl, or both isopropyl).

[0036] In some embodiments, R^1 is halo. In some embodiments, R^1 is at the 6,7, or 8 position.

[0037] In some embodiments, each R^1 is independently C_1 - C_8 alkyl (e.g., each R^1 is methyl). In some embodiments, each R^1 is independently halo (e.g., each R^1 is fluoro or each R^1 is chloro). In some embodiments, one R^1 is fluoro and the other is chloro. In some embodiments, one R^1 is chloro and the other is bromo.

[0038] In some embodiments, each R^1 is independently $-OR^d$. In some embodiments, each R^d is independently C_1 - C_8 alkyl (e.g., each R^d is methyl).

[0039] In some embodiments, one R^1 is halo (e.g., chloro) and the other is C_1 - C_8 alkyl (e.g., methyl). In some embodiments, one R^1 is halo (e.g., fluoro) and the other is heterocyclalkyl (e.g., $-CH_2$ -heterocycl). In some embodiments, the heterocycl is morpholino. In some embodiments, the heterocycl is pyrrolidiny. In some embodiments, the heterocycl is piperaziny. In some embodiments, the piperaziny is substituted with 1 R^6 . In some embodiments, R^6 is C_1 - C_8 alkyl (e.g., methyl).

[0040] In some embodiments, one R^1 is halo (e.g., fluoro or chloro) and the other is haloalkyl (e.g., trifluoromethyl).

[0041] In some embodiments, one R^1 is halo (e.g., chloro) and the other is haloalkoxy, (e.g., difluoromethoxy or trifluoromethoxy).

[0042] In some embodiments, one R^1 is halo (e.g., chloro) and the other is $-C(O)OR^a$. In some embodiments, R^a is hydrogen.

[0043] In some embodiments, one R^1 is halo (e.g., fluoro or chloro) and the other is $-C(Y)NR^bR^{b'}$. In some embodiments, Y is O. In some embodiments, R^b and $R^{b'}$ are both hydrogen. In some embodiments, one of R^b and $R^{b'}$ is hydrogen and the other is C_1 - C_8 alkyl (e.g., methyl).

[0044] In some embodiments, one R^1 is halo (e.g., chloro) and the other is $-NR^cC(Y)R^c$. In some embodiments, Y is O. In some embodiments, R^c is hydrogen and R^c is C_1 - C_8 alkyl (e.g., methyl).

[0045] In some embodiments, one R^1 is halo (e.g., fluoro or chloro) and the other is $-OR^d$. In some embodiments, R^d is hydrogen. In some embodiments, R^d is C_1 - C_8 alkyl (e.g., methyl).

[0046] In some embodiments, one R^1 is halo (e.g., fluoro or chloro) and the other is $-CN$.

[0047] In some embodiments, one R^1 is halo (e.g., chloro) and the other is $-NO_2$.

[0048] In some embodiments, one R^1 is $-C(Y)NR^bR^{b'}$ and the other is haloalkyl (e.g., trifluoromethyl). In some embodiments, R^b and $R^{b'}$ are both hydrogen.

[0049] In some embodiments, one R^1 is $-C(Y)NR^bR^{b'}$ and the other is haloalkoxy (e.g., trifluoromethoxy). In some embodiments, R^b and $R^{b'}$ are both hydrogen.

[0050] In some embodiments, one R^1 is $-C(Y)NR^bR^{b'}$ and the other is $-S(O)_qR^f$. In some embodiments, R^b and $R^{b'}$ are both hydrogen. In some embodiments, R^f is C_1 - C_8 alkyl (e.g., methyl).

[0051] In some embodiments, one R^1 is $-C(Y)NR^bR^{b'}$ and the other is $-CN$. In some embodiments, R^b and $R^{b'}$ are both hydrogen.

[0052] In some embodiments, one R^1 is $-OR^d$ and the other is C_1 - C_8 alkyl (e.g., methyl). In some embodiments, R^d is C_1 - C_8 alkyl (e.g., methyl).

[0053] In some embodiments, one R^1 is $-OR^d$ and the other is haloalkyl (e.g., trifluoromethyl). In some embodiments, R^d is C_1 - C_8 alkyl (e.g., methyl).

[0054] In some embodiments, one R^1 is $-OR^d$ and the other is $-C(O)OR^a$. In some embodiments, R^d is C_1 - C_8 alkyl (e.g., methyl). In some embodiments, R^a is hydrogen. In some embodiments, R^a is C_1 - C_8 alkyl (e.g., methyl).

[0055] In some embodiments, one R^1 is $-OR^d$ and the other is $-NR^cC(O)R^c$. In some embodiments, R^d is C_1 - C_8 alkyl (e.g., methyl). In some embodiments, R^c is hydrogen and R^c is C_1 - C_8 alkyl (e.g., methyl).

[0056] In some embodiments, one R^1 is haloalkyl (e.g., trifluoromethyl) and the other is $-CN$.

[0057] In some embodiments, each R^1 is independently halo (e.g., all three R^1 are fluoro or all three R^1 are chloro).

[0058] In some embodiments, two R^1 are independently halo (e.g., both are chloro) and the other is C_1 - C_8 alkyl (e.g., methyl). In some embodiments, two R^1 are independently halo (e.g., both are chloro) and the other is heteroaryl (e.g., pyrroly). In some embodiments, two R^1 are independently halo (e.g., both are fluoro) and the other is $-C(Y)NR^bR^{b'}$ (e.g., $-C(O)NH_2$). In some embodiments, two R^1 are independently C_1 - C_8 alkyl (e.g., both are methyl) and the other is halo (e.g., chloro or bromo).

[0059] In some embodiments, one R^1 is C_1 - C_8 alkyl (e.g., methyl), and two R^1 , together with the atoms to which they

are attached, are taken together to form a heterocyclyl ring.

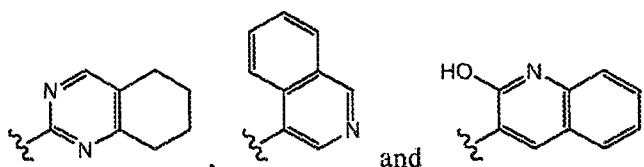
[0060] In some embodiments, one R^1 is $-OR^d$, and two R^1 , together with the atoms to which they are attached, are taken together to form a heterocyclyl ring. In some embodiments, R^d is C_1 - C_8 alkyl (e.g., methyl).

[0061] In some embodiments, B is a 6-membered heteroaryl.

[0062] In some embodiments, B is pyridyl. In some embodiments, B is 3-pyridyl. In some embodiments, m is 2. In some embodiments, m is 3.

[0063] In some embodiments, B is pyrazolyl. In some embodiments, m is 2.

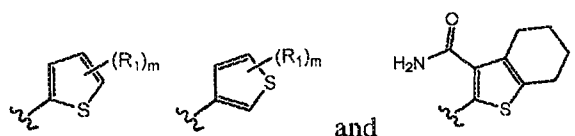
[0064] In some embodiments, B is selected from:



[0065] In some embodiments, B is a 5-membered heteroaryl (e.g., pyrazolyl). In some embodiments, m is 1.

[0066] In some embodiments, m is 2.

[0067] In some embodiments, B is thienyl. In some embodiments, B is selected from:



[0068] In some embodiments, m is 1. In some embodiments, m is 2.

[0069] In some embodiments, R^1 is $-C(Y)NR^bR^{b'}$. In some embodiments, Y is O. In some embodiments, R^b and $R^{b'}$ are both hydrogen.

[0070] In some embodiments, m is 2.

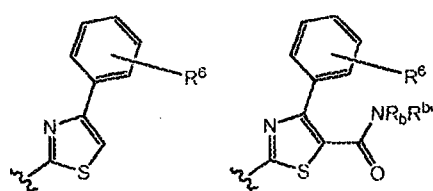
[0071] In some embodiments, one R^1 is C_1 - C_8 alkyl (e.g., methyl) and the other is $-C(Y)NR^bR^{b'}$. In some embodiments, Y is O. In some embodiments, R^b and $R^{b'}$ are both hydrogen.

[0072] In some embodiments, B is thiazolyl.

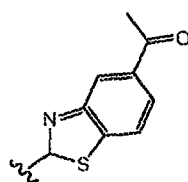
[0073] In some embodiments, m is 1.

[0074] In some embodiments, m is 2.

[0075] In some embodiments, B is:



[0076] In some embodiments, B is:



[0077] In some embodiments, E is aryl (e.g., phenyl).

[0078] In some embodiments, n is 1.

[0079] In some embodiments, R^2 is C_1 - C_8 alkyl (e.g., methyl).

[0080] In some embodiments, R^2 is heterocyclalkyl. In some embodiments, the alkyl is C_1 alkyl. In some embodiments, the alkyl is C_2 alkyl. In some embodiments, the alkyl is C_3 alkyl. In some embodiments, the heterocyclalkyl is morpholino. In some embodiments, the heterocyclalkyl is thiomorpholino.

[0081] In some embodiments, R^2 is haloalkyl (e.g., trifluoromethyl).

[0082] In some embodiments, R^2 is haloalkoxy (e.g., trifluoromethoxy).

[0083] In some embodiments, R^2 is -CN.

[0084] In some embodiments, R^2 is -NO₂.

[0085] In some embodiments, R^2 is -C(O)OR^a. In some embodiments, R^a is hydrogen. In some embodiments, R^a is C_1 - C_8 alkyl (e.g., methyl).

[0086] In some embodiments, R^2 is -C(Y)NR^bR^{b'}. In some embodiments, Y is O. In some embodiments, R^b and R^{b'} are each hydrogen. In some embodiments, R^b and R^{b'} are each independently C_1 - C_8 alkyl (e.g., R^b and R^{b'} are both methyl). In some embodiments, one of R^b and R^{b'} is hydrogen and the other is C_1 - C_8 alkyl (e.g., methyl or ethyl). In some embodiments, one of R^b and R^{b'} is hydrogen and the other is heterocyclalkyl (e.g., -CH₂-CH₂-morpholino). In some embodiments, one of R^b and R^{b'} is hydrogen and the other is haloalkyl (e.g., trifluoroethyl).

[0087] In some embodiments, R^2 is -NR^bR^{b'}. In some embodiments, R^b and R^{b'} are both hydrogen. In some embodiments, R^b and R^{b'} are each independently C_1 - C_8 alkyl (e.g., R^b and R^{b'} are both methyl). In some embodiments, one of R^b and R^{b'} is hydrogen and the other is C_1 - C_8 alkyl (e.g., methyl). In some embodiments, one of R^b and R^{b'} is hydrogen and the other is heterocyclalkyl (e.g., tetrahydropyranyl). In some embodiments, one of R^b and R^{b'} is hydrogen and the other is heterocyclalkyl. In some embodiments, the alkyl is C_1 alkyl. In some embodiments, the alkyl is C_2 alkyl. In some embodiments, the heterocyclalkyl is pyrrolidinyl. In some embodiments, the heterocyclalkyl is tetrahydrofuranyl. In some embodiments, the heterocyclalkyl is tetrahydropyranyl. In some embodiments, one of R^b and R^{b'} is hydrogen and the other is hydroxyalkyl. In some embodiments, the alkyl is C_2 alkyl. In some embodiments, one of R^b and R^{b'} is hydrogen and the other is alkoxyalkyl. In some embodiments, the alkyl is C_2 alkyl. In some embodiments, the alkyl is C_3 alkyl. In some embodiments, the alkoxy is methoxy. In some embodiments, one of R^b and R^{b'} is hydrogen and the other is -C(Y)R^e. In some embodiments, Y is O. In some embodiments, R^e is C_1 - C_8 alkyl (e.g., methyl). In some embodiments, R^e is heterocyclalkyl. In some embodiments, R^e is tetrahydropyranyl.

[0088] In some embodiments, R^2 is -OR^d.

[0089] In some embodiments, R^d is hydrogen. In some embodiments, R^d is C_1 - C_8 alkyl (e.g., methyl).

[0090] In some embodiments, R^d is ethyl. In some embodiments, R^d is C_3 alkyl (e.g., isopropyl or n-propyl). In some embodiments, R^d is optionally substituted heteroalkyl. In some embodiments, R^d is optionally substituted pyridinyl.

[0091] In some embodiments, R^d is n-propyl. In some embodiments, R^d is cyclyl (e.g., cyclopentyl).

[0092] In some embodiments, R^d is heteroalkyl (e.g., -CH₂-pyridyl).

[0093] In some embodiments, R^d is heterocyclalkyl. In some embodiments, the alkyl is C_1 alkyl. In some embodiments, the alkyl is C_2 alkyl. In some embodiments, the alkyl is C_3 alkyl. In some embodiments, the alkyl is C_4 alkyl. In some embodiments, the heterocyclalkyl is morpholino. In some embodiments, the heterocyclalkyl is piperidyl. In some embodiments, the heterocyclalkyl is tetrahydrofuranyl. In some embodiments, R^d is cyclylalkyl (e.g., -CH₂-cyclobutyl).

[0094] In some embodiments, R^d is alkoxyalkyl. In some embodiments, the alkyl is C_2 alkyl. In some embodiments, the alkyl is C_3 alkyl. In some embodiments, the alkoxy is methoxy.

[0095] In some embodiments, R^d is dialkylaminoalkyl. In some embodiments, the alkyl is C_2 alkyl. In some embodiments, the alkyl is C_3 alkyl. In some embodiments, the dialkylamino is dimethylamino.

[0096] In some embodiments, R^2 is -C(Y)R^e. In some embodiments, Y is O. In some embodiments, R^e is heterocyclalkyl. In some embodiments, R^e is piperidyl. In some embodiments, R^e is pyrrolidinyl. In some embodiments, R^e is piperazinyl. In some embodiments, R^e is morpholino. In some embodiments, R^e is thiomorpholino.

[0097] In some embodiments, n is 2.

[0098] In some embodiments, each R² is independently -OR^d.

[0099] In some embodiments, each R^d is C_1 - C_8 alkyl.

[0100] In some embodiments, each R² is methoxy. In some embodiments, one R² is methoxy and the other is ethoxy. In some embodiments, one R² is methoxy and the other is propoxy. In some embodiments, one R² is methoxy and the other is isopropoxy.

[0101] In some embodiments, one R² is methoxy and the other is ethoxy substituted with 1 R⁶. In some embodiments, R⁶ is -NR^bR^{b'}. In some embodiments, R^b and R^{b'} are each independently C_1 - C_8 alkyl (e.g., R^b and R^{b'} are both methyl). In some embodiments, R⁶ is -OR^d. In some embodiments, R^d is C_1 - C_8 alkyl (e.g., methyl).

[0102] In some embodiments, one R² is methoxy and the other is propoxy substituted with 1 R⁶. In some embodiments, R⁶ is -OR^d. In some embodiments, R^d is C_1 - C_8 alkyl (e.g., methyl).

[0103] In some embodiments, one R² is -OR^d and the other is C_1 - C_8 alkyl (e.g., methyl). In some embodiments, R^d is C_1 - C_8 alkyl (e.g., methyl or ethyl).

[0104] In some embodiments, one R² is -OR^d and the other is halo (e.g., chloro). In some embodiments, R^d is C_1 - C_8

alkyl (e.g., methyl).

[0105] In some embodiments, one R^2 is $-OR^d$ and the other is $-CN$. In some embodiments, R^d is C_1-C_8 alkyl (e.g., methyl).

[0106] In some embodiments, one R^2 is $-OR^d$ and the other is $-C(O)OR^a$. In some embodiments, R^d and R^a are both hydrogen.

[0107] In some embodiments, one R^2 is $-OR^d$ and the other is C_1-C_8 alkyl (e.g., methyl). In some embodiments, R^d is C_1-C_8 alkyl (e.g., methyl).

[0108] In some embodiments, one R^2 is $-OR^d$ and the other is $-C(Y)R^e$. In some embodiments, R^d is C_1-C_8 alkyl (e.g., methyl). In some embodiments, Y is O . In some embodiments, R^e is heterocyclyl (e.g., morpholino).

[0109] In some embodiments, one R^2 is halo (e.g., chloro or bromo) and the other is C_1-C_8 alkyl (e.g., methyl).

[0110] In some embodiments, one R^2 is C_1-C_8 alkyl (e.g., methyl) and the other is $-CN$.

[0111] In some embodiments, one R^2 is C_1-C_8 alkyl (e.g., methyl) and the other is heteroaryl (e.g., pyridyl). In some embodiments, the pyridyl is substituted with 1 R^6 . In some embodiments, R^6 is $-OR^d$. In some embodiments, R^d is C_1-C_8 alkyl (e.g., methyl).

[0112] In some embodiments, one R^2 is C_1-C_8 alkyl (e.g., methyl) and the other is heterocyclalkyl (e.g., $-CH_2$ -morpholino).

[0113] In some embodiments, p is 0.

[0114] In some embodiments, p is 1.

[0115] In some embodiments, R^3 is C_1-C_8 alkyl (e.g., methyl). In some embodiments, R^3 is halo (e.g., chloro). In some embodiments, R^3 is haloalkyl (e.g., trifluoromethyl).

[0116] In some embodiments, R^3 is $-OR^d$. In some embodiments, R^d is hydrogen. In some embodiments, R^d is C_1-C_8 alkyl (e.g., methyl).

[0117] In some embodiments, R^3 is $-NR^bR^{b'}$. In some embodiments, R^b and $R^{b'}$ are both hydrogen. In some embodiments, one of R^b and $R^{b'}$ is hydrogen and the other is C_1-C_8 alkyl (e.g., methyl).

[0118] In some embodiments, E is a 5-membered heteroaryl ring.

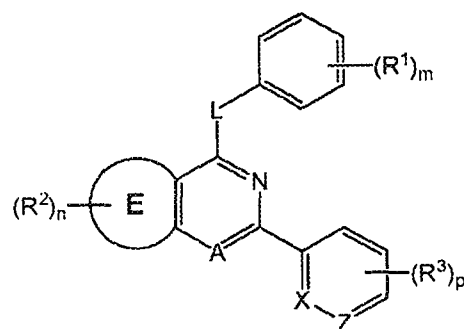
[0119] In some embodiments, E is a thiophene ring.

[0120] In some embodiments, E is a pyrrole ring.

[0121] In some embodiments, n is 1. In some embodiments, R^2 is C_1-C_8 alkyl (e.g., methyl). In some embodiments, E is an N -methylpyrrole ring.

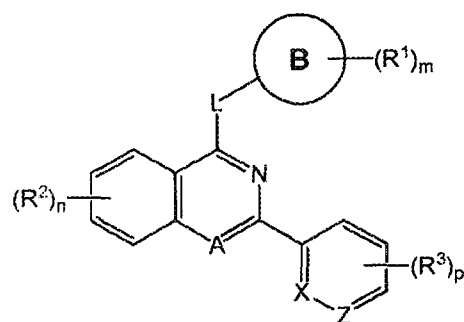
[0122] In some embodiments, L is NR^5 . In some embodiments, R^5 is hydrogen.

[0123] In some embodiments, the compound is:



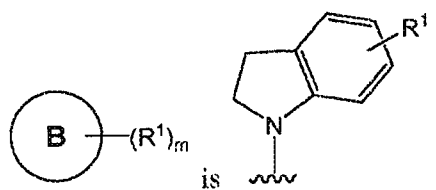
[0124] In some embodiments, R^1 is $-C(Y)NR^bR^{b'}$.

[0125] In some embodiments, the compound is:

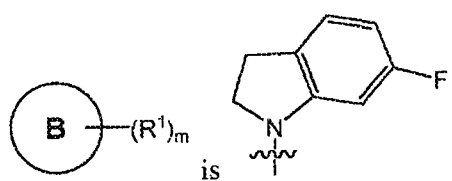


[0126] In some embodiments, R^1 is $-C(Y)NR^bR^{b'}$.

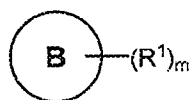
[0127] In some embodiments,



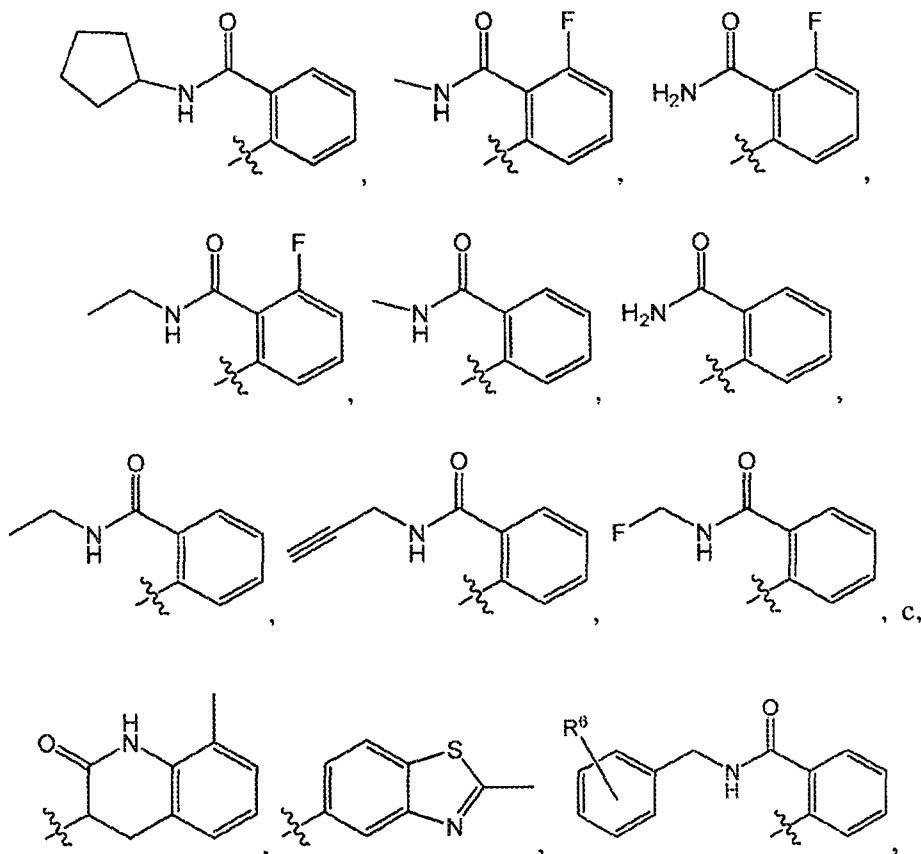
[0128] In some embodiments,



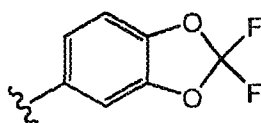
[0129] In some embodiments, L is NH, and



is selected from

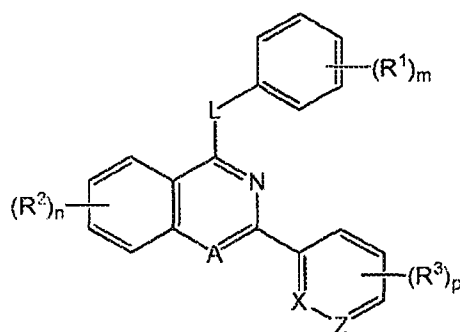


and



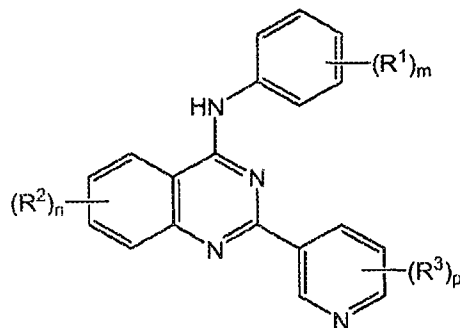
In some embodiments R^6 is halo.

[0130] In some embodiments, the compound is:



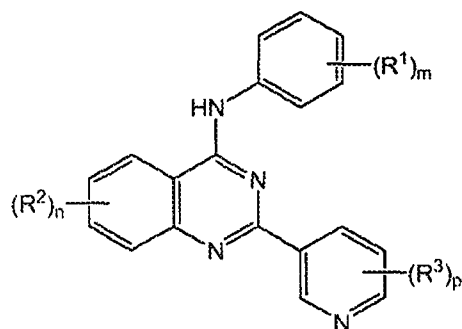
[0131] In some embodiments, R^1 is $-C(Y)NR^bR^{b'}$. In some embodiments, R^1 is halo. In some embodiments, R^2 is C_1-C_4 alkoxy. In some embodiments, m is 2 and two R^1 are 3,4-dichloro; 3,4-difluoro; 3,5-dichloro; 3,5-difluoro; 3-chloro,4-fluoro; or 3-chloro,5-fluoro. In some embodiments, R^2 is $-C(O)NR^bR^{b'}$ and R^3 is H. In some embodiments, R^b and $R^{b'}$ are H. In some embodiments, R^b and $R^{b'}$ are independently C_1-C_4 alkyl or halo-substituted C_1-C_4 alkyl. In some embodiments, R^b is methyl and $R^{b'}$ is trifluoroethyl. In some embodiments, R^1 is C_1-C_4 alkoxy or halo-substituted C_1-C_4 alkoxy. In some embodiments, n and p are zero.

[0132] In some embodiments, the compound is:



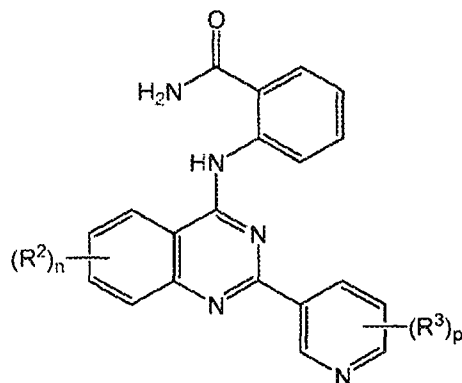
[0133] In some embodiments, R^1 is $-C(Y)NR^bR^{b'}$. In some embodiments, R^1 is halo. In some embodiments, R^2 is C_1-C_4 alkoxy. In some embodiments, m is 2 and two R^1 are 3,4-dichloro; 3,4-difluoro; 3,5-dichloro; 3,5-difluoro; 3-chloro,4-fluoro; or 3-chloro,5-fluoro. In some embodiments, R^2 is $-C(O)NR^bR^{b'}$ and R^3 is H. In some embodiments, R^b and $R^{b'}$ are H. In some embodiments, R^b and $R^{b'}$ are independently C_1-C_4 alkyl or halo-substituted C_1-C_4 alkyl. In some embodiments, R^b is methyl and $R^{b'}$ is trifluoroethyl. In some embodiments, R^1 is C_1-C_4 alkoxy or halo-substituted C_1-C_4 alkoxy. In some embodiments, n and p are zero.

[0134] In some embodiments, the compound is:



[0135] In some embodiments, R^1 is $-C(Y)NR^bR^{b'}$. In some embodiments, R^1 is halo. In some embodiments, R^2 is C_1-C_4 alkoxy. In some embodiments, m is 2 and two R^1 are 3,4-dichloro; 3,4-difluoro; 3,5-dichloro; 3,5-difluoro; 3-chloro,4-fluoro; or 3-chloro,5-fluoro. In some embodiments, R^b and $R^{b'}$ are H. In some embodiments, R^b and $R^{b'}$ are independently C_1-C_4 alkyl or halo-substituted C_1-C_4 alkyl. In some embodiments, R^b is methyl and $R^{b'}$ is trifluoroethyl. In some embodiments, R^1 is C_1-C_4 alkoxy or halo-substituted C_1-C_4 alkoxy. In some embodiments, n and p are zero.

[0136] In some embodiments, the compound is:



[0137] In some embodiments, R^1 is $-C(Y)NR^bR^{b'}$. In some embodiments, R^1 is halo. In some embodiments, R^2 is C_1-C_4 alkoxy. In some embodiments, m is 2 and two R^1 are 3,4-dichloro; 3,4-difluoro; 3,5-dichloro; 3,5-difluoro; 3-chloro,4-fluoro; or 3-chloro,5-fluoro. In some embodiments, R^2 is $-C(O)NR^bR^{b'}$ and R^3 is H. In some embodiments, R^b and $R^{b'}$ are H. In some embodiments, R^b and $R^{b'}$ are independently C_1-C_4 alkyl or halo-substituted C_1-C_4 alkyl. In some embodiments, R^b is methyl and $R^{b'}$ is trifluoroethyl. In some embodiments, R^1 is C_1-C_4 alkoxy or halo-substituted C_1-C_4 alkoxy. In some embodiments, n and p are zero.

Aspects and Embodiments of Compounds of Formula (I)

[0138] In one aspect, the invention features a composition comprising a compound of formula (I) and an acceptable carrier.

[0139] In one aspect, the invention features a pharmaceutical composition comprising a compound of formula (I) and a pharmaceutically acceptable carrier.

[0140] In one aspect, the invention features a kit comprising a composition comprising a compound of formula (I) and an acceptable carrier.

[0141] In one aspect, the invention features a kit comprising a pharmaceutical composition comprising a compound of formula (I) and a pharmaceutically acceptable carrier.

[0142] In one aspect, the invention features a dosage form comprising a composition comprising a compound of formula (I) and an acceptable carrier.

[0143] In one aspect, the invention features a dosage form comprising a pharmaceutical composition comprising a compound of formula (I) and a pharmaceutically acceptable carrier.

[0144] In one aspect, the invention features a method of treating a disorder that would benefit by the modulation of STEP (e.g., by activation or inhibition of STEP) in a subject, the method comprising administering to a subject in need thereof a compound of formula (I). In one aspect, the invention features a method of treating a disorder that would benefit by the inhibition of STEP, the method comprising administering to a subject in need thereof a compound of formula (I).

In some embodiments, the disorder is selected from schizophrenia, schizoaffective disorder, bipolar disorder, manic-depressive disorder, psychosis, mood and anxiety disorders, mania, drug or substance addiction, cognition disorders, learning disabilities, learning and memory disorders, aging and neurologic disorders associated with or linked with cognitive impairments; mild cognitive impairments (MCI), Alzheimer's disease, Alzheimer-related cognition disorders, Huntington's disease, Parkinson's disease, CADASIL syndrome (cerebral autosomal dominant arteriopathy with subcortical infarcts and leukoencephalopathy), amnesia, Wernicke-Korsakoff syndrome, Korsakoff syndrome, mild traumatic head injury (MBTI), traumatic head injury (TBI), fragile X syndrome, stroke, attention-deficit and hyperactivity disorder (ADHD), obsessive compulsive disorder (OCD), post-traumatic stress disorder (PTSD), loss of concentration, autism, cerebral palsy, encephalopathy, and narcolepsy. In some embodiments, the disorder affects learning and memory, neurogenesis, neuronal plasticity, pain perception, mood and anxiety, or neuroendocrine regulation. In some embodiments, the disorder is a cognitive deficit disorder. In some embodiments, the disorder involves pain perception or neuroendocrine regulation. In some embodiments, the disorder affects the central nervous system. In some embodiments the disorder is selected from the group consisting of schizophrenia; refractory, intractable or chronic schizophrenia; emotional disturbance; psychotic disorder; mood disorder; bipolar I type disorder; bipolar II type disorder; depression; endogenous depression; major depression; melancholy and refractory depression; dysthymic disorder; cyclothymic disorder; panic attack; panic disorder; agoraphobia; social phobia; obsessive-compulsive disorder; post-traumatic stress disorder; generalized anxiety disorder; acute stress disorder; hysteria; somatization disorder; conversion disorder, pain disorder; hypochondriasis; factitious disorder; dissociative disorder; sexual dysfunction; sexual desire disorder; sexual arousal disorder; erectile dysfunction; anorexia nervosa; bulimia nervosa; sleep disorder; adjustment disorder; alcohol abuse; alcohol intoxication; drug addiction; stimulant intoxication; narcotism; anhedonia; iatrogenic anhedonia; anhedonia of a psychic or mental cause; anhedonia associated with depression; anhedonia associated with schizophrenia; delirium; cognitive impairment; cognitive impairment associated with Alzheimer's disease, Parkinson's disease and other neurodegenerative diseases; cognitive impairment caused by Alzheimer's disease; Parkinson's disease and associated neurodegenerative diseases; cognitive impairment of schizophrenia; cognitive impairment caused by refractory, intractable or chronic schizophrenia; vomiting; motion sickness; obesity; migraine; pain (ache); mental retardation; autism disorder (autism); Tourette's disorder; tic disorder; attention-deficit/hyperactivity disorder; conduct disorder; and Down's syndrome.

[0145] In one aspect, the invention features a method of treating a condition that would benefit by the modulation of STEP (e.g., by activation or inhibition of STEP) in a subject, the method comprising administering to a subject in need thereof a compound of formula (I). In some embodiments, the condition is selected from decreased neurogenesis, cell resilience, or neuronal plasticity due to normal aging, neurodegenerative disorders of the CNS; Alzheimer's disease, Huntington's disease, fragile X syndrome, amyotrophic lateral sclerosis/Lou Gehrig's disease, stroke, Parkinson's disease, parkinsonism, dementia, Pick disease, Corticobasal degeneration, Multiple system atrophy, Progressive supranuclear palsy, traumatic brain injury, head trauma, mild traumatic head injury (MBTI), traumatic head injury (TBI), encephalopathy, intoxication related to ethanol, alcoholism, fetal alcohol syndrome, drug addiction or drug abuse.

[0146] In some embodiments, a compound of formula (I) is administered in combination with an additional therapeutic agent. In some embodiments, the additional therapeutic agent is an atypical antipsychotic. In some embodiments, the additional therapeutic agent is selected from the group consisting of aripiprazole, clozapine, ziprasidone, risperidone, quetiapine, olanzapine, amisulpride, asenapine, iloperidone, melperone, paliperidone, perospirone, sertindole and sulpiride. In some embodiments, the additional therapeutic agent is a typical antipsychotic. In some embodiments, the additional therapeutic agent is selected from the group consisting of haloperidol, molindone, loxapine, thioridazine, molindone, thiothixene, pimozide, fluphenazine, trifluoperazine, mesoridazine, chlorprothixene, chlorpromazine, perphenazine, triflupromazine and zuclopenthixol.

DETAILED DESCRIPTION

[0147] A compound or composition described herein can be used, e.g., in a method of treating schizophrenia or cognitive deficit. Many of the compounds described herein modulate STEP activity and can be used, e.g., to reduce or inhibit STEP activity, e.g., in a subject.

Definitions

[0148] The term "acyl" refers to an alkylcarbonyl, cycloalkylcarbonyl, arylcarbonyl, heterocyclylcarbonyl, or heteroarylcarbonyl substituent, any of which may be further substituted (e.g., by one or more substituents).

[0149] The term "alkenyl" refers to a straight or branched hydrocarbon chain containing 2-12 carbon atoms (unless otherwise noted) and having one or more double bonds. Examples of alkenyl groups include, but are not limited to, allyl, propenyl, 2-butenyl, 3-hexenyl and 3-octenyl groups. One of the double bond carbons may optionally be the point of attachment of the alkenyl substituent.

[0150] The term "alkenylene" refers to a divalent alkenyl, e.g. $-\text{CH}=\text{CH}-$, $-\text{CH}_2-\text{CH}=\text{CH}-$, and $-\text{CH}=\text{CH}-\text{CH}_2-$.

[0151] The term "alkynyl" refers to a straight or branched hydrocarbon chain containing 2-12 carbon atoms (unless otherwise noted) and characterized in having one or more triple bonds. Examples of alkynyl groups include, but are not limited to, ethynyl, propargyl, and 3-hexynyl. One of the triple bond carbons may optionally be the point of attachment of the alkynyl substituent.

[0152] The term "alkynylene" refers to a divalent alkynyl, e.g. $-\text{CH}\equiv\text{CH}-$, $-\text{CH}_2-\text{CH}\equiv\text{CH}-$, and $-\text{CH}\equiv\text{CH}-\text{CH}_2-$.

[0153] The terms "alkoxyl" or "alkoxy" as used herein refers to an alkyl group, as defined below, having an oxygen radical attached thereto. Representative alkoxy groups include methoxy, ethoxy, propyloxy, tert-butoxy and the like. The term "alkoxyalkyl" refers to an alkyl in which one or more hydrogen atoms are replaced by an alkoxy group.

[0154] An "ether" is two hydrocarbons covalently linked by an oxygen.

[0155] The term "alkyl" refers to the radical of saturated aliphatic groups, including straight-chain alkyl groups, and branched-chain alkyl groups. In preferred embodiments, a straight chain or branched chain alkyl has 12 or fewer carbon atoms in its backbone (unless otherwise noted) e.g., from 1-12, 1-8, 1-6, or 1-4. Exemplary alkyl moieties include methyl, ethyl, propyl (e.g., n-propyl or isopropyl), butyl (e.g., n-butyl, isobutyl or t-butyl), pentyl (e.g., n-pentyl, isopentyl or pentan-3-yl), hexyl and heptyl.

[0156] The term "alkylene" refers to a divalent alkyl, e.g., $-\text{CH}_2-$, $-\text{CH}_2\text{CH}_2-$, and $-\text{CH}_2\text{CH}_2\text{CH}_2-$.

[0157] The term "alkoxylene" refers to an alkylene wherein a CH_2 is substituted with an oxygen. For example, an aryl alkoxylene refers to a group with an alkylene attached to an aryl group through an oxygen, an optionally substituted heteroaryl alkoxylene refers to a group with an alkylene attached to an heteroaryl group through an oxygen.

[0158] The term "amino" refers to $-\text{NH}_2$.

[0159] The term "aminoalkyl" refers to an alkyl in which one or more hydrogen atoms are replaced by an amino group.

[0160] The terms "alkylamino" and "dialkylamino" refer to $-\text{NH}(\text{alkyl})$ and $-\text{N}(\text{alkyl})_2$ radicals respectively.

[0161] The term "aralkylamino" or "arylalkylamino" refers to a $-\text{NH}(\text{aralkyl})$ radical. The term "alkylaminoalkyl" refers to a $(\text{alkyl})\text{NH}-\text{alkyl}-$ radical; the term "dialkylaminoalkyl" refers to an $(\text{alkyl})_2\text{N}-\text{alkyl}-$ radical.

[0162] The term "amido" refers to a $-\text{NHC}(\text{O})-$ or $\text{C}(\text{O})\text{NH}_2$ substituent.

[0163] The term "aryl" refers to a 6-carbon monocyclic, 10-carbon bicyclic, or 14-carbon tricyclic aromatic ring system wherein 0, 1, 2, 3, or 4 atoms of each ring may be substituted by a substituent. Examples of aryl moieties include, but are not limited to, phenyl, naphthyl and the like. The term "arylalkyl" or "aralkyl" refers to alkyl substituted with an aryl. Exemplary aralkyls include but are not limited to benzyl, 1-phenylethyl, 2-phenylethyl, 3-phenylpropyl, 9-fluorenyl, benzhydryl, phenethyl, and trityl groups. The term "arylalkenyl" refers to an alkenyl substituted with an aryl. The term "arylalkynyl" refers to an alkynyl substituted with an aryl. Terms such as "arylC₂-C₆alkyl" are to be read as a further limitation on the size of the alkyl group. The term "arylalkoxy" refers to an alkoxy substituted with aryl. The term "arylenyl" refers to a divalent aryl (i.e., $-\text{Ar}-$).

[0164] The terms "cycloalkyl" or "cyclyl" as employed herein include saturated and partially unsaturated cyclic hydrocarbon groups having 3 to 12 carbons, preferably 3 to 8 carbons, and more preferably 3 to 6 carbons, wherein the cycloalkyl group may be optionally substituted. Exemplary cyclyl groups include, without limitation, cyclopropyl, cyclobutyl, cyclopentyl, cyclopentenyl, cyclohexyl, cyclohexenyl, cycloheptyl, and cyclooctyl. Cyclyl moieties also include both bridged and fused ring systems. Cyclyl groups also include those that are fused to additional ring systems, which may be saturated or unsaturated. A cyclyl group may thus be a bicyclic group in which one ring is saturated or partially unsaturated and the other is fully unsaturated (e.g., indanyl).

[0165] The term "cyclylalkyl" as used herein, refers to an alkyl group substituted with a cyclyl group. Cyclylalkyl includes groups in which more than one hydrogen atom of an alkyl group has been replaced by a cyclyl group.

[0166] The term "cycloalkylalkyl" as used herein, refers to an alkyl group substituted with a cycloalkyl group.

[0167] The term "halo" or "halogen" refers to any radical of fluorine, chlorine, bromine or iodine.

[0168] The term "haloalkyl" refers to an alkyl group that may have any number of hydrogens available on the group replaced with a halogen atom. Representative haloalkyl groups include but are not limited to: $-\text{CH}_2\text{Cl}$, $-\text{CH}_2\text{ClCF}_3$, $-\text{CHBr}_2$, $-\text{CF}_3$, $-\text{CH}_2\text{F}$, $-\text{CHF}_2$, and $-\text{CH}_2\text{CF}_3$. The term "fluoroalkyl" refers to an alkyl group that may have any number of hydrogens available on the group replaced with a fluorine atom. Representative fluoroalkyl groups include but are not limited to: $-\text{CH}_2\text{F}$, $-\text{CH}_2\text{FCF}_3$, $-\text{CHF}_2$ and $-\text{CF}_3$. The term "haloalkoxy" refers to an alkoxy group that may have any number of hydrogen atoms available on the alkyl group replaced with a halogen atom. Representative haloalkoxy groups include but are not limited to: $-\text{OCH}_2\text{Cl}$, $-\text{OCH}_2\text{ClCF}_3$, $-\text{OCHBr}_2$, $-\text{OCHF}_2$ or $-\text{OCF}_3$. The term "fluoroalkoxy" refers to an alkoxy group that may have any number of hydrogens available on the group replaced with a fluorine atom. Representative fluoroalkoxy groups include but are not limited to: $-\text{OCH}_2\text{F}$, $-\text{OCH}_2\text{FCF}_3$, $-\text{OCHF}_2$ or $-\text{OCF}_3$.

[0169] The term "heteroatom" as used herein means an atom of any element other than carbon or hydrogen. Preferred heteroatoms are nitrogen, oxygen, sulfur, phosphorus and silicon. A heteroatom may be present in any oxidation state (e.g., any oxidized form of nitrogen, sulfur, phosphorus or silicon) and any charged state (e.g., the quaternized form of any basic nitrogen), and includes a substitutable nitrogen of a heterocyclic ring, for example N (as in 3,4-dihydro-2H-pyrrolyl), NH (as in pyrrolidinyl) or NR^+ (as in N-substituted pyrrolidinyl).

[0170] The term "heteroaryl" refers to an aromatic 5-8 membered monocyclic, 8-12 membered bicyclic, or 11-14 membered tricyclic ring system having 1-3 heteroatoms if monocyclic, 1-6 heteroatoms if bicyclic, or 1-9 heteroatoms if tricyclic, said heteroatoms selected from O, N, or S (e.g., carbon atoms and 1-3, 1-6, or 1-9 heteroatoms of N, O, or S if monocyclic, bicyclic, or tricyclic, respectively), wherein 0, 1, 2, 3, or 4 atoms of each ring may be substituted by a substituent. Examples of heteroaryl groups include pyridyl, furyl or furanyl, imidazolyl, benzimidazolyl, pyrimidinyl, thiophenyl or thienyl, quinolinyl, indolyl, thiazolyl, oxazolyl and the like. The term "heteroarylalkyl" or the term "heteroaralkyl" refers to an alkyl substituted with a heteroaryl. The term "heteroarylalkenyl" refers to an alkenyl substituted with a heteroaryl. The term "heteroarylalkynyl" refers to an alkynyl substituted with a heteroaryl. The term "heteroarylalkoxy" refers to an alkoxy substituted with heteroaryl.

[0171] The term "heteroaryl" refers to a group having 5 to 14 ring atoms, preferably 5, 6, 9, or 10 ring atoms; having 6, 10, or 14 π electrons shared in a cyclic array; and having, in addition to carbon atoms, from one to five heteroatoms. A heteroaryl group may be mono-, bi-, tri-, or polycyclic, preferably mono-, bi-, or tricyclic, more preferably mono- or bicyclic. When a heteroaryl is substituted by a hydroxy group, it also includes its corresponding tautomer. The term "heteroaryl," as used herein, also includes groups in which a heteroaromatic ring is fused to one or more aryl rings. Nonlimiting examples of heteroaryl groups include thienyl, furanyl, pyrrolyl, imidazolyl, pyrazolyl, triazolyl, tetrazolyl, oxazolyl, isoxazolyl, oxadiazolyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyridazinyl, pyrimidinyl, pyrazinyl, indoliziny, purinyl, naphthyridinyl, pteridinyl, indolyl, isoindolyl, benzothienyl, benzofuranyl, dibenzofuranyl, indazolyl, benzimidazolyl, benzthiazolyl, quinolyl, isoquinolyl, cinnolinyl, phthalazinyl, quinazolinyl, quinoxalinyl, 4H-quinoliziny, carbazolyl, acridinyl, phenazinyl, phenothiazinyl, phenoxazinyl, tetrahydroquinoliny, tetrahydroisoquinoliny, and pyrido[2,3-b]-1,4-oxazin-3(4H)-one. The term "heteroaryl" may be used interchangeably with the terms "heteroaryl ring", "heteroaryl group," or "heteroaromatic," any of which terms include rings that are optionally substituted. A ring nitrogen atom of a heteroaryl may be oxidized to form the corresponding N-oxide compound. A nonlimiting example of such a heteroaryl having an oxidized ring nitrogen atom is N-oxopyridyl.

[0172] The term "heteroarylalkyl" or "heteroaralkyl" refers to an alkyl group substituted by a heteroaryl. Heteroaralkyl includes groups in which more than one hydrogen atom has been replaced by a heteroaryl group.

[0173] As used herein, the terms "heterocycle," "heterocyclyl" and "heterocyclic ring" are used interchangeably and refer to a stable 3- to 8-membered monocyclic or 7-10-membered bicyclic heterocyclic moiety that is either saturated or partially unsaturated, and having, in addition to carbon atoms, one or more, preferably one to four, heteroatoms, as defined above. When used in reference to a ring atom of a heterocycle, the term "nitrogen" includes a substituted nitrogen. As an example, in a saturated or partially unsaturated ring having 0-3 heteroatoms selected from oxygen, sulfur or nitrogen, the nitrogen may be N (as in 3,4-dihydro-2H-pyrrolyl), NH (as in pyrrolidinyl), or NR⁺ (as in N-substituted pyrrolidinyl). A heterocyclic ring can be attached to its pendant group at any heteroatom or carbon atom that results in a stable structure and any of the ring atoms can be optionally substituted. Examples of such saturated or partially unsaturated heterocyclic radicals include, without limitation, tetrahydrofuranyl, tetrahydropyranyl, tetrahydrothienyl, piperidinyl, decahydroquinoliny, oxazolidinyl, piperazinyl, dioxanyl, dioxolanyl, diazepinyl, oxazepinyl, thiazepinyl, morpholinyl, and thiomorpholinyl. A heterocyclyl group may be mono-, bi-, tri-, or polycyclic, preferably mono-, bi-, or tricyclic, more preferably mono- or bicyclic. Additionally, a heterocyclic ring also includes groups in which the heterocyclyl ring is fused to one or more aryl, heteroaryl or cyclyl rings. A ring nitrogen atom of a heterocyclic ring also may be oxidized to form the corresponding N-hydroxy compound.

[0174] The term "heterocyclylalkyl" refers to an alkyl group substituted by a heterocyclyl. Heterocyclylalkyl includes groups in which more than one hydrogen atom has been replaced by a heterocyclyl group.

[0175] The terms "heteraralkyl" and "heteroaralkyl", as used herein, refers to an alkyl group substituted with a heteroaryl group. Exemplary heteroaralkyl groups include but are not limited to methylpyridyl or methylpyrimidyl.

[0176] The term "heterocyclyl" or "heterocyclylalkyl" refers to a nonaromatic 5-8 membered monocyclic, 5-12 membered bicyclic, or 11-14 membered tricyclic ring system having 1-3 heteroatoms if monocyclic, 1-6 heteroatoms if bicyclic, or 1-9 heteroatoms if tricyclic, said heteroatoms selected from O, N, or S (e.g., carbon atoms and 1-3, 1-6, or 1-9 heteroatoms of N, O, or S if monocyclic, bicyclic, or tricyclic, respectively), wherein 0, 1, 2 or 3 atoms of each ring may be substituted by a substituent. Examples of heterocyclyl groups include piperazinyl, pyrrolidinyl, dioxanyl, morpholinyl, tetrahydrofuranyl, and include both bridged and fused ring systems. The term "heterocyclylalkyl" refers to an alkyl substituted with a heterocyclyl.

[0177] The term "heterocyclylalkyl", as used herein, refers to an alkyl group substituted with a heterocycle group.

[0178] The term "heteroalkyl," as used herein, refers to a saturate or unsaturated, straight or branched chain aliphatic group, wherein one or more of the carbon atoms in the chain are independently replaced by a heteroatom. Exemplary hetero atoms include O, S, and N.

[0179] In the case of aralkyl, heteroaralkyl, cyclylalkyl, heterocyclylalkyl etc., groups described as optionally substituted, it is intended that either or both aryl, heteroaryl, cyclyl, heterocyclyl and alkyl moieties may be independently optionally substituted or unsubstituted.

[0180] The term "hydroxyalkyl" refers to an alkyl in which one or more hydrogen atoms are replaced by a hydroxy group.

The term "oxo" refers to an oxygen atom (=O), which forms a carbonyl when attached to carbon, an N-oxide when attached to nitrogen, and a sulfoxide or sulfone when attached to sulfur.

The term "thioalkyl" as used herein refers to an -S(alkyl) group, where the point of attachment is through the sulfur atom and the alkyl group is as defined above. The term "thiono" or "thioxo" refers to a sulfur atom (=S), which forms a thioketone when attached to carbon.

[0181] The term "substituted" refers to moieties having substituents replacing a hydrogen on one or more carbons of the backbone. It will be understood that "substitution" or "substituted with" includes the implicit proviso that such substitution is in accordance with permitted valence of the substituted atom and the substituent, and that the substitution results in a stable compound, e.g., which does not spontaneously undergo transformation such as by rearrangement, cyclization, elimination, etc. As used herein, the term "substituted" is contemplated to include all permissible substituents of organic compounds. In a broad aspect, the permissible substituents include acyclic and cyclic, branched and unbranched, carbocyclic and heterocyclic, aromatic and non-aromatic substituents of organic compounds. The permissible substituents can be one or more and the same or different for appropriate organic compounds. For purposes of this invention, the heteroatoms such as nitrogen may have hydrogen substituents and/or any permissible substituents of organic compounds described herein which satisfy the valences of the heteroatoms.

[0182] The term "substituent" refers to a group "substituted" on a moiety described herein. Any atom on any substituent can be substituted. Substituents can include any substituents described herein. Exemplary substituents include, without limitation, alkyl (e.g., C1, C2, C3, C4, C5, C6, C7, C8, C9, C10, C11, C12 straight or branched chain alkyl), cycloalkyl, haloalkyl (e.g., perfluoroalkyl such as CF₃), aryl, heteroaryl, aralkyl, heteroaralkyl, heterocyclyl, alkenyl, alkynyl, cycloalkenyl, heterocycloalkenyl, alkoxy, haloalkoxy (e.g., perfluoroalkoxy such as OCF₃), halo, hydroxy, carboxy, carboxylate, cyano, nitro, amino, alkylamino, SO₃H, sulfate, phosphate, methylenedioxy (-O-CH₂-O- wherein oxygens are attached to vicinal atoms), ethylenedioxy, oxo, thioxo (e.g., C=S), imino (alkyl, aryl, aralkyl), S(O)_nalkyl (where n is 0-2), S(O)_n aryl (where n is 0-2), S(O)_n heteroaryl (where n is 0-2), S(O)_n heterocyclyl (where n is 0-2), amine (mono-, di-, alkyl, cycloalkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, and combinations thereof), ester (alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl), amide (mono-, di-, alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, and combinations thereof), sulfonamide (mono-, di-, alkyl, aralkyl, heteroaralkyl, and combinations thereof). In one aspect, the substituents on a group are independently any one single, or any subset of the aforementioned substituents. In another aspect, a substituent may itself be substituted with any one of the above substituents.

[0183] As used herein, the phrase "optionally substituted" is used interchangeably with the phrase "substituted or unsubstituted." In general, the term "substituted", whether preceded by the term "optionally" or not, means that a hydrogen radical of the designated moiety is replaced with the radical of a specified substituent, provided that the substitution results in a stable or chemically feasible compound. The term "substitutable", when used in reference to a designated atom, means that attached to the atom is a hydrogen radical, which hydrogen atom can be replaced with the radical of a suitable substituent. Unless otherwise indicated, an "optionally substituted" group may have a substituent at each substitutable position of the group, and when more than one position in any given structure may be substituted with more than one substituent selected from a specified group, the substituent may be either the same or different at every position. Combinations of substituents envisioned by this invention are preferably those that result in the formation of stable or chemically feasible compounds.

[0184] As used herein, the term "optionally substituted" means substituted or unsubstituted.

[0185] As used herein, the term "partially unsaturated" refers to a moiety that includes at least one double or triple bond between atoms. The term "partially unsaturated" encompasses rings, e.g., having one or more sites of unsaturation, but that are not completely unsaturated so as to be aryl or heteroaryl.

[0186] The term "chiral" refers to molecules which have the property of non-superimposability of the mirror image partner, while the term "achiral" refers to molecules which are superimposable on their mirror image partner. With respect to the nomenclature of a chiral center, terms "R" and "S" configuration are as defined by the IUPAC Recommendations. The term "enantiomers" refers to two stereoisomers of a compound which are non-superimposable mirror images of one another. An equimolar mixture of two enantiomers is called a "racemic mixture" or a "racemate." The term "isomers" or "stereoisomers" refers to compounds which have identical chemical constitution, but differ with regard to the arrangement of the atoms or groups in space. For example, isomers include *cis*- and *trans*-isomers, *E*- and *Z*-isomers, R- and S-enantiomers, diastereomers, (D)-isomers, (L)-isomers, racemic mixtures thereof, and other mixtures thereof. The term "diastereomers" refers to stereoisomers with two or more centers of dissymmetry and whose molecules are not mirror images of one another.

[0187] The term "administration" or "administering" includes routes of introducing the compounds, or a composition thereof, of the invention to a subject to perform their intended function. Examples of routes of administration that may be used include injection (subcutaneous, intravenous, parenterally, intraperitoneally, intrathecal), oral, inhalation, rectal and transdermal. The pharmaceutical compositions may be given by forms suitable for each administration route. For example, these compositions are administered in tablets or capsule form, by injection, inhalation, eye lotion, ointment, suppository, etc. administration by injection, infusion or inhalation; topical by lotion or ointment; and rectal by suppositories.

Oral administration is preferred. The injection can be bolus or can be continuous infusion. Depending on the route of administration, a compound described herein can be coated with or disposed in a selected material to protect it from natural conditions which may detrimentally affect its ability to perform its intended function. A compound or composition described herein can be administered alone, or in conjunction with either another agent as described above or with a pharmaceutically-acceptable carrier, or both. A compound or composition described herein can be administered prior to the administration of the other agent, simultaneously with the agent, or after the administration of the agent. Furthermore, a compound described herein can also be administered in a pro-drug form which is converted into its active metabolite, or more active metabolite *in vivo*.

[0188] The language "biological activities" of a compound described herein includes all activities elicited by a compound described herein in a responsive subject or cell. It includes genomic and non-genomic activities elicited by these compounds.

[0189] The terms "inhibit" and "inhibitor" as used herein means an agent that measurably slows or stops the production of STriatal-Enriched tyrosine Phosphatase (STEP), or decreases or inactivates STEP, or interferes with STEP-mediated biological pathways. Inhibitors of STEP include compounds of the invention, e.g., compounds of Formulas (I), (II), or (III). A compound can be evaluated to determine if it is an inhibitor by measuring either directly or indirectly the activity of STEP in the presence of the compound suspected to inhibit STEP. Exemplary methods of measure STEP inhibition are described in the EXAMPLES herein.

[0190] An "effective amount" or "an amount effective" refers to an amount of the compound or composition which is effective, upon single or multiple dose administrations to a subject and for periods of time necessary, in treating a cell, or curing, alleviating, relieving or improving a symptom of a disorder, e.g., a disorder described herein. An effective amount of a compound described herein may vary according to factors such as the disease state, age, and weight of the subject, and the ability of a compound described herein to elicit a desired response in the subject. Dosage regimens may be adjusted to provide the optimum therapeutic response. An effective amount is also one in which any toxic or detrimental effects (e.g., side effects) of a compound described herein are outweighed by the therapeutically beneficial effects. The term "effective amount" includes an amount effective, at dosages and for periods of time necessary, to achieve the desired result, e.g., modulate or regulate protein tyrosine phosphatases, e.g., STEP, in a subject and/or treat a disorder described herein such as a protein tyrosine phosphatase related disorder. Exemplary disorders include those related to cognition, learning and memory, neurogenesis. An effective amount may also affect neuronal plasticity, pain perception, mood and anxiety, and neuroendocrine regulation.

[0191] An effective amount of a compound described herein may vary according to factors such as the disease state, age, and weight of the subject, and the ability of a compound described herein to elicit a desired response in the subject. Dosage regimens may be adjusted to provide the optimum therapeutic response. An effective amount is also one in which any toxic or detrimental effects (e.g., side effects) of a compound described herein are outweighed by the therapeutically beneficial effects.

[0192] A therapeutically effective amount of a compound described herein (i.e., an effective dosage) may range from about 0.001 to 50 mg/kg body weight, preferably about 0.01 to 40 mg/kg body weight, more preferably about 0.1 to 35 mg/kg body weight, still more preferably about 1 to 30 mg/kg, and even more preferably about 10 to 30 mg/kg. The skilled artisan will appreciate that certain factors may influence the dosage required to effectively treat a subject, including but not limited to the severity of the disease or disorder, previous treatments, the general health and/or age of the subject, and other diseases present. Moreover, treatment of a subject with a therapeutically effective amount of a compound described herein can include a single treatment or, preferably, can include a series of treatments. In one example, a subject is treated with a compound described herein in the range of between about 0.1 to 20 mg/kg body weight, one time per week for between about 1 to 10 weeks, preferably between 2 to 8 weeks, more preferably between about 3 to 7 weeks, and even more preferably for about 4, 5, or 6 weeks. It will also be appreciated that the effective dosage of a compound described herein used for treatment may increase or decrease over the course of a particular treatment.

[0193] As used herein, an amount of a compound effective to prevent a disorder, or "a prophylactically effective amount" of the compound refers to an amount effective, upon single- or multiple-dose administration to the subject, in preventing or delaying the occurrence of the onset or recurrence of a disorder or a symptom of the disorder.

[0194] The language "improved biological properties" refers to any activity inherent in a compound described herein that enhances its effectiveness *in vivo*. In a preferred embodiment, this term refers to any qualitative or quantitative improved therapeutic property of a compound described herein, such as reduced off-target effects.

[0195] The term "modulate" refers to an increase or decrease, e.g., in the activity of an enzyme in response to exposure to a compound or composition described herein, e.g., the activation or inhibition of STEP, in at least a sub-population of cells in a subject such that a desired end result is achieved (e.g., a therapeutic result). In some embodiments, a compound as described herein inhibits a target described herein, e.g., STEP. In some embodiments, a compound as described herein is activates a target described herein, e.g., STEP.

[0196] As used herein, the term "subject" is intended to include human and non-human animals. Exemplary human subjects include a human patient having a disorder, e.g., a disorder described herein, or a normal subject. The term

"non-human animals" includes all vertebrates, e.g., non-mammals (such as chickens, amphibians, reptiles) and mammals, such as non-human primates, domesticated and/or agriculturally useful animals, e.g., sheep, dog, cat, cow, pig, etc.

[0197] As used herein, the term "treat" or "treating" is defined as applying or administering a compound or composition, alone or in combination with a second compound or composition, to a subject, e.g., a patient, or applying or administering the compound or composition to an isolated tissue or cell, e.g., cell line, from a subject, e.g., a patient, who has a disorder (e.g., a disorder as described herein), a symptom of a disorder, or a predisposition toward a disorder, with the purpose to cure, heal, alleviate, relieve, alter, remedy, ameliorate, improve or affect the disorder, one or more symptoms of the disorder or the predisposition toward the disorder (e.g., to prevent at least one symptom of the disorder or to delay onset of at least one symptom of the disorder).

[0198] The phrases "parenteral administration" and "administered parenterally" as used herein means modes of administration other than enteral and topical administration, usually by injection, and includes, without limitation, intravenous, intramuscular, intraarterial, intrathecal, intracapsular, intraorbital, intracardiac, intradermal, intraperitoneal, transtracheal, subcutaneous, subcuticular, intraarticular, subcapsular, subarachnoid, intraspinal and intrasternal injection and infusion.

[0199] The term "prodrug" or "pro-drug" includes compounds with moieties that can be metabolized *in vivo*. Generally, the prodrugs are metabolized *in vivo* by esterases or by other mechanisms to active drugs. Examples of prodrugs and their uses are well known in the art (See, e.g., Berge et al. (1977) "Pharmaceutical Salts", J. Pharm. Sci. 66:1-19). The prodrugs can be prepared *in situ* during the final isolation and purification of the compounds, or by separately reacting the purified compound in its free acid form or hydroxyl with a suitable esterifying agent. Hydroxyl groups can be converted into esters *via* treatment with a carboxylic acid. Examples of prodrug moieties include substituted and unsubstituted, branch or unbranched lower alkyl ester moieties, (e.g., propionic acid esters), lower alkenyl esters, di-lower alkyl-amino lower-alkyl esters (e.g., dimethylaminoethyl ester), acylamino lower alkyl esters (e.g., acetyloxymethyl ester), acyloxy lower alkyl esters (e.g., pivaloyloxymethyl ester), aryl esters (phenyl ester), aryl-lower alkyl esters (e.g., benzyl ester), substituted (e.g., with methyl, halo, or methoxy substituents) aryl and aryl-lower alkyl esters, amides, lower-alkyl amides, di-lower alkyl amides, and hydroxy amides. Preferred prodrug moieties are propionic acid esters and acyl esters. Prodrugs which are converted to active forms through other mechanisms *in vivo* are also included.

[0200] The language "a prophylactically effective amount" of a compound refers to an amount of a compound described herein any formula herein or otherwise described herein which is effective, upon single or multiple dose administration to the patient, in preventing or treating a disease or condition.

[0201] The language "reduced off-target effects" is intended to include a reduction in any undesired side effect elicited by a compound described herein when administered *in vivo*. In some embodiments, a compound described herein has little to no cardio and/or pulmonary toxicity (e.g., when administered to a subject). In some embodiments, a compound described herein has little to no hallucinogenic activity (e.g., when administered to a subject).

[0202] The term "selective" means a greater activity against a first target. In some embodiments a compound has a selectivity of at least 1.25-fold, at least 1.5 fold, at least 2-fold, at least 3-fold, at least 4-fold, at least 5-fold, at least 6-fold, at least 10-fold or at least 100-fold greater towards a first target relative to a second target. In some embodiments, a compound described herein, e.g., a compound of Formulas (I) is selective toward STEP relative to one or more other protein tyrosine phosphatases.

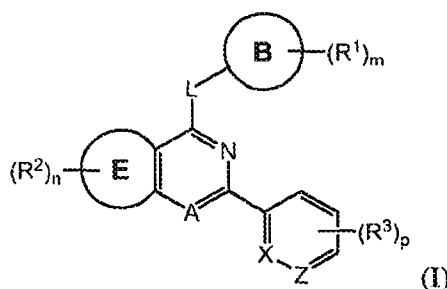
[0203] The term "subject" includes organisms which are capable of suffering from a serotonin-receptor-related disorder or who could otherwise benefit from the administration of a compound described herein of the invention, such as human and non-human animals. Preferred humans include human patients suffering from or prone to suffering from a serotonin-related disorder or associated state, as described herein. The term "non-human animals" of the invention includes all vertebrates, e.g., mammals, e.g., rodents, e.g., mice, and non-mammals, such as non-human primates, e.g., sheep, dog, cow, chickens, amphibians, reptiles, etc.

[0204] The phrases "systemic administration," "administered systemically," "peripheral administration" and "administered peripherally" as used herein mean the administration of a compound described herein(s), drug or other material, such that it enters the patient's system and, thus, is subject to metabolism and other like processes, for example, subcutaneous administration.

Compounds

[0205] The compounds described herein can be used for a variety of purposes, e.g., therapeutic purposes. Many of the compounds modulate STEP activity and can be used, for example to inhibit STEP, e.g., in a subject.

[0206] Exemplary compounds include a compound of formula (I):



wherein A, B, E, L, X, Z, R¹, R², R³, m, n and p are as defined above in the section relating to compound of Formula (I). In preferred embodiments, L is NH, B is aryl (e.g., phenyl) that may be optionally substituted, E is aryl (e.g., phenyl), A is N, X is CH and Z is N.

[0207] The present invention includes compounds that differ only in the presence of one or more isotopically enriched atoms. For example, compounds having the present structures except for the replacement of hydrogen by deuterium or tritium, the replacement of a carbon by a ¹³C- or ¹⁴C-enriched carbon, or the replacement of a fluorine by a ¹⁹F-enriched fluorine are within the scope of this invention. Such compounds are useful, for example, as analytical tools or probes in biological assays, or as bioactive agents.

[0208] In the compounds of the present invention, any atom not specifically designated as a particular isotope is meant to represent any stable isotope of that atom unless otherwise stated (e.g., hydrogen, ²H or deuterium and ³H or tritium). The formulas described herein may or may not indicate whether atoms at certain positions are isotopically enriched. When a structural formula is silent with respect to whether a particular position is isotopically enriched, it is to be understood that the isotopes at that particular position are present in natural abundance or, that the particular position is isotopically enriched with one or more naturally occurring stable isotopes. For example, the formula -CH₂- represents the following possible structures: -CH₂-, -CHD- or -CD₂-.

[0209] The variable "D" is defined as deuterium.

[0210] The terms "compound" or "compounds," when referring to a compound of this invention or a compound described herein, refers to a collection of molecules having an identical chemical structure, except that there may be isotopic variation among the constituent atoms of the molecules. Thus, it will be clear to those of skill in the art that a compound represented by a particular chemical structure containing indicated hydrogen atoms will contain lesser amounts of isotopologues having deuterium atoms at one or more of the designated hydrogen positions in that structure. Alternatively, a compound represented by a particular chemical structure containing indicated deuterium atoms will contain lesser amounts of isotopologues having hydrogen atoms at one or more of the designated deuterium positions in that structure. The relative amount of such isotopologues in a compound of this invention will depend on a number of factors including isotopic purity of deuterated reagents used to make the compound and the efficiency of incorporation of deuterium in the various synthetic steps used to prepare the compound. The relative amount of such isotopologues in total will be less than 55% of the compound. In other embodiments, the relative amount of such isotopologues in total will be less than 50%, less than 45%, less than 40%, less than 35%, less than 35%, less than 15%, less than 10%, less than 5%, less than 1% or less than 0.5% of the compound.

[0211] The term "isotopologue" refers to a species that differs from a specific compound of this invention only in the isotopic composition thereof. Isotopologues can differ in the level of isotopic enrichment at one or more positions and/or in the position(s) of isotopic enrichment.

[0212] The compounds of this invention may contain one or more asymmetric centers and thus occur as racemates and racemic mixtures, single enantiomers, individual diastereomers and diastereomeric mixtures. Described herein are enantiomerically enriched compounds (e.g., a compound resolved to an enantiomeric excess of 60%, 70%, 80%, 85%, 90%, 95%, 99% or greater). All such isomeric forms of these compounds are expressly included in the present invention. The compounds of this invention may also contain linkages (e.g., carbon-carbon bonds) or substituents that can restrict bond rotation, e.g. restriction resulting from the presence of a ring or double bond. Accordingly, all *cis/trans* and *E/Z* isomers are expressly included in the present invention. The compounds of this invention may also be represented in multiple tautomeric forms, in such instances, the invention expressly includes all tautomeric forms of the compounds described herein, even though only a single tautomeric form may be represented (e.g., alkylation of a ring system may result in alkylation at multiple sites, the invention expressly includes all such reaction products). All such isomeric forms of such compounds are expressly included in the present invention. All crystal forms of the compounds described herein are expressly included in the present invention.

[0213] Naturally occurring or synthetic isomers can be separated in several ways known in the art. Methods for separating a racemic mixture of two enantiomers include chromatography using a chiral stationary phase (see, e.g., "Chiral Liquid Chromatography," W.J. Lough, Ed. Chapman and Hall, New York (1989)). Enantiomers can also be separated

by classical resolution techniques. For example, formation of diastereomeric salts and fractional crystallization can be used to separate enantiomers. For the separation of enantiomers of carboxylic acids, the diastereomeric salts can be formed by addition of enantiomerically pure chiral bases such as brucine, quinine, ephedrine, strychnine, and the like. Alternatively, diastereomeric esters can be formed with enantiomerically pure chiral alcohols such as menthol, followed by separation of the diastereomeric esters and hydrolysis to yield the free, enantiomerically enriched carboxylic acid. For separation of the optical isomers of amino compounds, addition of chiral carboxylic or sulfonic acids, such as camphorsulfonic acid, tartaric acid, mandelic acid, or lactic acid can result in formation of the diastereomeric salts. For example a compound can be resolved to an enantiomeric excess (e.g., 60%, 70%, 80%, 85%, 90%, 95%, 99% or greater) via formation of diastereomeric salts, e.g. with a chiral base, e.g., (+) or (-) α -methylbenzylamine, or via high performance liquid chromatography using a chiral column. In some embodiments a product is purified directly on a chiral column to provide enantiomerically enriched compound.

[0214] Combinations of substituents and variables envisioned by this invention are only those that result in the formation of stable compounds. The term "stable", as used herein, refers to compounds which possess stability sufficient to allow manufacture and which maintains the integrity of the compound for a sufficient period of time to be useful for the purposes detailed herein (e.g., therapeutic administration to a subject).

[0215] Compounds of formulas (I) are described herein, for example as provided in the summary above. Exemplary compounds are shown in Tables 1-30 in the Examples section.

Synthetic Methods

[0216] A compound described herein may be prepared via a variety of synthetic methods. Representative syntheses are shown in the Examples section.

[0217] As can be appreciated by the skilled artisan, further methods of synthesizing the compounds of the formulae herein will be evident to those of ordinary skill in the art. Additionally, the various synthetic steps may be performed in an alternate sequence or order to give the desired compounds. Synthetic chemistry transformations and protecting group methodologies (protection and deprotection) useful in synthesizing the compounds described herein are known in the art and include, for example, those such as described in R. Larock, *Comprehensive Organic Transformations*, VCH Publishers (1989); T.W. Greene and P.G.M. Wuts, *Protective Groups in Organic Synthesis*, 2d. Ed., John Wiley and Sons (1991); L. Fieser and M. Fieser, *Fieser and Fieser's Reagents for Organic Synthesis*, John Wiley and Sons (1994); and L. Paquette, ed., *Encyclopedia of Reagents for Organic Synthesis*, John Wiley and Sons (1995), and subsequent editions thereof.

[0218] Additionally, the compounds disclosed herein can be prepared on a solid support. The term "solid support" refers a material to which a compound is attached to facilitate identification, isolation, purification, or chemical reaction selectivity of the compound. Such materials are known in the art and include, for example, beads, pellets, disks, fibers, gels, or particles such as cellulose beads, pore-glass beads, silica gels, polystyrene beads optionally cross-linked with divinylbenzene and optionally grafted with polyethylene glycol, poly-acrylamide beads, latex beads, dimethylacrylamide beads optionally cross-linked with N,N'-bis-acryloyl ethylene diamine, glass particles coated with hydrophobic polymer, and material having a rigid or semi-rigid surface. The solid supports optionally have functional groups such as amino, hydroxy, carboxy, or halo groups, (see, Obrecht, D. and Villagrodó, J.M., *Solid-Supported Combinatorial and Parallel Synthesis of Small-Molecular-Weight Compound Libraries*, Pergamon-Elsevier Science Limited (1998)), and include those useful in techniques such as the "split and pool" or "parallel" synthesis techniques, solid-phase and solution-phase techniques, and encoding techniques (see, for example, Czarnik, A.W., *Curr. Opin. Chem. Bio.*, (1997) 1,60).

[0219] A compound described herein may be modified by appending appropriate functionalities to enhance selective biological properties. Such modifications are known in the art and include those which increase biological penetration into a given biological compartment (e.g., brain, blood, lymphatic system, central nervous system), increase oral availability, increase solubility to allow administration by injection, alter metabolism and alter rate of excretion.

[0220] Included herein are pharmaceutically acceptable derivatives or prodrugs of the compounds described herein. A "pharmaceutically acceptable derivative or prodrug" means any pharmaceutically acceptable salt, ester, salt of an ester, or other derivative of a compound of this invention (for example an imide ester of an amide), which, upon administration to a recipient, is capable of providing (directly or indirectly) a compound described herein. Particularly favored derivatives and prodrugs are those that increase the bioavailability of the compounds of this invention when such compounds are administered to a mammal (e.g., by allowing an orally administered compound to be more readily absorbed into the blood) or which enhance delivery of the parent compound to a biological compartment (e.g., the brain or lymphatic system) relative to the parent species. In an exemplary embodiment, the prodrug is a derivative including a group that enhances aqueous solubility or active transport through the gut membrane is appended to the structure of formulae described herein. In another exemplary embodiment, the prodrug is suitable for treatment or prevention of those diseases and conditions that require the drug molecule to cross the blood brain barrier. In a preferred embodiment, the prodrug enters the brain, where it is converted into the active form of the drug molecule.

[0221] Pharmaceutically acceptable salts of the compounds of this invention include those derived from pharmaceutically acceptable inorganic and organic acids and bases. Examples of suitable acid salts include acetate, adipate, benzoate, benzenesulfonate, butyrate, citrate, digluconate, dodecylsulfate, formate, fumarate, glycolate, hemisulfate, heptanoate, hexanoate, hydrochloride, hydrobromide, hydroiodide, lactate, maleate, malonate, methanesulfonate, 2-naphthalenesulfonate, nicotinate, nitrate, palmoate, phosphate, picrate, pivalate, propionate, salicylate, succinate, sulfate, tartrate, tosylate and undecanoate. Salts derived from appropriate bases include alkali metal (e.g., sodium), alkaline earth metal (e.g., magnesium), ammonium and N-(alkyl)₄⁺ salts. This invention also envisions the quaternization of any basic nitrogen-containing groups of the compounds disclosed herein. Water or oil-soluble or dispersible products may be obtained by such quaternization.

Evaluating Compounds

[0222] A variety of methods can be used to evaluate a compound for ability to modulate STEP activity. Evaluation methods include *in vitro* assays (e.g., enzyme-based assays), *in vitro* cell-based signaling assays, and *in vivo* methods (e.g., testing in animal models). The evaluation methods can evaluate binding activity, phosphatase activity, or an activity downstream of STEP, such as the activity of ERK.

[0223] For example, a compound described herein may be evaluated using a fluorescence-based phosphatase assay. A phosphate-containing reagent may be used in the assay which, upon dephosphorylation by a phosphatase, generates a fluorescent product that may be detected using a fluorometer or fluorescence plate reader. Data may be expressed as percentage (%) inhibition of enzyme activity. For compounds showing enzymatic activation, data may be represented as percentage of inhibition but with negative values.

Composition and routes of administration

[0224] The invention also provides a pharmaceutical composition, comprising an effective amount of a compound described herein (e.g., a compound capable of treating or preventing a condition as described herein, e.g., a compound of any formula herein or otherwise described herein) and a pharmaceutically acceptable carrier.

[0225] The compositions delineated herein include the compounds delineated herein (e.g., a compound described herein), as well as additional therapeutic agents if present, in amounts effective for achieving a modulation of disease or disease symptoms, including those described herein.

[0226] The term "pharmaceutically acceptable carrier or adjuvant" refers to a carrier or adjuvant that may be administered to a patient, together with a compound of this invention, and which does not destroy the pharmacological activity thereof and is nontoxic when administered in doses sufficient to deliver a therapeutic amount of the compound.

[0227] Pharmaceutically acceptable carriers, adjuvants and vehicles that may be used in the pharmaceutical compositions of this invention include, but are not limited to, ion exchangers, alumina, aluminum stearate, lecithin, self-emulsifying drug delivery systems (SEDDS) such as d- α -tocopherol polyethylene glycol 1000 succinate, surfactants used in pharmaceutical dosage forms such as Tweens or other similar polymeric delivery matrices, serum proteins, such as human serum albumin, buffer substances such as phosphates, glycine, sorbic acid, potassium sorbate, partial glyceride mixtures of saturated vegetable fatty acids, water, salts or electrolytes, such as protamine sulfate, disodium hydrogen phosphate, potassium hydrogen phosphate, sodium chloride, zinc salts, colloidal silica, magnesium trisilicate, polyvinyl pyrrolidone, cellulose-based substances, polyethylene glycol, sodium carboxymethylcellulose, polyacrylates, waxes, polyethylene-polyoxypropylene-block polymers, polyethylene glycol and wool fat. Cyclodextrins such as α -, β -, and γ -cyclodextrin, or chemically modified derivatives such as hydroxyalkylcyclodextrins, including 2- and 3-hydroxypropyl- β -cyclodextrins, or other solubilized derivatives may also be advantageously used to enhance delivery of compounds of the formulae described herein.

[0228] The pharmaceutical compositions of this invention may be administered orally, parenterally, by inhalation spray, topically, rectally, nasally, buccally, vaginally or via an implanted reservoir, preferably by oral administration or administration by injection. The pharmaceutical compositions of this invention may contain any conventional non-toxic pharmaceutically-acceptable carriers, adjuvants or vehicles. In some cases, the pH of the formulation may be adjusted with pharmaceutically acceptable acids, bases or buffers to enhance the stability of the formulated compound or its delivery form. The term parenteral as used herein includes subcutaneous, intracutaneous, intravenous, intramuscular, intra-articular, intraarterial, intrasynovial, intrasternal, intrathecal, intralesional and intracranial injection or infusion techniques.

[0229] The pharmaceutical compositions may be in the form of a sterile injectable preparation, for example, as a sterile injectable aqueous or oleaginous suspension. This suspension may be formulated according to techniques known in the art using suitable dispersing or wetting agents (such as, for example, Tween 80) and suspending agents. The sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally acceptable diluent or solvent, for example, as a solution in 1,3-butanediol. Among the acceptable vehicles and solvents that may be employed are mannitol, water, Ringer's solution and isotonic sodium chloride solution. In addition, sterile, fixed oils

are conventionally employed as a solvent or suspending medium. For this purpose, any bland fixed oil may be employed including synthetic mono- or diglycerides. Fatty acids, such as oleic acid and its glyceride derivatives are useful in the preparation of injectables, as are natural pharmaceutically-acceptable oils, such as olive oil or castor oil, especially in their polyoxyethylated versions. These oil solutions or suspensions may also contain a long-chain alcohol diluent or dispersant, or carboxymethyl cellulose or similar dispersing agents which are commonly used in the formulation of pharmaceutically acceptable dosage forms such as emulsions and or suspensions. Other commonly used surfactants such as Tweens or Spans and/or other similar emulsifying agents or bioavailability enhancers which are commonly used in the manufacture of pharmaceutically acceptable solid, liquid, or other dosage forms may also be used for the purposes of formulation.

[0230] The pharmaceutical compositions of this invention may be orally administered in any orally acceptable dosage form including, but not limited to, capsules, tablets, emulsions and aqueous suspensions, dispersions and solutions. In the case of tablets for oral use, carriers which are commonly used include lactose and corn starch. Lubricating agents, such as magnesium stearate, are also typically added. For oral administration in a capsule form, useful diluents include lactose and dried corn starch. When aqueous suspensions and/or emulsions are administered orally, the active ingredient may be suspended or dissolved in an oily phase is combined with emulsifying and/or suspending agents. If desired, certain sweetening and/or flavoring and/or coloring agents may be added.

[0231] The pharmaceutical compositions of this invention may also be administered in the form of suppositories for rectal administration. These compositions can be prepared by mixing a compound of this invention with a suitable non-irritating excipient which is solid at room temperature but liquid at the rectal temperature and therefore will melt in the rectum to release the active components. Such materials include, but are not limited to, cocoa butter, beeswax and polyethylene glycols.

[0232] Topical administration of the pharmaceutical compositions of this invention is useful when the desired treatment involves areas or organs readily accessible by topical application. For application topically to the skin, the pharmaceutical composition should be formulated with a suitable ointment containing the active components suspended or dissolved in a carrier. Carriers for topical administration of the compounds of this invention include, but are not limited to, mineral oil, liquid petroleum, white petroleum, propylene glycol, polyoxyethylene polyoxypropylene compound, emulsifying wax and water. Alternatively, the pharmaceutical composition can be formulated with a suitable lotion or cream containing the active compound suspended or dissolved in a carrier with suitable emulsifying agents. Suitable carriers include, but are not limited to, mineral oil, sorbitan monostearate, polysorbate 60, cetyl esters wax, cetearyl alcohol, 2-octyldodecanol, benzyl alcohol and water. The pharmaceutical compositions of this invention may also be topically applied to the lower intestinal tract by rectal suppository formulation or in a suitable enema formulation. Topically-transdermal patches are also included in this invention.

[0233] The pharmaceutical compositions of this invention may be administered by nasal aerosol or inhalation. Such compositions are prepared according to techniques well-known in the art of pharmaceutical formulation and may be prepared as solutions in saline, employing benzyl alcohol or other suitable preservatives, absorption promoters to enhance bioavailability, fluorocarbons, and/or other solubilizing or dispersing agents known in the art.

[0234] When the compositions of this invention comprise a combination of a compound of the formulae described herein and one or more additional therapeutic agents, both the compound and the additional agent should be present at dosage levels of between about 1 to 100%, and more preferably between about 5 to 95% of the dosage normally administered in a monotherapy regimen. The additional agents may be administered separately, as part of a multiple dose regimen, from the compounds of this invention. Alternatively, those agents may be part of a single dosage form, mixed together with the compounds of this invention in a single composition.

[0235] The compounds described herein can, for example, be administered by injection, intravenously, intraarterially, subdermally, intraperitoneally, intramuscularly, or subcutaneously; or orally, buccally, nasally, transmucosally, topically, in an ophthalmic preparation, or by inhalation, with a dosage ranging from about 0.5 to about 100 mg/kg of body weight, alternatively dosages between 1 mg and 1000 mg/dose, every 4 to 120 hours, or according to the requirements of the particular drug. The methods herein contemplate administration of an effective amount of compound or compound composition to achieve the desired or stated effect. Typically, the pharmaceutical compositions of this invention will be administered from about 1 to about 6 times per day or alternatively, as a continuous infusion. Such administration can be used as a chronic or acute therapy. The amount of active ingredient that may be combined with the carrier materials to produce a single dosage form will vary depending upon the host treated and the particular mode of administration. A typical preparation will contain from about 5% to about 95% active compound (w/w). Alternatively, such preparations contain from about 20% to about 80% active compound.

[0236] Lower or higher doses than those recited above may be required. Specific dosage and treatment regimens for any particular patient will depend upon a variety of factors, including the activity of the specific compound employed, the age, body weight, general health status, sex, diet, time of administration, rate of excretion, drug combination, the severity and course of the disease, condition or symptoms, the patient's disposition to the disease, condition or symptoms, and the judgment of the treating physician.

[0237] Upon improvement of a patient's condition, a maintenance dose of a compound, composition or combination of this invention may be administered, if necessary. Subsequently, the dosage or frequency of administration, or both, may be reduced, as a function of the symptoms, to a level at which the improved condition is retained when the symptoms have been alleviated to the desired level. Patients may, however, require intermittent treatment on a long-term basis upon any recurrence of disease symptoms.

Methods of treatment

[0238] The compounds and compositions described herein can be administered to cells in culture, e.g. *in vitro* or *ex vivo*, or to a subject, e.g., *in vivo*, to treat, prevent, and/or diagnose a variety of disorders, including those described herein below.

[0239] The compounds and compositions described herein can be administered to a subject, for example using a method described herein, who is suffering from a disorder described herein, e.g., a disorder that would benefit from the modulation of STEP (e.g., activating or inhibiting STEP). The compounds and compositions described herein can be administered to a subject, for example using a method described herein, who is at risk for a disorder described herein, e.g., a disorder that would benefit from the modulation of STEP (e.g., activating or inhibiting STEP).

[0240] Inhibitors of STEP may increase phosphorylation of an NMDA-R. Thus, in some embodiments, a compound described herein, e.g., a compound that inhibits STEP, may be useful for treating a disorder in which increasing phosphorylation of an NMDA-R would be beneficial.

[0241] Inhibitors of STEP may activate an ERK1 or ERK2 kinase, for example, in the CNS. Thus, in some embodiments, a compound described herein, e.g., a compound that inhibits STEP, may be useful for treating a disorder in which activate an ERK1 or ERK2 kinase would be beneficial.

[0242] Compounds described herein may be useful in treating a variety of disorders, including disorders of the CNS. Exemplary disorders include schizophrenia, schizoaffective disorders, major depression, bipolar disorder, cognitive deficit, mild cognitive impairment (MCI), Alzheimer's disease (AD), attention-deficit/hyperactivity disorder (ADHD), dementia, generalized anxiety disorders, panic disorders, obsessive-compulsive disorders, phobias, post-traumatic stress syndrome, anorexia nervosa, drug addiction, ischemic stroke, head trauma or brain injury, Huntington's disease, Parkinson's disease, spinocerebellar degeneration, motor neuron diseases, epilepsy, neuropathic pain, chronic pain, neuropathies, autism and autistic disorders.

[0243] Compounds described herein may be useful for treating or preventing central nervous system disorders selected from the group consisting of schizophrenia; refractory, intractable or chronic schizophrenia; emotional disturbance; psychotic disorder; mood disorder; bipolar I type disorder; bipolar II type disorder; depression; endogenous depression; major depression; melancholy and refractory depression; dysthymic disorder; cyclothymic disorder; panic attack; panic disorder; agoraphobia; social phobia; obsessive-compulsive disorder; post-traumatic stress disorder; generalized anxiety disorder; acute stress disorder; hysteria; somatization disorder; conversion disorder; pain disorder; hypochondriasis; factitious disorder; dissociative disorder; sexual dysfunction; sexual desire disorder; sexual arousal disorder; erectile dysfunction; anorexia nervosa; bulimia nervosa; sleep disorder; adjustment disorder; alcohol abuse; alcohol intoxication; drug addiction; stimulant intoxication; narcotism; anhedonia; iatrogenic anhedonia; anhedonia of a psychic or mental cause; anhedonia associated with depression; anhedonia associated with schizophrenia; delirium; cognitive impairment; cognitive impairment associated with Alzheimer's disease, Parkinson's disease and other neurodegenerative diseases; cognitive impairment caused by Alzheimer's disease; Parkinson's disease and associated neurodegenerative diseases; cognitive impairment of schizophrenia; cognitive impairment caused by refractory, intractable or chronic schizophrenia; vomiting; motion sickness; obesity; migraine; pain (ache) ; mental retardation; autism disorder (autism); Tourette's disorder; tic disorder; attention-deficit/hyperactivity disorder; conduct disorder; and Down's syndrome.

[0244] Compounds described herein may be useful for treating or preventing disorders selected from schizophrenia, schizoaffective disorder, bipolar disorder, manic-depressive disorder, psychosis, mood and anxiety disorders, mania, drug or substance addiction, cognition disorders, learning disabilities, learning and memory disorders, aging and neurologic disorders associated with or linked with cognitive impairments; mild cognitive impairments (MCI), Alzheimer's disease, Alzheimer-related cognition disorders, Huntington's disease, Parkinson's disease, CADASIL syndrome (cerebral autosomal dominant arteriopathy with subcortical infarcts and leukoencephalopathy), amnesia, Wernicke-Korsakoff syndrome, Korsakoff syndrome, mild traumatic head injury (MBTI), traumatic head injury (TBI), fragile X syndrome, stroke, attention-deficit and hyperactivity disorder (ADHD), obsessive compulsive disorder (OCD), post-traumatic stress disorder (PTSD), loss of concentration, autism, cerebral palsy, encephalopathy, and narcolepsy. The disorder may affect learning and memory, neurogenesis, neuronal plasticity, pain perception, mood and anxiety, or neuroendocrine regulation. The disorder may be a cognitive deficit disorder. The disorder may involve pain perception or neuroendocrine regulation.

Schizophrenia

[0245] In some embodiments, a compound or composition described herein can be used in the treatment of schizophrenia. Schizophrenia is a psychiatric diagnosis that describes a mental disorder characterized by abnormalities in the perception or expression of reality. Distortions in perception may affect all five senses, including sight, hearing, taste, smell and touch, but most commonly manifests as auditory hallucinations, paranoid or bizarre delusions, or disorganized speech and thinking with significant social or occupational dysfunction. Onset of symptoms typically occurs in young adulthood, with approximately 0.4-0.6% of the population affected. Diagnosis is based on the patient's self-reported experiences and observed behavior.

[0246] The disorder is thought to mainly affect cognition, but it also usually contributes to chronic problems with behavior and emotion. People with schizophrenia are likely to have additional (comorbid) conditions, including major depression and anxiety disorders. Social problems, such as long-term unemployment, poverty and homelessness, are common. Furthermore, the average life expectancy of people with the disorder is 10 to 12 years less than those without, due to increased physical health problems and a higher suicide rate.

[0247] The Diagnostic and Statistical Manual of Mental Disorders (DSM) contains five sub-classifications of schizophrenia. These include Paranoid type (where delusions and hallucinations are present but thought disorder, disorganized behavior, and affective flattening are absent); Disorganized type (also known as hebephrenic schizophrenia, where thought disorder and flat affect are present together); Catatonic type (the subject may be almost immobile or exhibit agitated, purposeless movement; symptoms can include catatonic stupor and waxy flexibility); Undifferentiated type (psychotic symptoms are present but the criteria for paranoid, disorganized, or catatonic types have not been met); and Residual type (where positive symptoms are present at a low intensity only).

[0248] The International Statistical Classification of Diseases and Related Health Problems (10th Revision) defines two additional subtypes. These include Post-schizophrenic depression (a depressive episode arising in the aftermath of a schizophrenic illness where some low-level schizophrenic symptoms may still be present); and Simple schizophrenia (insidious and progressive development of prominent negative symptoms with no history of psychotic episodes.)

[0249] An agent for the treatment of schizophrenia may improve so-called positive symptoms in the acute period of schizophrenia such as hallucinations, delusions, excitations and the like. An agent for treating schizophrenia may also improve so-called negative symptoms that are observed in the chronic period of schizophrenia such as apathy, emotional depression, hyposychosis and the like.

Schizoaffective Disorder

[0250] Schizoaffective disorder is a psychiatric diagnosis that describes a mental disorder characterized by recurring episodes of elevated or depressed mood, or simultaneously elevated and depressed mood that alternate or occur together with distortions in perception. The perceptual distortion component of the disorder, called psychosis, may affect all five senses, including sight, hearing, taste, smell and touch, but most commonly manifest as auditory hallucinations, paranoid or bizarre delusions, or disorganized speech and thinking with significant social and occupational dysfunction. The elevated, depressed or simultaneously elevated and depressed mood episode components of the disorder, called mood disorder, are broadly recognized as depressive and bipolar types of the illness; the division is based on whether the individual has ever had a manic, hypomanic or mixed episode. Onset of symptoms usually begins in early adulthood and is rarely diagnosed in childhood (prior to age 13). The lifetime prevalence of the disorder is uncertain (due to studies using varying diagnostic criteria), although it is generally agreed to be less than 1 percent, and possibly in the range of 0.5 to 0.8 percent. Diagnosis is based on the patient's self-reported experiences and observed behavior. No laboratory test for schizoaffective disorder currently exists. As a group, people with schizoaffective disorder have a more favorable prognosis than people with schizophrenia, but a worse prognosis than those with mood disorders.

[0251] The disorder is thought to mainly affect cognition and emotion, but it also usually contributes to ongoing problems with behavior and motivation. People with schizoaffective disorder are likely to have additional (comorbid) conditions, including anxiety disorders and substance abuse. Social problems, such as long-term unemployment, poverty and homelessness, are common. Furthermore, the average life expectancy of people with the disorder is shorter than those without the disorder, due to increased physical health problems and a higher suicide rate.

Cognitive Deficit

[0252] Treatment using a compound or composition described herein may improve a cognitive deficit associated with a cognition-related disorder. Cognitive deficit is an inclusive term to describe any characteristic that acts as a barrier to cognitive performance. The term may describe deficits in global intellectual performance, such as mental retardation, it may describe specific deficits in cognitive abilities (learning disorders, dyslexia), or it may describe drug-induced cognitive/memory impairment, such as that seen with alcohol and the benzodiazepines. Cognitive deficits may be congenital

or caused by environmental factors such as brain injuries, neurological disorders, or mental illness.

[0253] Exemplary cognition-related disorders (e.g., cognitive dysfunction) include, without limitation, mild cognitive impairment (MCI), dementia, delirium, amnesic disorder, Alzheimer's disease, Parkinson's disease and Huntington's disease; memory disorders including memory deficits associated with depression, senile dementia, dementia of Alzheimer's disease; cognitive deficits or cognitive dysfunction associated with neurological conditions including, for example, Parkinson's disease (PD), Huntington's disease (HD), Alzheimer's disease, depression, schizophrenia and other psychotic disorders such as paranoia and manic-depressive illness; cognitive dysfunction in schizophrenia; disorders of attention and learning such as attention deficit disorders (e.g., attention deficit hyperactivity disorder (ADHD)) and dyslexia; cognitive dysfunction associated with developmental disorders such as Down's syndrome and Fragile X syndrome; loss of executive function; loss of learned information; vascular dementia; schizophrenia; cognitive decline; a neurodegenerative disorder; and other dementias, for example, dementia due to HIV disease, head trauma, Parkinson's disease, Huntington's disease, Pick's disease, Creutzfeldt-Jakob disease, or due to multiple etiologies. Cognition-related disorders also include, without limitation, cognitive dysfunction associated with MCI and dementias such as Lewy Body, vascular, and post stroke dementias. Cognitive dysfunction associated with surgical procedures, traumatic brain injury or stroke may also be treated in accordance with the embodiments described herein.

Major Depression

[0254] Major depression (also known as clinical depression, major depressive disorder, unipolar depression, or unipolar disorder) is a mental disorder characterized by a pervasive low mood, low self-esteem, and loss of interest or pleasure in normally enjoyable activities. Types of Major depressive disorder include, e.g., Atypical depression, Melancholic depression, Psychotic depression, Catatonic depression, Postpartum depression, and Seasonal affective disorder.

Bipolar Disorder

[0255] Bipolar disorder, also known as manic depressive disorder, manic depressive psychosis, manic depression or bipolar affective disorder, is a psychiatric diagnosis that describes a category of mood disorders defined by the presence of one or more episodes of abnormally elevated mood clinically referred to as mania or, if milder, hypomania. Individuals who experience manic episodes also commonly experience depressive episodes or symptoms, or mixed episodes in which features of both mania and depression are present at the same time. These episodes are usually separated by periods of "normal" mood, but in some individuals, depression and mania may rapidly alternate, known as rapid cycling. Extreme manic episodes can sometimes lead to psychotic symptoms such as delusions and hallucinations. The disorder has been subdivided into bipolar I, bipolar II, cyclothymia, and other types, based on the nature and severity of mood episodes experienced; the range is often described as the bipolar spectrum.

Anxiety Disorders

[0256] Anxiety disorder is a blanket term covering several different forms of abnormal and pathological fear and anxiety. Current psychiatric diagnostic criteria recognize a wide variety of anxiety disorders. Recent surveys have found that as many as 18% of Americans may be affected by one or more of them.

[0257] Generalized anxiety disorder is a common chronic disorder characterized by long-lasting anxiety that is not focused on any one object or situation. Those suffering from generalized anxiety experience non-specific persistent fear and worry and become overly concerned with everyday matters. Generalized anxiety disorder is the most common anxiety disorder to affect older adults.

[0258] In panic disorder, a person suffers from brief attacks of intense terror and apprehension, often marked by trembling, shaking, confusion, dizziness, nausea, difficulty breathing. These panic attacks, defined by the APA as fear or discomfort that abruptly arises and peaks in less than ten minutes, can last for several hours and can be triggered by stress, fear, or even exercise; although the specific cause is not always apparent. In addition to recurrent unexpected panic attacks, a diagnosis of panic disorder also requires that said attacks have chronic consequences: either worry over the attacks' potential implications, persistent fear of future attacks, or significant changes in behavior related to the attacks. Accordingly, those suffering from panic disorder experience symptoms even outside of specific panic episodes. Often, normal changes in heartbeat are noticed by a panic sufferer, leading them to think something is wrong with their heart or they are about to have another panic attack. In some cases, a heightened awareness (hypervigilance) of body functioning occurs during panic attacks, wherein any perceived physiological change is interpreted as a possible life threatening illness (i.e. extreme hypochondriasis).

[0259] Obsessive compulsive disorder is a type of anxiety disorder primarily characterized by repetitive obsessions (distressing, persistent, and intrusive thoughts or images) and compulsions (urges to perform specific acts or rituals). The OCD thought pattern may be likened to superstitions insofar as it involves a belief in a causative relationship where,

in reality, one does not exist. Often the process is entirely illogical; for example, the compulsion of walking in a certain pattern may be employed to alleviate the obsession of impending harm. And in many cases, the compulsion is entirely inexplicable, simply an urge to complete a ritual triggered by nervousness. In a minority of cases, sufferers of OCD may only experience obsessions, with no overt compulsions; a much smaller number of sufferers experience only compulsions.

[0260] The single largest category of anxiety disorders is that of Phobia, which includes all cases in which fear and anxiety is triggered by a specific stimulus or situation. Sufferers typically anticipate terrifying consequences from encountering the object of their fear, which can be anything from an animal to a location to a bodily fluid.

[0261] Post-traumatic stress disorder or PTSD is an anxiety disorder which results from a traumatic experience. Post-traumatic stress can result from an extreme situation, such as combat, rape, hostage situations, or even serious accident. It can also result from long term (chronic) exposure to a severe stressor, for example soldiers who endure individual battles but cannot cope with continuous combat. Common symptoms include flashbacks, avoidant behaviors, and depression.

Combination therapies

[0262] In some embodiments, the subject is being treated with an additional therapeutic agent. Such additional agents include atypical antipsychotics such as aripiprazole, clozapine, ziprasidone, risperidone, quetiapine, olanzapine, amisulpride, asenapine, iloperidone, melperone, paliperidone, perospirone, sertindole and sulpiride; and typical antipsychotics such as haloperidol, molindone, loxapine, thioridazine, molindone, thiothixene, pimozide, fluphenazine, trifluoperazine, mesoridazine, chlorprothixene, chlorpromazine, perphenazine, triflupromazine and zuclopenthixol.

Clinical outcomes

[0263] In some embodiments, treatment with a compound or composition described herein, for example, using a method described herein, improves one or more clinical outcomes. For example, in some embodiments, treatment with a compound or composition described herein may improve cognitive function. Elements of cognitive function include memory, orientation, attention, reasoning, language and praxis.

[0264] In some embodiments, clinical outcomes may be assessed using known methods. One such method is the Brief Psychiatric Rating Scale (BPRS), a multi-item inventory of general psychopathology traditionally used to evaluate the effects of drug treatment in schizophrenia. The BPRS psychosis cluster (conceptual disorganization, hallucinatory behavior, suspiciousness, and unusual thought content) is considered a particularly useful subset for assessing actively psychotic schizophrenic patients.

[0265] In some embodiments, clinical outcomes may be assessed using the 7-point Clinical Global Impression (CGI) rating scale, a commonly used measure of symptom severity, treatment response and the efficacy of treatments. The CGI reflects the impression of a skilled observer, fully familiar with the manifestations of schizophrenia, about the overall clinical state of the patient.

[0266] In some embodiments, clinical outcomes may be assessed using the 30-item Positive and Negative Symptoms Scale (PANSS). The name refers to the two types of symptoms in schizophrenia, as defined by the American Psychiatric Association: positive symptoms, which refer to an excess or distortion of normal functions (e.g. hallucinations and delusions), and negative symptoms, which represent a diminution or loss of normal functions.

[0267] In some embodiments, clinical outcomes may be assessed using the Scale for Assessing Negative Symptoms (SANS). SANS assesses five symptom complexes to obtain clinical ratings of negative symptoms in patients with schizophrenia. They are: affective blunting; alogia (impoverished thinking); avolition/apathy; anhedonia/asociality; and disturbance of attention. Assessments are conducted on a six-point scale.

[0268] The invention is further illustrated by the following examples which are intended to illustrate but not limit the scope of the invention.

EXAMPLES

Abbreviations:

[0269]

DCM: Dichloromethane
EA, EtOAc or AcOEt: Ethyl acetate
PE: Petroleum ether
DIPEA: Diisopropylethylamine
TEA: Triethylamine

	rt: Room temperature
	SOCl ₂ : Thionyl chloride
	POCl ₃ : Phosphorous oxychloride
	THF: Tetrahydrofuran
5	NaOAc: Sodium acetate
	MeOH: Methanol
	i-AmOH: Isoamyl alcohol
	NaH: Sodium hydride
	NaBH ₃ CN: Sodium cyanoborohydride
10	n-BuLi: <i>n</i> -Butyl lithium
	LHMDS: Lithium bis(trimethylsilyl)amide
	LDA: Lithium diisopropylamide
	i-PrOH: Isopropyl alcohol
	Na ₂ SO ₄ : Sodium sulfate
15	Mg ₂ SO ₄ : Magnesium sulfate
	MeCN: Acetonitrile
	NaOH: Sodium hydroxide
	EtOH: Ethanol
	CuI: Copper(I) iodide
20	Pd(PPh ₃) ₂ Cl ₂ : trans-Dichlorobis(triphenylphosphine)palladium(II)
	MsCl: Methanesulfonyl chloride
	BINAM: [1,1'-Binaphthalene]-2,2'-diamine
	Xphos: 2-Dicyclohexylphosphino-2',4',6'-triisopropylbiphenyl
	Sphos: 2-Dicyclohexylphosphino-2',6'-dimethoxybiphenyl
25	DavePhos: 2-(Dicyclohexylphosphino)-2'-(N,N-dimethylamino)biphenyl
	Cs ₂ CO ₃ : Cesium carbonate
	K ₂ CO ₃ : Potassium carbonate
	Mwave or μ W or mW: Microwave
	<i>t</i> -BuOH: <i>tert</i> -Butanol
30	K ₃ PO ₄ : Potassium phosphate
	Pd(APhos) ₂ Cl ₂ : Bis(di- <i>tert</i> -butyl(4-dimethylaminophenyl)phosphine)dichloro palladium(II)
	Pd(PPh ₃) ₄ : Tetrakis(triphenylphosphine)palladium (0)
	Pd(dppf) ₂ Cl ₂ : Dichloro[1,1'-bis(diphenylphosphino)ferrocene]palladium(II)
	PdOAc: Palladium(II) acetate
35	Pd ₂ dba ₃ : Tris(dibenzylideneacetone)dipalladium (0)
	Pd-118: Dichloro[1,1'-bis(di- <i>t</i> -butylphosphino)ferrocene]palladium(II)
	Xantphos: 9,9-Dimethyl-4,5-bis(diphenylphosphino)xanthene
	BINAP: ($\pm$)-2,2'-Bis(diphenylphosphino)-1,1'-binaphthalene
	EDCI: 1-Ethyl-3-(3-dimethylaminopropyl) carbodiimide
40	HOBt: Hydroxybenzotriazole
	NH ₄ OH: Ammonium hydroxide
	H ₂ O: Water
	Pd/C: Palladium on carbon
	DMF: <i>N,N</i> Dimethylformamide
45	KOCN: Potassium cyanate
	WSC-HCl or WSCDI: Water Soluble Carbodiimide hydrochloride
	HATU: O-(7-Azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate
	HBTU: O-(Benzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate
50	Py-Brop: Bromotripyrrolidino-phosphonium hexafluorophosphate
	BOP: Benzotriazole-1-yl-oxy-tris-(dimethylamino)-phosphonium hexafluoro phosphate
	DBU: diaza(1,3)bicyclo[5.4.0]undecene
	DMSO: Dimethyl sulfoxide
	LCMS: Liquid chromatography mass spectrometry
	HPLC: High performance liquid chromatography
55	DMA: N,N-dimethylacetamide
	h: hour
	TLC: Thin layer chromatography
	TFA: Trifluoroacetic acid

Et₃N: Triethylamine
 DIPEA: N,N-Diisopropylethylamine
 O.N: Overnight
 TBSO: tert-Butyldimethylsilyloxy
 5 DME: Dimethyl ether
 NMP: 1-methyl-2-pyrrolidinone
 PS-BEMP: 2-*tert*-Butylimino-2-diethylamino-1,3-dimethylperhydro-1,3,2-diazaphosphorine supported on Polystyrene
 PBr₃ : Phosphorus tribromide
 10 NaOtBu: Sodium tert-butoxide
 KI: Potassium iodide
 PPh₃ : Triphenylphosphine
 NMM: *N*-Methylmorpholine
 HCHO: Formaldehyde
 15 PG: Protecting group
 ISCO: Teledyne ISCO purification systems
 BINAM: 1,1'-binaphthyl-2,2'-diamine.

General Experimental:

20 **[0270]** All exemplified target compounds are fully analyzed and characterized (TLC, LCMS, ¹H-NMR) prior to submission for biological evaluation. Thin-layer chromatography was carried out on native silica 254F plates. Visualization was accomplished with ultraviolet or phosphomolybdic acid. ¹H-NMR spectra were recorded on multiple NMR spectrometers, either on 400 MHz on a Avance III 400 Ultra shield-plus TM digital Spectrometer or on 300 MHz using a Varian Mercury
 25 300Plus Spectrometer, designated by 400 MHz or 300 MHz, respectively. ¹H-NMR spectra were also recorded on a Bruker Spectrospin 300 MHz Spectrometer at 300.13 MHz in DMSO-d₆ with TMS as an internal standard and will be designated as Bruker 300 Hz. NMR assignments are based on a combination of the ¹H, ¹³C, ¹HCOSY, HMBC and HMQC spectra. Coupling constants are given in hertz (Hz). Anhydrous methylene chloride, tetrahydrofuran, and dimethylformamide were obtained by distillation, and other materials are reagent grade.

30 **[0271] LC-MS Methods** are listed here:

Method A: Mobile phase: A = 0.1% TFA/H₂O, B = 0.01% TFA/MeCN; Gradient: B = 5%-95% in 1.5 min; Flow rate: 2.0 mL/min; Column: sunfire-C₁₈, 50 x 4.6 mm, 3.5 μm;

35 **Method B:** Mobile phase: A = 10mM NH₄HCO₃/H₂O, B = MeCN; Gradient: B = 5%-95% in 1.5 min; Flow rate: 2.0 mL/min; Column: Xbridge-C₁₈, 50 x 4.6mm, 3.5 μm;

Method C: Mobile phase: A = 10 mM ammonium formate/H₂O/4.9% MeCN, B = MeCN; Gradient: B = 5%-100% in 2.0 min; Flow rate: 2.5 mL/min; Column: Atlantis T3 3μM 4.6x30mm

40 **Method D:** Mobile phase: A = 0.1% formic acid/H₂O/4.9% MeCN, B = MeCN; Gradient: B = 5%-100% in 2.0 min; Flow rate: 2.5 mL/min; Column: Atlantis T3 3μM 4.6x30mm

45 **Method E:** Mobile phase: A = 0.05% TFA/H₂O, B = 0.05% TFA/MeCN; Gradient: B = 5%-100% in 3.0 min; Flow rate: 0.8 mL/min; Column: CAPCELL PAK C18 (Shiseido, UG120, 3 mM, 2.0 mm I.D. x 50 mm).

[0272] Representative Conditions of PREP-HPLC are listed here:

PREP-HPLC Condition A (Basic Mobile Phase):

50 Instrument: Gilson 281
 Mobile Phase: A = 0.01% NH₄HCO₃/H₂O, B = MeCN
 Flow Rate: 40.0 mL/min
 Column: AGT Venusil XBP C₁₈, 10.0 μm, 30 mmx 100 mm

PREP-HPLC Condition B (Basic Mobile Phase):

55 Instrument: Gilson 281

Mobile Phase: A = $\text{NH}_3\text{-H}_2\text{O}$, 10 mmol/L, B = MeCN

Flow Rate: 40.0 mL/min

Column: Waters X-Bridge, 5.0 μm , 30 mm x 150 mm

PREP-HPLC Condition C (Basic Mobile Phase):

Instrument: Gilson 281

Mobile Phase: A = 0.01% $\text{NH}_4\text{HCO}_3/\text{H}_2\text{O}$, B = MeCN

Flow Rate: 30.0 mL/min

Column: Shimadzu PRC-ODS, 10.0 μm , 20 mm x 250 mm

Gradient: B = xx%-yy% 0.0 to 8.0 min

yy%-95% 8.0 to 8.2 min

95%-95% 8.2 to 11.0 min

[0273] The following table shows the relationship of representative value (xx%-yy%) of gradient and retention time on LC-MS of corresponding compound.

25%-30% 0.5-1.0 min

30%-50% 1.0-1.5 min

50%-70% 1.5-1.75 min

70%-90% 1.7-2.0 min

PREP-HPLC Condition D:

[0274] Instrument: Waters 600 pump, Waters 2996, Photodiode Array Detector, Waters Micromass ZQ, Gilson 215 Liquid Handler.

Mobile Phase: A = 0.05% TFA/ H_2O , B = MeCN

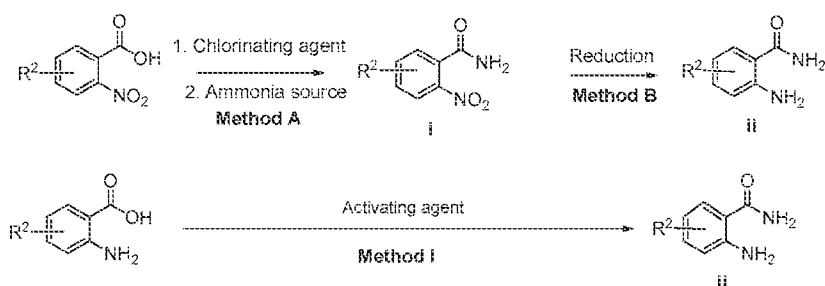
Flow Rate: 36.0 mL/min

Column: Shiseido CAPCELL PAK C18, UG120, 5 μm , 20 mm I.D. x 50 mm

Gradient: B = 5%-100% 0.0 to 4.0 min

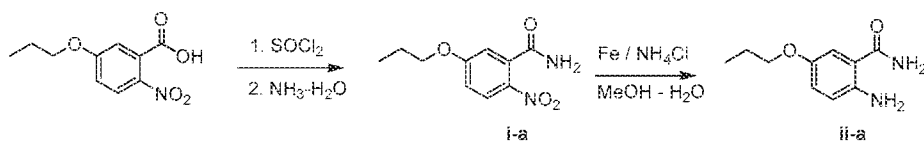
Scheme 1: General route for the synthesis of compounds with general formula ii

[0275]



Scheme 2: Representative synthesis of compounds of formula ii (see Scheme 1)

[0276]



[0277] Method A: 2-Nitro-5-propoxy-benzamide (i-a) A mixture of 2-nitro-5-propoxybenzoic acid (1.97 g, 8.75 mmol) and DMF (0.1 mL) in SOCl_2 (20 mL) was stirred at 65 °C for 2 h. After the reaction was completed, the mixture was

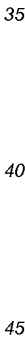
5 **[0278] Method B: 2-Amino-5-propoxy-benzamide (ii-a)** To a mixture of 2-nitro-5-propoxy-benzamide (1.20 g, 5.36 mmol) in MeOH-H₂O (v/v, 3:1, 60 mL) was added NH₄Cl (2.84 g, 53.6 mmol) and Fe (2.99 g, 53.6 mmol). The resulting mixture was stirred at 60 °C for 3 h. After the reaction was completed, the mixture was cooled to room temperature and the iron was filtered off. The filtrate was concentrated to 15 mL and the formed precipitate was collected and dried *in vacuo* to give 1.02 g of ii-a as a pale yellow solid (98%). LCMS m/z = 178.1 (M-16), 195.1 (M+1) (Method B) (retention time = 1.46 min).

[0279]



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[0281]



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- 35

F7: $\text{PBr}_3/\text{CH}_2\text{Cl}_2/\text{DMF}/60^\circ\text{C}$ **Method G for Coupling Conditions**

[0283]

G1: *i*-PrOH/85-100 °C

G2: THF/reflux

G3: *i*-AmOH/100-130°C

G4: MeOH/microwave/150°C

G5: *i*-AmOH/ microwave/150°CG6: THF/ Et_3N /refluxG7: THF- H_2O / NaOAc /rt-60°C

G8: NaH/THF

G9: *n*-BuLi/THF

G10: LHMDs/THF

G11: LDA/THF

G12: $\text{K}_2\text{CO}_3/\text{DMF}/60^\circ\text{C}$ G13: $\text{Cs}_2\text{CO}_3/\text{DMA}/80^\circ\text{C}$

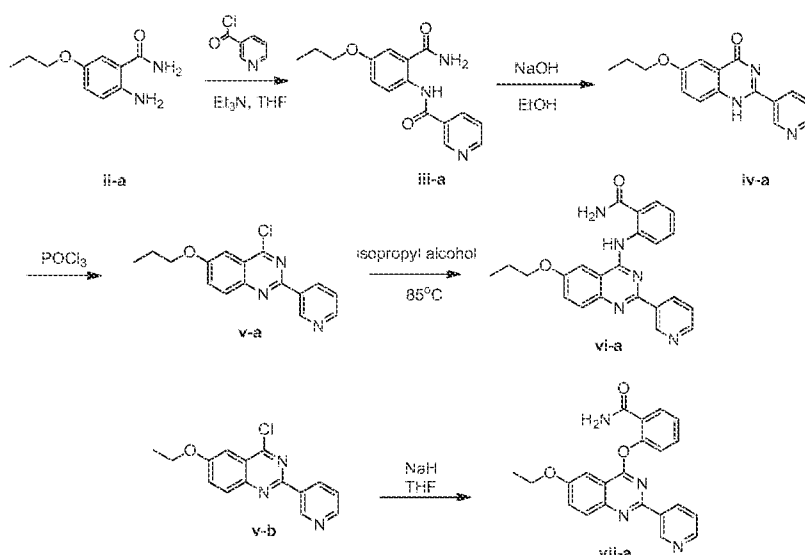
G14: NaOtBu/DMF/Microwave/100 °C

Method J for Coupling Conditions

[0284]

J1: $\text{Pd}(\text{PPh}_3)_4/ t\text{-BuOK/ Dioxane}$ J2: $\text{Pd}_2(\text{dba})_3/ \text{Xantphos/ Cs}_2\text{CO}_3/ \text{Dioxane}$ **Scheme 5: Representative synthesis of compounds of formula vi and vii (see Scheme 4)**

[0285]



[0286] Method C: N-(2-Carbamoyl-4-propoxy-phenyl)-nicotinamide (iii-a) To a solution of 2-amino-5-propoxybenzamide (760 mg, 3.91 mmol) in THF (15 mL) and Et_3N (1 mL) was added nicotinoyl chloride (607 mg, 4.30 mmol) in anhydrous THF (15 mL) dropwise. The resulting mixture was stirred at room temperature for 3 h. After the reaction was completed, the volatiles were removed. The residue was washed with H_2O (10 mL). The pH was adjusted to approximately 5 by adding dilute HCl (2N in water). The resulting solid was collected and dried *in vacuo* to give 1.00 g of iii-a as a pale yellow solid (89.0%). LCMS m/z = 300.1 (M+1) (Method B) (retention time = 1.60 min).

[0287] Method D: 2-benzamido-5-methoxy-3-methylbenzamide (iii-b) A 50 mL roundbottom flask was charged

with nicotinic acid (41 mg, 0.33 mmol, 1.0 eq.), 2-amino-5-methoxy-3-methylbenzamide (60 mg, 0.33 mmol, 1.0 eq.) and HBTU (190 mg, 0.50 mmol, 1.5 eq), which were suspended in 4 mL of DMF. DIPEA (86 mg, 0.66 mmol, 2.0 eq.) was added dropwise at room temperature and the reaction mixture was stirred overnight. The reaction mixture was added to water (10 mL) dropwise with stirring. The mixture was extracted with ethyl acetate. The ethyl acetate was evaporated and 55 mg of the orange solid (58.5% yield) was obtained. LCMS m/z = 286.1 ($M+1$) (Method B) (retention time = 1.24 min).

[0288] Method E: 6-Propoxy-2-pyridin-3-yl-1H-quinazolin-4-one (iv-a) A mixture of N-(2-carbamoyl-4-propoxy-phenyl)-nicotinamide (980 mg, 3.27 mmol) in EtOH (20 mL) was treated with NaOH (654 mg, 16.37 mmol). The resulting mixture was stirred at room temperature for 18 h. After the reaction was completed, the volatiles were removed *in vacuo*. The residue was partitioned between H₂O (50 mL) and ethyl acetate (50 mL). The aqueous layer was neutralized to pH 7 by slowly adding aq. citric acid and then a precipitate formed. The precipitate was collected and dried to give 1.00 g of iv-a as a grey solid (quantitative yield). LCMS m/z = 282.1 ($M+1$) (Method B) (retention time = 1.60 min).

Method F1: 4-Chloro-6-propoxy-2-pyridin-3-yl-quinazoline (v-a)

[0289] (This method is representative of method F1, F2, F3 and F4. These three methods can be implemented in a similar way except for substitution of the appropriate chlorinating reagent, solvent and temperature) To a mixture of 6-propoxy-2-pyridin-3-yl-1H-quinazolin-4-one (1.00 g, 3.56 mmol) in POCl₃ (10 mL) was added *N,N*-dimethylaniline (0.1 mL). The resulting mixture was stirred at 120 °C for 2 h. After the reaction was completed, POCl₃ was removed *in vacuo*, and the residue was added to ice-water slowly. The pH was adjusted to around 7 by slowly adding NaHCO₃ (sat.) at 0 °C. The resultant solid was collected and purified by chromatography on silica gel eluted with petroleum ether / ethyl acetate (v/v = 4: 1 to 1: 1) to give 580 mg of v-a as a pale yellow solid (54.7%).

[0290] Method F5: 4-chloro-6-methoxy-2-(pyridin-3-yl)quinazoline (v-c) In a sealed tube, phosphorus oxychloride (11 mL, 120 mmol) was added to 6-methoxy-2-(pyridin-3-yl)quinazolin-4(3H)-one (2.70 g, 10.66 mmol). The mixture was refluxed at 120°C for 12 h. After cooling, the remaining phosphorus oxychloride was removed *in vacuo* to leave a tan solid. This residue was added to an ice-water mixture (100 mL) with cooling and allowed to stir. The pH of the suspension was adjusted to about pH 9 via dropwise addition of 28% ammonium hydroxide, and stirring was continued for 30 mins. The resulting solid was filtered to give the desired product as a tan solid (2.55 g, 9.39 mmol, 88%). LC-MS m/z = 272.0 ($M+1$) (retention time = 2.05) ¹H NMR (300 MHz, DMSO) δ 9.55 (s, 1H), 8.81 - 8.64 (m, 2H), 8.09 (d, J = 9.2 Hz, 1H), 7.78 (dd, J = 9.2, 2.8 Hz, 1H), 7.61 (dd, J = 7.9, 4.8 Hz, 1H), 7.49 (d, J = 2.5 Hz, 1H), 4.00 (s, 3H).

[0291] Method F6: 6-Bromo-4-chloro-8-fluoro-2-(pyridin-3-yl)quinazoline (v-d) To a suspension of 6-bromo-8-fluoro-2-(pyridin-3-yl)quinazolin-4-ol (6.16 g, 0.0192 mol) in toluene (60 mL) was added phosphorus oxychloride (5.30 mL, 0.0579 mol) at room temperature. The mixture was refluxed for 3 h. The solvent was evaporated and water was added to the residue under cooling conditions. The suspension was stirred at room temperature for 30 min, the resulting precipitate was filtered and dried to give the title compound (6.5 g, quantitative). ¹H NMR (400 MHz, DMSO) δ 9.58 (d, J = 1.6 Hz, 1H), 8.82 (dd, J = 4.7, 1.5 Hz, 1H), 8.80 - 8.75 (m, 1H), 8.37 (dd, J = 9.7, 1.9 Hz, 1H), 8.34-8.29 (m, 1H), 7.67 (dd, J = 7.8, 4.6 Hz, 1H).

[0292] Method F7: 4-Bromo-6-methoxy-2-(pyridin-3-yl)quinazoline (v-e) To a sealed tube containing 6-methoxy-2-(pyridin-3-yl)quinazolin-4(3H)-one (1.30 g, 5.13 mmol) in dichloromethane (20 mL) was added 1 M phosphorus tribromide in dichloromethane (10.3 mL, 10.3 mmol) and DMF (2 mL). The reaction mixture was heated at 60°C for 4 h. After cooling, excess dichloromethane was evaporated leaving a tan residue. This solid was added to an ice-water mixture (100 mL) with cooling and allowed to stir. The pH of the suspension was adjusted to about pH 9 via dropwise addition of 28% ammonium hydroxide, and stirring was continued for 30 mins. The resulting solid was filtered to give the desired product as a tan solid (1.49 g, 4.71 mmol, 92%). LC-MS m/z = 318.3 ($M+2$) (retention time = 2.19).

Method G1: 2-(6-Propoxy-2-pyridin-3-yl-quinazolin-4-ylamino)-benzamide (vi-a)

[0293] (This method is representative of method G1, G2, and G3. These three methods can be implemented in a similar way except for substitution of the appropriate solvent and temperature) A mixture of 4-chloro-6-propoxy-2-(pyridin-3-yl)quinazoline (90 mg, 0.3 mmol) and 2-aminobenzamide (52 mg, 0.4 mmol) in i-PrOH (5 mL) was stirred at 85 °C for 18 h. The yellow precipitate was collected and washed with i-PrOH (10 mL). The solid was suspended in water (10 mL) and NH₃-H₂O (1 mL) was added. After filtration the solid was dried *in vacuo* to afford 31.0 mg of vi-a as a white solid (30.8%). LCMS m/z = 400.1 ($M+1$) (Method B) (retention time = 1.96 min). ¹H-NMR (400 MHz, DMSO-*d*₆): δ 9.59 (d, J = 2.0 Hz, 1H), 9.58 (d, J = 7.6 Hz, 1H), 8.69 - 8.74 (m, 2H), 8.48 (s, 1H), 7.97 (d, J = 6.8 Hz, 2H), 7.90 (d, J = 8.8 Hz, 1H), 7.74 (t, J = 7.6 Hz, 1H), 7.56 - 7.61 (m, 3H), 7.20 (t, J = 7.2 Hz, 1H), 4.16 (t, J = 6.4 Hz, 2H), 1.86 (dd, J = 14.0, 6.8 Hz, 2H), 1.07 (t, J = 7.2 Hz, 3H).

Method G8: 2-(6-ethoxy-2-(pyridin-3-yl)quinazolin-4-yloxy)benzamide (vii-a)

[0294] (The method G8 is representative of method G6, G7, G9, G10 and G11. These six methods can be implemented in a similar way except for substitution of the appropriate base, solvent and temperature) To a 2.5 dram reaction vial was first added sodium hydride 60% (0.028 g, 0.700 mmol) and salicylamide (0.072 g, 0.525 mmol) in DMF (2 mL). The mixture was allowed to stir at room temperature for 1 h. Then, 4-chloro-6-ethoxy-2-(pyridin-3-yl)quinazoline (0.100 g, 0.350 mmol) was added to the mixture, and the reaction was allowed to proceed at room temperature overnight. LC-MS analysis of the crude mixture showed about 85% of product formed and 10% remaining starting material. Water (30 mL) was added to the mixture, and the product was extracted with chloroform (3 x 15 mL). The combined organic layers were dried (Na₂SO₄), filtered, and concentrated. The crude product was purified via ISCO (silica gel, 97.5:2.5 CH₂Cl₂/MeOH; 12 g column) to afford 13.9 mg of the desired product as a white solid (10.3%) LCMS m/z = 387 (M+1) (Method C) (retention time = 2.05 min). ¹H NMR (300 MHz, DMSO) δ 11.47 (s, 2H), 9.39 (s, 1H), 8.66 (d, J = 2.8 Hz, 1H), 8.56 - 8.47 (m, 1H), 8.01 - 7.90 (m, 2H), 7.69 - 7.55 (m, 2H), 7.55 - 7.40 (m, 2H), 7.05 (t, J = 7.5 Hz, 1H), 6.97 (d, J = 8.2 Hz, 1H), 4.23 (q, J = 6.9 Hz, 2H), 1.44 (t, J = 6.9 Hz, 3H).

Method G13: 4-(4-chlorophenyl)-N-(6-methoxy-2-(pyridin-3-yl)quinazolin-4-yl)thiazol-2-amine (vi-c) (The method G13 is representative of method G12 also.

[0295] This method can be implemented in a similar way except for substitution of the appropriate base, solvent and temperature) To a suspension of 4-chloro-6-methoxy-2-(pyridine-3-yl)quinazoline (645.2 mg, 2.375 mmol) and 2-amino-4-(4-chlorophenyl)thiazole (1050 mg, 4.98 mmol) in DMA (40 mL) was added Cs₂CO₃ (2430 mg, 7.46 mmol) at room temperature. The mixture was stirred at 80°C for 9.5 h. Water was added and a precipitate formed which was collected by filtration and washed with H₂O Recrystallization from acetone/DMF/methanol gave 383.6 mg of the product in a 36% yield as yellow solid, >98% purity by ¹H NMR). ¹H NMR (400 MHz DMSO-d₆) δ 12.52 (s, 1H), 9.78 (d, J = 1.56 Hz, 1H), 8.91-8.88 (m, 1H), 8.74 (dd, J = 4.74, 1.60 Hz, 1H), 8.33 (brs, 1H), 8.06 (d, J = 8.56 Hz, 2H), 7.93 (d, J = 9.08 Hz, 1H), 7.89 (s, 1H), 7.66-7.59 (m, 2H), 7.55 (d, J = 8.56 Hz, 2H), 4.01 (s, 3H).

Method G14: 4-(6-methoxy-2-(pyridin-3-yl)quinazolin-4-ylamino)-1H-pyrazole-5-carboxamide, 2HCl (vi-d): (The method G14 is representative of method G4 and G5 also. This method can be implemented in a similar way except for substitution of the appropriate solvent and adjustment of the temperature)

[0296] To a microwave vial containing 4-bromo-6-methoxy-2-(pyridin-3-yl)quinazoline (150.0 mg, 0.47 mmol) in DMF (2 mL) was added 4-amino-1H-pyrazole-5-carboxamide (66.0 mg, 0.52 mmol) and sodium *tert*-butoxide (50 mg, 0.52 mmol). The reaction mixture was heated at 100°C for 15 mins by microwave irradiation. Water (50 mL) was added to the reaction mixture, and extracted with ethyl acetate (5 x 50 mL). The crude material was purified via ISCO (silica, 12 g column, 93% CH₂Cl₂ - 7% MeOH - 0.1% NH₄OH) giving the product as a yellow solid. The free base was then converted to the HCl salt to yield the final product as an orange solid (59.8 mg, 0.14 mmol, 22%). LC-MS m/z = 362.4 (M+1) (retention time = 1.57) ¹H NMR (300 MHz, DMSO) δ 11.28 (s, 1H), 9.62 (d, J = 1.7 Hz, 1H), 9.17 (d, J = 7.8 Hz, 1H), 8.93 (dd, J = 5.2, 1.3 Hz, 1H), 8.60 (s, 1H), 8.05 (s, 1H), 8.03 - 7.94 (m, 2H), 7.77 (s, 1H), 7.64 (dd, J = 9.2, 2.5 Hz, 1H), 7.40 (d, J = 2.6 Hz, 1H), 3.97 (s, 3H).

[0297] Method J1: 1-(6-Methoxy-2-(pyridin-3-yl)quinazolin-4-yl)-1H-benzo[d]imidazol-2(3H)-one (vi-b) To a 50-mL two-neck round bottom flask equipped with a reflux condenser was added a mixture of 4-chloro-6-methoxy-2-(pyridin-3-yl)quinazoline (50 mg, 0.18 mmol, 1 eq.) and 1H-benzo[d]imidazol-2(3H)-one (27 mg, 0.21 mmol, 1.1 eq.) in 5 mL of dry 1,4-dioxane. Pd(PPh₃)₄ (10.6 mg, 0.009 mmol, 0.05 eq) and *t*-BuOK (41 mg, 0.36 mmol, 2 eq.) were added. The resulting mixture was stirred at 100 °C under N₂ atmosphere overnight. After cooling, 20 mL of methanol was added. The mixture was filtered, the filtrate was concentrated *in vacuo* and then was purified with chromatography on silica gel (ethyl acetate/ petroleum ether from 1:4 to 1:2) to give the crude product, which was further purified with reverse phase HPLC to afford 4.8 mg of vi-e as a pale yellow solid (7%). MS m/z = 370.1 (M+1) (Method B) (retention time = 1.680 min). ¹H NMR (400 MHz, DMSO-d₆): δ 11.51 (s, 1H), 8.78 (d, J = 8.0 Hz, 2H), 8.17 (d, J = 9.2 Hz, 1H), 7.80 (dd, J = 9.2, 2.8 Hz, 1H), 7.70-7.64 (m, 1H), 7.46 (d, J = 2.8 Hz, 1H), 7.41 (d, J = 8.0 Hz, 1H), 7.20 (d, J = 4.0 Hz, 2H), 7.16 - 7.08 (m, 1H), 3.89 (s, 3H).

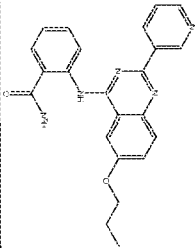
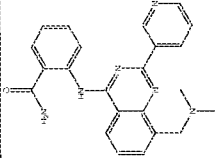
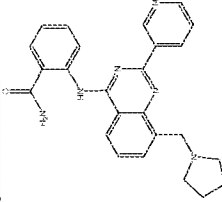
[0298] Method J2: 6-methoxy-N-(pyridin-2-yl)-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride (vi-e) To a suspension of 4-chloro-6-methoxy-2-(pyridine-3-yl)quinazoline (600 mg, 2.208 mmol) in dioxane (40 mL) under N₂, was added tris(dibenzylideneacetone)dipalladium(0) (103.1 mg, 0.113 mmol), 4,5-bis(diphenylphosphino)-9,9-dimethylxanthene (132.2 mg, 0.228 mmol) and cesium carbonate (1.1289 g, 3.46 mmol) at room temperature. 2-Aminopyridine (229 mg, 2.43 mmol) was added and the mixture was stirred at 100°C for 2h 30 min. Water was added and then a precipitate formed. The solid was collected and washed with water. The solid was dissolved in CH₂Cl₂. Purification was carried out using NH-silica gel to give the free base (649.3 mg). The free base was converted to the HCl salt by dissolving the

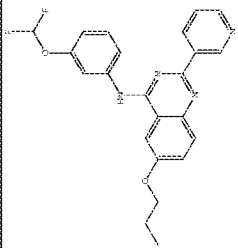
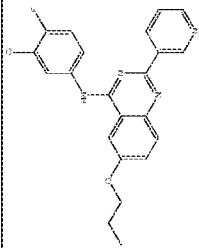
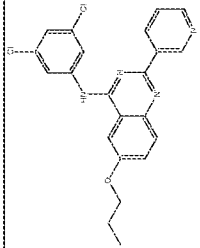
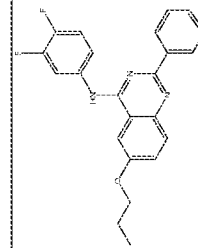
compound in CH_2Cl_2 / MeOH and 1.5 ml of 4N HCl in ethyl acetate was added and then a precipitate formed. The solid was collected and dried *in vacuo* (at 40 degrees on and 60 degrees for ca. 3 h) and then washed with methanol. The resulting solid was dried *in vacuo* at 60 degrees to give 682mg of the desired product as the HCl salt in a 77% yield as a pale yellow solid. ^1H NMR ($\text{DMSO}-d_6$) δ 11.14 (brs, 1H), 9.58 (s, 1H), 9.13 (dd, $J = 7.96$ Hz, 1H), 8.93 (d, $J = 5.24$ Hz, 1H), 8.54 (d, $J = 4.72$ Hz, 1H), 8.48 (d, $J = 8.32$ Hz, 1H), 8.28 (brs, 1H), 8.09 (brt, $J = 7.16$ Hz, 1H), 8.02-7.96 (m, 2H), 7.65 (dd, $J = 8.80, 2.48$ Hz, 1H), 7.34 (brt, $J = 6.52$ Hz, 1H), 4.01 (s, 3H). The 1H of 2HCl was not observed.

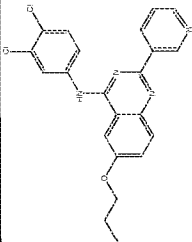
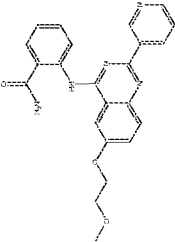
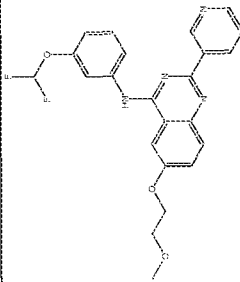
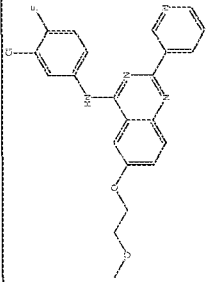
[0299] Method J3: N-(Biphenyl-4-yl)-6-methoxy-2-(pyridin-3-yl)quinazolin-4-amine hydrochloride (vi-f) To a suspension of N-(4-bromophenyl)-6-methoxy-2-(pyridin-3-yl)quinazolin-4-amine (548.7 mg, 1.347 mmol) and phenylboronic acid (270 mg, 2.21 mmol) in dioxane / H_2O (2/1) (30 mL) under N_2 was added Na_2CO_3 (485 mg, 4.58 mmol) and tetrakis(triphenylphosphine) palladium(0) (78 mg, 0.067 mmol) at room temperature. The mixture was stirred at 100°C for 1 h. Water was added and then a precipitate formed. The solid was dissolved in methanol/acetone at 60°C . The solution was filtrated through Celite to remove any extra palladium. The filtrate was concentrated down to give 493.4 mg of a solid residue in. The solid was added to CH_2Cl_2 followed by addition of 4N HCl in ethyl acetate (0.4 mL) at room temperature to form the HCl salt. The mixture was stirred at room temperature and the resulting solid was filtered and dried *in vacuo* to give 435.2 mg and a yield of 73% as the HCl salt. ^1H NMR ($\text{DMSO}-d_6$) δ 10.24 (br, 1H), 9.54 (s, 1H), 8.92 (d, $J = 7.52$ Hz, 1H), 8.82 (d, $J = 3.96$ Hz, 1H), 8.10 (d, $J = 2.44$ Hz, 1H), 8.03 (d, $J = 8.68$ Hz, 2H), 7.92 (d, $J = 9.08$ Hz, 1H), 7.84 (d, $J = 8.68$ Hz, 2H), 7.81 (m, 1H), 7.77-7.75 (m, 2H), 7.62 (dd, $J = 9.08, 2.44$ Hz, 1H), 7.50 (m, 2H), 7.38 (m, 1H), 4.01 (s, 3H). The 1H of HCl was not observed.

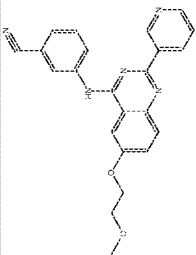
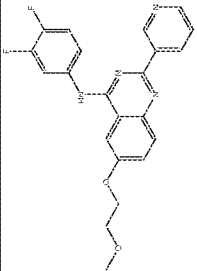
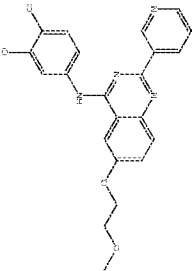
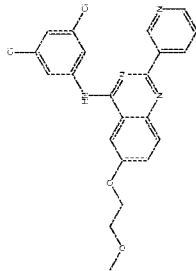
[0300] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 1 and 5** (prepared according to method **procedure A - J** as designated).

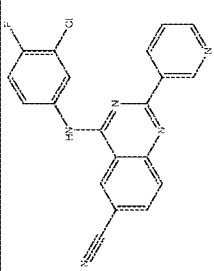
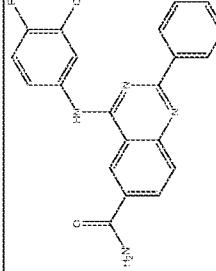
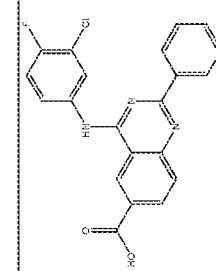
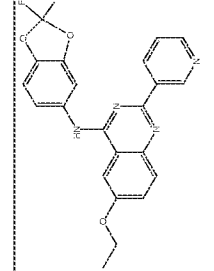
Table 1:

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1			399.45	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.59 (d, J = 2.0 Hz, 1H), 9.58 (d, J = 7.6 Hz, 1H), 8.69 – 8.74 (m, 2H), 8.46 (s, 1H), 7.97 (d, J = 6.8 Hz, 2H), 7.90 (d, J = 8.8 Hz, 1H), 7.74 (t, J = 7.6 Hz, 1H), 7.56 – 7.61 (m, 3H), 7.20 (t, J = 7.2 Hz, 1H), 4.16 (t, J = 6.4 Hz, 2H), 1.86 (dd, J = 7.2, 6.8 Hz, 2H), 1.07 (t, J = 7.2 Hz, 3H).	DMSO	400.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
2		HCl	396.46	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.20 (s, 1H), 9.79 (s, 1H), 9.02 – 9.08 (m, 2H), 8.87 (d, J = 3.2 Hz, 1H), 8.53 (s, 1H), 8.34 (d, J = 8.0 Hz, 1H), 8.19 (d, J = 6.8 Hz, 1H), 7.74 – 7.99 (m, 5H), 7.28 (t, J = 7.6 Hz, 1H), 4.92 (d, J = 4.0 Hz, 2H), 2.87 (d, J = 3.2 Hz, 6H).	DMSO	399.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
3		HCl	424.50	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.07 (s, 1H), 9.64 (d, J = 1.6 Hz, 1H), 9.15 (d, J = 8.0 Hz, 1H), 8.78 (ddd, J = 8.0, 4.0, 2.4 Hz, 1H), 8.73 (dd, J = 4.4, 1.2 Hz, 1H), 8.49 (s, 1H), 8.09 (d, J = 8.0 Hz, 1H), 7.92 – 7.69 (m, 3H), 7.67 – 7.74 (m, 2H), 7.62 (dd, J = 8.0, 4.8 Hz, 1H), 7.22 (t, J = 7.6 Hz, 1H), 4.29 (s, 2H), 2.61 (brs, 4H), 1.70 (brs, 4H).	DMSO	425.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

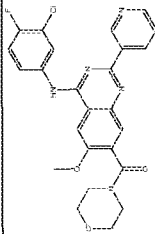
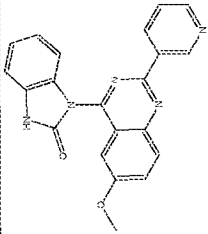
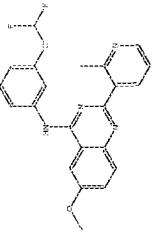
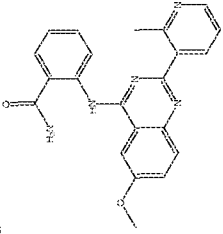
4			422.43	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.87 (s, 1H), 9.52 (s, 1H), 8.67 (d, J = 5.6 Hz, 1H), 7.99 (d, J = 2.4 Hz, 1H), 7.94 (s, 1H), 7.86 (d, J = 9.2 Hz, 1H), 7.80 (d, J = 8.0 Hz, 1H), 7.50 – 7.58 (m, 3H), 7.30 (t, J = 34.0 Hz, 1H), 7.00 (dd, J = 8.4, 2.0 Hz, 1H), 4.16 (t, J = 6.4 Hz, 2H), 1.86 (dd, J = 7.2, 6.4 Hz, 2H), 1.07 (t, J = 7.2 Hz, 3H).	DMSO	423.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
5		HCl	408.86	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.28 (s, 1H), 9.48 (s, 1H), 8.93 (d, J = 6.4 Hz, 1H), 8.85 (d, J = 4.4 Hz, 1H), 8.18 (dd, J = 7.6, 2.4 Hz, 1H), 8.06 (d, J = 2.0 Hz, 1H), 7.85 – 7.93 (m, 3H), 7.62 (dd, J = 8.6, 6.4 Hz, 1H), 7.55 (t, J = 8.8 Hz, 1H), 4.17 (t, J = 6.4 Hz, 3H), 1.86 (dd, J = 7.6, 6.4 Hz, 2H), 1.06 (t, J = 7.6 Hz, 3H).	DMSO	409.1, 411.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
6			425.31	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.93 (s, 1H), 9.51 (s, 1H), 8.68 – 8.71 (m, 2H), 8.16 (d, J = 2.4 Hz, 1H), 7.94 (d, J = 2.0 Hz, 1H), 7.86 (d, J = 8.8 Hz, 1H), 7.56 – 7.60 (m, 2H), 7.37 (s, 1H), 4.14 (t, J = 6.4 Hz, 2H), 1.84 (dd, J = 10.4 Hz, 7.2 Hz, 2H), 1.07 (t, J = 7.6 Hz, 3H).	DMSO	425.0, 427.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
7			392.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.06 (s, 1H), 9.49 (s, 1H), 8.77 – 8.81 (m, 2H), 8.09 (ddd, J = 10.0, 7.6, 2.4 Hz, 1H), 7.86 (d, J = 9.2 Hz, 1H), 7.69 – 7.74 (m, 2H), 7.54 – 7.61 (m, 2H), 4.15 (t, J = 6.4 Hz, 2H), 1.86 (dd, J = 14.0, 7.6 Hz, 2H), 1.06 (t, J = 7.2 Hz, 3H).	DMSO	393.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

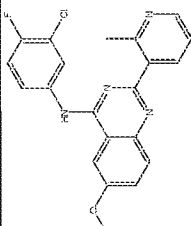
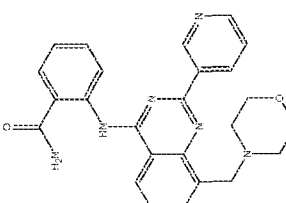
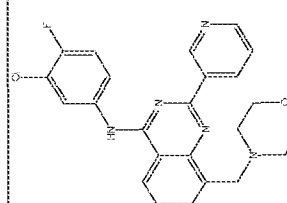
8			425.31	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.05 (s, 1H), 9.51 (s, 1H), 8.75 – 8.80 (m, 2H), 8.34 (d, J = 2.4 Hz, 1H), 7.96 – 8.00 (m, 2H), 7.88 (d, J = 9.2 Hz, 1H), 7.68 – 7.75 (m, 2H), 7.59 (dd, J = 9.2, 2.8 Hz, 1H), 4.15 (t, J = 6.4 Hz, 2H), 1.86 (dd, J = 7.2, 6.8 Hz, 2H), 1.06 (t, J = 7.6 Hz, 3H).	DMSO	425.0, 427.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
9			415.44	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.03 (s, 1H), 9.59 (d, J = 1.2 Hz, 1H), 9.15 (d, J = 8.4 Hz, 1H), 8.69 – 8.74 (m, 2H), 8.49 (s, 1H), 7.96 – 8.06 (m, 2H), 7.90 (d, J = 9.2 Hz, 1H), 7.56 – 7.63 (m, 3H), 7.20 (t, J = 8.0 Hz, 1H), 4.31 (t, J = 4.0 Hz, 2H), 3.79 (t, J = 4.8 Hz, 2H), 3.31 (s, 3H).	DMSO	416.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
10			438.43	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.86 (s, 1H), 9.53 (s, 1H), 8.66 – 8.68 (m, 2H), 8.02 (d, J = 2.8 Hz, 1H), 7.95 (t, J = 2.0 Hz, 1H), 7.86 (d, J = 8.8 Hz, 1H), 7.79 (d, J = 1.2 Hz, 1H), 7.48 – 7.60 (m, 3H), 7.30 (t, J = 7.4 Hz, 1H), 7.00 (dd, J = 8.0, 2.0 Hz, 1H), 4.33 (t, J = 4.8 Hz, 2H), 3.79 (t, J = 4.4 Hz, 2H), 3.37 (s, 3H).	DMSO	439.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
11		HCl	424.86	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.27 (s, 1H), 9.48 (s, 1H), 8.93 (d, J = 8.4 Hz, 1H), 8.85 (d, J = 4.8 Hz, 1H), 8.19 (dd, J = 6.8, 2.4 Hz, 1H), 8.10 (d, J = 2.4 Hz, 1H), 7.85 – 7.93 (m, 3H), 7.64 (dd, J = 9.2, 6.8 Hz, 1H), 7.55 (t, J = 8.8 Hz, 1H), 4.34 (t, J = 4.4 Hz, 2H), 3.78 (t, J = 4.4 Hz, 2H), 3.31 (s, 3H).	DMSO	425.1, 427.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

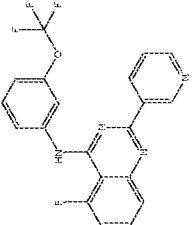
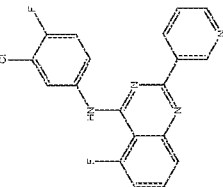
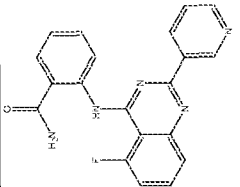
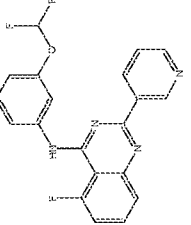
12			397.43	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.07 (s, 1H), 9.50 (s, 1H), 8.78 (dd, J = 17.6, 7.6 Hz, 1H), 8.39 (s, 1H), 8.26 (dd, J = 8.0, 2.0 Hz, 1H), 8.02 (d, J = 2.0 Hz, 1H), 7.90 (d, J = 9.2 Hz, 1H), 7.61 – 7.73 (m, 4H), 4.34 (t, J = 4.3 Hz, 2H), 3.79 (t, J = 4.4 Hz, 2H), 3.35 (s, 3H).	DMSO	398.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
13			408.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.86 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 8.67 (dd, J = 4.8, 1.6 Hz, 1H), 8.64 (dt, J = 8.0, 1.6 Hz, 1H), 8.12 (ddd, J = 13.2, 7.2, 2.4 Hz, 1H), 7.98 (d, J = 2.8 Hz, 1H), 7.86 (d, J = 9.2, 1H), 7.69 – 7.72 (m, 1H), 7.52 – 7.60 (m, 3H), 4.31 (t, J = 4.0 Hz, 2H), 3.76 – 3.80 (m, 2H), 3.37 (s, 3H).	DMSO	409.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
14			441.31	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.90 (s, 1H), 9.52 (d, J = 1.2 Hz, 1H), 8.64 – 8.68 (m, 1H), 8.37 (d, J = 2.8 Hz, 1H), 7.98 (d, J = 2.4 Hz, 1H), 7.96 (dd, J = 10.6, 2.4 Hz, 1H), 7.86 (d, J = 9.2 Hz, 1H), 7.74 (d, J = 9.2 Hz, 1H), 7.59 (dd, J = 8.8, 2.4 Hz, 1H), 7.55 (dd, J = 8.0, 4.8 Hz, 1H), 4.31 (t, J = 4.08 Hz, 2H), 3.78 (t, J = 4.4 Hz, 2H), 3.37 (s, 3H).	DMSO	441.0, 443.0 (M+1), 221.9 (M/2 +1)	Method A (TFA)	95	Method C, G1
15			441.31	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.93 (s, 1H), 9.53 (d, J = 1.6 Hz, 1H), 8.66 – 8.70 (m, 2H), 8.18 (s, 1H), 8.17 (s, 1H), 7.99 (d, J = 2.4 Hz, 1H), 7.89 (d, J = 8.8 Hz, 1H), 7.61 (dd, J = 8.8, 2.4 Hz, 1H), 7.56 (dd, J = 7.6, 4.8 Hz, 1H), 7.39 (t, J = 2.0 Hz, 1H), 4.33 (t, J = 4.3 Hz, 2H), 3.79 (t, J = 4.4 Hz, 2H), 3.37 (s, 3H).	DMSO	441.1, 443.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

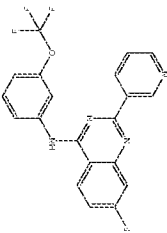
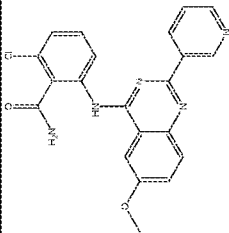
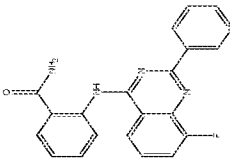
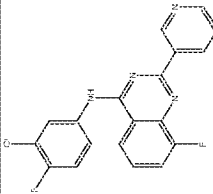
16		HCl	375.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.34 (s, 1H), 9.53 (d, J = 1.2 Hz, 1H), 9.16 (d, J = 1.2 Hz, 1H), 8.74 (dd, J = 4.0, 1.6 Hz, 1H), 8.66–8.69 (m, 2H), 8.20–8.28 (m, 2H), 8.00 (d, J = 8.4 Hz, 1H), 7.89–7.93 (m, 1H), 7.55–7.61 (m, 2H).	DMSO	376.0, 378.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
17			393.8	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.37 (s, 1H), 9.54 (d, J = 1.6 Hz, 1H), 9.09 (s, 1H), 8.69 (d, J = 10.0 Hz, 1H), 8.68 (d, J = 8.0 Hz, 1H), 8.23–8.27 (m, 2H), 8.11 (s, 1H), 8.78 (s, 2H), 7.50–7.53 (m, 3H).	DMSO	394.1, 396.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
18		HCl	394.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.81 (s, 1H), 9.48 (d, J = 0.8 Hz, 1H), 9.28 (d, J = 0.8 Hz, 1H), 8.69 (d, J = 10.0 Hz, 1H), 8.68 (d, J = 8.0 Hz, 1H), 8.23–8.27 (m, 2H), 8.11 (s, 1H), 8.78 (s, 2H), 7.50–7.56 (m, 3H).	DMSO	395.0, 397.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
19			422.38	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.11 (brs, 1H), 9.48 (s, 1H), 8.79 (d, J = 4.8 Hz, 1H), 8.83 (d, J = 8.0 Hz, 1H), 8.07 (d, J = 2.0 Hz, 1H), 8.02 (d, J = 2.4 Hz, 1H), 7.86–7.90 (m, 1H), 1.77 (d, J = 8.0 Hz, 1H), 7.52–7.62 (m, 3H), 4.27 (q, J = 7.2 Hz, 2H), 1.46 (t, J = 7.2 Hz, 3H).	DMSO	423.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

20		HCl	408.91	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.73 (s, 1H), 9.52 (d, J = 1.6 Hz, 1H), 9.32 (d, J = 1.6 Hz, 1H), 9.01 (d, J = 4.8 Hz, 1H), 8.91 (d, J = 5.2 Hz, 1H), 8.39 (dd, J = 8.4, 1.6 Hz, 1H), 8.17 (dd, J = 6.8, 2.0 Hz, 1H), 8.01 (d, J = 8.8 Hz, 1H), 7.90 – 7.95 (m, 2H), 7.55 (t, J = 9.2 Hz, 1H), 3.98 (s, 3H).	DMSO	409.1, 411.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
21			399.45	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.94 (s, 1H), 9.56 (s, 1H), 9.14 (d, J = 8.0 Hz, 1H), 8.83 – 8.66 (m, 2H), 8.46 (s, 1H), 8.06 – 7.91 (m, 2H), 7.76 – 7.58 (m, 2H), 7.35 (s, 1H), 7.27 (d, J = 2.0 Hz, 1H), 7.18 (t, J = 7.2 Hz, 1H), 4.15 (q, J = 6.8 Hz, 2H), 2.64 (s, 3H), 1.43 (t, J = 6.8 Hz, 3H).	DMSO	400.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
22			440.42	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.84 (s, 1H), 9.55 (d, J = 1.2 Hz, 1H), 8.73 – 8.60 (m, 2H), 8.14 (s, 1H), 7.91 (d, J = 8.4 Hz, 1H), 7.80 (d, J = 2.4 Hz, 1H), 7.60 (t, J = 8.0 Hz, 1H), 7.53 (dd, J = 7.6, 4.8 Hz, 1H), 7.45 (d, J = 1.6 Hz, 1H), 7.16 (d, J = 8.4 Hz, 1H), 4.23 (q, J = 6.8 Hz, 2H), 2.70 (s, 3H), 1.44 (t, J = 6.8 Hz, 3H).	DMSO	440.9 (M+1)	Method A (TFA)	95	Method D, G1
23			405.81	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.08 (s, 1H), 9.42 (s, 1H), 8.68 (d, J = 3.6 Hz, 1H), 8.55 (d, J = 8.4 Hz, 1H), 8.32 (s, 1H), 8.15 (dd, J = 6.8, 2.4 Hz, 1H), 8.04 (s, 1H), 7.93 – 7.78 (m, 1H), 7.54 (td, J = 8.3, 6.9 Hz, 2H), 4.08 (s, 3H).	DMSO	406.1, 408.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

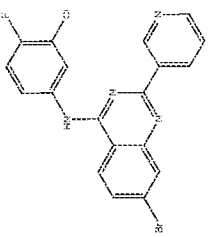
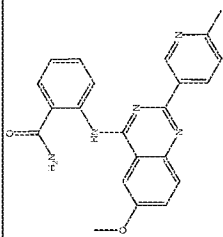
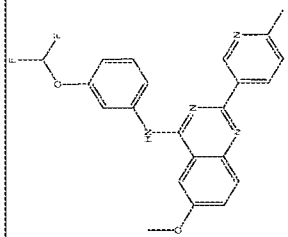
24			493.92	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.00 (s, 1H), 9.49 – 9.51 (m, 1H), 8.62 – 8.70 (m, 2H), 8.21 (dd, J = 6.8, 2.8 Hz, 1H), 8.03 (s, 1H), 7.87 – 7.91 (m, 1H), 7.75 (s, 1H), 7.53 – 7.59 (m, 2H), 4.02 (s, 3H), 3.64 – 3.73 (m, 4H), 3.49 – 3.58 (m, 2H), 3.14 – 3.22 (m, 2H).	DMSO	494.1, 496.1 (M+1)	Method B (NH ₄ HCO ₄)	95	Method C, G1
25		TFA	369.38	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.51 (s, 1H), 8.78 (d, J = 8.0 Hz, 2H), 8.17 (d, J = 9.2 Hz, 1H), 7.80 (dd, J = 9.2, 2.8 Hz, 1H), 7.70 – 7.64 (m, 1H), 7.46 (d, J = 2.8 Hz, 1H), 7.41 (d, J = 8.0 Hz, 1H), 7.20 (d, J = 4.0 Hz, 2H), 7.16 – 7.08 (m, 1H), 3.89 (s, 3H).	DMSO	370.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, J1
26			408.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.87 (s, 1H), 8.53 (dd, J = 4.7, 1.5 Hz, 1H), 8.23 (d, J = 6.8 Hz, 1H), 8.00 (d, J = 2.6 Hz, 1H), 7.8 (t, J = 2.1 Hz, 1H), 7.84 (d, J = 9.1 Hz, 1H), 7.73 (dd, J = 8.2, 1.2 Hz, 1H), 7.58 (dd, J = 9.1, 2.6 Hz, 1H), 7.41 – 7.25 (m, 3H), 7.24 (t, J = 74.0 Hz, 1H), 6.96 (dd, J = 8.0, 2.2 Hz, 1H), 3.99 (s, 3H), 2.75 (s, 3H).	DMSO	409.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
27			385.42	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.09 (s, 1H), 9.08 (d, J = 7.8 Hz, 1H), 8.55 (dd, J = 4.6, 1.6 Hz, 1H), 8.49 (s, 1H), 8.22 (dd, J = 7.8, 1.7 Hz, 1H), 8.01 (s, 1H), 7.95 (dd, J = 7.9, 1.2 Hz, 1H), 7.87 (d, J = 9.6 Hz, 1H), 7.66 – 7.58 (m, 3H), 7.40 (dd, J = 7.7, 4.8 Hz, 1H), 7.15 (dd, J = 11.5, 4.4 Hz, 1H), 3.99 (s, 3H), 2.77 (s, 3H).	DMSO	386.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

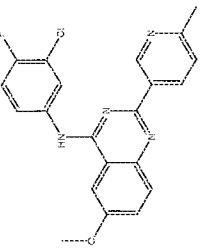
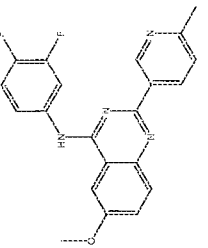
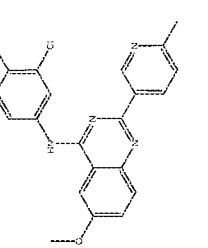
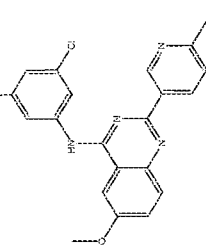
28		394.83		¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.10 (s, 1H), 8.63 (d, J = 4.1 Hz, 1H), 8.43 (d, J = 7.6 Hz, 1H), 8.15 (dd, J = 6.8, 2.5 Hz, 1H), 8.03 (d, J = 2.3 Hz, 1H), 7.88 – 7.78 (m, 2H), 7.58 (ddd, J = 12.5, 8.3, 4.1 Hz, 2H), 7.49 (t, J = 9.1 Hz, 1H), 3.99 (s, 3H), 2.80 (s, 3H).	DMSO	395.1, 397.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
29		440.5	HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.19 (s, 1H), 11.12 – 10.93 (m, 1H), 9.83 (d, J = 1.5 Hz, 1H), 9.21 (d, J = 7.8 Hz, 1H), 9.01 (d, J = 7.8 Hz, 1H), 8.93 – 8.89 (m, 1H), 8.53 (s, 1H), 8.36 (dd, J = 11.8, 7.9 Hz, 2H), 8.02 – 7.73 (m, 6H), 7.29 (d, J = 8.02 Hz, 1H), 5.01 (s, 2H), 3.95 – 3.81 (m, 4H), , 3.37 (s, 4H).	DMSO	441.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
30		449.91	HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.05 (s, 1H), 9.56 (s, 1H), 8.73 – 8.66 (m, 2H), 8.47 (d, J = 8.1 Hz, 1H), 8.28 (dd, J = 6.8, 2.5 Hz, 1H), 7.99 – 7.90 (m, 2H), 7.67 – 7.50 (m, 3H), 4.16 (s, 2H), 3.62 (s, 4H), 2.53 (s, 4H).	DMSO	450.1, 452.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

31			400.33	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.47 (d, J = 2.0 Hz, 2H), 8.70 (dd, J = 4.8, 1.6 Hz, 1H), 8.65 – 8.59 (m, 1H), 8.06 (s, 1H), 7.91 (dt, J = 8.1, 6.0 Hz, 1H), 7.82 (dd, J = 8.2, 1.2 Hz, 1H), 7.76 (d, J = 8.4 Hz, 1H), 7.59 (t, J = 8.2 Hz, 1H), 7.51 (ddd, J = 20.0, 10.0, 6.4 Hz, 2H), 7.21 (d, J = 8.4 Hz, 1H).	DMSO	401.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
32		HCl	368.77	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.53 (d, J = 11.3 Hz, 1H), 9.44 (d, J = 1.2 Hz, 1H), 8.84 – 8.79 (m, 2H), 8.09 (dd, J = 6.8, 2.6 Hz, 1H), 7.97 – 7.89 (m, 1H), 7.85 – 7.75 (m, 3H), 7.57 – 7.47 (m, 2H).	DMSO	369.1, 371.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
33		HCl	359.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.21 (d, J = 12.1 Hz, 1H), 9.53 (d, J = 1.2 Hz, 1H), 8.98 (d, J = 8.1 Hz, 1H), 8.89 (d, J = 4.2 Hz, 1H), 8.71 (d, J = 7.9 Hz, 1H), 8.35 (s, 1H), 7.97 – 7.76 (m, 5H), 7.72 – 7.65 (m, 1H), 7.50 (dd, J = 12.1, 7.5 Hz, 1H), 7.29 (t, J = 7.1 Hz, 1H).	DMSO	360.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
34			382.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.49 (d, J = 1.8 Hz, 1H), 9.42 (s, 1H), 8.71 (dd, J = 4.6, 1.3 Hz, 1H), 8.65 (d, J = 7.8 Hz, 1H), 7.90 (dd, J = 15.8, 9.7 Hz, 2H), 7.73 (dd, J = 19.0, 8.26 Hz, 2H), 7.56 – 7.12 (m, 4H), 7.04 (dd, J = 8.2, 1.8 Hz, 1H).	DMSO	383.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

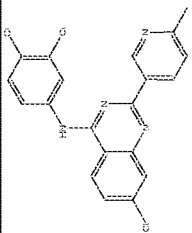
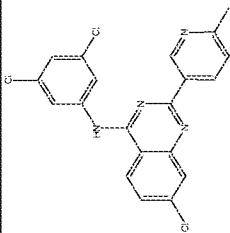
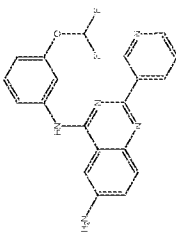
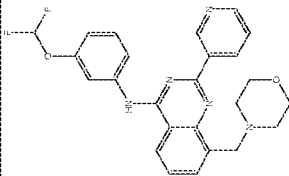
35		HCl	400.33	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.43 (s, 1H), 9.52 (s, 1H), 8.89 (d, J = 8.2 Hz, 1H), 8.85 (d, J = 4.2 Hz, 1H), 8.78 (dd, J = 9.2, 5.9 Hz, 1H), 8.09 (s, 1H), 7.95 (dd, J = 8.2, 1.2 Hz, 1H), 7.80 (dd, J = 7.9, 5.2 Hz, 1H), 7.72 – 7.58 (m, 3H), 7.21 (d, J = 8.4 Hz, 1H).	DMSO	401.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
36		HCl	405.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.74 – 10.48 (m, 1H), 9.36 (d, J = 1.6 Hz, 1H), 8.95 (d, J = 8.2 Hz, 1H), 8.89 – 8.85 (m, 1H), 8.06 (d, J = 1.7 Hz, 1H), 8.02 (d, J = 9.15 Hz, 1H), 7.93 – 7.86 (m, 2H), 7.73 – 7.63 (m, 3H), 7.60 – 7.52 (m, 2H), 3.98 (s, 3H).	DMSO	406.1, 408.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
37		HCl	359.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.61 (s, 1H), 9.00 (dd, J = 15.4, 8.2 Hz, 2H), 8.86 (d, J = 4.13 Hz, 1H), 8.52 (s, 1H), 8.04 – 7.94 (m, 3H), 7.86 – 7.70 (m, 4H), 7.26 (dd, J = 11.2, 4.2 Hz, 1H).	DMSO	360.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
38		HCl	368.77	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.22 (s, 1H), 9.52 (s, 1H), 8.73 (d, J = 3.6 Hz, 1H), 8.69 (td, J = 8.0, 1.8 Hz, 1H), 8.38 (d, J = 8.4 Hz, 1H), 8.25 (dd, J = 6.8, 2.6 Hz, 1H), 7.91 (ddd, J = 9.0, 4.2, 2.6 Hz, 1H), 7.79 (dd, J = 9.8, 7.9 Hz, 1H), 7.67 (dd, J = 8.2, 5.2 Hz, 1H), 7.61 – 7.53 (m, 2H).	DMSO	359.0, 371.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

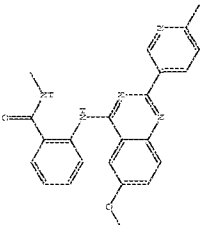
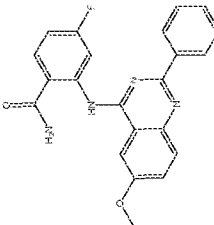
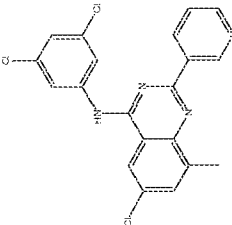
39		HCl	400.33	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.47 (s, 1H), 9.52 (d, J = 1.4 Hz, 1H), 9.04 (d, J = 8.0 Hz, 1H), 8.92 (dd, J = 5.2, 1.2 Hz, 1H), 8.52 (d, J = 8.4 Hz, 1H), 8.07 (s, 1H), 7.97 (d, J = 1.2 Hz, 2H), 7.86 – 7.80 (m, 1H), 7.74 – 7.68 (m, 1H), 7.62 (t, J = 8.2 Hz, 1H), 7.22 (d, J = 8.4 Hz, 1H).	DMSO	401.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
40			360.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.66 (d, J = 1.6 Hz, 1H), 8.92 (td, J = 7.9, 1.9 Hz, 1H), 8.71 (dd, J = 4.6, 1.6 Hz, 1H), 8.22 (d, J = 8.2 Hz, 1H), 7.87 (dd, J = 7.8, 1.9 Hz, 1H), 7.73 (dd, J = 9.2, 7.8 Hz, 1H), 7.59 (dt, J = 8.1, 5.0 Hz, 2H), 7.21 – 7.15 (m, 1H), 6.66 (d, J = 8.2 Hz, 1H), 6.54 (t, J = 7.4 Hz, 1H).	DMSO	360.9 (M+1), 382.9 (M+23)	Method B (NH ₄ HCO ₃)	95	Method C, G9
41			380.08	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.56 (brs, 1H), 9.62 (d, J = 1.2 Hz, 1H), 9.05 (d, J = 8.4 Hz, 1H), 9.00 (dd, J = 1.6, 4.8 Hz, 1H), 8.71 (d, J = 9.2 Hz, 1H), 8.31 (dd, J = 2.0, 7.2 Hz, 1H), 8.02 – 7.99 (m, 2H), 7.66 (t, J = 9.2 Hz, 1H), 7.54 (s, 1H), 7.50 (dd, J = 2.4, 9.2 Hz, 1H), 4.11 (s, 3H).	DMSO	381.1, 383.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
42			476.67	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.40 (s, 1H), 9.51 (s, 1H), 8.93 (d, J = 8.0 Hz, 1H), 8.88 (d, J = 4.8 Hz, 1H), 8.40 (d, J = 8.6 Hz, 1H), 8.35 (d, J = 1.6 Hz, 1H), 8.20 (dd, J = 4.4, 6.8 Hz, 1H), 8.04 (dd, J = 7.2, 8.8 Hz, 1H), 7.93 – 7.86 (m, 2H), 7.35 (t, J = 9.2 Hz, 1H).	DMSO	477.0, 499.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

43			429.67	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.46 (s, 1H), 9.49 (s, 1H), 8.95 (d, J = 8.1 Hz, 1H), 8.89 (d, J = 4.97 Hz, 1H), 8.60 (d, J = 8.9 Hz, 1H), 8.19 (dd, J = 6.8, 2.6 Hz, 1H), 8.14 (d, J = 1.9 Hz, 1H), 7.96 – 7.83 (m, 3H), 7.53 (t, J = 9.0 Hz, 1H).	DMSO	429.0, 430.9, 433.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
44		HCl	385.42	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.05 (s, 1H), 9.44 (d, J = 1.7 Hz, 1H), 9.16 – 8.91 (m, 2H), 8.03 – 7.89 (m, 3H), 7.88 – 7.82 (m, 1H), 7.75 – 7.68 (m, 1H), 7.67 – 7.60 (m, 2H), 7.25 (t, J = 7.6 Hz, 1H), 4.00 (s, 3H), 2.74 (s, 3H).	DMSO	385.0 (M+1), 193.4 (M/2+1)	Method A (TFA)	95	Method C, G1
45		HCl	408.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.37 – 10.12 (m, 1H), 9.35 (d, J = 1.8 Hz, 1H), 8.95 (dd, J = 4.6, 3.4 Hz, 1H), 8.03 (d, J = 2.3 Hz, 1H), 7.92 (d, J = 9.1 Hz, 1H), 7.87 (s, 1H), 7.84 – 7.74 (m, 2H), 7.62 (dd, J = 9.1, 2.6 Hz, 1H), 7.54 (t, J = 8.2 Hz, 1H), 7.32 (t, J = 74.0 Hz, 1H), 7.05 (dd, J = 8.3, 1.9 Hz, 1H), 4.00 (s, 3H), 2.72 (s, 3H).	DMSO	409.0 (M+1), 205.0 (M/2+1)	Method A (TFA)	95	Method C, G1

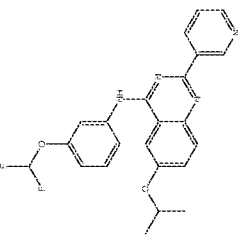
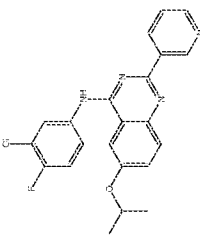
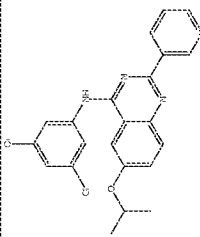
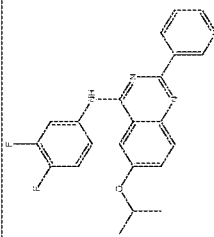
45		HCl	394.83	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.27 (brs, 1H), 9.32 (d, J = 1.7 Hz, 1H), 8.93 (d, J = 7.5 Hz, 1H), 8.19 (dd, J = 6.8, 2.5 Hz, 1H), 8.07 (d, J = 2.3 Hz, 1H), 7.88 – 7.92 (m, 2H), 7.84 (d, J = 8.1 Hz, 1H), 7.62 (dd, J = 9.2, 2.5 Hz, 1H), 7.54 (t, J = 9.1 Hz, 1H), 4.00 (s, 3H), 2.72 (s, 3H).	DMSO	395.0, 397.0 (M+1), 197.8, 198.8 (M/2+1)	Method A (TFA)	95	Method C, G1
47		HCl	378.37	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.58 – 10.30 (m, 1H), 9.32 (d, J = 1.9 Hz, 1H), 9.04 (d, J = 7.9 Hz, 1H), 8.17 – 8.04 (m, 2H), 7.95 (d, J = 9.0 Hz, 2H), 7.73 – 7.56 (m, 1H), 7.63 (dd, J = 9.2, 2.6 Hz, 1H), 7.60 – 7.50 (m, 1H), 4.01 (s, 3H), 2.76 (s, 3H).	DMSO	379.0 (M+1), 190.0 (M/2+1)	Method A (TFA)	95	Method C, G1
48		HCl	411.28	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.29 (s, 1H), 9.33 (d, J = 1.7 Hz, 1H), 9.04 – 8.89 (m, 1H), 8.31 (d, J = 2.4 Hz, 1H), 8.07 (d, J = 2.4 Hz, 1H), 7.97 (dd, J = 8.6, 2.5 Hz, 1H), 7.91 (d, J = 9.1 Hz, 1H), 7.85 (d, J = 8.6 Hz, 1H), 7.73 (d, J = 8.8 Hz, 1H), 7.62 (dd, J = 9.1, 2.5 Hz, 1H), 4.00 (s, 3H), 2.73 (s, 3H).	DMSO	411.0, 413.0 (M+1), 205.0, 206.9 (M/2+1)	Method A (TFA)	95	Method C, G1
49			411.28	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.30 (s, 1H), 9.31 (d, J = 1.5 Hz, 1H), 8.94 (d, J = 7.3 Hz, 1H), 8.13 (d, J = 1.7 Hz, 2H), 8.07 (d, J = 2.3 Hz, 1H), 7.90 (d, J = 9.2 Hz, 1H), 7.87 (d, J = 8.5 Hz, 1H), 7.61 (dd, J = 9.1, 2.6 Hz, 1H), 7.41 (d, J = 1.7 Hz, 1H), 4.00 (s, 3H), 2.72 (s, 3H).	DMSO	411.0, 413.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

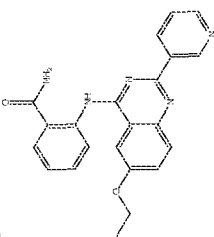
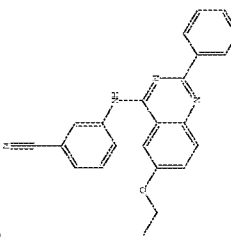
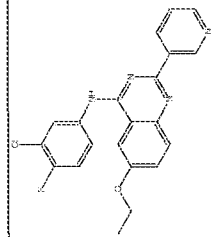
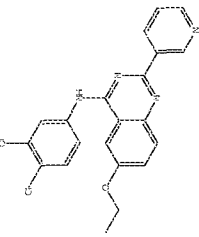
50		HCl	389.84	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.09 (s, 1H), 9.38 (s, 1H), 9.16 (d, J = 8.4 Hz, 1H), 8.83 (d, J = 8.3 Hz, 1H), 8.50 (s, 1H), 8.20 (d, J = 8.8 Hz, 1H), 8.05 – 7.99 (m, 2H), 7.97 (d, J = 7.9 Hz, 1H), 7.92 (s, 1H), 7.81 (dd, J = 8.8, 2.0 Hz, 1H), 7.68 (t, J = 7.8 Hz, 1H), 7.28 (t, J = 7.6 Hz, 1H), 2.82 (s, 3H).	DMSO	390.1, 392.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
51		HCl	412.82	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.49 (s, 1H), 9.35 (d, J = 1.7 Hz, 1H), 9.08 (dd, J = 8.3, 1.5 Hz, 1H), 8.75 (d, J = 9.0 Hz, 1H), 8.02 (d, J = 2.1 Hz, 1H), 7.97 (d, J = 8.3 Hz, 1H), 7.87 (s, 1H), 7.80 (dd, J = 9.0, 2.0 Hz, 2H), 7.53 (t, J = 8.2 Hz, 1H), 7.31 (t, J = 7.4 Hz, 1H), 7.06 (dd, J = 8.1, 2.0 Hz, 1H), 2.79 (s, 3H).	DMSO	413.0, 415.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
52			399.25	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.49 (s, 1H), 9.32 (d, J = 1.7 Hz, 1H), 9.03 (d, J = 8.4 Hz, 1H), 8.69 (d, J = 8.9 Hz, 1H), 8.20 (dd, J = 6.8, 2.6 Hz, 1H), 8.00 (d, J = 2.1 Hz, 1H), 7.96 – 7.86 (m, 2H), 7.80 (dd, J = 8.9, 2.1 Hz, 1H), 7.53 (t, J = 9.1 Hz, 1H), 2.76 (s, 3H).	DMSO	399.0, 401.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
53		HCl	382.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.55 (s, 1H), 9.32 (d, J = 1.7 Hz, 1H), 9.07 (dd, J = 8.2, 1.2 Hz, 1H), 8.73 (d, J = 8.9 Hz, 1H), 8.08 (ddd, J = 13.0, 7.5, 2.6 Hz, 1H), 8.00 (dd, J = 8.9, 5.2 Hz, 2H), 7.80 (dd, J = 8.9, 2.1 Hz, 1H), 7.71 (dd, J = 6.1, 2.9 Hz, 1H), 7.54 (dd, J = 19.7, 9.2 Hz, 1H), 2.78 (s, 3H).	DMSO	383.0, 385.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

54		HCl	415.7	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.63 (s, 1H), 9.30 (d, J = 1.7 Hz, 1H), 9.07 (dd, J = 6.3, 1.8 Hz, 1H), 8.75 (d, J = 9.0 Hz, 1H), 8.29 (d, J = 2.4 Hz, 1H), 8.01 (d, J = 8.4 Hz, 1H), 7.99 (d, J = 2.1 Hz, 1H), 7.96 (dd, J = 8.9, 2.5 Hz, 1H), 7.77 (dd, J = 8.9, 2.1 Hz, 1H), 7.70 (d, J = 8.8 Hz, 1H), 2.81 (s, 3H).	DMSO	415.0, 417.0, 419.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
55		HCl	415.7	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.54 (s, 1H), 9.32 (s, 1H), 9.01 (dd, J = 8.0, 1.8 Hz, 1H), 8.72 (d, J = 9.0 Hz, 1H), 8.11 (d, J = 1.8 Hz, 2H), 8.01 (d, J = 2.0 Hz, 1H), 7.97 (d, J = 8.5 Hz, 1H), 7.80 (dd, J = 8.9, 2.0 Hz, 1H), 2.78 (s, 3H).	DMSO	415.0, 417.0, 419.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
56			379.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.16 (s, 1H), 9.47 (s, 1H), 8.79 (dd, J = 8.9, 7.1 Hz, 2H), 7.91 (s, 1H), 7.84 – 7.77 (m, 2H), 7.76 – 7.68 (m, 1H), 7.56 (d, J = 2.1 Hz, 1H), 7.51 (t, J = 8.2 Hz, 1H), 7.39 (dd, J = 8.9, 2.2 Hz, 1H), 7.29 (t, J = 74.0 Hz, 1H), 7.00 (dd, J = 6.1, 2.1 Hz, 1H), 4.23 (brs, 2H).	DMSO	380.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
57		HCl	463.48	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.43 – 10.20 (m, 2H), 9.72 (s, 1H), 8.99 – 8.67 (m, 3H), 8.29 – 8.21 (m, 1H), 7.97 (s, 1H), 7.87 – 7.80 (m, 1H), 7.81 – 7.73 (m, 1H), 7.63 – 7.46 (m, 2H), 7.30 (t, J = 74.0 Hz, 1H), 7.08 – 6.98 (m, 1H), 5.00 (s, 2H), 4.05 – 3.65 (m, 4H), 3.39 – 3.36 (m, 4H).	DMSO	464.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

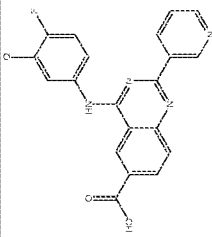
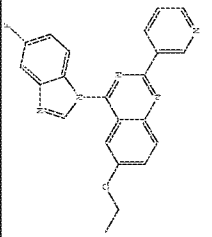
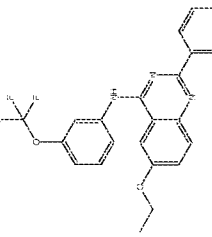
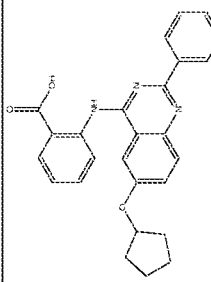
58		HCl	399.45	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.53 (s, 1H), 9.30 (d, J = 1.6 Hz, 1H), 9.03 – 8.96 (m, 1H), 8.94 – 8.88 (m, 1H), 8.71 (d, J = 8.4 Hz, 1H), 7.91 – 7.83 (m, 3H), 7.67 – 7.55 (m, 3H), 7.25 (t, J = 7.2 Hz, 1H), 3.98 (s, 3H), 2.80 – 2.74 (m, 6H).	DMSO	400.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
59		HCl	389.38	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.27 (s, 1H), 9.23 (s, 1H), 8.98 (d, J = 5.2 Hz, 1H), 8.91 (d, J = 8.0 Hz, 1H), 8.54 (dd, J = 12.0, 2.6 Hz, 1H), 8.48 (s, 1H), 8.07 (dd, J = 8.0, 5.6 Hz, 1H), 7.97 – 7.93 (m, 2H), 7.66 (d, J = 9.0 Hz, 1H), 7.37 (dd, J = 9.0, 2.4 Hz, 1H), 7.13 (d, J = 2.0 Hz, 1H), 6.91 (dt, J = 8.4, 2.6 Hz, 1H), 3.80 (s, 3H).	DMSO	390.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
60			415.70	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.13 (s, 1H), 9.48 (s, 1H), 8.87 (dd, J = 14.0, 6.2 Hz, 2H), 8.54 (d, J = 1.7 Hz, 1H), 8.10 (d, J = 1.8 Hz, 2H), 7.89 – 7.78 (m, 2H), 7.37 (t, J = 1.8 Hz, 1H), 2.70 (s, 3H).	DMSO	414.9, 416.9, 419.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

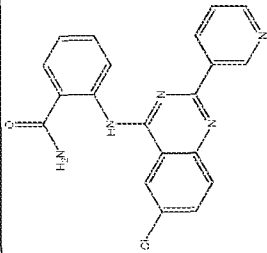
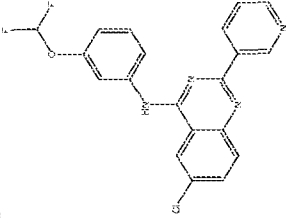
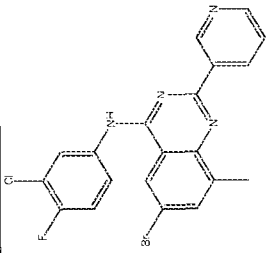
61			415.70	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.10 (s, 1H), 9.44 (s, 1H), 8.86 (d, J = 7.8 Hz, 1H), 8.79 (d, J = 4.5 Hz, 1H), 8.50 (d, J = 1.7 Hz, 1H), 8.23 (d, J = 2.4 Hz, 1H), 7.89 (dd, J = 8.8, 2.4 Hz, 1H), 7.83 – 7.71 (m, 2H), 7.62 (d, J = 8.8 Hz, 1H), 2.62 (s, 3H).	DMSO	414.9, 416.9, 418.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
62			412.82	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.02 (s, 1H), 9.58 (d, J = 1.8 Hz, 1H), 8.76 – 8.67 (m, 2H), 8.59 (d, J = 1.8 Hz, 1H), 7.96 (s, 1H), 7.82 (d, J = 6.6 Hz, 2H), 7.54 (dd, J = 16.4, 10.0, 6.4 Hz, 2H), 7.30 (t, J = 74.0 Hz, 1H), 7.01 (dd, J = 8.0, 2.0 Hz, 1H), 2.74 (s, 3H).	DMSO	413.0, 415.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
63			389.84	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.15 (s, 1H), 9.58 (s, 1H), 9.03 (d, J = 0.8 Hz, 1H), 8.78 – 8.84 (m, 2H), 8.50 (s, 1H), 7.95 – 8.03 (m, 3H), 7.80 (s, 1H), 7.70 – 7.74 (m, 2H), 7.23 (d, J = 7.2 Hz, 1H), 2.70 (s, 3H).	DMSO	391.0, 392.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
64			371.82	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.31 (s, 1H), 9.48 (d, J = 1.2 Hz, 1H), 9.01 (d, J = 8.4 Hz, 1H), 8.90 (d, J = 4.0 Hz, 1H), 8.61 (d, J = 1.6 Hz, 1H), 8.33 (s, 1H), 8.27 (dd, J = 6.0, 2.0 Hz, 1H), 7.95 – 7.63 (m, 3H), 2.70 (s, 3H).	DMSO	372.0, 373.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

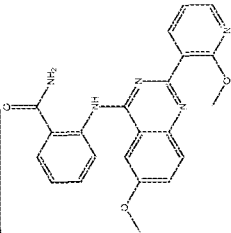
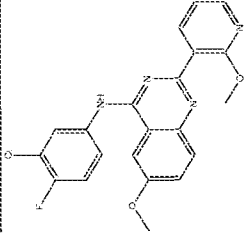
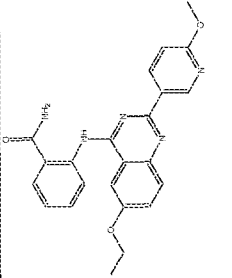
69		HCl	422.43	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.63 (s, 1H), 9.51 (s, 1H), 9.09 (d, J = 7.6 Hz, 1H), 8.91 (d, J = 7.6 Hz, 1H), 8.23 (s, 1H), 8.03 – 7.97 (m, 1H), 7.83 – 7.80 (m, 2H), 7.64 (dd, J = 9.2, 2.4 Hz, 1H), 7.55 (t, J = 8.1 Hz, 1H), 7.32 (t, J = 7.4 Hz, 1H), 7.08 (d, J = 7.4 Hz, 1H), 5.02 – 4.97 (m, 1H), 1.39 (d, J = 6.0 Hz, 6H).	DMSO	423.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
70		HCl	408.86	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.61 (s, 1H), 9.48 (s, 1H), 9.05 (d, J = 7.2 Hz, 1H), 8.93 (d, J = 5.2 Hz, 1H), 8.19 – 8.15 (m, 2H), 8.01 – 7.92 (m, 3H), 7.63 – 7.53 (m, 2H), 5.01 – 4.94 (m, 1H), 1.39 (d, J = 6.0 Hz, 6H).	DMSO	409.1, 411.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
71		HCl	425.31	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.92 (s, 1H), 9.52 (d, J = 1.2 Hz, 1H), 8.70 – 8.66 (m, 2H), 8.17 (d, J = 1.6 Hz, 2H), 7.98 (d, J = 2.4 Hz, 1H), 7.88 (d, J = 8.8 Hz, 1H), 7.60 – 7.56 (m, 2H), 7.39 (t, J = 1.8 Hz, 1H), 4.94 – 4.88 (m, 1H), 1.39 (d, J = 6.0 Hz, 6H).	DMSO	424.9, 426.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
72		HCl	392.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.28 (s, 1H), 9.49 (d, J = 1.2 Hz, 1H), 8.93 (d, J = 8.0 Hz, 1H), 8.86 – 8.84 (m, 1H), 8.10 – 8.05 (m, 2H), 7.93 – 7.85 (m, 2H), 7.71 – 7.69 (m, 2H), 7.61 – 7.52 (m, 1H), 4.97 – 4.91 (m, 1H), 1.39 (d, J = 6.0 Hz, 6H).	DMSO	393.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

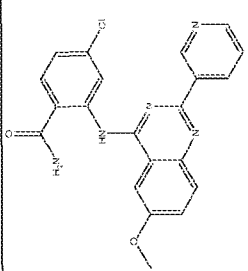
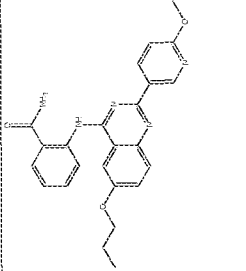
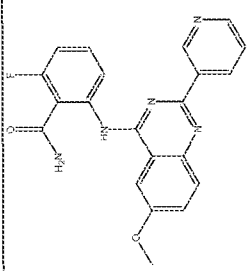
73		HCl	395.42	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.03 (s, 1H), 9.53 (d, J = 1.2 Hz, 1H), 9.10 (d, J = 8.4 Hz, 1H), 8.93 – 8.90 (m, 2H), 8.49 (s, 1H), 8.01 – 7.91 (m, 4H), 7.69 (t, J = 7.6 Hz, 1H), 7.59 – 7.54 (m, 2H), 7.24 (t, J = 7.6 Hz, 1H), 4.21 (q, J = 6.8 Hz, 2H), 1.46 (t, J = 6.9 Hz, 3H).	DMSO	386.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
74		HCl	367.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.35 (s, 1H), 9.49 (s, 1H), 8.95 (d, J = 7.6 Hz, 1H), 8.85 (d, J = 4.8 Hz, 1H), 8.37 (s, 1H), 8.28 (d, J = 7.6 Hz, 1H), 8.08 (s, 1H), 7.94 – 7.85 (m, 2H), 7.73 – 7.60 (m, 3H), 4.23 (q, J = 6.8 Hz, 2H), 1.46 (t, J = 6.8 Hz, 3H).	DMSO	388.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
75		HCl	394.83	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.64 (s, 1H), 9.44 (s, 1H), 8.99 (d, J = 8.4 Hz, 1H), 8.91 (d, J = 4.4 Hz, 1H), 8.17 (d, J = 3.2 Hz, 2H), 7.59 – 7.50 (m, 2H), 4.26 (q, J = 6.8 Hz, 2H), 1.44 (t, J = 6.8 Hz, 3H).	DMSO	395.1 397.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
76		HCl	411.28	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.48 (s, 1H), 9.47 (s, 1H), 9.02 (d, J = 7.6 Hz, 1H), 8.91 (d, J = 4.4 Hz, 1H), 8.28 (d, J = 1.6 Hz, 1H), 8.13 (s, 1H), 8.01 – 7.91 (m, 3H), 7.72 (d, J = 8.8 Hz, 1H), 7.59 – 7.57 (m, 1H), 4.26 (q, J = 6.8 Hz, 2H), 1.44 (t, J = 6.8 Hz, 3H).	DMSO	411.0 413.0 415.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

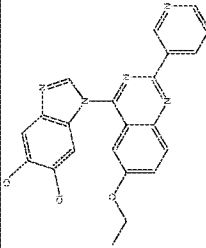
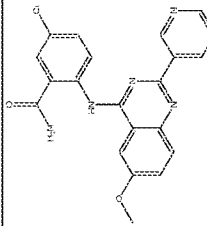
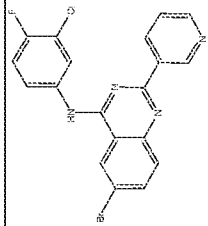
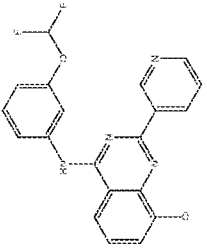
77		HCl	411.28	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.23 (s, 1H), 9.48 (s, 1H), 8.93 (d, J = 8.0 Hz, 1H), 8.85 (d, J = 4.8 Hz, 1H), 8.14 (s, 2H), 8.05 (d, J = 1.6 Hz, 1H), 7.92 – 7.85 (m, 2H), 7.61 (dd, J = 9.1, 1.8 Hz, 1H), 7.41 (s, 1H), 4.26 (q, J = 6.8 Hz, 2H), 1.45 (t, J = 6.8 Hz, 3H).	DMSO	411.0 413.1 415.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
78		HCl	378.37	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.39 (s, 1H), 9.49 (s, 1H), 8.98 (d, J = 8.0 Hz, 1H), 8.88 (d, J = 4.8 Hz, 1H), 8.11 – 8.05 (m, 2H), 7.95 – 7.91 (m, 2H), 7.72 – 7.52 (m, 3H), 4.28 (q, J = 6.8 Hz, 2H), 1.45 (t, J = 6.8 Hz, 3H).	DMSO	379.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
79		HCl	429.47	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.99 (s, 1H), 9.59 (d, J = 1.6 Hz, 1H), 9.14 (d, J = 7.6 Hz, 1H), 8.75 – 8.69 (m, 2H), 8.48 (s, 1H), 7.99 – 7.96 (m, 2H), 7.90 (d, J = 8.8 Hz, 1H), 7.76 – 7.72 (m, 1H), 7.61 – 7.56 (m, 3H), 7.22 – 7.19 (m, 1H), 4.24 (t, J = 6.0 Hz, 2H), 3.56 (t, J = 6.0 Hz, 2H), 3.28 (s, 3H), 2.11 – 2.04 (m, 2H).	DMSO	430.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
80		HCl	380.80	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.67 (s, 1H), 9.18 (d, J = 2.0 Hz, 1H), 8.90 (d, J = 8.4 Hz, 1H), 8.66 (dd, J = 8.8, 2.4 Hz, 1H), 8.35 (d, J = 8.4 Hz, 1H), 8.14 – 8.08 (m, 2H), 7.90 – 7.81 (m, 2H), 7.62 (t, J = 9.2 Hz, 1H), 7.09 (d, J = 8.8 Hz, 1H), 3.98 (s, 3H).	DMSO	380.9, 382.9 (M+1)	Method A (TFA)	95	Method C, G1

61		HCl	394.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.81 (s, 1H), 9.49 (d, J = 1.2 Hz, 1H), 9.28 (d, J = 0.8 Hz, 1H), 9.15 (d, J = 6.0 Hz, 1H), 9.02 (d, J = 5.2 Hz, 1H), 8.37 – 8.34 (m, 1H), 8.12 – 8.09 (m, 2H), 8.01 (d, J = 8.4 Hz, 1H), 7.95 – 7.91 (m, 1H), 7.53 (t, J = 9.0 Hz, 1H).	DMSO	395.0, 397.0 (M+1)	Method B (NH ₄ HCO ₂)	95	Method C, G1
82			385.39	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.63 (d, J = 1.6 Hz, 1H), 9.13 (s, 1H), 8.80 – 8.75 (m, 2H), 8.21 (d, J = 9.2 Hz, 1H), 7.97 – 7.94 (m, 1H), 7.82 (dd, J = 9.2, 2.8 Hz, 1H), 7.73 (dd, J = 9.2, 2.4 Hz, 1H), 7.64 – 7.61 (m, 1H), 7.38 – 7.32 (m, 2H), 4.18 (q, J = 6.8 Hz, 2H), 3.19 (t, J = 7.0 Hz, 3H).	DMSO	386.1 (M+1)	Method B (NH ₄ HCO ₂)	95	Method C, G10
83		HCl	426.39	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.43 (s, 1H), 9.49 (d, J = 1.6 Hz, 1H), 8.99 (d, J = 8.0 Hz, 1H), 8.89 (dd, J = 9.2, 1.6 Hz, 1H), 8.13 (d, J = 2.4 Hz, 1H), 8.05 (s, 1H), 7.97 – 7.89 (m, 3H), 7.65 – 7.61 (m, 2H), 7.23 (d, J = 8.4 Hz, 1H), 4.29 (q, J = 6.8 Hz, 2H), 1.46 (t, J = 7.2 Hz, 3H).	DMSO	427.1 (M+1)	Method B (NH ₄ HCO ₂)	95	Method C, G1
84			426.47	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.00 (s, 1H), 9.55 (s, 1H), 9.12 (d, J = 8.0 Hz, 1H), 8.94 (d, J = 4.8 Hz, 1H), 8.84 (d, J = 8.4 Hz, 1H), 8.11 – 8.09 (m, 1H), 8.02 – 7.99 (m, 1H), 7.96 (d, J = 9.2 Hz, 1H), 7.82 – 7.11 (m, 1H), 7.65 (s, 1H), 7.62 – 7.59 (m, 1H), 7.30 (t, J = 7.2 Hz, 1H), 5.06 – 5.03 (m, 1H), 2.15 – 2.09 (m, 2H), 1.86 – 1.74 (m, 4H), 1.68 – 1.65 (m, 2H).	DMSO	427.0 (M+1)	Method B (NH ₄ HCO ₂)	95	Method C, G1

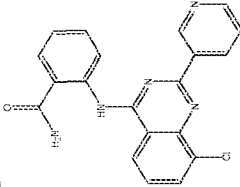
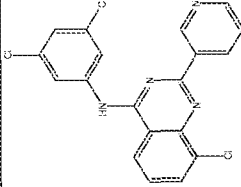
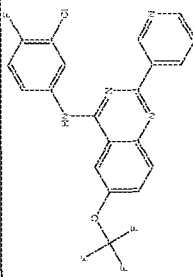
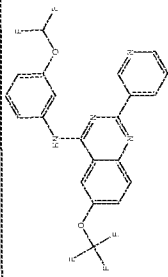
85		375.81	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.09 (s, 1H), 9.58 (s, 1H), 9.12 (d, J = 8.4 Hz, 1H), 8.94 (d, J = 4.4 Hz, 1H), 8.88 (d, J = 8.4 Hz, 1H), 8.50 (s, 1H), 8.25 (s, 1H), 8.01 – 7.96 (m, 5H), 7.71 (t, J = 8.0 Hz, 1H), 7.27 (t, J = 8.0 Hz, 1H).	DMSO	376.0, 378.0 (M+1), 397.9 (M+23), 188.4 (M/2+1)	Method A (TFA)	95	Method C, G1
86		398.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.10 (s, 1H), 9.54 (s, 1H), 8.78 (s, 1H), 8.71 – 8.67 (m, 2H), 7.96 (s, 1H), 7.91 (d, J = 8.0 Hz, 2H), 7.82 (d, J = 8.0 Hz, 1H), 7.56 – 7.51 (m, 2H), 7.30 (t, J = 74.0 Hz, 1H), 7.01 (dd, J = 2.0, 8.0 Hz, 1H).	DMSO	399.0, 401.0 (M+1), 199.9, 200.8 (M/2+1)	Method A (TFA)	95	Method C, G1
87		443.7	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.09 (s, 1H), 9.39 (s, 1H), 8.93 (d, J = 8.4 Hz, 1H), 8.90 (d, J = 5.6 Hz, 1H), 8.62 (d, J = 1.6 Hz, 1H), 8.11 (dd, J = 6.8, 2.4 Hz, 1H), 7.95 – 7.91 (m, 1H), 7.89 – 7.85 (m, 1H), 7.80 (s, 1H), 7.45 (t, J = 8.8 Hz, 1H), 2.66 (s, 3H).	DMSO	443.0, 445.0, 447.0 (M+1)	Method B (NH ₄ HCO ₃)	98	Method C, G1

88			401.42	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.19 (s, 1H), 9.43 (d, J = 8.0 Hz, 1H), 8.53 (s, 1H), 8.31 (dd, J = 4.8, 1.5 Hz, 1H), 8.26 (dd, J = 1.6, 7.2 Hz, 1H), 8.03 (s, 1H), 7.96 – 7.89 (m, 1H), 7.86 (d, J = 9.2 Hz, 1H), 7.65 – 7.54 (m, 3H), 7.18 – 7.12 (m, 2H), 4.02 (s, 3H), 3.98 (s, 3H).	DMSO	402.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
89			410.83	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.79 (s, 1H), 8.52 (dd, J = 2.4, 7.2 Hz, 1H), 8.28 (dd, J = 5.6, 2.4 Hz, 1H), 8.12 (dd, J = 2.4, 5.2 Hz, 1H), 7.98 – 7.94 (m, 2H), 7.82 (d, J = 9.2 Hz, 1H), 7.56 (dd, J = 9.2, 2.8 Hz, 1H), 7.47 (t, J = 9.2 Hz, 1H), 7.14 – 7.11 (m, 1H), 4.00 (s, 3H), 3.99 (s, 3H).	DMSO	411.1, 413.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
90			415.44	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.03 (s, 1H), 9.21 (d, J = 2.2 Hz, 1H), 9.16 (d, J = 7.9 Hz, 1H), 8.67 (dd, J = 8.7, 2.4 Hz, 1H), 8.50 (s, 1H), 8.02 (s, 1H), 7.99 – 7.93 (m, 1H), 7.84 (d, J = 8.9 Hz, 1H), 7.78 – 7.72 (m, 1H), 7.56 (dd, J = 11.2, 2.1 Hz, 2H), 7.19 (t, J = 7.2 Hz, 1H), 6.99 (d, J = 8.8 Hz, 1H), 4.24 (q, J = 6.9 Hz, 2H), 3.96 (s, 3H), 1.46 (t, J = 6.9 Hz, 3H).	DMSO	416.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

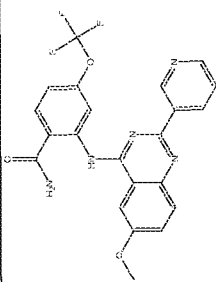
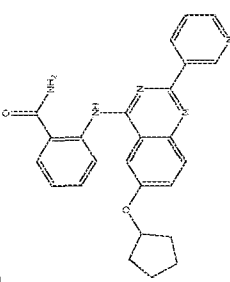
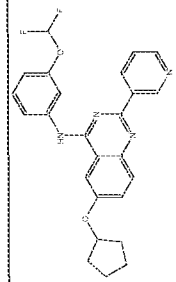
91		HCl	405.84	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.28 (s, 1H), 9.55 (d, J = 1.7 Hz, 1H), 9.31 (d, J = 2.1 Hz, 1H), 8.90 (d, J = 8.3 Hz, 1H), 8.83 (dd, J = 5.0, 1.4 Hz, 1H), 8.58 (s, 1H), 8.12 (s, 1H), 8.01 (d, J = 8.6 Hz, 1H), 7.93 (d, J = 9.1 Hz, 1H), 7.82 (dd, J = 7.9, 5.0 Hz, 1H), 7.63 (dd, J = 9.1, 2.6 Hz, 1H), 7.54 (d, J = 2.5 Hz, 1H), 7.30 (dd, J = 8.5, 2.2 Hz, 1H), 3.98 (s, 3H).	DMSO	406.0, 408.0 (M+1)	Method A (TFA)	95	Method C, G2
92			429.47	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.99 (s, 1H), 9.21 (d, J = 2.0 Hz, 1H), 9.14 (d, J = 8.4 Hz, 1H), 8.67 (dd, J = 8.7, 2.3 Hz, 1H), 8.49 (s, 1H), 8.05 – 7.90 (m, 2H), 7.85 (d, J = 9.0 Hz, 1H), 7.75 (t, J = 7.2 Hz, 1H), 7.67 – 7.47 (m, 2H), 7.20 (t, J = 7.5 Hz, 1H), 6.99 (d, J = 8.8 Hz, 1H), 4.15 (t, J = 6.3 Hz, 2H), 3.96 (s, 3H), 1.07 (t, J = 7.4 Hz, 3H), 1.86 (dd, J = 14.0, 6.7 Hz, 2H).	DMSO	430.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
93		HCl	389.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.34 (s, 1H), 9.52 (d, J = 1.7 Hz, 1H), 8.99 (d, J = 7.8 Hz, 1H), 8.85 (dd, J = 5.2, 1.3 Hz, 1H), 8.22 (d, J = 8.1 Hz, 1H), 8.11 – 8.00 (m, 2H), 7.95 (d, J = 9.1 Hz, 1H), 7.87 (dd, J = 7.9, 5.3 Hz, 1H), 7.77 – 7.59 (m, 3H), 7.27 – 7.16 (m, 1H), 3.99 (s, 3H).	DMSO	390.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G3

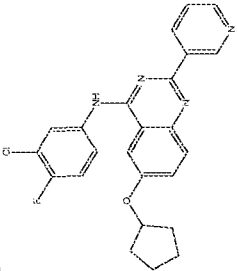
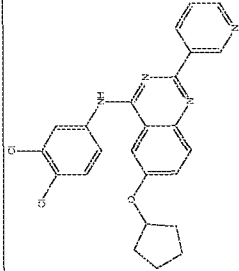
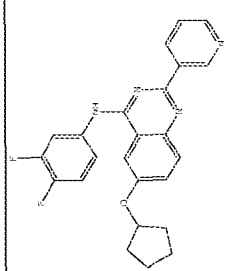
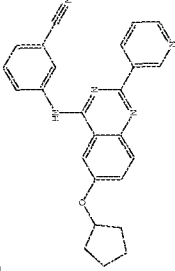
94			436.29	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.62 (s, 1H), 9.17 (s, 1H), 8.77 (d, J = 8.0 Hz, 1H), 8.76 (d, J = 4.0 Hz, 1H), 8.26 – 8.07 (m, 3H), 7.83 (dd, J = 9.1, 2.5 Hz, 1H), 7.63 (dd, J = 7.5, 5.0 Hz, 1H), 7.36 (d, J = 2.4 Hz, 1H), 4.18 (q, J = 6.8 Hz, 2H), 1.39 (t, J = 6.9 Hz, 3H).	DMSO	436.1, 438.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G10
95			405.84	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.93 (s, 1H), 9.58 (s, 1H), 9.16 (d, J = 9.0 Hz, 1H), 8.70 – 8.74 (m, 2H), 8.59 (s, 1H), 8.13 (s, 1H), 8.05 (d, J = 2.3 Hz, 1H), 7.92 (d, J = 9.1 Hz, 1H), 7.82 (dd, J = 8.9, 2.1 Hz, 1H), 7.65 – 7.49 (m, 3H), 3.98 (s, 3H).	DMSO	406.1, 408.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G3
96			429.67	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.45 (s, 1H), 9.47 (s, 1H), 9.02 – 8.92 (m, 3H), 8.18 – 8.16 (m, 1H), 8.06 – 8.04 (m, 1H), 7.96 – 7.84 (m, 3H), 7.54 – 7.50 (m, 1H).	DMSO	429.9, 431.0, 433.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
97			398.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.25 (s, 1H), 9.58 (s, 1H), 8.85 – 8.76 (m, 2H), 8.60 (d, J = 7.2 Hz, 1H), 8.12 – 8.10 (m, 1H), 7.92 (s, 1H), 7.83 – 7.80 (m, 1H), 7.71 – 7.64 (m, 2H), 7.56 – 7.52 (m, 1H), 7.30 (t, J = 74.0 Hz, 1H), 7.05 – 7.03 (m, 1H).	DMSO	399.0, 401.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

66

101			375.09	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.18 (s, 1H), 9.61 (s, 1H), 9.02 (d, J = 8.4 Hz, 1H), 8.93 (d, J = 8.4 Hz, 1H), 8.83 (d, J = 4.0 Hz, 1H), 8.52 (s, 1H), 8.16 – 8.08 (m, 2H), 7.98 – 7.94 (m, 2H), 7.80 – 7.67 (m, 3H), 7.26 (t, J = 7.6 Hz, 1H).	DMSO	376.0, 378.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
102			401.68	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.47 (s, 1H), 9.49 (d, J = 2.0 Hz, 1H), 9.00 – 8.98 (m, 1H), 8.91 (dd, J = 5.2, 1.6 Hz, 1H), 8.64 – 8.62 (m, 1H), 8.12 – 8.09 (m, 3H), 7.94 (dd, J = 8.0, 5.2 Hz, 1H), 7.67 (t, J = 8.0 Hz, 1H), 7.41 (t, J = 2.0 Hz, 1H).	DMSO	400.9, 402.9, 404.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
103			434.77	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.17 (d, J = 1.2 Hz, 1H), 9.51 (s, 1H), 8.71 – 8.62 (m, 3H), 8.25 – 8.22 (m, 1H), 8.04 – 8.00 (m, 1H), 7.92 – 7.87 (m, 2H), 7.58 – 7.53 (m, 2H).	DMSO	435.1, 437.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
104			448.35	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.16 (s, 1H), 9.55 (d, J = 1.2 Hz, 1H), 8.72 – 8.68 (m, 3H), 8.05 (d, J = 9.2 Hz, 1H), 7.92 – 7.95 (m, 2H), 7.80 (dd, J = 8.0, 1.2 Hz, 1H), 7.57 – 7.53 (m, 2H), 7.31 (t, J = 74.4 Hz, 1H), 7.03 (dd, J = 8.0, 1.2 Hz, 1H).	DMSO	448.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

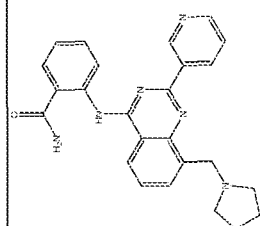
105			425.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.24 (s, 1H), 9.59 (s, 1H), 9.06 (d, J = 8.0 Hz, 1H), 8.74 – 8.73 (m, 2H), 8.54 (s, 1H), 8.10 – 7.91 (m, 5H), 7.77 – 7.73 (m, 1H), 7.61 – 7.58 (m, 1H), 7.26 (t, J = 7.6 Hz, 1H).	DMSO	426.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
106			448.35	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.23 (s, 1H), 9.55 (d, J = 1.6 Hz, 1H), 8.76 (d, J = 9.1 Hz, 1H), 8.74 – 8.67 (m, 2H), 7.94 (t, J = 2.0 Hz, 1H), 7.82 – 7.78 (m, J = 10.3 Hz, 2H), 7.68 (dd, J = 9.0, 1.6 Hz, 1H), 7.59 – 7.50 (m, 2H), 7.30 (t, J = 74.0 Hz, 1H), 7.03 (dd, J = 8.2, 2.2 Hz, 1H).	DMSO	449.0 (M+1) 225.0 (M/2+1)	Method A (TFA)	95	Method C, G1
107			425.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.17 (s, 1H), 9.59 (s, 1H), 9.04 (d, J = 8.0 Hz, 1H), 8.74 – 8.70 (m, 2H), 8.51 (s, 1H), 8.30 (d, J = 9.0 Hz, 1H), 8.02 – 7.90 (m, 2H), 7.86 – 7.69 (m, 3H), 7.65 – 7.56 (m, 1H), 7.25 (t, J = 9.8 Hz, 1H).	DMSO	426.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
108			434.77	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.26 (s, 1H), 9.52 (s, 1H), 8.81 – 8.59 (m, 3H), 8.24 (dd, J = 6.9, 2.5 Hz, 1H), 7.89 – 7.52 (m, 1H), 7.79 (s, 1H), 7.70 (d, J = 9.1 Hz, 1H), 7.65 – 7.50 (m, 2H).	DMSO	435.0, 437.0 (M+1) 218.1 (M/2+1)	Method A (TFA)	95	Method C, G1

109			455.39	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.35 (s, 1H), 9.58 (s, 1H), 9.27 (s, 1H), 8.83 – 8.70 (m, 2H), 8.62 (s, 1H), 8.18 (s, 1H), 8.11 (d, J = 8.8 Hz, 1H), 7.92 (d, J = 9.1 Hz, 1H), 7.68 – 7.58 (m, 2H), 7.54 (d, J = 2.5 Hz, 1H), 7.19 (dd, J = 8.4, 2.17 Hz, 1H), 3.98 (s, 3H).	DMSO	456.0 (M+1) 228.5 (M/2+1)	Method A (TFA)	95	Method C, J1
110			425.48	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.08 (s, 1H), 9.59 (s, 1H), 9.07 (d, J = 7.6 Hz, 1H), 9.02 (d, J = 8.0 Hz, 1H), 8.89 (d, J = 4.0 Hz, 1H), 8.50 (s, 1H), 7.92 – 7.99 (m, 4H), 7.73 (t, J = 7.6 Hz, 1H), 7.59 – 7.60 (m, 2H), 7.25 (t, J = 7.6 Hz, 1H), 2.12 – 2.15 (m, 2H), 5.02 – 5.03 (m, 1H), 1.67 – 1.86 (m, 6H).	DMSO	426.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
111			448.46	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.26 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 8.95 (d, J = 7.2 Hz, 1H), 8.55 (d, J = 4.4 Hz, 1H), 8.08 (s, 1H), 7.94 (d, J = 8.8 Hz, 1H), 7.85 (d, J = 9.6 Hz, 2H), 7.79 (d, J = 8.8 Hz, 1H), 7.60 – 7.63 (m, 1H), 7.55 (t, J = 8.0 Hz, 1H), 7.31 (t, J = 74.0 Hz, 1H), 7.05 (d, J = 8.4 Hz, 1H), 5.14 (s, 1H), 2.03 – 2.09 (m, 2H), 1.61 – 1.81 (m, 6H).	DMSO	449.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

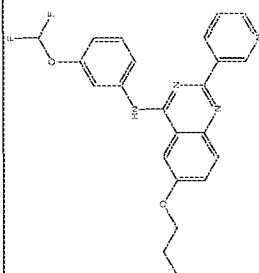
112		434.89	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.47 (s, 1H), 9.47 (s, 1H), 8.97 (d, J = 7.6 Hz, 1H), 8.89 (d, J = 4.4 Hz, 1H), 8.16 – 8.19 (m, 1H), 7.53 – 7.61 (m, 2H), 8.10 (s, 1H), 7.90 – 7.96 (m, 3H), 5.15 (s, 1H), 2.09 – 2.10 (m, 2H), 1.67 – 1.78 (m, 6H).	DMSO	435.0, 437.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
113		451.35	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.23 (s, 1H), 9.49 (s, 1H), 8.85 (d, J = 7.6 Hz, 1H), 8.80 (d, J = 4.0 Hz, 1H), 8.32 (s, 1H), 8.03 (s, 1H), 7.97 – 7.99 (m, 1H), 7.88 (d, J = 8.8 Hz, 1H), 7.77 – 7.79 (m, 1H), 7.75 (d, J = 8.8 Hz, 1H), 7.55 – 7.58 (m, 1H), 5.13 (s, 1H), 2.08 (d, J = 8.4 Hz, 2H), 1.79 (d, J = 9.6 Hz, 4H).	DMSO	451.0, 453.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
114		418.44	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.24 (s, 1H), 9.49 (s, 1H), 8.92 (d, J = 6.0 Hz, 1H), 8.40 (d, J = 6.4 Hz, 1H), 8.04 – 8.11 (m, 2H), 7.92 (d, J = 9.2 Hz, 1H), 7.94 – 7.91 (m, 1H), 7.69 – 7.71 (m, 1H), 7.52 – 7.61 (m, 2H), 5.13 (s, 1H), 2.06 – 2.11 (m, 2H), 1.65 – 1.81 (m, 6H).	DMSO	419.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
115		407.47	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.70 (s, 1H), 9.45 (s, 1H), 8.98 (d, J = 6.8 Hz, 1H), 8.89 (d, J = 6.0 Hz, 1H), 8.37 (s, 1H), 8.32 (d, J = 6.8 Hz, 1H), 8.18 (s, 1H), 7.93 (t, J = 9.2 Hz, 2H), 7.70 (t, J = 7.2 Hz, 2H), 7.56 (d, J = 6.8 Hz, 1H), 5.19 (s, 1H), 2.09 (s, 2H), 1.77 (m, 4H), 1.66 (m, 2H).	DMSO	408.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

116			401.68	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.81 (s, 1H), 9.47 (s, 1H), 8.71 (d, J = 6.4 Hz, 1H), 8.62 (d, J = 7.6 Hz, 1H), 8.26 (s, 1H), 7.88 (t, J = 7.2 Hz, 3H), 7.75 – 7.73 (m, 2H), 7.57 – 7.56 (m, 1H).	DMSO	401.0, 403.0 405.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
117			368.77	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.74 (s, 1H), 9.45 (d, J = 1.2 Hz, 1H), 8.70 (dd, J = 4.4, 1.6 Hz, 1H), 8.60 (d, J = 8.0 Hz, 1H), 8.04 – 8.00 (m, 1H), 7.88 – 7.85 (m, 2H), 7.74 (dd, J = 8.4, 1.6 Hz, 1H), 7.65 – 7.63 (m, 1H), 7.57 – 7.56 (m, 2H).	DMSO	369.1, 371.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
118			436.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.33 (s, 1H), 8.87 – 8.20 (m, 2H), 8.27 (d, J = 1.6 Hz, 1H), 7.99 (d, J = 8.4 Hz, 1H), 7.90 – 7.72 (m, 3H), 7.39 – 7.36 (m, 2H), 3.44 (brs, 4H), 2.05 (brs, 4H).	DMSO	435.9, 437.9, 439.8 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
119			403.43	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.38 (s, 1H), 9.43 (d, J = 1.2 Hz, 1H), 8.85 – 8.80 (m, 2H), 8.07 – 8.05 (m, 1H), 7.90 (d, J = 8.0 Hz, 1H), 7.81 (t, J = 6.8 Hz, 1H), 7.69 – 7.67 (m, 1H), 7.58 – 7.83 (m, 1H), 7.43 – 7.41 (m, 2H), 3.45 (brs, 4H), 2.06 (brs, 4H).	DMSO	403.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

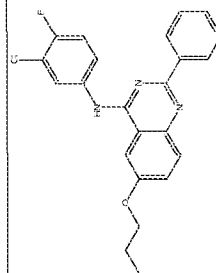
5		Method C, G1					
	95						
10		Method A (TFA)					
	358.0 (M+1)						
15							
	DMSO						
20							
	1H-NMR (400 MHz, DMSO-d6): δ 10.25 (s, 1H), 9.52 (s, 1H), 8.67–8.76 (m, 3H), 8.44 (s, 1H), 8.27 (m, 1H), 7.96 (s, 2H), 7.66–7.69 (m, 2H), 7.57–7.59 (m, 1H).						
25							
	357.8						
30							
35							
40							
45							
50							
55							



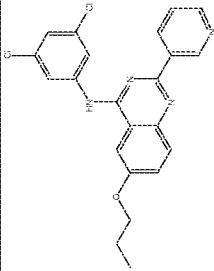
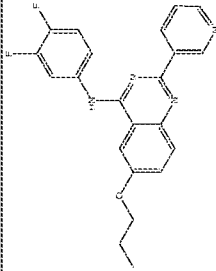
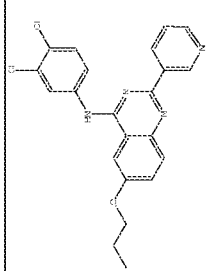
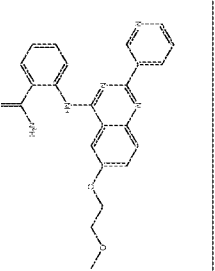
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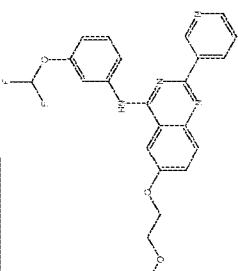
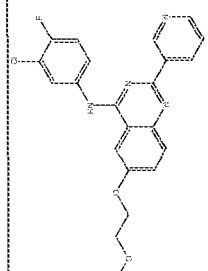
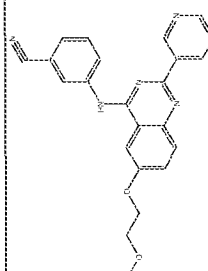
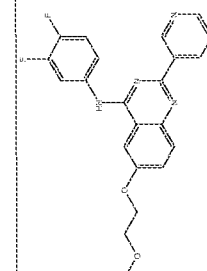


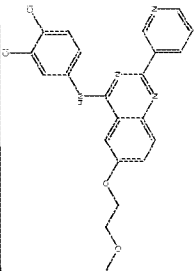
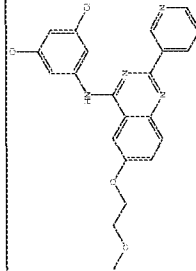
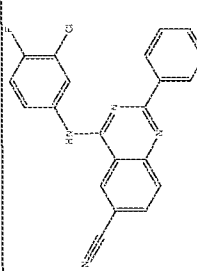
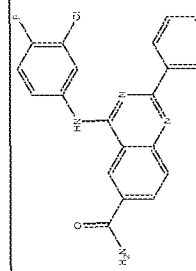
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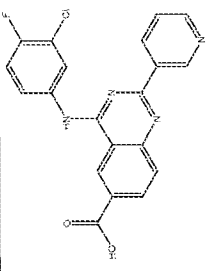
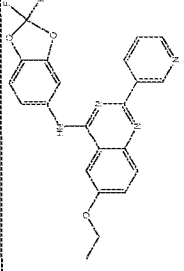
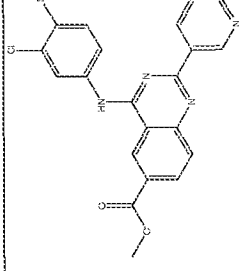
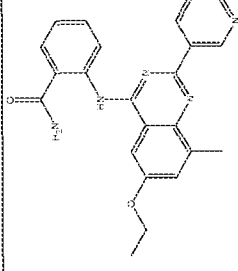


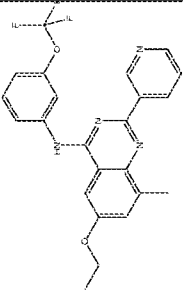
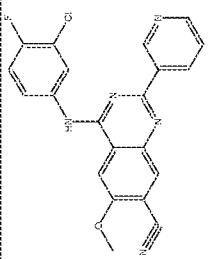
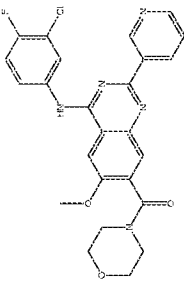
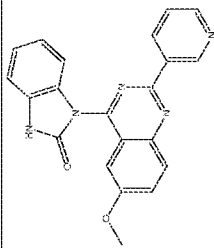
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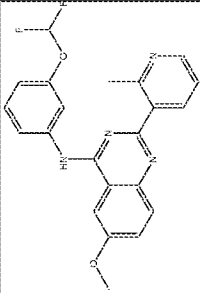
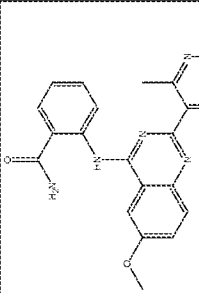
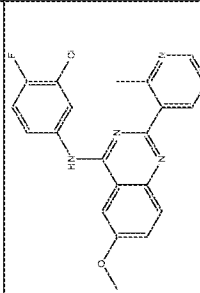
126			364.35	1H-NMR (400 MHz, DMSO-d6): δ 10.65 (s, 1H), 9.47 (s, 1H), 9.02 (d, J = 7.6 Hz, 1H), 8.92 (s, 1H), 8.21 (s, 1H), 8.08 – 8.10 (m, 1H), 7.95 – 7.97 (m, 2H), 7.74 – 7.75 (m, 1H), 7.54 – 7.62 (m, 2H), 4.00 (s, 3H).	DMSO	365.1 (M+1), 183.1 (M/2+1)	Method A (TFA)	95	Method C, G1
127			397.26	1H-NMR (400 MHz, DMSO-d6): δ 10.56 (s, 1H), 9.48 (s, 1H), 9.02 (d, J = 8.0 Hz, 1H), 8.91 – 8.92 (m, 1H), 8.28 (d, J = 2.4 Hz, 1H), 8.16 – 8.17 (m, 1H), 7.92 – 8.03 (m, 3H), 7.72 (d, J = 8.8 Hz, 1H), 7.58 – 7.61 (m, 1H), 4.00 (s, 3H).	DMSO	397.0, 399.0 (M+1), 200.1 (M/2+1)	Method A (TFA)	95	Method C, G1
128			407.45	1H-NMR (400 MHz, DMSO-d6): δ 10.32 (s, 1H), 9.55 (s, 1H), 8.92 – 8.84 (m, 2H), 8.16 – 8.09 (m, 3H), 7.95 – 7.89 (m, 3H), 7.83 – 7.79 (m, 1H), 7.62 – 7.58 (m, 1H), 7.36 (s, 2H), 3.40 (s, 3H).	DMSO	408.1 (M+1)	Method A (TFA)	95	Method C, G1
129			362.81	1H-NMR (400 MHz, DMSO-d6): δ 10.06 (s, 1H), 9.51 (s, 1H), 8.82 – 8.85 (m, 2H), 8.10 – 8.11 (m, 1H), 8.05 (d, J = 2.0 Hz, 1H), 7.98 – 7.93 (m, 2H), 7.75 – 7.77 (m, 1H), 7.59 – 7.61 (m, 1H), 7.52 (t, J = 8.0 Hz, 1H), 7.27 – 7.28 (m, 1H), 3.99 (s, 3H).	DMSO	363.0, 365.0 (M+1), 182.1 (M/2+1)	Method A (TFA)	95	Method C, G1

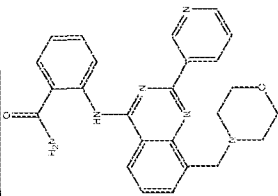
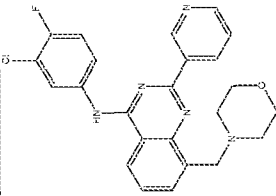
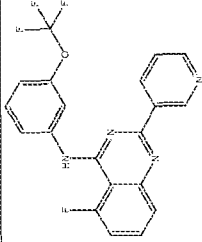
130			380.8	1H-NMR (300 MHz, DMSO-d6): δ 10.02 (s, 1H), 9.50 (s, 1H), 7.78 – 8.79 (m, 2H), 8.19 – 8.24 (m, 1H), 7.86 – 8.00 (m, 3H), 7.52 – 7.70 (m, 3H), 3.98 (s, 3H).	DMSO	381.1, 383.1 (M+1), 191.1 (M/2+1)	Method A (TFA)	95	Method C, G1
131			371.39	1H-NMR (300 MHz, DMSO-d6): δ 13.08 (s, 1H), 9.59 (s, 1H), 9.07 (d, J = 8.1 Hz, 1H), 8.96 (d, J = 8.1 Hz, 1H), 8.93 (d, J = 3.9 Hz, 1H), 8.49 (s, 1H), 7.89 – 7.99 (m, 3H), 7.80 – 7.84 (m, 1H), 7.72 (t, J = 7.8 Hz, 1H), 7.60 – 7.63 (m, 2H), 7.23 (t, J = 7.5 Hz, 1H), 3.98 (s, 3H).	DMSO	371.9 (M+1)	Method A (TFA)	95	Method C, G1
132			362.81	1H-NMR (400 MHz, DMSO-d6): δ 10.12 (s, 1H), 9.28 (s, 1H), 8.69 (d, J = 4.0 Hz, 1H), 8.59 (d, J = 8.4 Hz, 1H), 8.00 (d, J = 2.8 Hz, 1H), 7.87 (d, J = 9.2 Hz, 1H), 7.68 – 7.70 (m, 2H), 7.57 – 7.63 (m, 2H), 7.48 – 7.53 (m, 1H), 7.40 – 7.44 (m, 1H), 3.86 (s, 3H).	DMSO	363.0, 365.1 (M+1), 182.1 (M/2+1)	Method A (TFA)	95	Method C, G1
133			348.35	1H-NMR (400 MHz, DMSO-d6): δ 9.46 (s, 1H), 9.11 (s, 1H), 8.79 – 8.76 (m, 2H), 8.00 – 7.94 (m, 1H), 7.63 – 7.73 (m, 3H), 7.62 – 7.48 (m, 3H), 3.00 (s, 3H).	DMSO	349.1 (M+1), 175.1 (M/2+1)	Method A (TFA)	95	Method C, G1

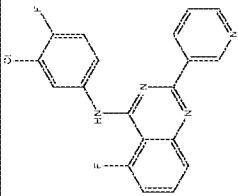
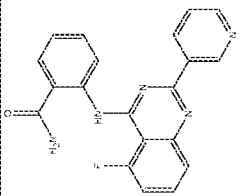
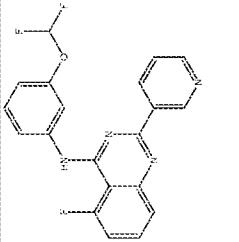
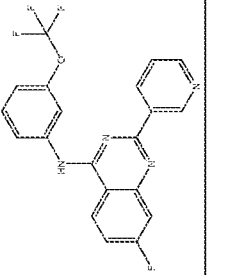
134		381.26	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.48 (s, 1H), 9.13 (s, 1H), 8.75 – 8.77 (m, 2H), 8.18 (s, 1H), 7.72 – 7.85 (m, 5H), 7.49 – 7.50 (m, 1H), 3.00 (s, 3H).	DMSO	381.0, 383.0, 385.0 (M+1)	Method A (TFA)	95	Method C, G1
135		364.8	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.43 (m, 2H), 8.94 – 8.89 (m, 2H), 8.08 – 8.06 (m, 1H), 7.92 – 7.81 (m, 4H), 7.57 – 7.53 (m, 2H), 3.03 (s, 3H).	DMSO	365.0, 367.0 (M+1), 183.1 (M/2+1)	Method A (TFA)	95	Method C, G1
136		337.38	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.52 – 9.44 (m, 2H), 8.98 (d, J = 8.0 Hz, 1H), 8.92 (d, J = 4.4 Hz, 1H), 8.36 – 8.19 (m, 2H), 8.02 – 7.84 (m, 3H), 7.73 – 7.69 (m, 2H), 7.55 (d, J = 6.8 Hz, 1H), 3.04 (s, 3H).	DMSO	338.1 (M+1), 169.6 (M/2+1)	Method A (TFA)	95	Method C, G1
137		364.35	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.97 (s, 1H), 9.51 (s, 1H), 8.69 – 8.66 (m, 2H), 8.16 – 8.08 (m, 2H), 7.75 – 7.73 (m, 1H), 7.61 – 7.53 (m, 3H), 7.40 (d, J = 7.6 Hz, 1H), 4.01 (s, 3H).	DMSO	183.1 (M/2+1), 365.1 (M+1)	Method A (TFA)	95	Method C, G1

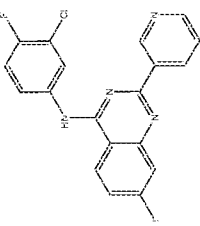
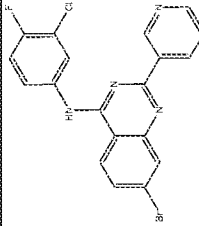
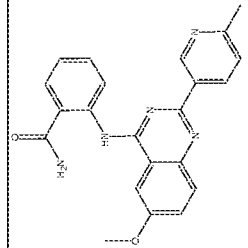
138			397.26	¹H-NMR (400 MHz, DMSO-d₆): δ 10.01 (s, 1H), 9.53 (s, 1H), 8.67–8.70 (m, 2H), 8.41 (d, J = 2.4 Hz, 1H), 8.10 (d, J = 8.4 Hz, 1H), 7.95–7.99 (m, 1H), 7.73 (d, J = 8.8 Hz, 1H), 7.56–7.63 (m, 2H), 7.42 (d, J = 8.4 Hz, 1H), 4.02 (s, 3H).	DMSO	397.0, 399.0 (M+1)	Method A (TFA)	95	Method C, G1
139			380.8	¹H-NMR (400 MHz, DMSO-d₆): δ 10.21 (s, 1H), 9.48 (s, 1H), 8.99 (d, J = 8.0 Hz, 1H), 8.89 (d, J = 4.8 Hz, 1H), 8.15–8.22 (m, 2H), 7.92–7.95 (m, 2H), 7.62 (t, J = 8.0 Hz, 1H), 7.52 (t, J = 8.8 Hz, 1H), 7.43 (d, J = 8.0 Hz, 1H), 4.02 (s, 3H).	DMSO	191.1 (M/2+1) 381.1, 383.1 (M+1)	Method A (TFA)	95	Method C, G1
140			401.68	¹H-NMR (400 MHz, DMSO-d₆): δ 10.11 (s, 1H), 8.50 (m, 1H), 8.63–8.70 (m, 3H), 8.14 (d, J = 2.0 Hz, 2H), 7.92 (m, 2H), 7.57 (m, 1H), 7.38 (t, J = 1.6 Hz, 1H).	DMSO	401.0, 403.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
141			348.35	¹H-NMR (400 MHz, DMSO-d₆): δ 9.46 (s, 1H), 9.11 (s, 1H), 8.76–8.70 (m, 2H), 7.95–7.99 (m, 1H), 7.78–7.60 (m, 3H), 7.54–7.44 (m, 3H), 3.00 (s, 3H).	DMSO	175.1 (M/2+1) 349.1 (M+1)	Method A (TFA)	95	Method C, G1

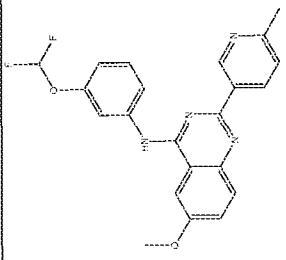
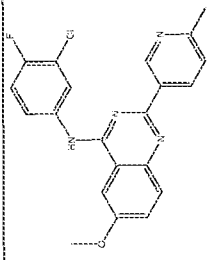
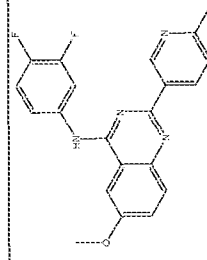
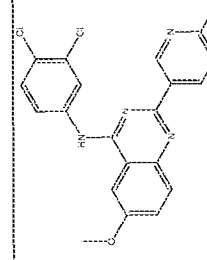
142			397.26	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.30 (s, 1H), 8.39 (s, 1H), 8.83 (s, 2H), 8.03 – 8.12 (m, 3H), 7.79 – 7.81 (m, 2H), 7.50 – 7.51 (m, 1H), 7.32 (s, 1H), 3.96 (s, 3H).	DMSO	397.0, 399.0, 401.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
143			452.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.86 (s, 1H), 9.51 (s, 1H), 8.65 – 8.66 (m, 2H), 8.34 – 8.35 (m, 1H), 7.95 – 7.96 (m, 1H), 7.72 – 7.80 (m, 4H), 7.54 (m, 1H), 3.84 (m, 4H), 3.56 (m, 4H).	DMSO	452.1, 454.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
144			435.88	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.83 (s, 1H), 9.49 (s, 1H), 8.62 – 8.65 (m, 2H), 8.21 – 8.23 (m, 1H), 7.89 – 7.91 (m, 1H), 7.78 – 7.81 (m, 1H), 7.72 – 7.73 (m, 2H), 7.51 – 7.54 (m, 2H), 3.84 (t, J = 4.0 Hz, 4H), 3.36 (m, 4H).	DMSO	436.1, 438.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
145			419.43	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.50 (s, 1H), 9.45 (s, 1H), 8.83 – 8.90 (m, 2H), 8.06 – 8.08 (m, 1H), 7.70-7.93 (m, 5H), 7.56 (dd, J = 19.6, 9.6 Hz, 1H), 3.82 – 3.83 (m, 4H), 3.39 – 3.41 (m, 4H).	DMSO	420.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

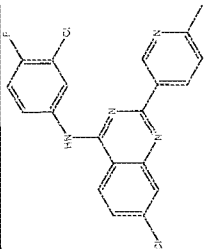
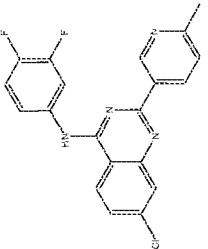
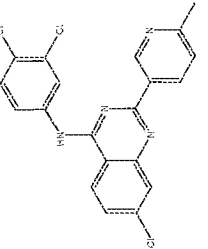
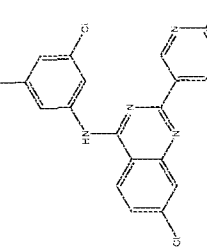
146			408.46	¹H-NMR (400 MHz, DMSO-d₆): δ 10.54 (s, 1H), 9.44 (s, 1H), 8.90 (d, J = 8.0 Hz, 1H), 8.84 (d, J = 4.4 Hz, 1H), 8.34 (s, 1H), 8.27 (d, J = 7.2 Hz, 1H), 7.67 – 7.92 (m, 6H), 3.82 – 3.83 (m, 4H), 3.39 – 3.40 (m, 4H).	DMSO	409.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
147			452.34	¹H-NMR (400 MHz, DMSO-d₆): δ 10.42 (s, 1H), 9.42 (s, 1H), 8.85 – 8.87 (m, 2H), 8.11 (s, 2H), 7.73 – 7.88 (m, 4H), 7.38 (s, 1H), 3.82 – 3.83 (m, 4H), 3.39 – 3.40 (m, 4H).	DMSO	452.1, 454.1, 456.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
148			426.47	¹H-NMR (400 MHz, DMSO-d₆): δ 12.93 (s, 1H), 9.57 (m, 1H), 9.12 (d, J = 8.4 Hz, 1H), 8.68 – 8.72 (m, 2H), 8.56 (s, 1H), 8.02 (d, J = 6.8 Hz, 1H), 7.96 (s, 1H), 7.72 – 7.84 (m, 3H), 7.56 – 7.57 (m, 1H), 7.40 (m, 1H), 7.19 (t, J = 8.0 Hz, 1H), 3.84 (t, J = 4.4 Hz, 4H), 3.35 (t, J = 4.4 Hz, 4H).	DMSO	427.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

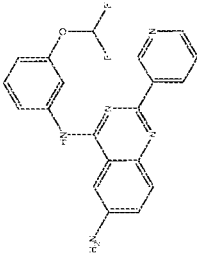
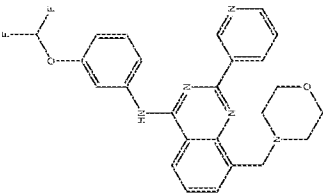
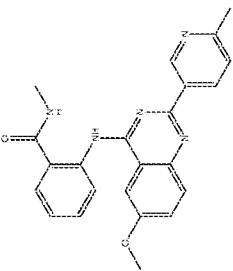
Method C, G1	Method C, G1	Method C, G1
95	95	95
Method B (NH ₄ HCO ₃)	Method B (NH ₄ HCO ₃)	Method B (NH ₄ HCO ₃)
394.1, 396.1 (M+1)	378.2 (M+1)	410.1, 412.0, 414.1 (M+1)
DMSO	DMSO	DMSO
¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.71 (s, 1H), 9.48 (s, 1H), 8.62 – 8.63 (m, 2H), 8.23 (m, 1H), 7.92 (m, 1H), 7.77 (d, J = 9.6 Hz, 1H), 7.51 – 7.56 (m, 3H), 7.42 (m, 1H), 3.12 (s, 6H).	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.72 (s, 1H), 9.48 (m, 1H), 8.60 – 8.63 (m, 2H), 8.08 – 8.14 (m, 1H), 7.77 (d, J = 9.2 Hz, 1H), 7.69 – 7.71 (m, 1H), 7.50 – 7.57 (m, 3H), 7.42 – 7.43 (m, 1H), 3.12 (s, 6H).	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.72 (s, 1H), 9.49 (s, 1H), 8.60 – 8.64 (m, 2H), 8.17 (s, 2H), 7.76 (d, J = 8.8 Hz, 1H), 7.51 – 7.53 (m, 2H), 7.37 (s, 1H), 7.32 (s, 1H), 3.11 (s, 6H).
393.84	377.39	410.3
		
149	150	151

152			410.3	¹H-NMR (400 MHz, DMSO-d₆): δ 10.44 (s, 1H), 9.45 (s, 1H), 8.83 – 8.91 (m, 2H), 8.26 – 8.27 (m, 1H), 7.72 – 7.99 (m, 4H), 7.57 – 7.61 (m, 2H), 3.14 (s, 6H).	DMSO	410.1, 412.1, 414.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
153			366.42	¹H-NMR (400 MHz, DMSO-d₆): δ 9.82 (s, 1H), 9.48 (s, 1H), 8.55 – 8.64 (m, 2H), 8.41 (s, 1H), 8.28 (d, J = 8.0 Hz, 1H), 7.79 (d, J = 9.6 Hz, 1H), 7.69 (t, J = 8.0 Hz, 1H), 7.50 – 7.61 (m, 3H), 7.43 (m, 1H), 3.12 (s, 6H).	DMSO	367.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
154			435.23	¹H-NMR (400 MHz, DMSO-d₆): δ 10.57 (s, 1H), 9.53 (s, 1H), 8.86 – 8.94 (m, 3H), 8.32 – 8.33 (m, 1H), 8.21 (s, 1H), 7.97 – 8.01 (m, 2H), 7.84 – 7.88 (m, 1H), 7.73 (d, J = 8.8 Hz, 1H).	DMSO	435.1, 437.1, 439.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
155			402.32	¹H-NMR (400 MHz, DMSO-d₆): δ 10.32 (s, 1H), 9.51 – 9.52 (m, 1H), 8.65 – 8.78 (m, 3H), 8.19 (s, 1H), 8.11 (m, 1H), 7.97 (d, J = 8.0 Hz, 1H), 7.71 (m, 1H), 7.55 – 7.59 (m, 2H).	DMSO	403.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

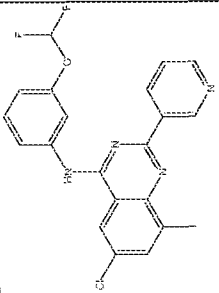
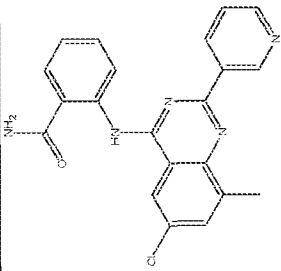
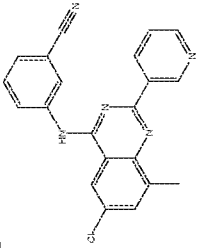
162			375.81	1H-NMR (400 MHz, DMSO-d6): δ 13.13 (s, 1H), 9.53 (s, 1H), 9.01 (d, J = 8.0 Hz, 1H), 8.68 – 8.72 (m, 2H), 8.50 (s, 1H), 8.10 (d, J = 8.8, 1H), 7.88 – 7.97 (m, 3H), 7.70 – 7.72 (m, 2H), 7.57 – 7.59 (m, 1H), 7.22 (t, J = 7.6, 1H).	DMSO	376.0, 378.0 (M+1)	Method B (NH4HCO3)	95	Method C, G1
163			376.8	1H-NMR (400 MHz, DMSO-d6): δ 12.07(s, 1H), 9.56 (s, 1H), 8.84 (d, J = 8.4 Hz, 1H), 8.71 – 8.73 (m, 2H), 8.34 (s, 1H), 7.96 – 8.09(m, 3H), 7.59 – 7.80 (m, 2H), 7.29 (t, J = 7.2 Hz, 1H).	DMSO	377.0, 379.0 (M+1)	Method B (NH4HCO3)	95	Method C, G1
164			394.37	1H-NMR (400 MHz, DMSO-d6): δ 9.90 (s, 1H), 9.53 (s, 1H), 8.67 (d, J = 5.2 Hz, 2H), 7.99 (d, J = 2.4 Hz, 1H), 7.94 (m, 1H), 7.87 (d, J = 9.2 Hz, 1H), 7.80 (d, J = 8.4 Hz, 1H), 7.49 – 7.59 (m, 3H), 7.31 (t, J = 74.0 Hz, 1H), 7.00 (d, J = 8.0 Hz, 1H), 3.99 (s, 3H).	DMSO	395.1 (M+1)	Method B (NH4HCO3)	95	Method C, G1

165		HCl	456.51	¹H-NMR (400 MHz, DMSO-d₆): δ 9.42 (s, 1H), 8.98 (d, J = 8.4 Hz, 1H), 8.68 – 8.71 (m, 2H), 7.94 (d, J = 7.6 Hz, 1H), 7.70 (t, J = 8.0 Hz, 1H), 7.57 (t, J = 5.6 Hz, 1H), 7.53 (s, 1H), 7.38 (s, 1H), 7.19 (t, J = 7.2 Hz, 1H), 4.52 (s, 2H), 4.00 (s, 3H), 3.63 (s, 2H), 2.89 (s, 6H).	DMSO	459.1 (M+1) 230.2 (M/2+1)	Method A (TFA)	95	Method C, G1
166			467.92	¹H-NMR (400 MHz, DMSO-d₆): δ 9.71 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 8.62 – 8.67 (m, 2H), 8.20 (dd, J = 6.8, 2.4 Hz, 1H), 7.86 – 7.89 (m, 2H), 7.51 – 7.55 (m, 2H), 7.31 (s, 1H), 4.24 (t, J = 6.0 Hz, 2H), 3.98 (s, 3H), 2.77 (t, J = 5.6 Hz, 2H), 2.29 (s, 6H).	DMSO	468.1 (M+1) 234.6 (M/2+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
167		HCl	428.82	¹H-NMR (400 MHz, DMSO-d₆): δ 10.41 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 8.98 (d, J = 8.4 Hz, 1H), 8.88 (dd, J = 5.2, 1.6 Hz, 1H), 8.13 (d, J = 2.8 Hz, 1H), 8.00 (t, J = 2.0 Hz, 1H), 7.89 – 7.95 (m, 3H), 7.62 (dd, J = 9.2, 3.2 Hz, 1H), 7.36 (t, J = 73.6 Hz, 1H), 7.16 (d, J = 2.0 Hz, 1H), 4.01 (s, 3H).	DMSO	429.1, 431.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
168			385.42	¹H-NMR (400 MHz, DMSO-d₆): δ 13.03 (s, 1H), 9.62 (d, J = 1.6 Hz, 1H), 9.20 (d, J = 8.4 Hz, 1H), 8.74 – 8.77 (m, 1H), 8.70 (dd, J = 4.4, 1.6 Hz, 1H), 8.48 (s, 1H), 7.96 – 7.99 (m, 2H), 7.72 – 7.76 (m, 1H), 7.57 – 7.60 (m, 1H), 7.47 (m, 1H), 7.41 (m, 1H), 7.17 – 7.21 (m, 1H), 3.95 (s, 3H), 2.72 (s, 3H).	DMSO	386.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

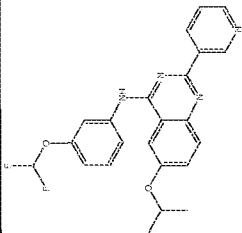
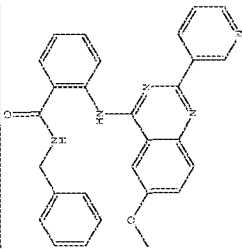
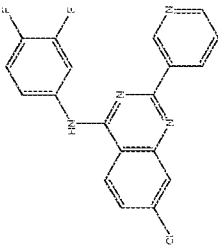
172		HCl	382.34	¹H-NMR (400 MHz, DMSO-d₆): δ 10.30 (s, 1H), 9.53 (d, J = 1.2 Hz, 1H), 8.98 (d, J = 8.0 Hz, 1H), 8.88 (d, J = 4.4 Hz, 1H), 8.62 (dd, J = 10.0, 2.8 Hz, 1H), 8.01 – 8.05 (m, 1H), 7.84 – 7.92 (m, 4H), 7.50 – 7.56 (m, 1H), 7.32 (t, J = 74.0 Hz, 1H), 7.03 (dd, J = 8.0, 2.0 Hz, 1H).	DMSO	383.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
173			348.35	¹H-NMR (400 MHz, DMSO-d₆): δ 10.09 (s, 1H), 9.51 (s, 1H), 8.79 – 8.76 (m, 2H), 8.38 (s, 1H), 8.16 – 8.11 (m, 1H), 7.86 – 7.66 (m, 4H), 7.59 – 7.50 (m, 1H), 2.57 (s, 3H).	DMSO	349.1 (M+1), 175.1 (M/2+1)	Method A (TFA)	95	Method C, G1
174			364.8	¹H-NMR (300 Hz, CD₃OD): δ 9.50 (d, J = 1.2 Hz, 1H), 9.14 (d, J = 8.4 Hz, 1H), 8.94 (d, J = 4.8 Hz, 1H), 8.35 (s, 1H), 8.09 – 7.99 (m, 2H), 7.93 (s, 2H), 7.78 – 7.73 (m, 1H), 7.39 (t, J = 9.0 Hz, 1H), 2.64 (s, 3H).	CD ₃ OD	365.1, 367.1 (M+1)	Method A (TFA)	95	Method C, G1
175			381.26	¹H-NMR (400 MHz, DMSO-d₆): δ 10.13 (s, 1H), 9.51 (s, 1H), 8.84 – 8.79 (m, 2H), 8.36 – 8.34 (m, 2H), 7.96 (dd, J = 8.8, 2.5 Hz, 1H), 7.85 – 7.70 (m, 4H), 2.55 (s, 3H).	DMSO	381.0, 383.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

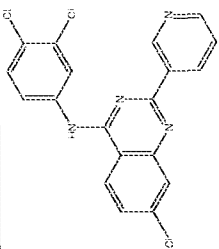
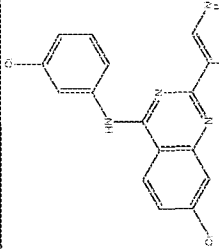
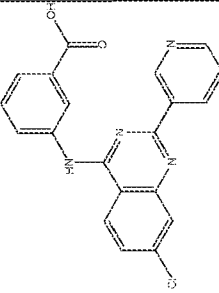
176			346.81	¹H-NMR (300 Hz, CD₃OD): δ 9.48 (d, J = 1.2 Hz, 1H), 8.99 – 8.95 (m, 1H), 8.82 – 8.80 (m, 1H), 8.30 – 8.29 (m, 1H), 7.90 – 7.81 (m, 5H), 7.52 – 7.46 (m, 2H), 2.62 (s, 3H).	CD ₃ OD	347.0, 349.1 (M+1)	Method A (TFA)	95	Method C, G1
177			346.81	¹H-NMR (400 MHz, DMSO-d₆): δ 10.12 (s, 1H), 9.52 (s, 1H), 8.84 – 8.78 (m, 2H), 8.42 (s, 1H), 8.16 – 8.15 (m, 1H), 7.94 – 7.91 (m, 1H), 7.85 (d, J = 8.4 Hz, 1H), 7.79 (d, J = 8.1 Hz, 1H), 7.75 – 7.71 (m, 1H), 7.50 (t, J = 8.1 Hz, 1H), 7.27 – 7.23 (m, 1H), 2.56 (s, 3H).	DMSO	347.1, 349.1 (M+1)	Method A (TFA)	95	Method C, G1
178			360.37	¹H-NMR (400 MHz, DMSO-d₆): δ 10.60 (s, 1H), 9.50 (d, J = 2.1 Hz, 1H), 9.00 (d, J = 8.1 Hz, 1H), 8.90 (dd, J = 5.1, 1.2 Hz, 1H), 8.55 (s, 1H), 8.43 (s, 1H), 8.25 (d, J = 7.8 Hz, 1H), 7.96 – 7.90 (m, 2H), 7.84 (d, J = 8.1 Hz, 1H), 7.27 (t, J = 8.2 Hz, 1H), 7.57 (d, J = 7.8 Hz, 1H), 2.62 (s, 3H).	DMSO	331.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

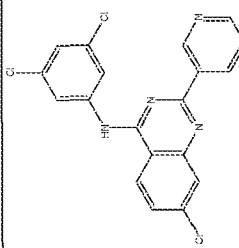
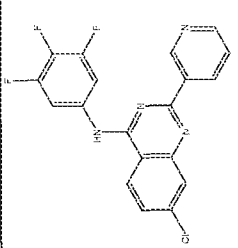
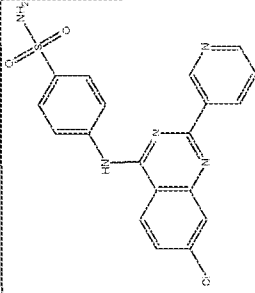
89

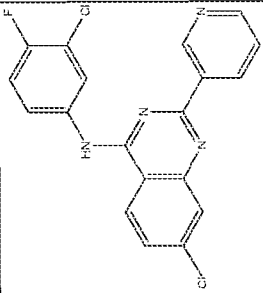
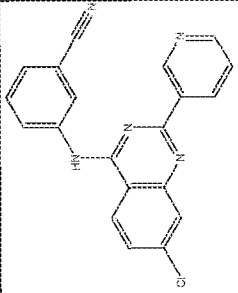
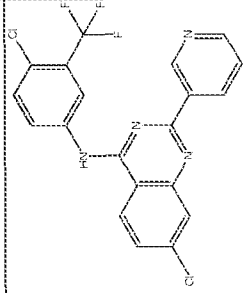
182			390.82	¹H-NMR (400 MHz, DMSO-d₆): δ 12.25 (s, 1H), 9.54 (s, 1H), 9.25 (s, 1H), 8.84 – 8.80 (m, 2H), 8.09 (d, J = 8.0 Hz, 1H), 7.96 (s, 1H), 7.88 (d, J = 8.0 Hz, 1H), 7.81 (d, J = 9.2 Hz, 1H), 7.77 – 7.75 (m, 1H), 7.29 (d, J = 8.4 Hz, 1H), 2.56 (s, 3H).	DMSO	391.1, 393.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
183			366.34	¹H-NMR (400 MHz, DMSO-d₆): δ 9.98 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 8.69 (dd, J = 4.8, 1.6 Hz, 1H), 8.66 – 8.63 (m, 1H), 8.30 (s, 1H), 7.99 – 7.95 (m, 2H), 7.82 (d, J = 8.4 Hz, 1H), 7.76 (dd, J = 8.6, 1.4 Hz, 1H), 7.58 – 7.55 (m, 1H), 2.55 (s, 3H).	DMSO	367.1 (M+1), 184.1 (M/2+1)	Method A (TFA)	95	Method C, G1
184			415.7	¹H-NMR (400 MHz, DMSO-d₆): δ 10.09 (s, 1H), 9.51 (s, 1H), 8.75 – 8.72 (m, 2H), 8.38 (s, 2H), 8.32 (s, 1H), 7.83 (d, J = 8.6 Hz, 1H), 7.77 (d, J = 8.0 Hz, 1H), 7.66 – 7.64 (m, 1H), 2.55 (s, 3H).	DMSO	414.8, 416.8 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

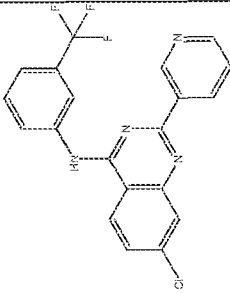
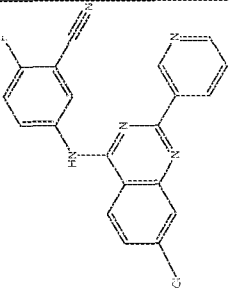
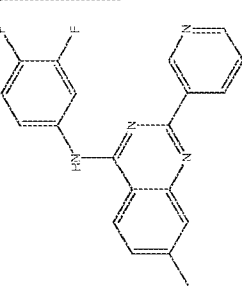
185			354.4	1H-NMR (400 MHz, DMSO-d6): δ 10.26 (s, 1H), 9.56 (s, 1H), 8.93 (d, J = 8.0 Hz, 1H), 8.81 (d, J = 4.0 Hz, 1H), 8.65 (s, 1H), 8.48 (s, 1H), 8.21 (d, J = 8.0 Hz, 1H), 7.86 (d, J = 8.4 Hz, 1H), 7.88 – 7.89 (m, 3H), 7.65 (t, J = 7.6 Hz, 1H), 2.65 (s, 3H), 2.58 (s, 3H).	DMSO	355.1 (M+1), 178.1 (M/2+1)	Method A (TFA)	95	Method C, G1
186			356.38	1H-NMR (400 MHz, DMSO-d6): δ 12.18 (s, 1H), 9.57 (s, 1H), 9.07 (d, J = 8.4 Hz, 1H), 8.71 (d, J = 5.6 Hz, 2H), 8.10 (d, J = 8.0 Hz, 1H), 7.99 (s, 1H), 7.86 – 7.76 (m, 3H), 7.58 (t, J = 6.2 Hz, 1H), 7.23 (t, J = 7.4 Hz, 1H), 2.55 (s, 3H).	DMSO	357.1 (M+1)	Method A (TFA)	95	Method C, G1
187			337.38	1H-NMR (400 MHz, DMSO-d6): δ 10.10 (s, 1H), 9.51 (s, 1H), 8.65 – 8.70 (m, 2H), 8.45 (s, 1H), 8.38 (s, 1H), 8.28 (d, J = 8.0 Hz, 1H), 7.83 (d, J = 8.8 Hz, 1H), 7.76 (d, J = 8.0 Hz, 1H), 7.59 (t, J = 7.8 Hz, 1H), 7.62 (d, J = 7.2 Hz, 1H), 7.54 – 7.57 (m, 1H), 2.56 (s, 3H).	DMSO	338.1 (M+1)	Method B (NH4HCO3)	95	Method C, G1
188			355.39	1H-NMR (400 MHz, DMSO-d6): δ 13.11 (s, 1H), 9.61 (s, 1H), 9.09 (d, J = 8.8 Hz, 1H), 8.89 (d, J = 8.0 Hz, 1H), 8.81 (s, 1H), 8.50 (s, 1H), 8.01 – 7.98 (m, 2H), 7.96 (dd, J = 7.8, 1.0 Hz, 1H), 7.86 (d, J = 8.8 Hz, 1H), 7.78 – 7.72 (m, 3H), 7.23 (t, J = 7.4 Hz, 1H), 2.56 (s, 3H).	DMSO	356.2 (M+1)	Method B (NH4HCO3)	95	Method C, G1

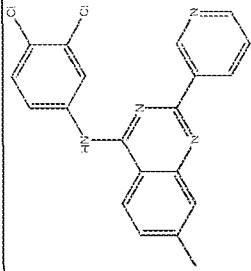
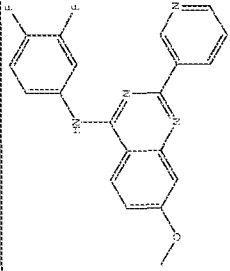
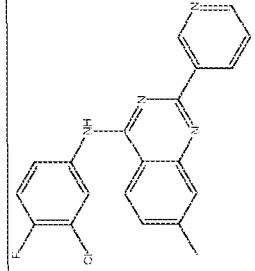
189			385.42	1H-NMR (400 MHz, DMSO-d6): δ 12.82 (s, 1H), 9.58 (s, 1H), 9.03 (d, J = 8.4 Hz, 1H), 8.92 (d, J = 4.0 Hz, 1H), 8.74 – 8.69 (m, 2H), 7.92 – 7.87 (m, 2H), 7.72 (t, J = 7.6 Hz, 1H), 7.66 – 7.56 (m, 3H), 7.23 (t, J = 7.6 Hz, 1H), 4.00 (s, 3H), 2.84 (d, J = 4.4 Hz, 3H).	DMSO	386.2 (M+1)	Method B (NH4HCO3)	95	Method C, G1
190			461.51	1H-NMR (400 MHz, DMSO-d6): δ 12.40 (s, 1H), 9.57 (d, J = 1.2 Hz, 1H), 9.48 (t, J = 6.0 Hz, 1H), 8.89 (d, J = 8.8 Hz, 1H), 8.73 – 8.68 (m, 2H), 7.96 (dd, J = 8.0, 1.2 Hz, 1H), 7.89 (d, J = 8.8 Hz, 1H), 7.76 – 7.72 (m, 1H), 7.61 – 7.55 (m, 3H), 7.34 – 7.20 (m, 6H), 4.52 (d, J = 5.6 Hz, 2H), 3.93 (s, 3H).	DMSO	462.2 (M+1)	Method B (NH4HCO3)	95	Method C, G1
191			368.77	1H NMR (300 MHz, DMSO-d6): δ 10.32 (s, 1H), 9.55 – 9.45 (m, 1H), 8.93 – 8.76 (m, 2H), 8.63 (d, J = 6.9 Hz, 1H), 8.10 (ddd, J = 13.2, 7.5, 2.6 Hz, 1H), 7.97 (d, J = 2.1 Hz, 1H), 7.80 – 7.67 (m, 3H), 7.62 – 7.49 (m, 1H).	DMSO	369.0, 371.0 (M+1)	Method B (NH4HCO3)	97	Method C, G1

192			401.68	¹H NMR (300 MHz, DMSO-d₆): δ 10.38 (s, 1H), 9.58 – 9.43 (m, 1H), 8.90 – 8.79 (m, 2H), 8.65 (d, J = 9.0 Hz, 1H), 8.33 (d, J = 2.4 Hz, 1H), 8.01 – 7.90 (m, 2H), 7.82 – 7.68 (m, 3H).	DMSO	401.0, 403.0, 405.0 (M+1)	Method B (NH ₄ HCO ₃)	97	Method C, G1
193			367.23	¹H NMR (300 MHz, DMSO-d₆): δ 10.19 (s, 1H), 9.53 (d, J = 2.1 Hz, 1H), 8.78 – 8.56 (m, 3H), 8.17 (t, J = 2.0 Hz, 1H), 7.96 (d, J = 2.1 Hz, 1H), 7.93 – 7.85 (m, 1H), 7.75 (dd, J = 8.9, 2.2 Hz, 1H), 7.58 (ddd, J = 8.0, 4.8, 0.7 Hz, 1H), 7.51 (t, J = 8.1 Hz, 1H), 7.26 (ddd, J = 8.0, 2.0, 0.6 Hz, 1H).	DMSO	367.0, 369.0 (M+1)	Method B (NH ₄ HCO ₃)	98	Method C, G1
194			376.80	¹H NMR (300 MHz, DMSO-d₆): δ 13.11 (s, 1H), 10.27 (s, 1H), 9.58 (d, J = 2.1 Hz, 1H), 8.87 – 8.62 (m, 4H), 8.21 – 8.10 (m, 1H), 7.96 (d, J = 2.1 Hz, 1H), 7.82 – 7.68 (m, 2H), 7.64 – 7.53 (m, 2H).	DMSO	377.0, 379.0 (M+1)	Method A (TFA)	95	Method C, G1

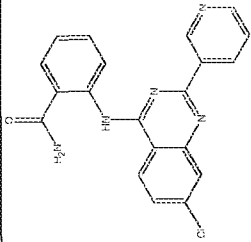
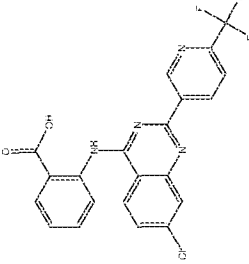
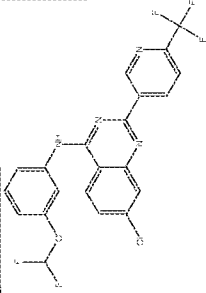
195			401.68	¹H-NMR (300 MHz, DMSO-d₆): δ 10.32 (s, 1H), 9.52 – 9.49 (m, 1H), 8.85 – 8.73 (m, 2H), 8.61 (d, J = 9.0 Hz, 1H), 8.18 – 8.08 (m, 2H), 7.99 – 7.95 (m, 1H), 7.77 (dd, J = 8.9, 2.2 Hz, 1H), 7.71 (dd, J = 7.8, 5.0 Hz, 1H), 7.42 – 7.39 (m, 1H).	DMSO	401.0, 403.0, 405.0 (M+1)	Method B (NH ₄ HCO ₃)	97	Method C, G1
196			386.76	¹H-NMR (300 MHz, DMSO-d₆): δ 10.43 (s, 1H), 9.49 (d, J = 1.9 Hz, 1H), 8.95 – 8.82 (m, 2H), 8.64 (d, J = 9.0 Hz, 1H), 8.01 – 7.82 (m, 4H), 7.78 (dd, J = 8.9, 2.2 Hz, 1H).	DMSO	387.0, 389.1 (M+1)	Method B (NH ₄ HCO ₃)	97	Method C, G1
197			411.86	¹H-NMR (400 MHz, DMSO-d₆): δ 10.50 (brs, 1H), 9.56 (s, 1H), 9.08 – 8.93 (m, 1H), 8.92 – 8.82 (m, 1H), 8.73 (d, J = 8.9 Hz, 1H), 8.21 – 8.11 (m, 2H), 8.02 (d, J = 1.9 Hz, 1H), 7.98 – 7.77 (m, 4H), 7.38 (s, 2H).	DMSO	412.1, 414.1 (M+1)	Method B (NH ₄ HCO ₃)	97	Method C, G1

198		365.22	¹H NMR (400 MHz, DMSO-d₆): δ 10.25 (s, 1H), 9.51 (d, J = 1.4 Hz, 1H), 8.77 – 8.69 (m, 2H), 8.60 (d, J = 9.0 Hz, 1H), 8.27 – 8.20 (m, 1H), 7.97 (d, J = 2.1 Hz, 1H), 7.90 (ddd, J = 9.0, 4.3, 2.6 Hz, 1H), 7.76 (dd, J = 8.9, 2.1 Hz, 1H), 7.64 (dd, J = 8.0, 4.9 Hz, 1H), 7.55 (t, J = 9.1 Hz, 1H).	DMSO	365.0, 367.0 (M+1)	Method B (NH ₄ HCO ₃)	100	Method C, G1
199		357.80	¹H NMR (400 MHz, DMSO-d₆): δ 10.36 (s, 1H), 9.53 – 9.49 (m, 1H), 8.81 – 8.70 (m, 2H), 8.64 (d, J = 8.9 Hz, 1H), 8.45 – 8.36 (m, 1H), 8.29 – 8.21 (m, 1H), 7.96 (d, J = 2.1 Hz, 1H), 7.77 (dd, J = 8.9, 2.1 Hz, 1H), 7.74 – 7.61 (m, 3H).	DMSO	357.9, 359.9 (M+1)	Method B (NH ₄ HCO ₃)	99	Method C, G1
200		435.23	¹H NMR (400 MHz, DMSO-d₆): δ 10.44 (s, 1H), 9.53 (s, 1H), 8.84 – 8.74 (m, 2H), 8.69 – 8.61 (m, 2H), 8.26 (dd, J = 8.8, 2.6 Hz, 1H), 7.99 (d, J = 2.1 Hz, 1H), 7.83 (d, J = 8.8 Hz, 1H), 7.78 (dd, J = 8.9, 2.1 Hz, 1H), 7.70 (dd, J = 7.7, 5.3 Hz, 1H).	DMSO	435.0, 437.0 (M+1)	Method B (NH ₄ HCO ₃)	99	Method C, G1

201			400.78		¹H-NMR (400 MHz, DMSO-d₆): δ 10.45 (s, 1H), 9.53 (d, J = 1.5 Hz, 1H), 8.85 (d, J = 8.1 Hz, 1H), 8.82 (dd, J = 5.0, 1.4 Hz, 1H), 8.69 (d, J = 9.0 Hz, 1H), 8.49 (s, 1H), 8.21 (d, J = 8.0 Hz, 1H), 7.99 (d, J = 2.1 Hz, 1H), 7.83 – 7.67 (m, 3H), 7.56 (d, J = 7.8 Hz, 1H).	DMSO	401.1, 403.0 (M+1)	Method B (NH ₄ HCO ₃)	99	Method C, G1
202			375.79		¹H-NMR (400 MHz, DMSO-d₆): δ 10.47 (s, 1H), 9.49 (s, 1H), 8.87 – 8.77 (m, 2H), 8.63 (d, J = 8.9 Hz, 1H), 8.42 (dd, J = 5.8, 2.7 Hz, 1H), 8.26 (ddd, J = 9.1, 4.9, 2.8 Hz, 1H), 7.98 (d, J = 2.1 Hz, 1H), 7.83 – 7.72 (m, 2H), 7.66 (t, J = 9.1 Hz, 1H).	DMSO	375.9, 377.9 (M+1)	Method B (NH ₄ HCO ₃)	99	Method C, G1
203			348.35		¹H-NMR (400 MHz, DMSO-d₆): δ 10.42 (s, 1H), 9.51 (s, 1H), 8.91 (d, J = 6.8 Hz, 1H), 8.86 (d, J = 4.4 Hz, 1H), 8.57 (d, J = 8.3 Hz, 1H), 8.11 (ddd, J = 13.0, 7.5, 2.5 Hz, 1H), 7.90 – 7.68 (m, 3H), 7.65 – 7.46 (m, 2H), 2.56 (s, 3H).	DMSO	343.1 (M+1)	Method B (NH ₄ HCO ₃)	100	Method C, G1

204			381.26	1H-NMR (400 MHz, DMSO-d6): δ 10.95 (s, 1H), 9.54 (s, 1H), 8.75 – 8.65 (m, 2H), 8.46 (d, J = 8.5 Hz, 1H), 8.42 (d, J = 2.4 Hz, 1H), 7.98 (dd, J = 8.8, 2.4 Hz, 1H), 7.72 (d, J = 9.1 Hz, 2H), 7.57 (dd, J = 7.9, 4.9 Hz, 1H), 7.53 (d, J = 8.5 Hz, 1H), 2.55 (s, 3H).	DMSO	381.0, 383.0 (M+1)	Method B (NH4HCO3)	100	Method C, G1
205			364.35	1H-NMR (400 MHz, DMSO-d6): δ 9.86 (s, 1H), 9.53 (s, 1H), 8.79 – 8.58 (m, 2H), 8.49 (d, J = 9.0 Hz, 1H), 8.13 (ddd, J = 9.2, 7.1, 1.7 Hz, 1H), 7.84 – 7.65 (m, 1H), 7.67 – 7.46 (m, 2H), 7.41 – 7.23 (m, 2H), 3.97 (s, 3H).	DMSO	365.1 (M+1)	Method B (NH4HCO3)	95	Method C, G1
206			364.80	1H-NMR (400 MHz, DMSO-d6): δ 10.32 (s, 1H), 9.50 (d, J = 1.4 Hz, 1H), 8.92 – 8.77 (m, 2H), 8.53 (d, J = 8.5 Hz, 1H), 8.24 (dd, J = 6.8, 2.5 Hz, 1H), 7.98 – 7.87 (m, 1H), 7.83 – 7.70 (m, 2H), 7.60 – 7.50 (m, 2H), 2.56 (s, 3H).	DMSO	365.1, 367.1 (M+1)	Method B (NH4HCO3)	100	Method C, G1

207			381.26		1H-NMR (400 MHz, DMSO-d6): δ 10.21 (s, 1H), 9.53 (s, 1H), 8.87 – 8.73 (m, 2H), 8.50 (d, J = 8.5 Hz, 1H), 8.17 (d, J = 1.7 Hz, 2H), 7.77 (s, 1H), 7.72 (dd, J = 7.8, 5.1 Hz, 1H), 7.57 (dd, J = 8.4, 0.9 Hz, 1H), 7.39 (t, J = 1.6 Hz, 1H), 2.56 (s, 3H).	DMSO	381.1, 383.0 (M+1)	Method B (NH4HCO3)	99	Method C, G1
208			337.38		1H-NMR (400 MHz, DMSO-d6): δ 10.12 (s, 1H), 9.53 (d, J = 1.6 Hz, 1H), 8.77 – 8.63 (m, 2H), 8.53 – 8.42 (m, 2H), 8.28 (d, J = 8.3 Hz, 1H), 7.75 (s, 1H), 7.69 (t, J = 7.9 Hz, 1H), 7.62 (d, J = 7.7 Hz, 1H), 7.60 – 7.51 (m, 2H), 2.55 (s, 3H).	DMSO	338.2 (M+1)	Method B (NH4HCO3)	100	Method C, G1
209			355.39		1H-NMR (400 MHz, DMSO-d6): δ 13.09 (s, 1H), 9.60 (s, 1H), 9.02 (d, J = 8.4 Hz, 1H), 8.96 (d, J = 8.4 Hz, 1H), 8.85 (d, J = 4.0 Hz, 1H), 8.49 (s, 1H), 8.13 (d, J = 8.4 Hz, 1H), 7.97 (d, J = 7.8 Hz, 1H), 7.92 (s, 1H), 7.83 (dd, J = 8.0, 5.2 Hz, 1H), 7.80 (s, 1H), 7.73 (t, J = 7.4 Hz, 1H), 7.61 (d, J = 8.0 Hz, 1H), 7.25 (t, J = 7.6 Hz, 1H), 2.56 (s, 3H).	DMSO	356.1 (M+1)	Method B (NH4HCO3)	99	Method C, G1

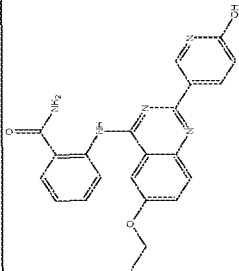
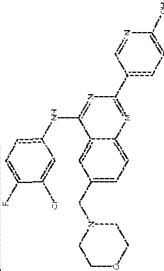
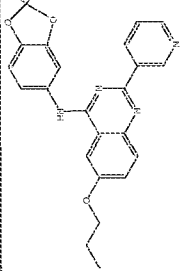
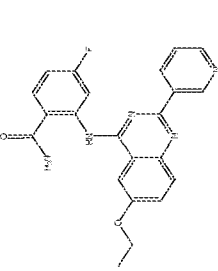
210			375.81	1H-NMR (400 MHz, DMSO-d6): δ 13.13 (s, 1H), 9.59 (s, 1H), 9.03 – 8.89 (m, 3H), 8.50 (s, 1H), 8.21 – 8.19 (d, J = 8.6 Hz, 1H), 7.99 – 7.87 (m, 4H), 7.82 – 7.79 (dd, J = 8.8, 2.3 Hz, 1H), 7.74 – 7.70 (m, 1H), 7.26 – 7.24 (m, 1H).	DMSO	376.1, 378.1 (M+1)	Method B (NH4HCO3)	98	Method C, G1
211			444.79	1H-NMR (400 MHz, DMSO-d6): δ 13.68 (brs, 1H), 11.98 (s, 1H), 9.49 (s, 1H), 8.63 – 8.64 (m, 2H), 8.11 – 7.90 (m, 3H), 7.81 (d, J = 2.0 Hz, 1H), 7.75 – 7.62 (m, 2H), 7.22 (t, J = 7.6 Hz, 1H).	DMSO	444.9, 446.9 (M+1)	Method B (NH4HCO3)	100	Method C, G1
212		HCl	466.79	1H-NMR (400 MHz, DMSO-d6): δ 10.36 (s, 1H), 9.62 (s, 1H), 8.90 (dd, J = 8.2, 1.3 Hz, 1H), 8.69 (d, J = 9.0 Hz, 1H), 8.05 (d, J = 8.2 Hz, 1H), 7.99 (d, J = 2.1 Hz, 1H), 7.92 – 7.86 (m, 1H), 7.86 – 7.79 (m, 1H), 7.77 (dd, J = 8.9, 2.1 Hz, 1H), 7.53 (t, J = 8.2 Hz, 1H), 7.30 (t, J = 7.3 Hz, 1H), 7.04 (dd, J = 8.1, 2.1 Hz, 1H).	DMSO	466.9, 468.9 (M+1)	Method B (NH4HCO3)	100	Method C, G1

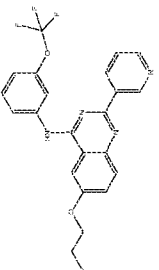
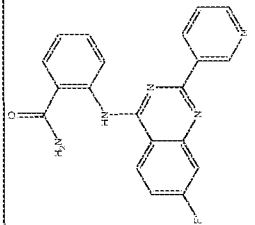
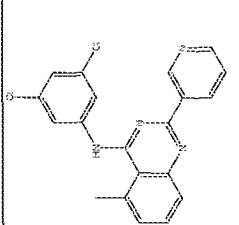
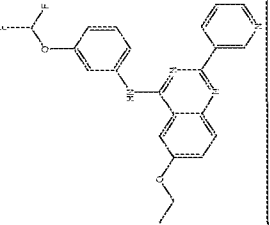
213			469.67	1H-NMR (400 MHz, DMSO-d6): δ 10.31 (s, 1H), 9.57 (s, 1H), 8.85 (dd, J = 8.1, 1.3 Hz, 1H), 8.60 (d, J = 9.0 Hz, 1H), 8.35 – 8.23 (m, 1H), 8.06 (d, J = 8.2 Hz, 1H), 8.00 – 7.90 (m, 2H), 7.80 – 7.65 (m, 2H).	DMSO	468.9, 470.8 (M+1)	Method B (NH4HCO3)	100	Method C, G1
214			436.77	1H-NMR (400 MHz, DMSO-d6): δ 10.35 (s, 1H), 9.56 (s, 1H), 8.84 (d, J = 8.3 Hz, 1H), 8.64 (d, J = 9.0 Hz, 1H), 8.14 – 7.99 (m, 2H), 7.95 (d, J = 2.1 Hz, 1H), 7.81 – 7.65 (m, 2H), 7.59 – 7.48 (m, 1H).	DMSO	436.9, 438.9 (M+1)	Method B (NH4HCO3)	100	Method C, G1
215			440.37	1H-NMR (400 MHz, DMSO-d6): δ 13.81 (brs, 1H), 11.97 (s, 1H), 9.57 (s, 1H), 9.01 – 8.91 (m, 1H), 8.84 (d, J = 8.3 Hz, 1H), 8.06 (dd, J = 7.9, 1.4 Hz, 1H), 8.01 (d, J = 8.2 Hz, 1H), 7.82 (d, J = 9.1 Hz, 1H), 7.79 – 7.70 (m, 1H), 7.52 (dd, J = 9.1, 2.6 Hz, 1H), 7.49 – 7.44 (m, 1H), 7.20 (t, J = 7.6 Hz, 1H), 3.94 (s, 3H).	DMSO	441.1 (M+1)	Method B (NH4HCO3)	100	Method C, G1

216			396.80		1H-NMR (400 MHz, DMSO-d6): δ 11.89 (s, 1H), 9.80 (s, 1H), 8.34 (dd, J = 9.6, 2.6 Hz, 1H), 8.22 (s, 1H), 8.11 (dd, J = 6.9, 2.6 Hz, 1H), 7.88 (d, J = 2.7 Hz, 1H), 7.84 – 7.77 (m, 1H), 7.74 (d, J = 9.1 Hz, 1H), 7.56 – 7.43 (m, 2H), 6.44 (d, J = 9.6 Hz, 1H), 3.95 (s, 3H).	DMSO	397.1, 399.1 (M+1)	Method B (NH4HCO3)	100	Method D, G1
217			410.37		1H-NMR (400 MHz, DMSO-d6): δ 11.89 (s, 1H), 9.80 (s, 1H), 8.36 (dd, J = 9.6, 2.6 Hz, 1H), 8.25 (d, J = 2.2 Hz, 1H), 7.92 (d, J = 2.6 Hz, 1H), 7.80 – 7.67 (m, 3H), 7.53 – 7.46 (m, 2H), 7.27 (t, J = 74.0 Hz, 1H), 6.98 (dd, J = 8.2, 2.0 Hz, 1H), 6.43 (d, J = 9.6 Hz, 1H), 3.96 (s, 3H).	DMSO	411.1 (M+1)	Method B (NH4HCO3)	100	Method D, G1
218			387.39		1H-NMR (400 MHz, DMSO-d6): δ 12.98 (s, 1H), 12.58 – 12.05 (m, 1H), 8.55 (s, 2H), 8.41 (s, 1H), 8.33 (dd, J = 9.7, 2.5 Hz, 1H), 8.11 – 7.98 (m, 1H), 7.97 – 7.86 (m, 2H), 7.79 – 7.64 (m, 3H), 7.44 – 7.29 (m, 1H), 6.52 (d, J = 9.7 Hz, 1H), 3.98 (s, 3H).	DMSO	388.2 (M+1)	Method B (NH4HCO3)	100	Method D, G1
219			389.38		1H-NMR (400 MHz, DMSO-d6): δ 12.73 (s, 1H), 9.56 (s, 1H), 9.10 (dd, J = 8.9, 5.4 Hz, 1H), 8.79 – 8.63 (m, 2H), 8.50 (s, 1H), 8.08 (s, 1H), 7.95 – 7.76 (m, 2H), 7.68 – 7.45 (m, 4H), 3.97 (s, 3H).	DMSO	390.1 (M+1)	Method B (NH4HCO3)	98	Method C, G1

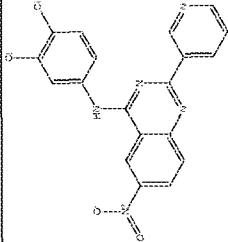
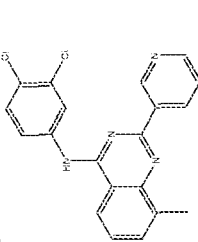
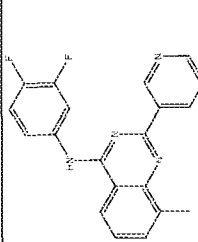
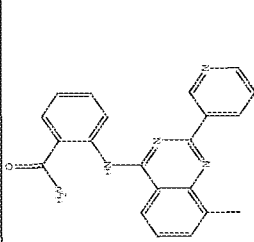
220			369.80	¹H-NMR (400 MHz, DMSO-d₆): δ 14.87 (brs, 1H), 10.08 (brs, 1H), 8.85 – 7.66 (m, 6H), 7.69 – 6.63 (m, 3H), 3.96 (s, 3H).	DMSO	397.1, 399.1 (M+1)	Method B (NH ₄ HCO ₃)	100	Method C, G1
221			387.39	¹H-NMR (400 MHz, DMSO-d₆): δ 14.80 (brs, 1H), 13.19 (s, 1H), 8.68 (d, J = 7.3 Hz, 1H), 8.44 (s, 1H), 8.38 (d, J = 8.0 Hz, 1H), 8.13 (d, J = 9.1 Hz, 1H), 8.10 – 8.03 (m, 1H), 7.95 (d, J = 7.8 Hz, 1H), 7.90 (s, 1H), 7.86 (s, 1H), 7.78 – 7.65 (m, 2H), 7.43 (t, J = 7.5 Hz, 1H), 6.78 (t, J = 6.8 Hz, 1H), 3.99 (s, 3H).	DMSO	388.1 (M+1)	Method B (NH ₄ HCO ₃)	100	Method C, G1
222			410.37	¹H-NMR (400 MHz, DMSO-d₆): δ 15.33 (brs, 1H), 13.47 (brs, 1H), 11.69 (s, 1H), 8.62 (dd, J = 7.5, 2.1 Hz, 1H), 8.45 (d, J = 2.3 Hz, 1H), 8.17 (d, J = 9.2 Hz, 1H), 8.07 (dd, J = 6.2, 2.1 Hz, 1H), 7.81 – 7.68 (m, 3H), 7.61 (t, J = 8.1 Hz, 1H), 7.32 (t, J = 74.0 Hz, 1H), 7.26 – 7.16 (m, 1H), 6.80 – 6.69 (m, 1H), 4.03 (s, 3H).	DMSO	411.1 (M+1)	Method B (NH ₄ HCO ₃)	100	Method C, G1

223			401.22	¹H-NMR (400 MHz, DMSO-d₆): δ 11.22 (brs, 1H), 8.47 (d, J = 8.6 Hz, 1H), 8.30 (dd, J = 9.6, 2.3 Hz, 1H), 8.27 – 8.16 (m, 1H), 8.11 (dd, J = 6.8, 2.4 Hz, 1H), 7.90 – 7.73 (m, 2H), 7.60 (dd, J = 8.8, 1.8 Hz, 1H), 7.47 (t, J = 9.1 Hz, 1H), 6.44 (d, J = 9.6 Hz, 1H).	DMSO	401.0, 403.0 (M+1)	Method B (NH ₄ HCO ₃)	100	Method C, G1
224			414.79	¹H-NMR (400 MHz, DMSO-d₆): δ 12.00 (s, 1H), 10.07 (s, 1H), 8.54 (d, J = 8.9 Hz, 1H), 8.34 (dd, J = 9.6, 2.6 Hz, 1H), 8.30 (d, J = 2.3 Hz, 1H), 7.87 – 7.69 (m, 3H), 7.62 (dd, J = 8.9, 2.1 Hz, 1H), 7.48 (t, J = 8.2 Hz, 1H), 7.26 (t, J = 74.0 Hz, 1H), 7.00 (dd, J = 8.1, 2.0 Hz, 1H), 6.44 (d, J = 9.6 Hz, 1H).	DMSO	415.1, 417.1 (M+1)	Method B (NH ₄ HCO ₃)	100	Method C, G1
225			432.78	¹H-NMR (400 MHz, DMSO-d₆): δ 12.06 (s, 1H), 10.15 (s, 1H), 8.55 (d, J = 9.0 Hz, 1H), 8.34 (dd, J = 9.6, 2.5 Hz, 1H), 8.29 (s, 1H), 7.97 (s, 1H), 7.88 (d, J = 8.0 Hz, 1H), 7.84 (d, J = 2.1 Hz, 1H), 7.65 (dd, J = 8.8, 2.1 Hz, 1H), 7.57 (t, J = 8.2 Hz, 1H), 7.17 (d, J = 7.8 Hz, 1H), 6.44 (d, J = 9.6 Hz, 1H).	DMSO	433.0, 435.0 (M+1)	Method B (NH ₄ HCO ₃)	100	Method C, G1
226			428.36	¹H-NMR (400 MHz, DMSO-d₆): δ 12.26 (brs, 1H), 8.47 – 8.06 (m, 3H), 8.06 – 7.76 (m, 3H), 7.67 – 7.59 (m, 2H), 7.28 (s, 1H), 6.47 (d, J = 9.7 Hz, 1H), 3.96 (s, 3H).	DMSO	429.1 (M+1)	Method B (NH ₄ HCO ₃)	94	Method D, G1

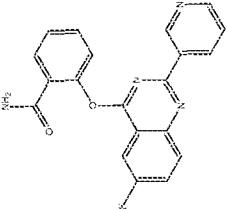
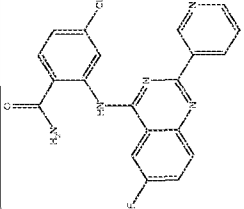
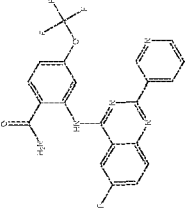
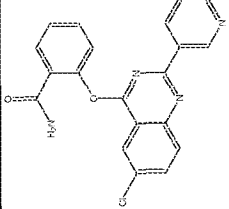
227		401.42	¹H-NMR (400 MHz, DMSO-d₆): δ 12.97 (s, 1H), 12.27 (s, 1H), 8.73 – 8.56 (m, 1H), 8.53 (s, 1H), 8.42 (s, 1H), 8.33 (dd, J = 9.7, 2.6 Hz, 1H), 8.04 – 7.86 (m, 3H), 7.74 – 7.59 (m, 3H), 7.34 (t, J = 8.7 Hz, 1H), 6.52 (d, J = 9.7 Hz, 1H), 4.23 (q, J = 6.9 Hz, 2H), 1.45 (t, J = 6.9 Hz, 3H).	DMSO	402.2 (M+1)	Method B (NH ₄ HCO ₃)	100	Method D, G1
228		465.91	¹H-NMR (400 MHz, DMSO-d₆): δ 8.43 – 8.31 (m, 2H), 8.27 (s, 1H), 8.14 (dd, J = 6.7, 2.3 Hz, 1H), 7.89 – 7.70 (m, 3H), 7.47 (t, J = 9.1 Hz, 1H), 6.44 (d, J = 9.6 Hz, 1H), 3.62 (d, J = 9.3 Hz, 6H), 2.42 (s, 4H).	DMSO	466.1, 468.1 (M+1)	Method B (NH ₄ HCO ₃)	97	Method D, G1
229		436.41	¹H-NMR (400 MHz, DMSO-d₆): δ 10.26 (brs, 1H), 9.47 (s, 1H), 8.91 (d, J = 7.2 Hz, 1H), 8.84 (d, J = 5.2 Hz, 1H), 8.08 – 8.06 (m, 2H), 7.91 (d, J = 7.6 Hz, 1H), 7.85 (t, J = 8 Hz, 1H), 7.65 – 7.60 (m, 2H), 7.54 – 7.52 (m, 1H), 4.17 (t, J = 7.6 Hz, 2H), 1.86 (q, J = 6.8 Hz, 2H), 1.08 – 1.03 (m, 3H).	DMSO	423.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
230		403.41	¹H-NMR (400 MHz, DMSO-d₆): δ 13.42 (s, 1H), 9.53 (s, 1H), 9.10 (d, J = 12.8 Hz, 1H), 8.70 – 8.65 (m, 2H), 8.50 (s, 1H), 8.07 – 8.04 (m, 2H), 7.86 (d, J = 8.8 Hz, 1H), 7.60 – 7.55 (m, 2H), 7.48 (s, 1H), 7.06 – 7.01 (m, 1H), 4.21 (q, J = 6.8 Hz, 2H), 1.45 (t, J = 7.2 Hz, 3H).	DMSO	403.9 (M+1)	Method A (TFA)	95	Method C, G1

231		440.42	1H-NMR (400 MHz, DMSO-d6): δ 10.57 (s, 1H), 9.48 (s, 1H), 9.01 (d, J = 7.6 Hz, 1H), 8.90 (d, J = 5.2 Hz, 1H), 8.19 (s, 1H), 8.04 (s, 1H), 7.98 (d, J = 8.8 Hz, 2H), 7.93 (t, J = 5.6 Hz, 1H), 7.64 – 7.60 (m, 2H), 7.22 (d, J = 8.4 Hz, 1H), 4.19 (t, J = 6.4 Hz, 1H), 1.87 – 1.81 (m, 2H), 1.06 (t, J = 7.2 Hz, 3H).	DMSO	440.9 (M+1)	Method A (TFA)	95	Method C, G1
232		359.12	1H-NMR (400 MHz, DMSO-d6): δ 13.10 (s, 1H), 9.54 (s, 1H), 8.96 (d, J = 8.4 Hz, 1H), 8.82 – 8.80 (m, 2H), 8.52 (s, 1H), 8.22 – 8.17 (m, 1H), 7.97 – 7.93 (m, 2H), 7.70 – 7.59 (m, 4H), 7.21 (t, J = 8.0 Hz, 1H).	DMSO	360.1 (M+1)	Method B (NH4HCO3)	95	Method C, G1
233		381.26	1H-NMR (400 MHz, DMSO-d6): δ 9.48 (s, 1H), 9.08 (s, 1H), 8.67 – 8.70 (m, 1H), 8.64 (d, J = 8.0 Hz, 1H), 7.98 (s, 2H), 7.76 – 7.78 (m, 2H), 7.55 – 7.57 (m, 1H), 7.43 – 7.46 (m, 1H), 7.36 (s, 1H), 2.99 (s, 3H).	DMSO	381.0, 383.0, 385.0 (M+1)	Method B (NH4HCO3)	95	Method C, G1
234		408.4	1H-NMR (400 MHz, DMSO-d6): δ 10.38 (s, 1H), 9.51 (s, 1H), 9.00 (d, J = 7.2 Hz, 1H), 8.88 (d, J = 4.8 Hz, 1H), 8.13 (s, 1H), 7.96 (d, J = 7.2 Hz, 1H), 7.86 – 7.91 (m, 2H), 7.86 (s, 1H), 7.80 (d, J = 8.4 Hz, 1H), 7.63 (d, J = 2.4 Hz, 1H), 7.62 (dd, J = 9.2, 2.0 Hz, 1H), 7.55 (t, J = 8.4 Hz, 1H), 7.31 (t, J = 74.0 Hz, 1H), 7.05 (d, J = 8.0 Hz, 1H), 4.28 (q, J = 6.8 Hz, 2H), 1.46 (t, J = 6.8 Hz, 3H).	DMSO	409.2 (M+1)	Method B (NH4HCO3)	95	Method C, G1

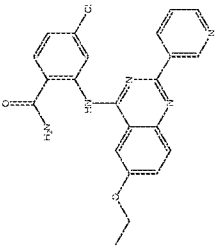
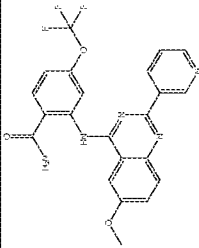
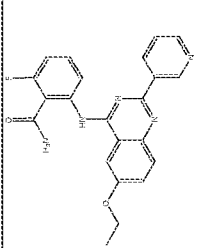
235		412.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.81 (s, 1H), 9.49 (s, 1H), 9.02 (d, J = 8.4 Hz, 1H), 8.91 (dd, J = 5.2, 1.2 Hz, 1H), 8.31 (d, J = 2.0 Hz, 1H), 8.06 (s, 1H), 8.01 (d, J = 8.8 Hz, 2H), 7.94 – 7.91 (m, 1H), 7.64 – 7.60 (m, 2H), 7.22 (d, J = 8.4 Hz, 1H), 4.02 (s, 3H).	DMSO	412.9 (M+1)	Method A (TFA)	95	Method C, G1
236		395.46	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.64 (s, 1H), 8.90 (s, 1H), 8.77 (dd, J = 13.5, 6.2 Hz, 2H), 8.19 (d, J = 9.2 Hz, 1H), 7.81 (dd, J = 9.2, 2.4 Hz, 1H), 7.72 – 7.56 (m, 3H), 7.33 (d, J = 2.0 Hz, 1H), 4.15 (q, J = 6.8 Hz, 2H), 2.39 (s, 3H), 2.35 (s, 3H), 1.38 (t, J = 6.8 Hz, 3H).	DMSO	396.2 (M+1)	Method B (NH ₄ HCO ₃)	98	Method C, J1
237		376.8	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.64 (brs, 1H), 11.31 (brs, 1H), 9.45 (s, 1H), 8.82 – 8.69 (m, 1H), 8.61 (s, 1H), 8.15 (d, J = 2.0 Hz, 1H), 8.04 – 7.91 (m, 1H), 7.81 (dd, J = 6.9, 2.1 Hz, 1H), 7.70 – 7.55 (m, 1H), 7.47 (s, 1H), 7.06 (s, 1H), 6.98 (d, J = 8.2 Hz, 1H).	DMSO	376.9, 378.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G8

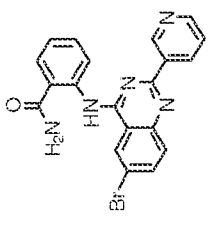
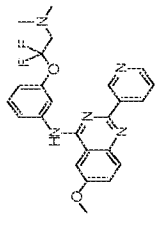
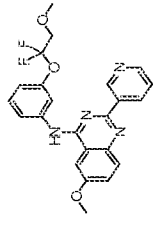
238		412.23	1H-NMR (400 MHz, DMSO-d6): δ 10.70 (s, 1H), 9.62 (d, J = 2.4 Hz, 1H), 9.51 (d, J = 1.6 Hz, 1H), 8.76 – 8.79 (m, 2H), 8.57 (dd, J = 6.8, 2.0 Hz, 1H), 8.29 (d, J = 2.4 Hz, 1H), 8.01 (d, J = 8.8 Hz, 1H), 7.94 (dd, J = 6.8, 2.0 Hz, 1H), 7.71 (m, 2H).	DMSO	412.0, 414.0, 416.0 (M+1)	Method B (NH4HCO3)	95	Method C, G1
239		381.26	1H-NMR (400 MHz, DMSO-d6): 10.04 (s, 1H), 9.59 (d, J = 1.2 Hz, 1H), 8.74 – 8.70 (m, 2H), 8.42 – 8.39 (m, 2H), 7.98 (dd, J = 2.4 Hz, 1H), 7.79 (d, J = 7.2 Hz, 1H), 7.72 (d, J = 8.8 Hz, 1H), 7.59 – 7.55 (m, 2H), 2.75 (s, 3H).	DMSO	381, 383 (M+1)	Method B (NH4HCO3)	95	Method C, G1
240		348.35	1H-NMR (400 MHz, DMSO-d6): 10.00 (s, 1H), 9.57 (d, J = 1.2 Hz, 1H), 8.72 – 8.70 (m, 2H), 8.38 (d, J = 8.4 Hz, 1H), 8.17 – 8.12 (m, 1H), 7.79 – 7.74 (m, 2H), 7.59 – 7.50 (m, 3H), 2.74 (s, 3H).	DMSO	349.2 (M+1)	Method B (NH4HCO3)	95	Method C, G1
241		355.39	1H-NMR (400 MHz, DMSO-d6): 11.10 (s, 1H), 9.60 (d, J = 1.6 Hz, 1H), 9.16 (d, J = 8.4 Hz, 1H), 8.82 – 8.72 (m, 2H), 8.50 (s, 1H), 8.04 (d, J = 8.4 Hz, 1H), 7.97 – 7.92 (m, 2H), 7.77 – 7.72 (m, 2H), 7.63 – 7.60 (m, 2H), 7.24 – 7.19 (m, 1H), 2.76 (s, 3H).	DMSO	356.2 (M+1)	Method B (NH4HCO3)	95	Method C, G1

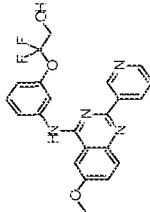
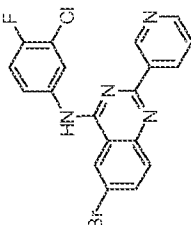
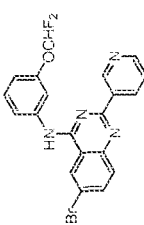
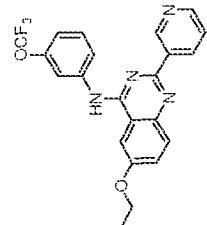
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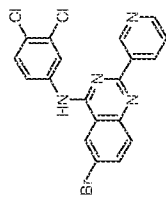
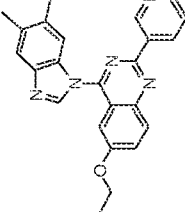
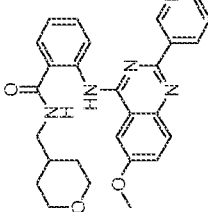
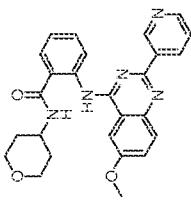
245			360.34	¹H-NMR (400 MHz, DMSO-d₆): δ 11.95 (s, 1H), 9.54 (s, 1H), 9.29 (s, 1H), 8.06 (d, J = 0.9 Hz, 1H), 7.99 (d, J = 8.5 Hz, 2H), 7.82 (dd, J = 26.4, 8.4 Hz, 2H), 7.58 (dd, J = 7.9, 4.8 Hz, 1H), 7.28 (d, J = 8.0 Hz, 1H), 8.72 (d, J = 3.6 Hz, 1H), 8.67 (d, J = 7.8 Hz, 1H), 8.64 – 8.53 (m, 1H).	DMSO	360.8 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G8
246			393.8	¹H-NMR (400 MHz, DMSO-d₆): δ 13.29 (s, 1H), 9.54 (s, 1H), 9.29 (s, 1H), 8.06 (d, J = 0.9 Hz, 1H), 7.99 (d, J = 8.5 Hz, 2H), 7.82 (dd, J = 26.4, 8.4 Hz, 2H), 7.58 (dd, J = 7.9, 4.8 Hz, 1H), 7.28 (d, J = 8.0 Hz, 1H), 8.72 (d, J = 3.6 Hz, 1H), 8.67 (d, J = 7.8 Hz, 1H), 8.64 – 8.53 (m, 1H).	DMSO	394.0, 396.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, J1
247				¹H-NMR (400 MHz, DMSO-d₆): δ 13.36 (brs, 1H), 9.54 (s, 1H), 9.36 – 9.08 (m, 1H), 8.12 (d, J = 8.8 Hz, 2H), 8.08 – 7.96 (m, 1H), 7.94 – 7.78 (m, 2H), 7.57 (dd, J = 7.8, 4.8 Hz, 1H), 7.29 – 7.16 (m, 1H), 8.94 – 8.49 (m, 3H).	DMSO	444.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, J1
248			376.8	¹H-NMR (400 MHz, DMSO-d₆): δ 9.57 (s, 1H), 8.71 (d, J = 5.6 Hz, 2H), 8.40 (d, J = 1.8 Hz, 1H), 8.01 – 7.97 (m, 2H), 7.89 (dd, J = 7.8, 1.5 Hz, 1H), 7.63 – 7.51 (m, 1H), 7.29 (t, J = 7.4 Hz, 1H), 6.79 (d, J = 8.4 Hz, 1H), 6.76 – 6.74 (m, 1H).	DMSO	377.0, 379.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G8

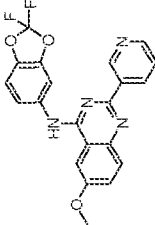
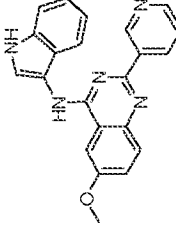
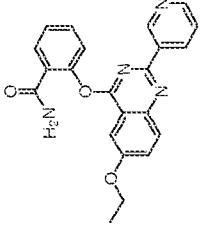
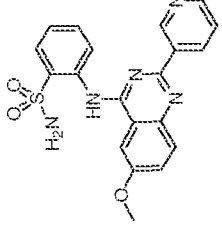
249		426.35	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.70 (d, J = 1.5 Hz, 1H), 8.74 (td, J = 8.0, 1.9 Hz, 1H), 8.71 (dd, J = 4.7, 1.7 Hz, 1H), 8.50 (d, J = 9.0 Hz, 1H), 7.88 (dd, J = 7.8, 1.9 Hz, 1H), 7.76 (d, J = 1.1 Hz, 1H), 7.20 – 7.11 (m, 1H), 7.64 – 7.56 (m, 2H), 6.70 – 6.60 (m, 1H), 6.52 (dd, J = 10.8, 3.9 Hz, 1H).	DMSO	427.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G8
250		360.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.69 (s, 1H), 8.83 (d, J = 7.4 Hz, 1H), 8.71 (d, J = 3.9 Hz, 1H), 8.49 (t, J = 7.2 Hz, 1H), 7.85 (d, J = 7.8 Hz, 1H), 7.63 – 7.51 (m, 3H), 7.14 (t, J = 7.0 Hz, 1H), 6.61 (d, J = 8.0 Hz, 1H), 6.46 (t, J = 7.2 Hz, 1H).	DMSO	360.8 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G8
251		400.43	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.92 (s, 1H), 9.04 (d, J = 8.3 Hz, 1H), 8.66 (dd, J = 7.7, 1.7 Hz, 1H), 8.48 (s, 1H), 8.10 (dd, J = 4.6, 1.7 Hz, 1H), 8.01 – 7.94 (m, 2H), 7.89 (d, J = 8.9 Hz, 1H), 7.72 (t, J = 7.7 Hz, 1H), 7.55 (m, 2H), 7.20 (t, J = 7.5 Hz, 1H), 6.74 (dd, J = 7.7, 4.7 Hz, 1H), 4.24 (q, J = 6.9 Hz, 2H), 1.46 (t, J = 6.9 Hz, 3H).	DMSO	401.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
252		483.44	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.28 (s, 1H), 9.54 (s, 1H), 9.28 (s, 1H), 8.70 – 8.66 (m, 2H), 8.59 (s, 1H), 8.11 (d, J = 8.8 Hz, 2H), 7.90 (d, J = 8.9 Hz, 1H), 7.64 – 7.49 (m, 3H), 7.18 (d, J = 9.2 Hz, 1H), 4.14 (t, J = 6.2 Hz, 2H), 1.86 (q, J = 6.8 Hz, 2H), 1.07 (t, J = 7.2 Hz, 3H).	DMSO	484.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, J1

253					¹H-NMR (400 MHz, DMSO-d₆): δ 13.24 (s, 1H), 9.56 (s, 1H), 9.39 (d, J = 2.1 Hz, 1H), 8.77 - 8.73 (m, 2H), 8.55 (s, 1H), 8.09 (s, 1H), 8.00 (d, J = 8.6 Hz, 1H), 7.90 (d, J = 9.1 Hz, 1H), 7.65 (dd, J = 7.9, 4.9 Hz, 1H), 7.59 (dd, J = 9.1, 2.5 Hz, 1H), 4.24 (q, J = 6.9 Hz, 2H), 1.46 (t, J = 6.9 Hz, 3H).	DMSO	420.0, 422.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
254		469.42			¹H-NMR (400 MHz, DMSO-d₆): δ 13.34 (s, 1H), 9.60 (s, 1H), 9.35 (s, 1H), 8.77 (d, J = 3.5 Hz, 1H), 8.72 (d, J = 7.9 Hz, 1H), 8.67 (s, 1H), 8.22 (s, 1H), 8.17 (d, J = 8.8 Hz, 1H), 7.94 (d, J = 9.1 Hz, 1H), 7.69 - 7.58 (m, 2H), 7.55 (s, 1H), 7.24 (d, J = 8.5 Hz, 1H), 4.29 (q, J = 7.0 Hz, 2H), 1.53 (t, J = 6.9 Hz, 3H).	DMSO	470.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
255		403.41			¹H-NMR (400 MHz, DMSO-d₆): δ 11.43 (s, 1H), 9.53 (s, 1H), 8.68 - 8.66 (m, 2H), 8.56 (d, J = 8.3 Hz, 1H), 8.23 (s, 1H), 8.12 (s, 1H), 7.88 (d, J = 9.1 Hz, 1H), 7.72 - 7.64 (m, 1H), 7.61 - 7.47 (m, 3H), 7.17 - 7.08 (m, 1H), 4.23 (q, J = 6.9 Hz, 2H), 1.46 (t, J = 6.9 Hz, 3H).	DMSO	404.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

Number	Product	Salt Type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	Retention Time (min)	LCMS Protocol	Purity percent	Method of Coupling
256		HCl	456.723	¹ H NMR (300 MHz, DMSO) δ 9.55 (s, 1H), 9.09 (d, J = 8.0 Hz, 1H), 9.01 – 8.76 (m, 2H), 8.50 (s, 1H), 8.37 (s, 1H), 8.13 – 7.91 (m, 4H), 7.85 (d, J = 8.9 Hz, 1H), 7.69 (t, J = 7.7 Hz, 1H), 7.25 (t, J = 7.6 Hz, 1H).	DMSO	420 (M+1)	1.89	Method D	100	Method G1
257		HCl	524.39	¹ H NMR (300 MHz, DMSO) δ 10.68 (s, 1H), 9.50 (s, 1H), 9.08 (d, J = 8.1 Hz, 1H), 8.92 (d, J = 5.2 Hz, 1H), 8.28 (s, 1H), 8.11 (s, 1H), 8.06 – 7.93 (m, 2H), 7.87 (d, J = 7.6 Hz, 1H), 7.70 – 7.51 (m, 2H), 7.19 (d, J = 7.6 Hz, 1H), 4.28 (t, J = 9.1 Hz, 2H), 4	DMSO	452 (M+1)	1.34	Method D	100	Method G1
258		HCl	474.89	¹ H NMR (300 MHz, DMSO) δ 10.36 (s, 1H), 9.52 (s, 1H), 9.00 (d, J = 8.2 Hz, 1H), 8.89 (d, J = 4.4 Hz, 1H), 8.12 (s, 1H), 8.02 – 7.78 (m, 4H), 7.64 (d, J = 9.2 Hz, 1H), 7.54 (t, J = 8.2 Hz, 1H), 7.07 (d, J = 7.7 Hz, 1H), 4.08 – 3.87 (m, 5H), 3.46 (s, 3H).	DMSO	439 (M+1)	2.21	Method C	98	Method G1

259		HCl	460.86	¹H NMR (300 MHz, DMSO) δ 10.61 (s, 1H), 9.51 (s, 1H), 9.10 (d, J = 8.2 Hz, 1H), 8.94 (d, J = 4.9 Hz, 1H), 8.22 (s, 1H), 8.08 – 7.94 (m, 2H), 7.93 – 7.78 (m, 2H), 7.65 (d, J = 9.1 Hz, 1H), 7.53 (t, J = 8.2 Hz, 1H), 7.08 (d, J = 7.3 Hz, 1H), 4.01 (s, 3H), 3	DMSO	425 (M+1)	1.64	Method C	98	Method G1	
260		HCl	429.67	No Data			428.9 (M+1)	2.35	Method C	100	Method G1
261		HCl	443.24	¹H NMR (300 MHz, DMSO) δ 10.35 (s, 1H), 9.52 (d, J = 1.5 Hz, 1H), 8.97 (dd, J = 6.6, 1.8 Hz, 2H), 8.91 – 8.83 (m, 1H), 8.07 (dd, J = 8.9, 2.0 Hz, 1H), 7.85 (dd, J = 14.6, 8.2 Hz, 4H), 7.54 (dd, J = 11.7, 4.6 Hz, 1H), 7.31 (s, 1H), 7.10 – 6.99 (m, 1H).	DMSO	442.96 (M+1)	2.36	Method C	100	Method G1	
262		HCl	426.39	¹H NMR (300 MHz, DMSO) δ 10.71 (s, 1H), 9.47 (s, 1H), 9.01 (d, J = 8.2 Hz, 1H), 8.90 (d, J = 4.0 Hz, 1H), 8.23 (s, 1H), 8.08 – 7.86 (m, 4H), 7.61 (t, J = 8.2 Hz, 2H), 7.22 (d, J = 7.6 Hz, 1H), 4.28 (d, J = 7.0 Hz, 2H), 1.44 (t, J = 6.9 Hz, 3H).	DMSO	427.1 (M+1)	2.47	Method C	93	Method G1	

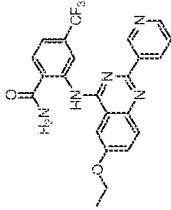
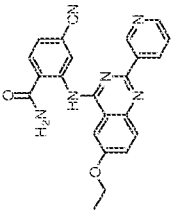
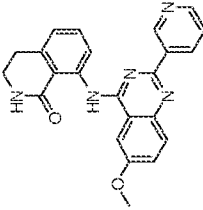
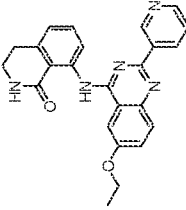
263		HCl	446.13	¹ H NMR (300 MHz, DMSO) δ : 10.35 (s, 1H), 9.47 (s, 1H), 8.90 (s, 3H), 8.27 (s, 1H), 8.02 (d, J = 8.9 Hz, 1H), 7.94 (d, J = 8.8 Hz, 1H), 7.61 (d, J = 9.0 Hz, 2H), 7.66 (d, J = 8.8 Hz, 1H).	DMSO	494.0 (M+1)	2.29	Method C	91	Method G1
264			395.46	¹ H NMR (300 MHz, DMSO) δ : 9.63 (s, 1H), 8.88 (s, 1H), 8.83 – 8.71 (m, 2H), 8.18 (d, J = 9.2 Hz, 1H), 7.90 (dd, J = 9.2, 2.3 Hz, 1H), 7.69 – 7.57 (m, 3H), 7.32 (d, J = 2.1 Hz, 1H), 4.14 (q, J = 6.6 Hz, 2H), 2.39 (s, 3H), 2.35 (s, 3H), 1.37 (t, J = 6.9 Hz, 3).	DMSO	396.10 (M+1)	2.3	Method C	100	Method G12
265		HCl	469.53	¹ H NMR (300 MHz, DMSO) δ : 12.13 (s, 1H), 9.54 (s, 1H), 8.99 (d, J = 7.4 Hz, 1H), 8.85 (s, 2H), 8.61 (d, J = 8.2 Hz, 1H), 7.95 (d, J = 9.1 Hz, 1H), 7.86 (d, J = 7.3 Hz, 2H), 7.68 (dd, J = 15.1, 7.8 Hz, 3H), 7.32 (d, J = 6.8 Hz, 1H), 4.00 (s, 3H), 3.71 (d, J).	DMSO	470.1 (M+1)	2.15	Method C	100	Method G1
266		HCl	455.51	¹ H NMR (300 MHz, DMSO) δ : 12.07 (s, 1H), 9.52 (s, 1H), 8.82 – 8.65 (m, 4H), 7.86 (dd, J = 8.1, 5.0 Hz, 2H), 7.73 – 7.55 (m, 4H), 7.27 (t, J = 7.2 Hz, 1H), 3.98 (m, 1H), 3.82 (d, J = 10.4 Hz, 2H), 3.35 (t, J = 11.7 Hz, 2H), 1.68 (d, J = 10.3 Hz, 2H), 1.55 –	DMSO	456.1 (M+1)	2.09	Method C	100	Method G1

267		HCl	406.36	¹ H NMR (300 MHz, DMSO) δ 10.57 (s, 1H), 9.45 (d, J = 1.5 Hz, 1H), 8.99 (d, J = 8.0 Hz, 1H), 8.89 (d, J = 3.9 Hz, 1H), 8.16 (d, J = 2.3 Hz, 1H), 8.05 (d, J = 2.0 Hz, 1H), 7.94 (t, J = 7.4 Hz, 2H), 7.63 (ddd, J = 9.3, 6.9, 2.3 Hz, 2H), 7.52 (d, J = 8.7 Hz, 1H).	DMSO	409.1 (M+1)	2.3	Method C	100	Method G1
268		HCl	367.4	¹ H NMR (300 MHz, DMSO) δ 11.12 (s, 1H), 9.81 (s, 1H), 9.43 – 9.37 (m, 1H), 8.58 (ddd, J = 8.0, 4.8, 1.4 Hz, 2H), 8.09 (d, J = 2.5 Hz, 1H), 7.89 (d, J = 2.2 Hz, 1H), 7.80 (s, 1H), 7.72 (d, J = 7.9 Hz, 1H), 7.53 – 7.41 (m, 3H), 7.17 (t, J = 7.5 Hz, 1H), 7.0	DMSO	368.1 (M+1)	1.98	Method C	100	Method G1
269			386.4	¹ H NMR (300 MHz, DMSO) δ 11.47 (s, 2H), 9.39 (s, 1H), 8.66 (d, J = 2.8 Hz, 1H), 8.56 – 8.47 (m, 1H), 8.01 – 7.90 (m, 2H), 7.69 – 7.55 (m, 2H), 7.55 – 7.40 (m, 2H), 7.05 (t, J = 7.5 Hz, 1H), 6.97 (d, J = 8.2 Hz, 1H), 4.23 (q, J = 6.9 Hz, 2H), 1.44 (t, J =	DMSO	387.0 (M+1)	2.05	Method C	100	Method G6
270			407.45	¹ H NMR (300 MHz, DMSO) δ 10.52 (s, 1H), 9.44 (s, 1H), 8.92 (d, J = 7.4 Hz, 1H), 8.86 (d, J = 5.1 Hz, 1H), 8.46 (d, J = 6.4 Hz, 1H), 8.01 (d, J = 8.7 Hz, 2H), 7.92 – 7.74 (m, 6H), 7.63 (dd, J = 9.2, 2.4 Hz, 1H), 7.46 (t, J = 7.8 Hz, 1H), 3.98 (s, 4H).	DMSO	408.0 (M+1)	1.92	Method C	100	Method G1, with two drops of conc. HCl

271		HCl	365.42	¹ H NMR (300 MHz, DMSO) δ 9.75 (s, 1H), 9.66 (s, 1H), 9.39 (s, 1H), 8.61 (d, J = 3.1 Hz, 1H), 8.54 (d, J = 8.2 Hz, 1H), 7.85 (s, 1H), 7.82 (s, 2H), 7.71 (d, J = 7.4 Hz, 1H), 7.54 (d, J = 9.1 Hz, 1H), 7.46 (dd, J = 7.5, 5.2 Hz, 1H), 7.35 – 7.23 (m, 2H), 3.9 (m, 2H), 3.9	DMSO	386.1 (M+1)	1.79	Method C	96	Method G2
272		HCl	413.47	¹ H NMR (300 MHz, DMSO) δ 12.30 (s, 1H), 9.57 (d, J = 2.1 Hz, 1H), 8.89 (d, J = 8.0 Hz, 1H), 8.77 – 8.67 (m, 2H), 8.63 (d, J = 7.7 Hz, 1H), 7.94 – 7.83 (m, 2H), 7.71 (t, J = 7.1 Hz, 1H), 7.66 – 7.52 (m, 3H), 7.25 (t, J = 7.4 Hz, 1H), 4.24 – 4.06 (m, 1H), 3	DMSO	414.2 (M+1)	2.36	Method C	100	Method G1
273		HCl	387.46	¹ H NMR (300 MHz, DMSO) δ 10.96 (s, 1H), 10.29 (s, 1H), 9.85 (s, 1H), 9.51 (d, J = 2.0 Hz, 1H), 8.71 – 8.62 (m, 2H), 8.47 (dd, J = 8.2, 0.9 Hz, 1H), 7.89 (d, J = 9.1 Hz, 1H), 7.67 – 7.62 (m, 2H), 7.60 (d, J = 2.5 Hz, 1H), 7.57 (d, J = 2.6 Hz, 1H), 7.55 – 7	DMSO	388.1 (M+1)	2.01	Method C	95	Method G1
274		HCl	399.45	¹ H NMR (300 MHz, DMSO) δ 12.45 (s, 1H), 9.57 (d, J = 1.3 Hz, 1H), 8.96 (dd, J = 8.4, 0.9 Hz, 1H), 8.89 (t, J = 5.2 Hz, 1H), 8.75 – 8.66 (m, 2H), 7.94 – 7.84 (m, 2H), 7.72 (t, J = 7.9 Hz, 1H), 7.65 – 7.52 (m, 3H), 7.24 (t, J = 7.6 Hz, 1H), 4.00 (s, 3H), 3	DMSO	400.3 (M+1)	2.21	Method C	100	Method G1

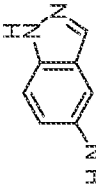
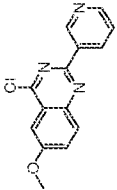
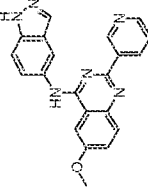
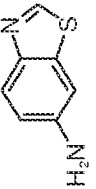
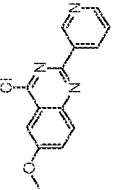
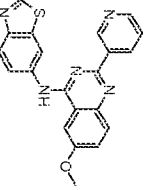
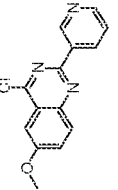
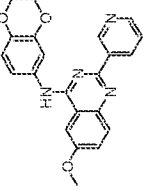
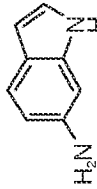
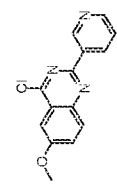
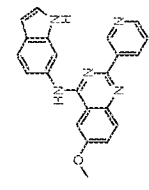
275		HCl	453.42	¹ H NMR (300 MHz, DMSO) δ 11.70 (s, 1H), 9.54 (d, J = 2.0 Hz, 1H), 9.40 (t, J = 6.1 Hz, 1H), 8.79 (d, J = 7.7 Hz, 1H), 8.73 – 8.63 (m, 2H), 7.89 (d, J = 9.8 Hz, 2H), 7.76 (t, J = 7.1 Hz, 1H), 7.64 – 7.50 (m, 3H), 7.30 (t, J = 7.1 Hz, 1H), 4.16 – 4.01 (m, 2H)	DMSO	454.0 (M+1)	2.28	Method C	100	Method G1
276			421.25	¹ H NMR (300 MHz, DMSO) δ 9.26 (d, J = 2.2 Hz, 1H), 8.71 – 8.62 (m, 2H), 8.46 (dt, J = 8.0, 1.9 Hz, 1H), 8.21 (dd, J = 8.9, 2.3 Hz, 1H), 8.03 (d, J = 8.9 Hz, 1H), 7.89 (s, 1H), 7.78 (dd, J = 7.6, 1.6 Hz, 1H), 7.72 – 7.62 (m, 1H), 7.56 – 7.45 (m, 3H), 7.32	DMSO	423.0 (M+1)	1.85	Method C	94	Method G1
277		HCl	447.49	¹ H NMR (300 MHz, DMSO) δ 11.25 (s, 1H), 10.46 (s, 1H), 9.45 (s, 1H), 8.93 (d, J = 7.9 Hz, 1H), 8.82 (d, J = 3.9 Hz, 1H), 8.22 (d, J = 8.1 Hz, 1H), 7.97 – 7.79 (m, 4H), 7.73 (t, J = 7.1 Hz, 1H), 7.62 (dd, J = 9.1, 2.5 Hz, 1H), 7.53 – 7.39 (m, 3H), 7.19 (t, 1H)	DMSO	448.1 (M+1)	2.67	Method C	100	Method G1
278		HCl	399.45	¹ H NMR (300 MHz, DMSO) δ 12.59 (s, 1H), 9.57 (d, J = 1.4 Hz, 1H), 9.03 (dd, J = 8.4, 0.9 Hz, 1H), 8.91 (d, J = 4.6 Hz, 1H), 8.76 – 8.64 (m, 2H), 7.88 (t, J = 7.8 Hz, 2H), 7.72 (t, J = 7.9 Hz, 1H), 7.63 – 7.53 (m, 3H), 7.22 (dd, J = 7.9, 1.1 Hz, 1H), 4.28	DMSO	400.1 (M+1)	2.5	Method C	100	Method G1

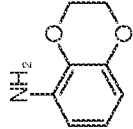
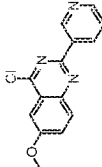
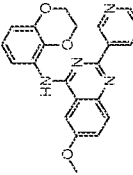
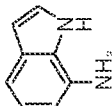
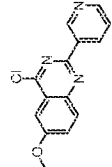
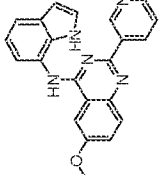
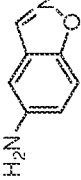
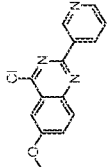
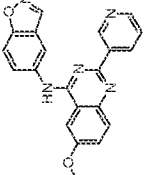
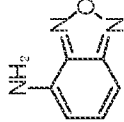
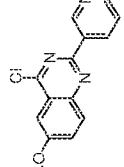
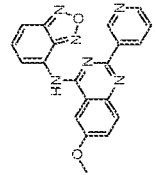
279		HCl	399.45	¹ H NMR (300 MHz, DMSO) δ 9.99 (s, 1H), 9.37 (d, J = 1.4 Hz, 1H), 8.61 (dd, J = 4.7, 1.7 Hz, 1H), 8.58 – 8.51 (m, 1H), 7.84 (dd, J = 9.0, 5.3 Hz, 3H), 7.65 – 7.42 (m, 4H), 7.38 (t, J = 7.5 Hz, 1H), 3.95 (s, 3H), 2.74 (s, 6H).	DMSO	400.1 (M+1)	2.2	Method C	100	Method G1
280		HCl	439.51	¹ H NMR (300 MHz, DMSO) δ 12.05 (s, 1H), 9.51 (d, J = 1.6 Hz, 1H), 9.04 (d, J = 8.1 Hz, 1H), 8.88 (dd, J = 5.2, 1.3 Hz, 1H), 8.60 (d, J = 7.0 Hz, 1H), 8.44 (d, J = 8.0 Hz, 1H), 8.00 – 7.87 (m, 2H), 7.83 (dd, J = 7.8, 1.4 Hz, 1H), 7.77 (d, J = 1.9 Hz, 1H).	DMSO	440.1 (M+1)	2.52	Method C	100	Method G1
281		HCl	403.41	¹ H NMR (300 MHz, DMSO) δ 10.98 (s, 1H), 9.50 (s, 1H), 8.72 – 8.55 (m, 3H), 8.38 (d, J = 8.2 Hz, 1H), 7.87 (d, J = 8.9 Hz, 1H), 7.72 – 7.45 (m, 4H), 7.19 – 7.07 (m, 1H), 3.96 (s, J = 5.2 Hz, 3H), 2.73 (d, J = 4.5 Hz, 3H).	DMSO	404.1 (M+1)	2.18	Method C	100	Method G1
282		HCl	417.44	¹ H NMR (300 MHz, DMSO) δ 10.96 (s, 1H), 9.50 (d, J = 1.9 Hz, 1H), 8.71 – 8.55 (m, 3H), 8.38 (d, J = 8.2 Hz, 1H), 7.86 (d, J = 9.0 Hz, 1H), 7.66 (dd, J = 14.8, 8.3 Hz, 1H), 7.60 – 7.47 (m, 3H), 7.21 – 7.05 (m, 1H), 4.24 (q, J = 6.9 Hz, 2H), 2.73 (d, J = 4.5 Hz, 3H).	DMSO	419.1 (M+1)	2.38	Method C	100	Method G1
283		HCl	439.39	¹ H NMR (300 MHz, DMSO) δ 13.10 (s, 1H), 9.88 (d, J = 1.1 Hz, 1H), 9.52 (d, J = 1.4 Hz, 1H), 8.75 – 8.58 (m, 3H), 8.26 (s, 1H), 8.15 (d, J = 8.2 Hz, 1H), 7.85 (d, J = 9.1 Hz, 1H), 7.59 – 7.47 (m, 3H), 7.45 (d, J = 2.5 Hz, 1H), 3.95 (s, 3H).	DMSO	440.1 (M+1)	2.25	Method C	100	Method G1

284		HCl	453.42	¹ H NMR (300 MHz, DMSO) δ 13.08 (s, 1H), 9.70 (s, 1H), 9.55 (d, J = 1.3 Hz, 1H), 8.77 - 8.60 (m, 3H), 8.26 (s, 1H), 8.15 (d, J = 8.5 Hz, 1H), 7.88 (d, J = 9.1 Hz, 1H), 7.62 - 7.39 (m, 4H), 4.21 (q, J = 7.1 Hz, 2H), 1.45 (t, J = 6.9 Hz, 3H).	DMSO	454.1 (M+1)	2.43	Method C	95	Method G1
285		HCl	410.43	¹ H NMR (300 MHz, DMSO) δ 12.87 (s, 1H), 9.50 (d, J = 1.5 Hz, 2H), 8.73 - 8.59 (m, 3H), 8.26 (s, 1H), 8.07 (d, J = 8.2 Hz, 1H), 7.85 (d, J = 9.1 Hz, 1H), 7.64 (dd, J = 8.1, 1.5 Hz, 1H), 7.59 - 7.49 (m, 2H), 7.42 (d, J = 2.4 Hz, 1H), 4.19 (q, J = 6.9 Hz, 2H).	DMSO	411.1 (M+1)	2.16	Method C	95	Method G1
286		HCl	397.43	¹ H NMR (300 MHz, DMSO) δ 13.63 (s, 1H), 9.55 (s, 1H), 9.13 (d, J = 8.3 Hz, 1H), 8.75 - 8.56 (m, 3H), 7.84 (d, J = 9.8 Hz, 1H), 7.69 - 7.48 (m, 4H), 7.01 (d, J = 7.4 Hz, 1H), 3.94 (s, 3H), 3.46 - 3.36 (m, 2H), 2.96 (t, J = 5.7 Hz, 2H).	DMSO	398.3 (M+1)	2.12	Method C	100	Method G1
287		HCl	411.46	¹ H NMR (300 MHz, DMSO) δ 13.61 (s, 1H), 9.56 (s, 1H), 9.14 (d, J = 8.1 Hz, 1H), 8.75 - 8.60 (m, 3H), 7.84 (d, J = 9.8 Hz, 1H), 7.70 - 7.46 (m, 4H), 7.02 (d, J = 7.2 Hz, 1H), 4.19 (q, J = 6.9 Hz, 3H), 3.48 - 3.37 (m, 2H), 2.96 (t, J = 6.1 Hz, 2H), 1.44 (t,	DMSO	412.4 (M+1)	2.4	Method C	100	Method G1

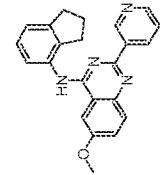
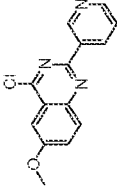
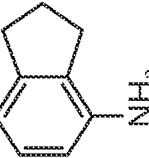
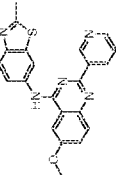
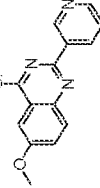
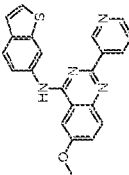
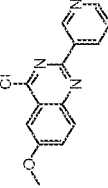
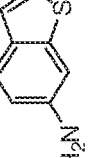
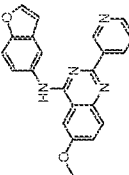
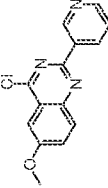
288		HCl	383.4	¹ H NMR (300 MHz, DMSO) δ 11.62 (s, 1H), 9.60 (d, J = 1.5 Hz, 1H), 9.02 (s, 1H), 8.88 (d, J = 8.0 Hz, 1H), 8.80 - 8.65 (m, 2H), 7.90 (d, J = 9.0 Hz, 1H), 7.75 (t, J = 7.9 Hz, 1H), 7.66 - 7.51 (m, 3H), 7.27 (d, J = 7.6 Hz, 1H), 4.47 (s, 2H), 3.97 (s, 3H).	DMSO	384.1 (M+1)	2.01	Method C	95	Method G1
289		HCl	421.4	¹ H NMR (300 MHz, DMSO) δ 12.07 (s, 1H), 9.48 (d, J = 1.6 Hz, 1H), 8.76 - 8.55 (m, 3H), 8.34 (s, 1H), 8.14 (s, 1H), 7.87 (d, J = 9.1 Hz, 1H), 7.62 - 7.49 (m, 2H), 7.38 (d, J = 2.5 Hz, 1H), 7.14 (ddd, J = 11.5, 9.0, 2.6 Hz, 1H), 4.19 (q, J = 6.9 Hz, 2H), 1.	DMSO	422.2 (M+1)	2.35	Method C	100	Method G1
290		HCl	412.44	¹ H NMR (300 MHz, DMSO) δ 9.82 (s, 1H), 9.50 (d, J = 2.0 Hz, 1H), 8.68 - 8.58 (m, 2H), 7.98 (d, J = 2.5 Hz, 1H), 7.85 (d, J = 1.8 Hz, 1H), 7.80 (d, J = 9.1 Hz, 1H), 7.55 - 7.43 (m, 3H), 7.23 (d, J = 8.4 Hz, 1H), 3.96 (s, 3H), 3.37 (d, J = 4.2 Hz, 6H).	DMSO	413.0 (M+1)	1.79	Method C		Method G1
291		HCl	426.47	¹ H NMR (300 MHz, DMSO) δ 9.78 (s, 1H), 9.50 (d, J = 2.0 Hz, 1H), 8.68 - 8.57 (m, 2H), 7.97 (d, J = 2.5 Hz, 1H), 7.85 (d, J = 1.8 Hz, 1H), 7.79 (d, J = 9.1 Hz, 1H), 7.55 - 7.43 (m, 3H), 7.22 (d, J = 8.4 Hz, 1H), 4.22 (q, J = 6.9 Hz, 2H), 3.37 (d, J = 4.3 Hz, 6H).	DMSO	427.1 (M+1)	1.92	Method C		Method G1

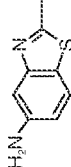
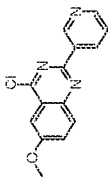
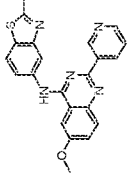
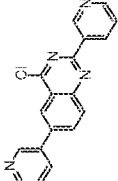
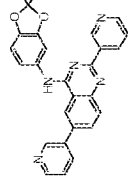
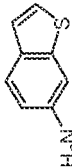
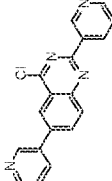
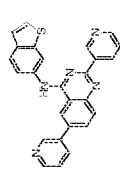
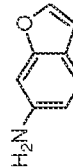
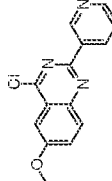
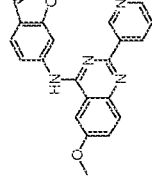
292			404.39	¹ H NMR (300 MHz, DMSO) δ 9.36 (d, J = 1.4 Hz, 1H), 8.66 (d, J = 2.7 Hz, 1H), 8.52 (d, J = 7.8 Hz, 1H), 7.97 (d, J = 8.9 Hz, 2H), 7.64 (d, J = 13.5 Hz, 2H), 7.50 (dd, J = 7.7, 4.2 Hz, 1H), 6.86 (t, J = 6.8 Hz, 1H), 6.70 (dd, J = 11.8, 1.7 Hz, 1H), 4.22 (q, 2H).	DMSO	405.1 (M+1)	1.98	Method C	95	Method G6
293			406.82	¹ H NMR (300 MHz, DMSO) δ 9.23 (d, J = 1.4 Hz, 1H), 8.63 (d, J = 3.6 Hz, 1H), 8.43 (d, J = 8.0 Hz, 1H), 8.00 (d, J = 9.1 Hz, 1H), 7.92 (s, 1H), 7.81 - 7.65 (m, 4H), 7.55 (dd, J = 8.4, 2.1 Hz, 1H), 7.48 (dd, J = 6.7, 3.8 Hz, 1H), 7.37 (s, 1H), 3.97 (s, 3H).	DMSO	407.1 (M+1)	1.88	Method C	100	Method G8
294		HCl	406.46	¹ H NMR (300 MHz, DMSO) δ 10.11 (s, 1H), 9.39 (d, J = 1.4 Hz, 1H), 8.63 (dd, J = 4.7, 1.7 Hz, 1H), 8.59 - 8.53 (m, 1H), 8.50 (dd, J = 8.2, 0.8 Hz, 1H), 8.03 (dd, J = 8.0, 1.4 Hz, 1H), 7.96 - 7.84 (m, 2H), 7.65 - 7.46 (m, 4H), 3.94 (s, 3H), 3.25 (s, 3H).	DMSO	407.1 (M+1)	2.13	Method C	95	Method G1

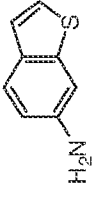
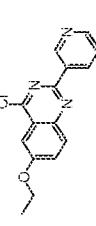
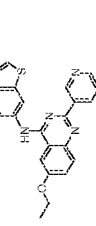
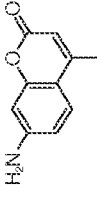
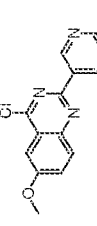
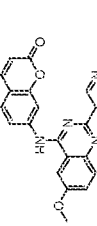
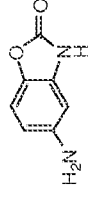
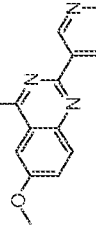
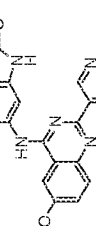
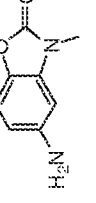
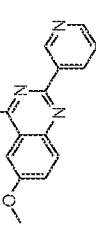
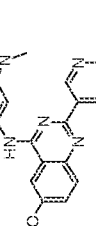
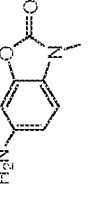
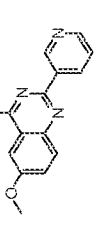
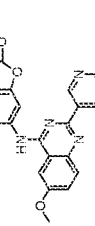
Number	Starting Material 1	Starting Material 2	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
295					¹ H NMR (400 MHz, DMSO) δ 13.12 (s, 1H), 9.90 (s, 1H), 9.47 (d, J = 1.4 Hz, 1H), 8.70 – 8.54 (m, 2H), 8.20 – 8.15 (m, 1H), 8.14 (s, 1H), 8.01 (d, J = 2.6 Hz, 1H), 7.88 – 7.75 (m, 2H), 7.65 (d, J = 8.9 Hz, 1H), 7.59 – 7.44 (m, 2H), 3.96 (s, 3H).	DMSO	>98	G1
296					¹ H NMR (400 MHz, DMSO) δ 10.04 (s, 1H), 9.54 – 9.47 (m, 1H), 9.36 (s, 1H), 6.72 (d, J = 2.0 Hz, 1H), 8.69 – 8.59 (m, 2H), 8.19 (d, J = 8.8 Hz, 1H), 8.06 – 7.97 (m, 2H), 7.86 (d, J = 9.1 Hz, 1H), 7.61 – 7.47 (m, 2H), 3.99 (s, 3H).	DMSO	>98	G1
297					¹ H NMR (400 MHz, DMSO) δ 9.64 (s, 1H), 9.55 – 9.45 (m, 1H), 8.72 – 8.60 (m, 2H), 7.95 (d, J = 2.7 Hz, 1H), 7.82 (d, J = 9.1 Hz, 1H), 7.58 – 7.48 (m, 3H), 7.34 (dd, J = 8.7, 2.5 Hz, 1H), 6.96 (d, J = 8.7 Hz, 1H), 4.39 – 4.23 (m, 4H), 3.97 (s, 3H).	DMSO	>98	G1
298					¹ H NMR (400 MHz, DMSO) δ 11.21 (s, 1H), 9.79 (s, 1H), 9.53 (d, J = 1.4 Hz, 1H), 8.75 – 8.56 (m, 2H), 8.07 (s, 1H), 8.03 (d, J = 2.7 Hz, 1H), 7.82 (d, J = 9.1 Hz, 1H), 7.61 (d, J = 8.5 Hz, 1H), 7.56 – 7.47 (m, 2H), 7.43 (dd, J = 6.5, 1.8 Hz, 1H), 7.39 – 7.34 (m, 1H), 6.50 – 6.41 (m, 1H), 3.99 (s, 3H).	DMSO	>98	G1

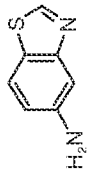
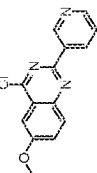
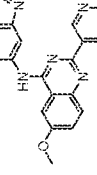
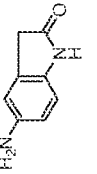
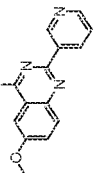
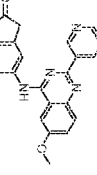
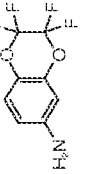
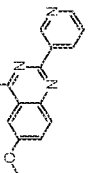
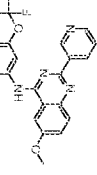
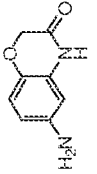
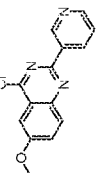
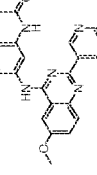
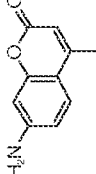
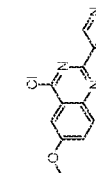
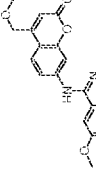
299				¹ H NMR (400 MHz, DMSO) δ 9.54 (s, 1H), 9.42 – 9.34 (m, 1H), 8.61 (dd, J = 4.7, 1.7 Hz, 1H), 8.57 – 8.47 (m, 1H), 7.93 (d, J = 2.7 Hz, 1H), 7.81 (d, J = 9.1 Hz, 1H), 7.56 – 7.44 (m, 2H), 7.19 (dd, J = 7.9, 1.5 Hz, 1H), 7.00 – 6.91 (m, 1H), 6.86 (dd, J = 8.2, 1.6 Hz, 1H), 4.34 – 4.17 (m, 4H), 3.95 (s, 3H).	DMSO	>98	G1
300				¹ H NMR (400 MHz, DMSO) δ 10.76 (s, 1H), 10.01 (s, 1H), 9.17 – 9.06 (m, 1H), 8.53 (dd, J = 4.7, 1.7 Hz, 1H), 8.36 – 8.27 (m, 1H), 8.05 (d, J = 2.7 Hz, 1H), 7.84 (d, J = 9.1 Hz, 1H), 7.60 – 7.43 (m, 2H), 7.36 (ddd, J = 8.0, 4.8, 0.8 Hz, 1H), 7.32 – 7.28 (m, 1H), 7.25 (d, J = 6.8 Hz, 1H), 7.16 – 7.04 (m, 1H), 6.54 (dd, J = 3.0, 1.9 Hz, 1H), 3.96 (s, 3H).	DMSO	>98	G1
301				¹ H NMR (400 MHz, DMSO) δ 10.02 (s, 1H), 9.48 (d, J = 1.5 Hz, 1H), 9.33 (d, J = 1.0 Hz, 1H), 8.69 – 8.59 (m, 2H), 8.37 (d, J = 1.7 Hz, 1H), 8.10 (dd, J = 9.0, 2.1 Hz, 1H), 8.00 (d, J = 2.7 Hz, 1H), 7.90 (d, J = 9.0 Hz, 1H), 7.86 (d, J = 9.1 Hz, 1H), 7.57 (dd, J = 9.1, 2.7 Hz, 1H), 7.54 – 7.47 (m, 1H), 3.99 (s, 3H).	DMSO	>98	G1
302				¹ H NMR (400 MHz, DMSO) δ 10.52 (s, 1H), 9.16 (dd, J = 2.2, 0.8 Hz, 1H), 8.60 (dd, J = 4.7, 1.7 Hz, 1H), 8.45 – 8.36 (m, 1H), 8.03 (d, J = 2.7 Hz, 1H), 7.97 – 7.90 (m, 2H), 7.88 (d, J = 7.0 Hz, 1H), 7.77 (dd, J = 9.0, 7.1 Hz, 1H), 7.61 (dd, J = 9.1, 2.7 Hz, 1H), 7.44 (ddd, J = 8.0, 4.8, 0.8 Hz, 1H), 4.01 (s, 3H).	DMSO	>98	G1

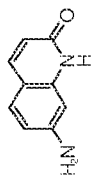
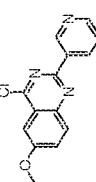
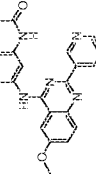
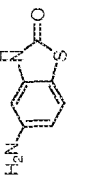
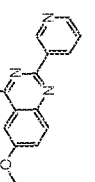
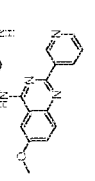
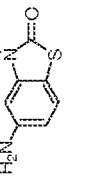
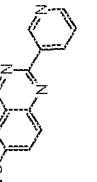
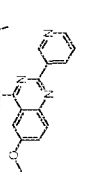
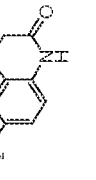
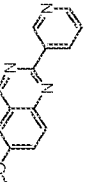
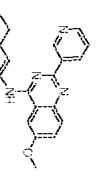
303				¹ H NMR (400 MHz, DMSO) δ 9.86 (s, 1H), 9.60 – 9.56 (m, 1H), 8.72 – 8.67 (m, 1H), 8.65 (dd, J = 4.8, 1.7 Hz, 1H), 8.26 – 8.21 (m, 1H), 8.05 (d, J = 2.7 Hz, 1H), 7.83 (d, J = 9.1 Hz, 1H), 7.62 (d, J = 8.4 Hz, 1H), 7.56 – 7.50 (m, 2H), 7.46 (dd, J = 8.5, 1.8 Hz, 1H), 7.35 (d, J = 3.1 Hz, 1H), 6.46 (dd, J = 3.1, 0.8 Hz, 1H), 3.99 (s, 3H), 3.86 (s, 3H).	DMSO	>98	G1
304				¹ H NMR (400 MHz, DMSO) δ 9.96 (s, 1H), 9.51 – 9.45 (m, 1H), 8.68 – 8.59 (m, 2H), 8.57 – 8.52 (m, 1H), 8.02 – 7.86 (m, 3H), 7.87 – 7.80 (m, 1H), 7.58 – 7.46 (m, 2H), 3.98 (s, 3H), 2.83 (s, 3H).	DMSO	>98	G1
305				¹ H NMR (400 MHz, DMSO) δ 9.94 (s, 1H), 9.51 – 9.43 (m, 1H), 8.68 – 8.58 (m, 2H), 8.39 (d, J = 2.0 Hz, 1H), 7.99 (d, J = 2.7 Hz, 1H), 7.87 – 7.80 (m, 2H), 7.78 (d, J = 8.6 Hz, 1H), 7.54 (dd, J = 9.1, 2.7 Hz, 1H), 7.51 (ddd, J = 8.0, 4.8, 0.8 Hz, 1H), 4.19 (s, 3H), 3.98 (s, 3H).	DMSO	>98	G1
306				¹ H NMR (400 MHz, DMSO) δ 9.72 (s, 1H), 9.52 – 9.47 (m, 1H), 8.67 – 8.60 (m, 2H), 7.98 (d, J = 2.7 Hz, 1H), 7.82 (d, J = 9.1 Hz, 1H), 7.79 – 7.74 (m, 1H), 7.68 – 7.62 (m, 1H), 7.55 – 7.48 (m, 2H), 7.32 (d, J = 8.1 Hz, 1H), 3.97 (s, 3H), 2.99 – 2.88 (m, 4H), 2.15 – 2.04 (m, 2H).	DMSO	>98	G1

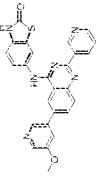
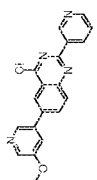
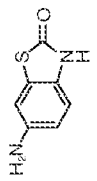
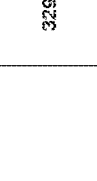
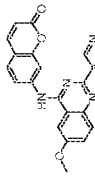
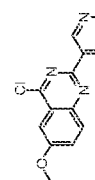
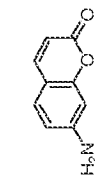
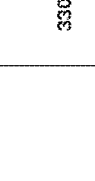
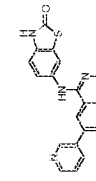
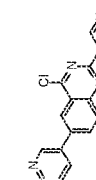
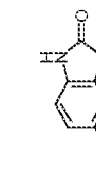
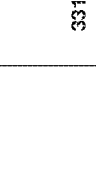
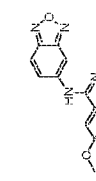
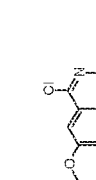
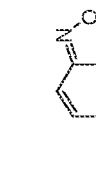
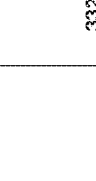
307				¹ H NMR (400 MHz, CDCl ₃) δ 9.74 (s, 1H), 9.37 – 9.31 (m, 1H), 8.60 (dd, J = 4.7, 1.7 Hz, 1H), 8.53 – 8.46 (m, 1H), 7.95 (d, J = 2.7 Hz, 1H), 7.82 (d, J = 9.1 Hz, 1H), 7.52 (dd, J = 9.1, 2.7 Hz, 1H), 7.47 (ddd, J = 8.0, 4.8, 0.8 Hz, 1H), 7.34 (d, J = 7.0 Hz, 1H), 7.31 – 7.24 (m, 1H), 7.22 (d, J = 7.1 Hz, 1H), 3.96 (s, 3H), 2.99 (t, J = 7.3 Hz, 2H), 2.82 (t, J = 7.4 Hz, 2H), 2.06 – 1.96 (m, 2H).	DMSO	>98	G1
306				¹ H NMR (400 MHz, DMSO) δ 10.45 (s, 1H), 9.50 – 9.44 (m, 1H), 8.94 – 8.87 (m, 1H), 8.83 (dd, J = 5.1, 1.5 Hz, 1H), 8.54 (d, J = 2.0 Hz, 1H), 8.13 (d, J = 2.4 Hz, 1H), 8.02 (d, J = 8.7 Hz, 1H), 7.97 – 7.89 (m, 2H), 7.83 (dd, J = 8.0, 5.2 Hz, 1H), 7.63 (dd, J = 9.1, 2.7 Hz, 1H), 4.01 (s, 3H), 2.83 (s, 3H).	DMSO	>98	G1
309				¹ H NMR (400 MHz, DMSO) δ 10.64 (s, 1H), 9.47 (d, J = 1.5 Hz, 1H), 8.94 (d, J = 8.1 Hz, 1H), 8.86 (dd, J = 5.2, 1.5 Hz, 1H), 8.54 – 8.48 (m, 1H), 8.23 – 8.16 (m, 1H), 8.04 – 7.94 (m, 2H), 7.92 – 7.83 (m, 2H), 7.78 (d, J = 5.4 Hz, 1H), 7.65 (dd, J = 9.1, 2.6 Hz, 1H), 7.51 (dd, J = 5.4, 0.7 Hz, 1H), 4.02 (s, 3H).	DMSO	>98	G1
310				¹ H NMR (400 MHz, DMSO) δ 10.86 (s, 1H), 9.43 (d, J = 1.6 Hz, 1H), 8.94 (d, J = 7.5 Hz, 1H), 8.89 (dd, J = 5.2, 1.5 Hz, 1H), 8.30 – 8.20 (m, 1H), 8.13 – 7.99 (m, 3H), 7.94 – 7.84 (m, 1H), 7.74 (d, J = 1.3 Hz, 2H), 7.67 (dd, J = 9.1, 2.5 Hz, 1H), 7.08 (d, J = 2.2 Hz, 1H), 4.02 (s, 3H).	DMSO	>98	G1

311				HCl	¹ H NMR (400 MHz, DMSO) δ 10.53 (s, 1H), 9.48 (d, J = 1.5 Hz, 1H), 8.93 (d, J = 7.7 Hz, 1H), 8.85 (dd, J = 5.2, 1.5 Hz, 1H), 8.44 (d, J = 1.9 Hz, 1H), 8.20–8.15 (m, 1H), 8.13 (d, J = 8.7 Hz, 1H), 7.97 (d, J = 9.1 Hz, 1H), 7.92 (dd, J = 8.7, 2.0 Hz, 1H), 7.86 (dd, J = 7.8, 5.1 Hz, 1H), 7.64 (dd, J = 9.1, 2.6 Hz, 1H), 4.02 (s, 3H), 2.85 (s, 3H).	DMSO	>98	G1
312				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.85 (s, 1H), 9.54 (d, J = 1.5 Hz, 1H), 9.47 (s, 1H), 9.32 (s, 1H), 9.03 (d, J = 8.0 Hz, 1H), 8.93–8.87 (m, 1H), 8.86–8.78 (m, 2H), 8.44 (dd, J = 8.8, 1.9 Hz, 1H), 8.14 (d, J = 2.1 Hz, 1H), 8.10 (d, J = 8.7 Hz, 1H), 7.99–7.89 (m, 2H), 7.78–7.71 (m, 1H), 7.55 (d, J = 8.7 Hz, 1H).	DMSO	>98	G1
313				2 HCl	¹ H NMR (400 MHz, DMSO) δ 11.07 (s, 1H), 9.54 (d, J = 1.6 Hz, 1H), 9.51 (d, J = 2.0 Hz, 1H), 9.46–9.39 (m, 1H), 9.10–9.01 (m, 1H), 8.97–8.88 (m, 2H), 8.86 (dd, J = 5.3, 1.3 Hz, 1H), 8.63–8.55 (m, 1H), 8.48 (dd, J = 8.8, 1.9 Hz, 1H), 8.15 (d, J = 8.7 Hz, 1H), 8.04–7.90 (m, 4H), 7.79 (d, J = 5.4 Hz, 1H), 7.52 (dd, J = 5.4, 0.7 Hz, 1H).	DMSO	>98	G1
314				HCl	¹ H NMR (400 MHz, DMSO) δ 10.49 (s, 1H), 9.48 (d, J = 1.5 Hz, 1H), 8.93 (d, J = 7.9 Hz, 1H), 8.85 (dd, J = 5.2, 1.5 Hz, 1H), 8.24 (s, 1H), 8.16 (d, J = 2.5 Hz, 1H), 8.05 (d, J = 2.2 Hz, 1H), 7.96 (d, J = 9.1 Hz, 1H), 7.87 (dd, J = 8.1, 5.3 Hz, 1H), 7.77 (d, J = 8.4 Hz, 1H), 7.72 (dd, J = 8.4, 1.8 Hz, 1H), 7.64 (dd, J = 9.1, 2.6 Hz, 1H), 7.03 (dd, J = 2.2, 1.0 Hz, 1H), 4.02 (s, 3H).	DMSO	>98	G1

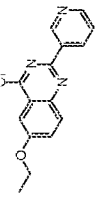
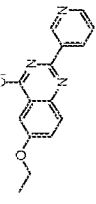
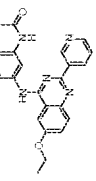
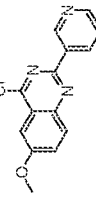
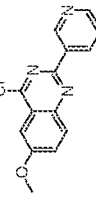
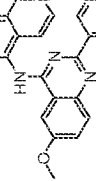
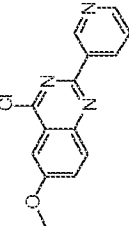
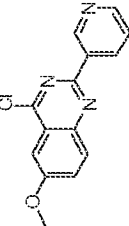
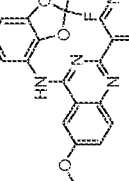
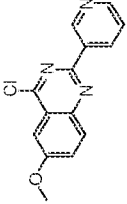
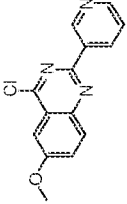
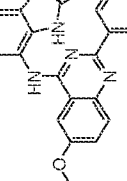
315				HCl	¹ H NMR (400 MHz, DMSO) δ 10.72 (s, 1H), 9.46 (d, J = 1.5 Hz, 1H), 8.96 (d, J = 8.2 Hz, 1H), 8.87 (dd, J = 5.2, 1.5 Hz, 1H), 8.50 (d, J = 1.8 Hz, 1H), 8.21 (d, J = 2.4 Hz, 1H), 8.06 – 7.96 (m, 2H), 7.93 – 7.84 (m, 2H), 7.78 (d, J = 5.4 Hz, 1H), 7.63 (dd, J = 9.1, 2.6 Hz, 1H), 7.51 (dd, J = 5.4, 0.7 Hz, 1H), 4.29 (q, J = 7.0 Hz, 2H), 1.46 (t, J = 7.0 Hz, 3H).	DMSO	>98	G1
316					¹ H NMR (400 MHz, DMSO) δ 10.04 (s, 1H), 9.60 – 9.52 (m, 1H), 8.75 – 8.63 (m, 2H), 8.16 (d, J = 2.0 Hz, 1H), 8.04 – 7.95 (m, 2H), 7.93 – 7.83 (m, 2H), 7.63 – 7.53 (m, 2H), 6.32 (d, J = 1.2 Hz, 1H), 4.01 (s, 3H), 2.47 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)
317					¹ H NMR (400 MHz, DMSO) δ 11.82 (s, 1H), 10.06 (s, 1H), 9.53 – 9.46 (m, 1H), 8.81 – 8.70 (m, 2H), 8.04 (d, J = 2.6 Hz, 1H), 7.88 (d, J = 9.1 Hz, 1H), 7.81 (d, J = 2.1 Hz, 1H), 7.67 (dd, J = 7.9, 5.0 Hz, 1H), 7.57 (dd, J = 9.1, 2.7 Hz, 1H), 7.53 (dd, J = 8.7, 2.1 Hz, 1H), 7.39 (d, J = 8.6 Hz, 1H), 4.00 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)
318					¹ H NMR (400 MHz, DMSO) δ 9.89 (s, 1H), 9.57 – 9.48 (m, 1H), 8.72 – 8.58 (m, 2H), 8.01 (d, J = 2.1 Hz, 1H), 7.99 (d, J = 2.7 Hz, 1H), 7.85 (d, J = 9.1 Hz, 1H), 7.59 – 7.49 (m, 3H), 7.43 (d, J = 8.6 Hz, 1H), 3.98 (s, 3H), 3.41 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)
319					¹ H NMR (400 MHz, DMSO) δ 9.88 (s, 1H), 9.52 – 9.44 (m, 1H), 8.71 – 8.59 (m, 2H), 8.02 – 7.91 (m, 2H), 7.83 (d, J = 9.1 Hz, 1H), 7.66 (dd, J = 8.4, 2.0 Hz, 1H), 7.58 – 7.48 (m, 2H), 7.36 (d, J = 8.4 Hz, 1H), 3.96 (s, 3H), 3.40 (s, 3H).	DMSO	>96	G1 (0.1 N HCl added)

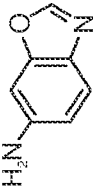
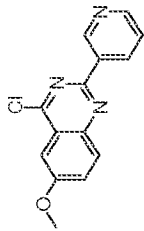
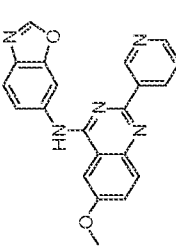
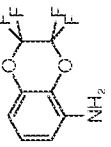
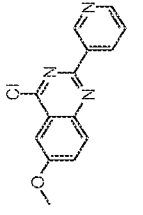
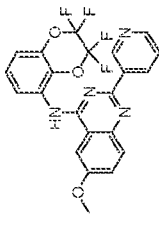
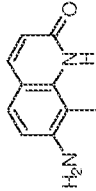
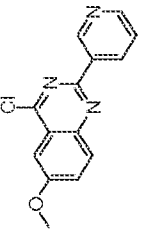
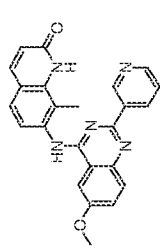
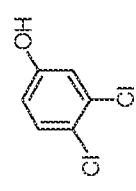
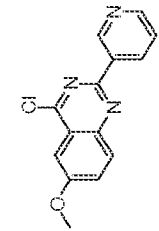
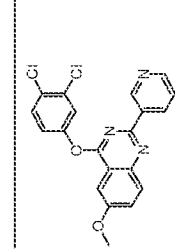
320				¹ H NMR (400 MHz, DMSO) δ 10.01 (s, 1H), 9.58–9.51 (m, 1H), 9.45 (s, 1H), 8.75 (d, J = 1.9 Hz, 1H), 8.71–8.63 (m, 2H), 8.25 (d, J = 8.7 Hz, 1H), 8.08–8.00 (m, 2H), 7.86 (d, J = 9.1 Hz, 1H), 7.61–7.50 (m, 2H), 4.00 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)
321				¹ H NMR (400 MHz, DMSO) δ 10.43 (s, 1H), 9.72 (s, 1H), 9.51–9.45 (m, 1H), 8.67–8.57 (m, 2H), 7.96 (d, J = 2.7 Hz, 1H), 7.81 (d, J = 9.1 Hz, 1H), 7.73–7.68 (m, 1H), 7.64 (dd, J = 8.3, 2.1 Hz, 1H), 7.56–7.48 (m, 2H), 6.93 (d, J = 8.3 Hz, 1H), 3.97 (s, 3H), 3.58 (s, 2H).	DMSO	>98	G1 (0.1 N HCl added)
322				¹ H NMR (400 MHz, DMSO) δ 9.95 (s, 1H), 9.54–9.45 (m, 1H), 8.67 (dd, J = 4.7, 1.7 Hz, 1H), 8.66–8.60 (m, 1H), 8.15 (d, J = 2.4 Hz, 1H), 7.94 (d, J = 2.6 Hz, 1H), 7.89–7.83 (m, 2H), 7.63–7.52 (m, 3H), 3.96 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)
323				¹ H NMR (400 MHz, DMSO) δ 10.99 (s, 1H), 9.75 (s, 1H), 9.55–9.49 (m, 1H), 8.69–8.62 (m, 2H), 7.96 (d, J = 2.7 Hz, 1H), 7.83 (d, J = 9.1 Hz, 1H), 7.58–7.49 (m, 3H), 7.35 (dd, J = 8.7, 2.5 Hz, 1H), 7.06 (d, J = 8.6 Hz, 1H), 4.62 (s, 2H), 3.97 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)
324				¹ H NMR (400 MHz, DMSO) δ 10.24 (s, 1H), 9.58–9.51 (m, 1H), 8.92–8.84 (m, 1H), 8.80 (dd, J = 5.0, 1.5 Hz, 1H), 8.16 (d, J = 2.0 Hz, 1H), 8.07 (d, J = 2.6 Hz, 1H), 7.98 (dd, J = 8.8, 2.1 Hz, 1H), 7.91 (d, J = 9.1 Hz, 1H), 7.83–7.72 (m, 2H), 7.61 (dd, J = 9.1, 2.6 Hz, 1H), 6.37 (s, 1H), 4.75 (d, J = 1.2 Hz, 2H), 4.01 (s, 3H), 3.47 (s, 3H).	HCl	>98	G1 (0.1 N HCl added)

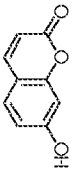
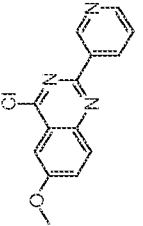
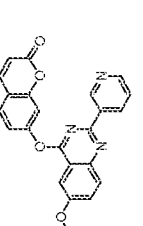
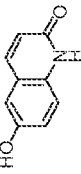
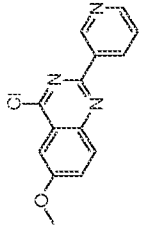
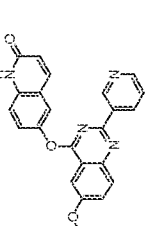
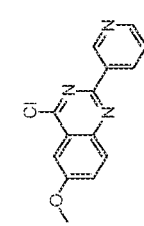
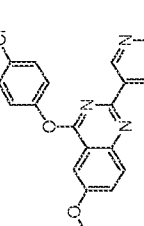
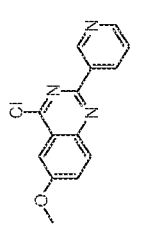
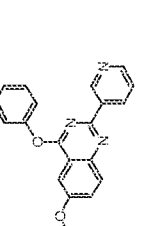
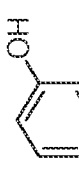
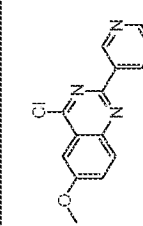
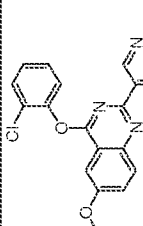
325				HCl	¹ H NMR (400 MHz, DMSO) δ 12.01 (s, 1H), 10.15 (s, 1H), 9.58 (d, J = 1.5 Hz, 1H), 8.94 – 8.86 (m, 1H), 8.78 (dd, J = 5.0, 1.5 Hz, 1H), 8.07 (d, J = 2.6 Hz, 1H), 8.04 – 7.98 (m, 1H), 7.91 (dd, J = 9.3, 2.2 Hz, 2H), 7.78 – 7.69 (m, 2H), 7.66 (dd, J = 8.5, 2.0 Hz, 1H), 7.61 (dd, J = 9.1, 2.7 Hz, 1H), 6.44 (d, J = 9.5 Hz, 1H), 4.01 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)
326				HCl	¹ H NMR (400 MHz, DMSO) δ 12.19 (s, 1H), 10.21 (s, 1H), 9.58 – 9.51 (m, 1H), 8.94 – 8.86 (m, 1H), 8.81 (dd, J = 5.1, 1.6 Hz, 1H), 8.10 (d, J = 2.6 Hz, 1H), 7.96 – 7.88 (m, 2H), 7.79 (dd, J = 7.9, 5.1 Hz, 1H), 7.68 – 7.56 (m, 3H), 4.00 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)
327				HCl	¹ H NMR (400 MHz, DMSO) δ 10.43 (s, 1H), 9.57 – 9.50 (m, 1H), 9.07 – 8.98 (m, 1H), 8.88 (dd, J = 5.2, 1.4 Hz, 1H), 8.16 (d, J = 2.5 Hz, 1H), 8.06 (d, J = 1.9 Hz, 1H), 8.00 – 7.89 (m, 2H), 7.75 (d, J = 8.5 Hz, 1H), 7.67 (dd, J = 8.5, 2.0 Hz, 1H), 7.63 (dd, J = 9.1, 2.6 Hz, 1H), 4.02 (s, 3H), 3.47 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)
328				HCl	¹ H NMR (400 MHz, DMSO) δ 10.68 (s, 1H), 10.24 (s, 1H), 9.45 (d, J = 1.6 Hz, 1H), 9.05 – 8.95 (m, 1H), 8.92 (dd, J = 5.2, 1.5 Hz, 1H), 8.25 – 8.16 (m, 1H), 8.09 – 7.99 (m, 1H), 7.98 – 7.89 (m, 1H), 7.71 – 7.60 (m, 3H), 6.96 (d, J = 8.3 Hz, 1H), 4.00 (s, 3H), 2.99 (t, J = 7.5 Hz, 2H), 2.57 – 2.52 (m, 2H).	DMSO	>98	G1 (0.1 N HCl added)

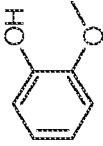
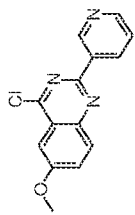
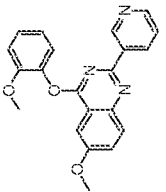
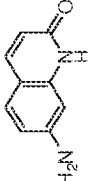
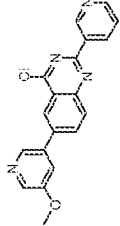
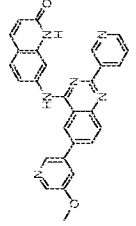
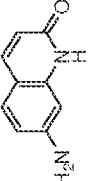
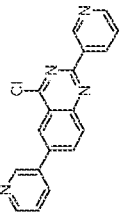
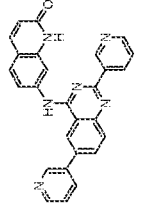
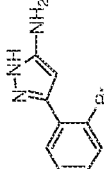
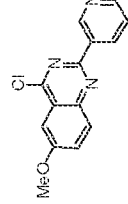
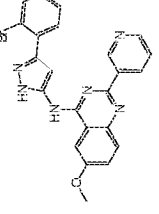
329					2 HCl	¹ H NMR (400 MHz, DMSO) δ 12.05 (s, 1H), 10.98 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 9.32 (s, 1H), 9.03 (d, J = 8.2 Hz, 1H), 8.98 (d, J = 1.7 Hz, 1H), 8.92 (dd, J = 5.3, 1.4 Hz, 1H), 8.53 (d, J = 2.6 Hz, 1H), 8.46 (dd, J = 8.8, 1.8 Hz, 1H), 8.30 (s, 1H), 8.13 (d, J = 2.1 Hz, 1H), 8.09 (d, J = 8.7 Hz, 1H), 7.95 (dd, J = 7.7, 5.3 Hz, 1H), 7.82 (dd, J = 8.6, 2.1 Hz, 1H), 7.26 (d, J = 8.6 Hz, 1H), 4.96 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)
330					HCl	¹ H NMR (400 MHz, DMSO) δ 10.40 (s, 1H), 9.54 (d, J = 1.5 Hz, 1H), 9.05 (s, 1H), 8.97 (m, 1H), 8.87 (dd, J = 5.2, 1.5 Hz, 1H), 8.15 (m, 3H), 8.00 (m, 3H), 7.81 (d, J = 8.5 Hz, 1H), 7.64 (dd, J = 9.1, 2.7 Hz, 1H), 6.43 (d, J = 9.4 Hz, 1H), 4.02 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)
331					2 HCl	¹ H NMR (400 MHz, DMSO) δ 12.05 (s, 1H), 10.77 (s, 1H), 9.51 (d, J = 1.5 Hz, 1H), 9.43 (m, 1H), 9.35 (m, 1H), 9.31 (m, 1H), 9.04 (m, 1H), 8.93 (m, 1H), 8.88 (dd, J = 5.2, 1.5 Hz, 1H), 8.81 (dd, J = 5.1, 1.4 Hz, 1H), 8.73 (d, J = 8.4 Hz, 1H), 8.42 (dd, J = 8.8, 1.8 Hz, 1H), 8.13 (d, J = 2.1 Hz, 1H), 8.09 (d, J = 8.7 Hz, 1H), 7.94 (m, 2H), 7.83 (m, 2H), 7.80 (dd, J = 8.6, 2.2 Hz, 1H), 7.27 (d, J = 8.6 Hz, 1H).	DMSO	>98	G1 (0.1 N HCl added)
332					HCl	¹ H NMR (400 MHz, DMSO) δ 10.57 (s, 1H), 9.55 (d, J = 1.6 Hz, 1H), 9.09 (s, 1H), 8.90 (dd, J = 5.3, 1.5 Hz, 1H), 8.65 (m, 1H), 8.59 (m, 1H), 8.24 (m, 2H), 8.16 (m, 2H), 8.14 (dd, J = 9.6, 0.8 Hz, 1H), 8.03 (m, 2H), 7.92 (m, 2H), 7.85 (dd, J = 9.1, 2.7 Hz, 1H), 4.03 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)

333			HCl	¹ H NMR (400 MHz, DMSO) δ 11.81 (s, 1H), 10.31 (s, 1H), 9.58 (d, J = 1.5 Hz, 1H), 9.08 (d, J = 7.9 Hz, 1H), 8.87 (dd, J = 5.2, 1.5 Hz, 1H), 8.13 (d, J = 2.6 Hz, 1H), 8.04 (d, J = 2.0 Hz, 1H), 7.98 – 7.88 (m, 2H), 7.81 (d, J = 8.8 Hz, 1H), 7.69 (dd, J = 8.8, 2.1 Hz, 1H), 7.63 (dd, J = 9.1, 2.7 Hz, 1H), 6.36 (s, 1H), 4.02 (s, 3H), 2.46 (d, J = 1.1 Hz, 3H).	DMSO	>98	G1 (0.1 N HCl added)
334			HCl	¹ H NMR (400 MHz, DMSO) δ 10.51 (s, 1H), 9.53 (d, J = 1.6 Hz, 1H), 9.08 – 9.02 (m, 1H), 8.90 (dd, J = 5.3, 1.5 Hz, 1H), 8.63 – 8.59 (m, 1H), 8.22 – 8.10 (m, 3H), 8.00 – 7.92 (m, 2H), 7.63 (dd, J = 9.1, 2.6 Hz, 1H), 4.30 (q, J = 7.0 Hz, 2H), 1.46 (t, J = 7.0 Hz, 3H).	DMSO	>98	G1 (0.1 N HCl added)
335			HCl	¹ H NMR (400 MHz, DMSO) δ 11.99 (s, 1H), 10.34 (s, 1H), 9.62 – 9.52 (m, 1H), 9.11 (d, J = 8.2 Hz, 1H), 8.90 (d, J = 4.4 Hz, 1H), 8.13 (d, J = 2.6 Hz, 1H), 8.06 (d, J = 1.8 Hz, 1H), 8.00 – 7.88 (m, 3H), 7.75 (d, J = 8.6 Hz, 1H), 7.68 – 7.60 (m, 2H), 6.46 (d, J = 9.5 Hz, 1H), 4.29 (q, J = 7.0 Hz, 2H), 1.46 (t, J = 7.0 Hz, 3H).	DMSO	>98	G1 (0.1 N HCl added)
336			HCl	¹ H NMR (400 MHz, DMSO) δ 11.02 (s, 1H), 10.49 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 9.02 (d, J = 8.1 Hz, 1H), 8.90 (dd, J = 5.2, 1.5 Hz, 1H), 6.15 (d, J = 2.3 Hz, 1H), 8.00 (d, J = 9.1 Hz, 1H), 7.93 (dd, J = 7.8, 5.4 Hz, 1H), 7.63 (dd, J = 9.1, 2.6 Hz, 1H), 7.54 (d, J = 2.4 Hz, 1H), 7.35 (dd, J = 8.7, 2.5 Hz, 1H), 7.08 (d, J = 6.6 Hz, 1H), 4.64 (s, 2H), 4.28 (q, J = 6.9 Hz, 2H), 1.45 (t, J = 7.0 Hz, 3H).	DMSO	>98	G1 (0.1 N HCl added)

337				HCl	¹ H NMR (400 MHz, DMSO) δ 10.36 (s, 1H), 10.25 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 8.98 (d, J = 8.1 Hz, 1H), 8.86 (dd, J = 5.2, 1.5 Hz, 1H), 8.10 (d, J = 2.4 Hz, 1H), 7.94 (d, J = 9.1 Hz, 1H), 7.87 (dd, J = 7.7, 5.2 Hz, 1H), 7.61 (dd, J = 9.1, 2.6 Hz, 1H), 7.47 (d, J = 1.9 Hz, 1H), 7.36 (dd, J = 8.1, 2.1 Hz, 1H), 7.28 (d, J = 8.1 Hz, 1H), 4.27 (q, J = 7.0 Hz, 2H), 2.93 (t, J = 7.5 Hz, 2H), 2.56 – 2.52 (m, 2H), 1.45 (t, J = 7.0 Hz, 3H).	DMSO	>98	G1 (0.1 N HCl added)
338					¹ H NMR (400 MHz, DMSO) δ 10.25 (s, 1H), 9.50 – 9.39 (m, 1H), 9.03 (dd, J = 2.1, 0.7 Hz, 1H), 8.52 (dd, J = 4.7, 1.7 Hz, 1H), 8.48 (d, J = 6.0 Hz, 1H), 8.31 – 8.24 (m, 1H), 8.16 (d, J = 8.2 Hz, 1H), 8.12 (d, J = 2.7 Hz, 1H), 8.04 (d, J = 7.3 Hz, 1H), 7.91 – 7.78 (m, 3H), 7.58 (dd, J = 9.1, 2.7 Hz, 1H), 7.35 (ddd, J = 8.0, 4.8, 0.7 Hz, 1H), 4.01 (s, 3H).	DMSO	>98	J2
339					¹ H NMR (400 MHz, DMSO) δ 10.23 (s, 1H), 9.37 (dd, J = 2.1, 0.7 Hz, 1H), 8.62 (dd, J = 4.7, 1.7 Hz, 1H), 8.57 – 8.49 (m, 1H), 7.93 (d, J = 2.7 Hz, 1H), 7.89 (d, J = 9.1 Hz, 1H), 7.58 (dd, J = 9.1, 2.7 Hz, 1H), 7.47 (ddd, J = 8.0, 4.8, 0.8 Hz, 1H), 7.40 – 7.30 (m, 3H), 3.96 (s, 3H).	DMSO	>98	J2
340					¹ H NMR (400 MHz, DMSO) δ 11.31 (s, 1H), 9.56 (s, 1H), 9.54 (dd, J = 2.1, 0.7 Hz, 1H), 8.80 – 8.66 (m, 2H), 8.14 (d, J = 9.2 Hz, 1H), 7.77 (dd, J = 9.2, 2.8 Hz, 1H), 7.58 (ddd, J = 8.0, 4.8, 0.8 Hz, 1H), 7.36 (d, J = 2.8 Hz, 1H), 7.03 – 6.94 (m, 1H), 6.68 (dd, J = 7.8, 0.8 Hz, 1H), 6.56 (dd, J = 8.2, 0.8 Hz, 1H), 3.86 (s, 3H).	DMSO	>98	J2

341					¹ H NMR (400 MHz, DMSO) δ 10.00 (s, 1H), 9.54 – 9.49 (m, 1H), 8.75 (s, 1H), 8.69 – 8.62 (m, 2H), 8.50 (d, J = 1.7 Hz, 1H), 8.02 (d, J = 2.7 Hz, 1H), 7.91 – 7.85 (m, 2H), 7.83 (dd, J = 8.6, 1.9 Hz, 1H), 7.60 – 7.51 (m, 2H), 4.00 (s, 3H).	DMSO	>98	J2
342					¹ H NMR (400 MHz, DMSO) δ 10.08 (s, 1H), 9.34 – 9.27 (m, 1H), 8.60 (dd, J = 4.7, 1.7 Hz, 1H), 8.51 – 8.44 (m, 1H), 7.94 (d, J = 2.7 Hz, 1H), 7.88 (d, J = 9.1 Hz, 1H), 7.58 (dd, J = 9.1, 2.7 Hz, 1H), 7.56 – 7.51 (m, 1H), 7.51 – 7.47 (m, 2H), 7.45 (ddd, J = 8.0, 4.8, 0.8 Hz, 1H), 3.98 (s, 3H).	DMSO	>98	J2
343					¹ H NMR (400 MHz, DMSO) δ 10.99 (s, 1H), 9.93 (s, 1H), 9.26 (d, J = 1.4 Hz, 1H), 8.59 (dd, J = 4.8, 1.7 Hz, 1H), 8.50 – 8.43 (m, 1H), 8.01 (d, J = 2.7 Hz, 1H), 7.98 (d, J = 9.5 Hz, 1H), 7.85 (d, J = 9.1 Hz, 1H), 7.65 (d, J = 8.4 Hz, 1H), 7.55 (dd, J = 9.1, 2.7 Hz, 1H), 7.49 – 7.42 (m, 1H), 7.36 (d, J = 8.3 Hz, 1H), 6.55 (dd, J = 9.5, 1.7 Hz, 1H), 3.96 (s, 3H), 2.33 (s, 3H).	DMSO	>98	J2
344				HCl	¹ H NMR (400 MHz, DMSO) δ 9.34 – 9.28 (m, 1H), 8.81 – 8.75 (m, 1H), 8.74 – 8.65 (m, 1H), 8.06 (d, J = 9.2 Hz, 1H), 7.93 (d, J = 2.7 Hz, 1H), 7.85 (d, J = 8.8 Hz, 1H), 7.80 – 7.72 (m, 2H), 7.69 (d, J = 2.8 Hz, 1H), 7.59 (dd, J = 8.8, 2.7 Hz, 1H), 4.00 (s, 3H).	DMSO	>98	G8 using DMF instead of THF

345				HCl	¹ H NMR (400 MHz, DMSO) δ 9.35 – 9.29 (m, 1H), 8.97 – 8.78 (m, 2H), 8.19 (d, J = 9.4 Hz, 1H), 8.09 (d, J = 9.2 Hz, 1H), 7.94 (d, J = 8.5 Hz, 1H), 7.77 (dd, J = 8.0, 5.3 Hz, 1H), 7.72 (dd, J = 9.1, 2.9 Hz, 1H), 7.67 (d, J = 2.3 Hz, 1H), 7.54 (dd, J = 8.5, 2.3 Hz, 1H), 6.56 (d, J = 9.6 Hz, 1H), 4.02 (s, 3H).	DMSO	>98	G12 at room temperature
346				HCl	¹ H NMR (400 MHz, DMSO) δ 11.98 (s, 1H), 9.29 (d, J = 1.7 Hz, 1H), 8.88 – 8.78 (m, 2H), 8.08 (d, J = 9.2 Hz, 1H), 7.97 (d, J = 9.6 Hz, 1H), 7.88 (dd, J = 8.1, 5.4 Hz, 1H), 7.81 (d, J = 2.6 Hz, 1H), 7.78 – 7.72 (m, 2H), 7.67 (dd, J = 8.9, 2.6 Hz, 1H), 7.46 (d, J = 8.9 Hz, 1H), 6.60 (d, J = 9.5 Hz, 1H), 4.02 (s, 3H).	DMSO	>98	G12 at room temperature
347				HCl	¹ H NMR (400 MHz, DMSO) δ 9.33 – 9.26 (m, 1H), 8.87 – 8.67 (m, 2H), 8.06 (d, J = 9.1 Hz, 1H), 7.91 – 7.79 (m, 1H), 7.75 (dd, J = 9.1, 2.9 Hz, 1H), 7.71 (d, J = 2.8 Hz, 1H), 7.67 – 7.61 (m, 2H), 7.56 – 7.51 (m, 2H), 4.01 (s, 3H).	DMSO	>98	G12 at room temperature
348				HCl	¹ H NMR (400 MHz, DMSO) δ 9.28 (d, J = 2.0 Hz, 1H), 8.85 – 8.67 (m, 2H), 8.06 (d, J = 9.0 Hz, 1H), 7.98 – 7.77 (m, 1H), 7.77 – 7.69 (m, 2H), 7.62 – 7.54 (m, 2H), 7.51 – 7.45 (m, 2H), 7.44 – 7.37 (m, 1H), 4.02 (d, J = 3.9 Hz, 3H).	DMSO	>98	G12 at room temperature
349				HCl	¹ H NMR (400 MHz, DMSO) δ 9.23 (dd, J = 2.1, 0.7 Hz, 1H), 8.80 (dd, J = 5.2, 1.6 Hz, 1H), 8.72 – 8.64 (m, 1H), 8.09 (d, J = 9.1 Hz, 1H), 7.94 – 7.71 (m, 4H), 7.66 (dd, J = 8.1, 1.6 Hz, 1H), 7.51 – 7.54 (m, 1H), 7.52 – 7.45 (m, 1H), 4.02 (s, 3H).	DMSO	>98	G12 at room temperature

350				HCl	¹ H NMR (400 MHz, DMSO) δ 9.22 (d, J = 1.7 Hz, 1H), 8.87 – 8.79 (m, 1H), 8.77 – 8.67 (m, 1H), 8.06 (d, J = 9.2 Hz, 1H), 7.92 – 7.79 (m, 1H), 7.74 (dd, J = 9.1, 2.9 Hz, 1H), 7.70 (d, J = 2.8 Hz, 1H), 7.45 – 7.37 (m, 2H), 7.33 – 7.27 (m, 1H), 7.17 – 7.09 (m, 1H), 4.01 (s, 3H), 3.71 (s, 3H).	DMSO	>98	G12 at room temperature
351				2 HCl	¹ H NMR (400 MHz, DMSO) δ 12.03 (s, 1H), 10.69 (s, 1H), 9.65 (d, J = 1.8 Hz, 1H), 9.26 – 9.21 (m, 1H), 9.20 – 9.12 (m, 1H), 8.99 – 8.88 (m, 2H), 8.52 (d, J = 2.6 Hz, 1H), 8.49 – 8.42 (m, 1H), 8.25 – 8.17 (m, 1H), 8.15 – 8.08 (m, 2H), 8.02 – 7.96 (m, 1H), 7.93 (d, J = 9.4 Hz, 1H), 7.77 (d, J = 8.5 Hz, 1H), 7.72 (dd, J = 8.5, 1.9 Hz, 1H), 6.47 (d, J = 9.6 Hz, 1H), 4.04 (s, 3H).	DMSO	>98	G1
352				2 HCl	¹ H NMR (400 MHz, DMSO) δ 12.04 (s, 1H), 10.92 (s, 1H), 9.65 (d, J = 1.7 Hz, 1H), 9.58 – 9.50 (m, 1H), 9.46 – 9.39 (m, 1H), 9.27 – 9.18 (m, 1H), 9.01 – 8.92 (m, 2H), 8.88 (dd, J = 5.3, 1.2 Hz, 1H), 8.48 (dd, J = 8.8, 1.9 Hz, 1H), 8.20 – 8.10 (m, 2H), 8.09 – 8.01 (m, 2H), 7.93 (d, J = 9.5 Hz, 1H), 7.81 – 7.72 (m, 2H), 6.47 (d, J = 9.5 Hz, 1H).	DMSO	>98	G1
353				HCl	¹ H NMR (300 MHz, DMSO) δ 10.90 (s, 1H), 9.63 (s, 1H), 8.89 (d, J = 8.0 Hz, 1H), 8.76 (d, J = 4.0 Hz, 1H), 8.19 (d, J = 2.1 Hz, 1H), 7.89 (d, J = 9.1 Hz, 1H), 7.83 (d, J = 7.9 Hz, 1H), 7.78 – 7.65 (m, 2H), 7.63 – 7.49 (m, 2H), 7.47 – 7.33 (m, 2H), 3.98 (s, 3H).	DMSO	>98	G1

354			HCl	¹H NMR (300 MHz, DMSO) δ 11.44 (s, 1H), 9.58 (d, J = 1.8 Hz, 1H), 9.16 (d, J = 8.2 Hz, 1H), 8.96 (dd, J = 5.3, 1.3 Hz, 1H), 8.29 (d, J = 2.5 Hz, 1H), 8.13 (d, J = 9.2 Hz, 1H), 7.99 (dd, J = 8.0, 5.4 Hz, 1H), 7.72 – 7.49 (m, 2H), 7.44 – 7.27 (m, 3H), 7.12 (s, 1H), 3.99 (s, 3H), 2.52 (s, 3H).	DMSO	>98	G1
355			HCl	¹H NMR (300 MHz, DMSO) δ 11.38 (s, 1H), 9.53 (s, 1H), 9.18 (d, J = 8.1 Hz, 1H), 8.97 (d, J = 5.3 Hz, 1H), 8.21 (s, 1H), 8.14 – 8.02 (m, 1H), 7.92 (d, J = 9.1 Hz, 1H), 7.60 (d, J = 9.1 Hz, 1H), 6.90 (s, 1H), 3.99 (s, 3H).	DMSO	>98	G1
356			HCl	¹H NMR (300 MHz, DMSO) δ 11.30 (s, 1H), 9.59 (d, J = 1.8 Hz, 1H), 9.19 (d, J = 8.2 Hz, 1H), 8.97 (dd, J = 5.3, 1.2 Hz, 1H), 8.24 (d, J = 2.5 Hz, 1H), 8.11 – 7.93 (m, 2H), 7.76 – 7.54 (m, 4H), 7.40 (t, J = 7.6 Hz, 1H), 7.26 (s, 1H), 7.21 (d, J = 7.4 Hz, 1H), 3.98 (s, 3H), 2.41 (s, 3H).	DMSO	>98	G1
357			HCl	¹H NMR (300 MHz, DMSO) δ 11.21 (s, 1H), 9.59 (d, J = 1.7 Hz, 1H), 9.22 (d, J = 8.1 Hz, 1H), 9.02 – 8.95 (m, 1H), 8.23 (d, J = 2.5 Hz, 1H), 8.13 – 7.96 (m, 3H), 7.85 (d, J = 8.7 Hz, 2H), 7.72 (d, J = 8.7 Hz, 2H), 7.60 (dd, J = 9.1, 2.6 Hz, 1H), 7.31 (s, 1H), 3.98 (s, 3H).	DMSO	>98	G1
358			HCl	¹H NMR (300 MHz, DMSO) δ 10.87 (s, 1H), 9.65 – 9.55 (m, 1H), 8.93 (d, J = 8.0, 1.3 Hz, 1H), 8.31 (dd, J = 5.0, 1.6 Hz, 1H), 8.18 (d, J = 2.6 Hz, 1H), 7.94 – 7.82 (m, 4H), 7.78 (dd, J = 8.0, 5.0 Hz, 1H), 7.60 – 7.46 (m, 4H), 7.44 – 7.35 (m, 1H), 7.32 (s, 1H), 3.98 (s, 3H).	DMSO	>98	G1

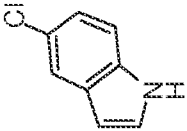
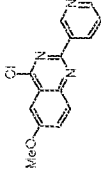
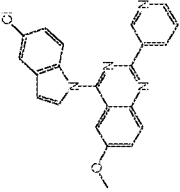
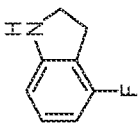
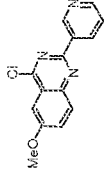
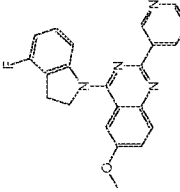
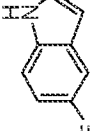
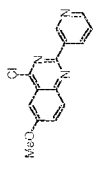
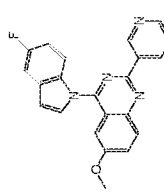
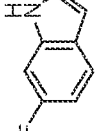
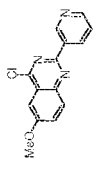
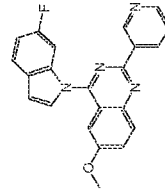
359			HCl	¹H NMR (300 MHz, DMSO) δ 11.29 (s, 1H), 9.59 (s, 1H), 9.20 (d, J = 8.1 Hz, 1H), 9.01 – 8.94 (m, 1H), 8.25 (s, 1H), 8.12 – 7.98 (m, 2H), 7.90 – 7.75 (m, 2H), 7.62 (d, J = 9.1 Hz, 1H), 7.21 (s, 1H), 7.08 (d, J = 8.6 Hz, 2H), 3.99 (s, 3H), 3.84 (s, 3H).	DMSO	>98	G1
360			HCl	¹H NMR (300 MHz, DMSO) δ 11.29 (s, 1H), 9.60 (s, 1H), 9.21 (d, J = 8.0 Hz, 1H), 8.99 (d, J = 5.3 Hz, 1H), 8.26 (d, J = 2.2 Hz, 1H), 8.11 – 7.99 (m, 2H), 7.77 (d, J = 8.0 Hz, 2H), 7.62 (dd, J = 9.1, 2.4 Hz, 1H), 7.32 (d, J = 8.1 Hz, 2H), 7.26 (s, 1H), 3.99 (s, 3H), 2.37 (s, 3H).	DMSO	>98	G1
361			HCl	¹H NMR (300 MHz, DMSO) δ 11.00 (s, 1H), 9.55 (d, J = 1.7 Hz, 1H), 8.96 (dt, J = 8.1, 1.7 Hz, 1H), 8.86 (dd, J = 5.1, 1.5 Hz, 1H), 8.19 (d, J = 2.6 Hz, 1H), 7.97 (d, J = 9.2 Hz, 1H), 7.83 (dt, J = 9.5, 4.8 Hz, 1H), 7.55 (dd, J = 9.1, 2.6 Hz, 1H), 6.72 (s, 1H), 3.95 (d, J = 7.0 Hz, 3H), 2.35 (d, J = 3.5 Hz, 3H).	DMSO	>98	G1
362			HCl	¹H NMR (300 MHz, DMSO) δ 11.21 (s, 1H), 9.58 (d, J = 1.5 Hz, 1H), 9.21 – 9.06 (m, 1H), 8.95 (d, J = 4.3 Hz, 1H), 8.22 (d, J = 2.3 Hz, 1H), 8.09 – 7.92 (m, 2H), 7.70 – 7.51 (m, 3H), 7.24 – 7.17 (m, 1H), 7.13 (s, 1H), 3.99 (s, 3H).	DMSO	>98	G1
363			HCl	¹H NMR (300 MHz, DMSO) δ 11.07 (s, 1H), 9.60 (s, 1H), 9.13 (d, J = 6.5 Hz, 1H), 8.93 (d, J = 5.1 Hz, 1H), 8.23 (s, 1H), 8.05 – 7.85 (m, 4H), 7.67 – 7.54 (m, 1H), 7.43 – 7.23 (m, 3H), 3.99 (s, 3H).	DMSO	>98	G1

364			2 HCl		¹ H NMR (300 MHz, DMSO) δ 11.16 (s, 1H), 9.57 (d, J = 1.5 Hz, 1H), 9.14 (d, J = 8.1 Hz, 1H), 8.93 (d, J = 5.2 Hz, 1H), 8.18 (d, J = 2.4 Hz, 1H), 8.03 – 7.99 (m, 4H), 7.61 – 7.54 (m, 2H), 7.32 (s, 1H), 3.97 (s, 3H).	DMSO	>98	G1
365			2 HCl		¹ H NMR (300 MHz, DMSO) δ 11.29 (s, 1H), 9.58 (d, J = 1.6 Hz, 1H), 9.27 (d, J = 8.2 Hz, 1H), 9.06 – 8.99 (m, 1H), 8.18 (d, J = 2.5 Hz, 1H), 8.10 (dd, J = 8.1, 5.5 Hz, 1H), 8.02 (d, J = 9.1 Hz, 1H), 7.91 (dd, J = 10.7, 1.8 Hz, 1H), 7.80 – 7.65 (m, 2H), 7.57 (dd, J = 9.1, 2.6 Hz, 1H), 7.29 (s, 1H), 3.96 (s, 3H).	DMSO	>98	G1
366			2 HCl		¹ H NMR (300 MHz, DMSO) δ 11.17 (s, 1H), 9.60 (s, 1H), 9.24 (d, J = 8.2 Hz, 1H), 8.93 (d, J = 5.1 Hz, 1H), 8.22 (d, J = 2.5 Hz, 1H), 8.11 – 8.04 (m, 1H), 8.02 – 7.91 (m, 2H), 7.82 – 7.68 (m, 2H), 7.62 (dd, J = 9.1, 2.5 Hz, 1H), 7.36 (s, 1H), 4.01 (d, J = 11.0 Hz, 3H).	DMSO	>98	G1
367			2 HCl		¹ H NMR (300 MHz, DMSO) δ 11.53 (s, 1H), 9.58 (d, J = 1.8 Hz, 1H), 9.20 (d, J = 8.2 Hz, 1H), 8.98 (dd, J = 5.3, 1.3 Hz, 1H), 8.25 (d, J = 2.5 Hz, 1H), 8.12 (d, J = 9.1 Hz, 1H), 8.04 – 7.95 (m, 1H), 7.80 – 7.68 (m, 2H), 7.63 – 7.57 (m, 1H), 7.56 – 7.48 (m, 1H), 7.31 (s, 1H), 3.99 (d, J = 6.7 Hz, 3H).	DMSO	>98	G1

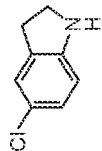
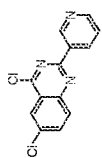
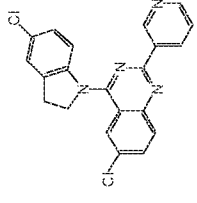
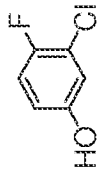
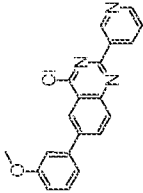
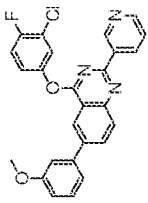
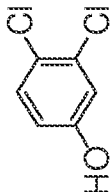
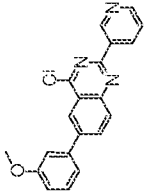
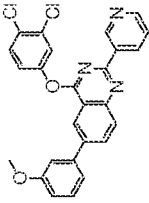
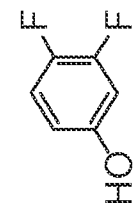
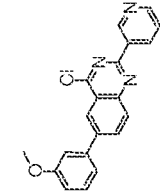
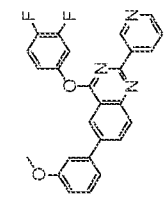
368			2 HCl		¹ H NMR (300 MHz, DMSO) δ 11.24 (s, 1H), 9.59 (d, J = 1.7 Hz, 1H), 9.26 (d, J = 8.3 Hz, 1H), 9.00 (d, J = 4.2 Hz, 1H), 8.23 (d, J = 2.6 Hz, 1H), 8.16 (d, J = 2.0 Hz, 1H), 8.09 (dd, J = 8.1, 5.4 Hz, 1H), 8.01 (d, J = 9.2 Hz, 1H), 7.91 (dd, J = 8.4, 2.1 Hz, 1H), 7.77 (d, J = 8.4 Hz, 1H), 7.62 (dd, J = 9.1, 2.5 Hz, 1H), 7.37 (s, 1H), 3.98 (s, 3H).	DMSO	>98	G1
369			HCl		¹ H NMR (300 MHz, DMSO) δ 11.33 (s, 1H), 9.62 (s, 1H), 9.27 (d, J = 8.4 Hz, 1H), 9.01 (d, J = 5.2 Hz, 1H), 8.29 (m, 3H), 8.13 (m, 2H), 7.79 (m, 2H), 7.63 (dd, J = 9.1, 2.6 Hz, 1H), 7.42 (s, 1H), 3.99 (s, 3H).	DMSO	>98	G1
370			HCl		¹ H NMR (300 MHz, DMSO) δ 11.31 (s, 1H), 9.55 (d, J = 1.7 Hz, 1H), 9.07 (d, J = 7.9 Hz, 1H), 8.92 (d, J = 4.0 Hz, 1H), 8.26 (d, J = 2.5 Hz, 1H), 8.04 (d, J = 9.1 Hz, 1H), 7.98 (m, 5H), 7.63 (dd, J = 9.2, 2.5 Hz, 1H), 7.13 (s, 1H), 4.00 (s, 3H).	DMSO	>98	G1
371			2 HCl		¹ H NMR (300 MHz, CDCl ₃) δ 11.29 (s, 1H), 9.57 (d, J = 1.8 Hz, 1H), 9.07 (d, J = 8.1 Hz, 1H), 8.93 (dd, J = 5.2, 1.3 Hz, 1H), 8.25 (d, J = 2.5 Hz, 1H), 8.07 (m, 2H), 7.92 (dd, J = 8.0, 5.3 Hz, 1H), 7.67 (m, 4H), 7.28 (s, 1H), 3.99 (s, 3H).	DMSO	>98	G1

372			2 HCl	¹ H NMR (300 MHz, DMSO) δ 11.32 (s, 1H), 9.80 (d, J = 1.7 Hz, 1H), 9.27 (d, J = 8.1 Hz, 1H), 9.03 (d, J = 5.4 Hz, 1H), 8.22 (d, J = 2.5 Hz, 1H), 8.12 (dd, J = 8.1, 5.7 Hz, 1H), 8.03 (d, J = 8.8 Hz, 3H), 7.59 (dd, J = 9.1, 2.5 Hz, 1H), 7.52 (d, J = 8.1 Hz, 2H), 7.32 (s, 1H), 3.98 (d, J = 6.7 Hz, 3H).	DMSO	>98	G1
373			2 HCl	¹ H NMR (300 MHz, DMSO) δ 11.38 (s, 1H), 9.61 (d, J = 1.6 Hz, 1H), 9.26 (d, J = 8.2 Hz, 1H), 9.02 (d, J = 5.3 Hz, 1H), 8.23 (d, J = 2.4 Hz, 1H), 8.12 (dd, J = 8.1, 5.7 Hz, 1H), 8.03 (d, J = 8.8 Hz, 3H), 7.59 (dd, J = 9.1, 2.5 Hz, 1H), 7.52 (d, J = 8.1 Hz, 2H), 7.32 (s, 1H), 3.98 (d, J = 6.7 Hz, 3H).	DMSO	>98	G1
374			2 HCl	¹ H NMR (300 MHz, DMSO) δ 11.29 (s, 1H), 9.56 (s, 1H), 9.12 (d, J = 7.7 Hz, 1H), 8.94 (d, J = 4.9 Hz, 1H), 8.26 (d, J = 2.3 Hz, 1H), 8.07 (dd, J = 7.90 (m, 2H), 7.71 (m, 2H), 7.51 (s, 1H), 7.47 (m, 1H), 7.17 (s, 1H), 3.99 (s, 3H), 2.53 (s, 3H).	DMSO	>98	G1
375			2 HCl	¹ H NMR (300 MHz, DMSO) δ 13.13 (s, 1H), 11.86 (m, 1H), 9.66 (d, J = 1.4 Hz, 1H), 9.36 (d, J = 8.2 Hz, 1H), 9.00 (d, J = 5.4 Hz, 1H), 8.38 (d, J = 2.2 Hz, 1H), 8.14 (dd, J = 8.1, 5.6 Hz, 1H), 8.01 (d, J = 7.5 Hz, 2H), 7.89 (d, J = 9.1 Hz, 1H), 7.78 (s, 1H), 7.64 (m, 9H), 5.23 (s, 2H).	DMSO	>98	J NaO ^t Bu, Pd(OAc) ₂ , Tri-t-butylphosphonium tetrafluoroborate, Toluene
376			2 HCl	¹ H NMR (300 MHz, DMSO) δ 12.49 (s, 1H), 9.63 (s, 1H), 9.27 (d, J = 8.1 Hz, 1H), 8.96 (d, J = 5.4 Hz, 1H), 8.34 (s, 1H), 8.07 (dd, J = 8.1, 5.6 Hz, 1H), 7.99 (d, J = 8.3 Hz, 2H), 7.85 (d, J = 9.1 Hz, 1H), 7.80 (s, 1H), 7.64 (m, 9H), 5.21 (s, 2H).	DMSO	>98	J NaO ^t Bu, Pd(OAc) ₂ , Tri-t-butylphosphonium tetrafluoroborate, Toluene

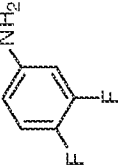
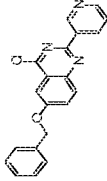
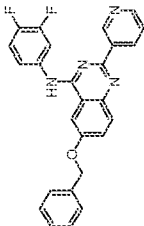
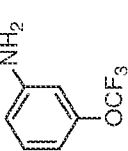
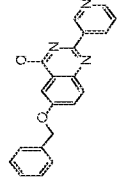
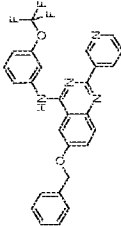
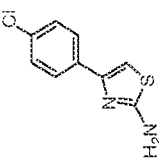
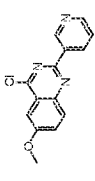
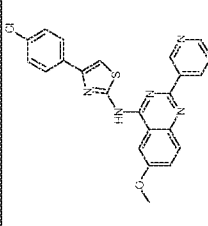
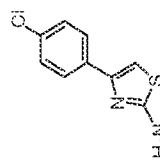
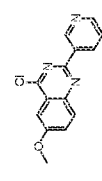
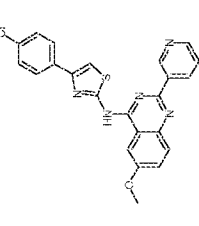
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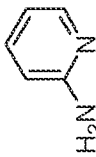
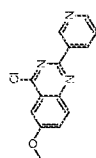
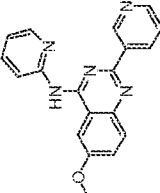
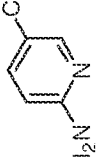
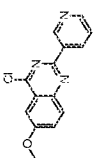
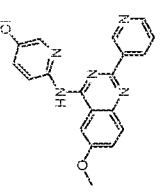
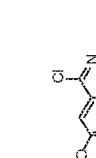
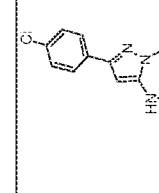
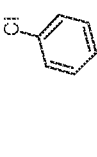
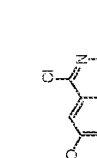
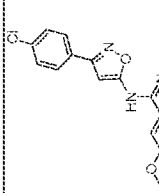
381				HCl	¹H NMR (300 MHz, DMSO) δ 9.66 (d, J = 2.0 Hz, 1H), 9.15 – 9.06 (m, 1H), 8.91 (dd, J = 5.2, 1.5 Hz, 1H), 8.27 – 8.17 (m, 2H), 8.01 (d, J = 8.9 Hz, 1H), 7.94 (dd, J = 8.2, 5.2 Hz, 1H), 7.88 – 7.78 (m, 2H), 7.42 (d, J = 2.7 Hz, 1H), 7.35 (dd, J = 8.8, 2.2 Hz, 1H), 6.96 (d, J = 3.5 Hz, 1H), 3.91 (s, 3H).	DMSO	>98	G8 at 50°C
382				HCl	¹H NMR (300 MHz, DMSO) δ 9.56 (d, J = 1.6 Hz, 1H), 9.26 (d, J = 8.2 Hz, 1H), 9.05 – 8.98 (m, 1H), 8.20 – 8.02 (m, 2H), 7.69 (dd, J = 9.2, 2.7 Hz, 1H), 7.53 – 7.40 (m, 2H), 7.32 (dd, J = 14.0, 8.0 Hz, 1H), 6.92 (t, J = 8.5 Hz, 1H), 4.71 (t, J = 7.9 Hz, 2H), 3.91 (s, 3H), 3.28 (t, J = 7.8 Hz, 2H).	DMSO	>98	G8 at 50°C
383				2 HCl	¹H NMR (300 MHz, DMSO) δ 9.66 (d, J = 1.5 Hz, 1H), 9.23 – 9.09 (m, 1H), 8.93 (dd, J = 5.3, 1.5 Hz, 1H), 8.31 – 8.16 (m, 2H), 8.08 – 7.93 (m, 2H), 7.82 (dd, J = 9.2, 2.8 Hz, 1H), 7.56 (dd, J = 9.4, 2.5 Hz, 1H), 7.43 (d, J = 2.7 Hz, 1H), 7.19 (td, J = 9.2, 2.7 Hz, 1H), 7.02 – 6.99 (m, 1H), 3.91 (s, 3H).	DMSO	>96	G8 at 50°C
384				2 HCl	¹H NMR (300 MHz, DMSO) δ 9.66 (d, J = 1.6 Hz, 1H), 9.23 – 9.09 (m, 1H), 8.94 (dd, J = 5.3, 1.5 Hz, 1H), 8.26 – 8.14 (m, 2H), 8.00 (dd, J = 8.1, 5.3 Hz, 1H), 7.90 – 7.73 (m, 3H), 7.45 (d, J = 2.7 Hz, 1H), 7.25 – 7.10 (m, 1H), 6.98 (d, J = 3.5 Hz, 1H), 3.92 (s, 3H).	DMSO	>96	G8 at 50°C

385			2 HCl	¹H NMR (300 MHz, DMSO) δ 9.54 (d, J = 1.8 Hz, 1H), 9.21 (dt, J = 8.2, 1.6 Hz, 1H), 8.98 (dd, J = 5.5, 1.3 Hz, 1H), 8.20 – 8.00 (m, 2H), 7.68 (dd, J = 9.2, 2.7 Hz, 1H), 7.55 – 7.31 (m, 3H), 7.00 – 6.78 (m, 1H), 4.69 (t, J = 7.9 Hz, 2H), 4.19 (q, J = 7.0 Hz, 2H), 3.22 (t, J = 7.9 Hz, 2H), 1.41 (t, J = 6.9 Hz, 3H).	DMSO	>98	G8 at 50°C
386			2 HCl	¹H NMR (300 MHz, DMSO) δ 9.42 (d, J = 1.5 Hz, 1H), 9.06 (d, J = 8.1 Hz, 1H), 8.93 (d, J = 5.4 Hz, 1H), 8.09 – 7.99 (m, 2H), 7.68 (dd, J = 9.1, 2.7 Hz, 1H), 7.53 (d, J = 2.6 Hz, 1H), 7.30 – 7.16 (m, 3H), 4.66 (t, J = 7.9 Hz, 2H), 3.98 (s, 3H), 3.29 (t, J = 7.7 Hz, 2H).	DMSO	>98	G8 at 50°C
387			2 HCl	¹H NMR (300 MHz, DMSO) δ 9.37 (d, J = 1.4 Hz, 1H), 9.11 (d, J = 8.2 Hz, 1H), 8.99 (d, J = 4.9 Hz, 1H), 8.14 (dd, J = 8.2, 5.6 Hz, 1H), 8.06 (d, J = 9.2 Hz, 1H), 7.70 (dd, J = 9.2, 2.6 Hz, 1H), 7.51 (d, J = 2.7 Hz, 1H), 7.45 – 7.36 (m, 2H), 7.26 – 7.15 (m, 1H), 4.83 (d, J = 9.5 Hz, 1H), 4.49 – 4.38 (m, 2H), 4.00 (s, 3H), 3.48 – 3.28 (m, 1H), 3.27 – 3.05 (m, 1H).	DMSO	>98	G8 at 50°C
388			2 HCl	¹H NMR (400 MHz, DMSO) δ 9.56 (d, J = 1.9 Hz, 1H), 9.31 – 9.19 (m, 1H), 9.00 (dd, J = 5.5, 1.3 Hz, 1H), 8.21 – 8.02 (m, 2H), 7.73 – 7.64 (m, 2H), 7.47 (dd, J = 5.4, 2.4 Hz, 2H), 7.32 (dd, J = 8.6, 2.3 Hz, 1H), 4.67 (t, J = 8.0 Hz, 2H), 4.18 (q, J = 7.0 Hz, 2H), 3.26 (t, J = 7.8 Hz, 2H), 1.41 (t, J = 6.9 Hz, 3H).	DMSO	>98	G8 at 50°C

389				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.54 (d, J = 1.8 Hz, 1H), 9.25 (dd, J = 8.2, 1.6 Hz, 1H), 9.04 (dd, J = 5.5, 1.1 Hz, 1H), 8.26 (d, J = 2.1 Hz, 1H), 8.16 (dd, J = 8.0, 5.6 Hz, 1H), 8.11 – 7.96 (m, 2H), 7.91 (d, J = 8.6 Hz, 1H), 7.47 (d, J = 2.1 Hz, 1H), 7.34 (dd, J = 8.6, 2.3 Hz, 1H), 4.68 (t, J = 7.9 Hz, 2H), 3.25 (t, J = 7.8 Hz, 2H).	DMSO	>98	G8 at 50°C
390				HCl	¹ H NMR (400 MHz, DMSO) δ 9.37 (d, J = 1.4 Hz, 1H), 8.82 (dd, J = 5.1, 1.5 Hz, 1H), 8.76 (dd, J = 8.1 Hz, 1H), 8.62 (d, J = 1.9 Hz, 1H), 8.46 (dd, J = 8.8, 2.1 Hz, 1H), 8.20 (d, J = 8.7 Hz, 1H), 7.91 (dd, J = 6.3, 2.6 Hz, 1H), 7.80 (dd, J = 6.0, 5.1 Hz, 1H), 7.71 – 7.59 (m, 2H), 7.52 – 7.38 (m, 3H), 7.10 – 7.01 (m, 1H), 3.89 (s, 3H).	DMSO	>98	G8 with DMF instead of THF
391				HCl	¹ H NMR (400 MHz, DMSO) δ 9.38 (d, J = 1.4 Hz, 1H), 8.82 (d, J = 5.1 Hz, 1H), 8.76 (d, J = 7.9 Hz, 1H), 8.63 (d, J = 1.8 Hz, 1H), 8.46 (dd, J = 8.8, 2.2 Hz, 1H), 8.20 (d, J = 8.7 Hz, 1H), 7.97 (d, J = 2.7 Hz, 1H), 7.87 (d, J = 8.8 Hz, 1H), 7.84 – 7.76 (m, 1H), 7.64 (dd, J = 8.8, 2.7 Hz, 1H), 7.52 – 7.39 (m, 3H), 7.10 – 7.02 (m, 1H), 3.88 (s, 3H).	DMSO	>98	G8 with DMF instead of THF
392				HCl	¹ H NMR (400 MHz, DMSO) δ 9.39 (t, J = 3.2 Hz, 1H), 8.93 – 8.82 (m, 2H), 8.62 (d, J = 1.8 Hz, 1H), 8.47 (dd, J = 8.8, 2.2 Hz, 1H), 8.21 (d, J = 9.0 Hz, 1H), 7.91 (dd, J = 6.1, 5.3 Hz, 1H), 7.84 – 7.77 (m, 1H), 7.73 – 7.63 (m, 1H), 7.53 – 7.40 (m, 4H), 7.09 – 7.03 (m, 1H), 3.87 (d, J = 6.5 Hz, 3H).	DMSO	>98	G8 with DMF instead of THF

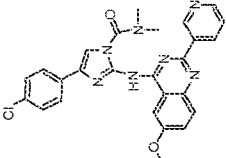
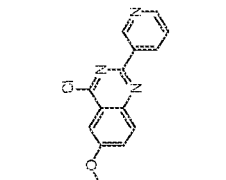
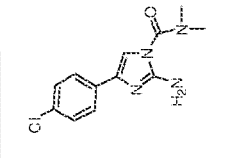
393			HCl	¹ H NMR (400 MHz, DMSO) δ 11.71 (s, 1H), 11.10 (s, 1H), 9.42 (d, J = 1.9 Hz, 1H), 8.90 – 8.69 (m, 2H), 8.62 (d, J = 1.7 Hz, 1H), 8.40 (dd, J = 8.8, 1.9 Hz, 1H), 8.14 (d, J = 8.7 Hz, 1H), 7.92 (d, J = 7.7 Hz, 1H), 7.84 – 7.69 (m, 1H), 7.57 – 7.40 (m, 3H), 7.11 – 7.02 (m, 2H), 6.96 (d, J = 8.3 Hz, 1H), 3.89 (s, 3H).	DMSO	>98	G8 with DMF instead of THF
394			HCl	¹ H NMR (400 MHz, DMSO) δ 9.43 – 9.29 (m, 1H), 8.75 – 8.64 (m, 2H), 8.60 – 8.48 (m, 1H), 8.47 – 8.32 (m, 1H), 8.21 – 8.01 (m, 2H), 7.86 – 7.74 (m, 2H), 7.64 (d, J = 8.6 Hz, 1H), 7.59 – 7.34 (m, 4H), 7.13 – 7.01 (m, 1H), 6.87 (d, J = 8.7 Hz, 1H), 3.90 (t, J = 5.3 Hz, 3H).	DMSO	>98	G8 with DMF instead of THF
395			HCl	¹ H NMR (400 MHz, DMSO) δ 9.24 (dd, J = 2.2, 0.8 Hz, 1H), 8.76 (dd, J = 5.0, 1.7 Hz, 1H), 8.61 – 8.54 (m, 1H), 8.49 (dd, J = 1.8, 1.1 Hz, 1H), 8.21 – 8.12 (m, 2H), 7.75 (dd, J = 8.0, 1.5 Hz, 1H), 7.70 – 7.62 (m, 2H), 7.62 – 7.53 (m, 1H), 7.53 – 7.44 (m, 1H).	DMSO	>96	G8 with DMF instead of THF
396			HCl	¹ H NMR (400 MHz, DMSO) δ 11.71 (s, 1H), 11.10 (s, 1H), 9.42 (d, J = 1.9 Hz, 1H), 8.90 – 8.69 (m, 2H), 8.62 (d, J = 1.7 Hz, 1H), 8.40 (dd, J = 8.8, 1.9 Hz, 1H), 8.14 (d, J = 8.7 Hz, 1H), 7.92 (d, J = 7.7 Hz, 1H), 7.84 – 7.69 (m, 1H), 7.57 – 7.40 (m, 3H), 7.11 – 7.02 (m, 2H), 6.96 (d, J = 8.3 Hz, 1H), 3.89 (s, 3H).	DMSO	>98	G12

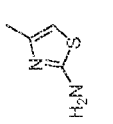
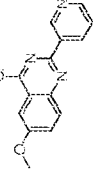
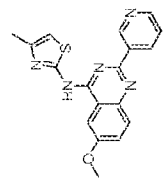
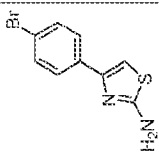
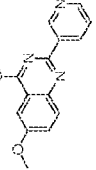
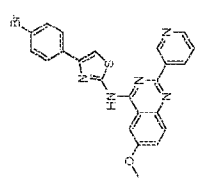
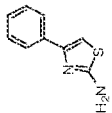
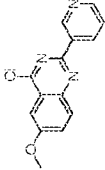
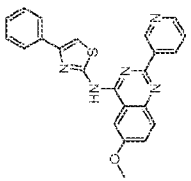
397				HCl	¹ H NMR (400 MHz, DMSO-d ₆) δ 10.40 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 8.99 (d, J = 8.2 Hz, 1H), 8.88 (dd, J = 5.3, 1.5 Hz, 1H), 8.30 (d, J = 2.5 Hz, 1H), 8.14 – 8.06 (m, 1H), 7.99 – 7.88 (m, 2H), 7.78 – 7.66 (m, 2H), 7.61 – 7.51 (m, 3H), 7.50 – 7.34 (m, 3H), 5.34 (s, 2H).	DMSO	>98	Method G1
398					¹ H NMR (400 MHz, DMSO-d ₆) δ 9.96 (s, 1H), 9.52 (d, J = 1.4 Hz, 1H), 8.98 (dd, J = 9.9 Hz, 1H), 8.65 (d, J = 9.9 Hz, 1H), 8.16 (dd, J = 2.5, 5.4 Hz, 2H), 7.68 – 7.41 (m, 2H), 7.78 – 7.66 (m, 8H), 7.39 – 7.19 (m, 1H), 5.30 (s, 2H).	DMSO	>98	Method G1
399					¹ H NMR (DMSO-d ₆) ppm 12.52 (s, 1H), 9.78 (d, J = 1.56 Hz, 1H), 8.91-8.88 (m, 1H), 8.74 (dd, J = 4.74, 1.60 Hz, 1H), 8.33 (brs, 1H), 8.06 (d, J = 8.56 Hz, 2H), 7.93 (d, J = 9.08 Hz, 1H), 7.89 (s, 1H), 7.66-7.59 (m, 2H), 7.55 (d, J = 8.56 Hz, 2H), 4.01 (s, 3H).	DMSO	>98	G13
400				2 HCl	¹ H NMR (DMSO-d ₆) ppm 12.80-12.40 (br, 1H), 9.77 (d, J = 1.84 Hz, 1H), 9.32 (d, J = 8.04 Hz, 1H), 8.97 (dd, J = 5.36, 1.40 Hz, 1H), 8.34 (d, J = 2.68 Hz, 1H), 8.09-8.06 (m, 3H), 7.97 (d, J = 9.12 Hz, 1H), 7.91 (s, 1H), 7.64 (dd, J = 9.12, 2.68 Hz, 1H), 7.55 (d, J = 8.60 Hz, 2H), 4.01 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13

401				2 HCl	1H NMR (DMSO-d6) ppm 11.14 (brs, 1H), 9.58 (s, 1H), 9.13 (dd, J = 7.96 Hz, 1H), 8.93 (d, J = 5.24 Hz, 1H), 8.54 (d, J = 4.72 Hz, 1H), 8.48 (d, J = 8.32 Hz, 1H), 8.28 (brs, 1H), 8.09 (brt, J = 7.16 Hz, 1H), 8.02-7.96 (m, 2H), 7.65 (dd, J = 8.80, 2.48 Hz, 1H), 7.34 (brt, J = 6.52 Hz, 1H), 4.01 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	J2
402				2 HCl	1H NMR (DMSO-d6) ppm 10.89 (s, 1H), 9.57 (d, J = 1.60 Hz, 1H), 9.04 (d, J = 8.04 Hz, 1H), 8.88 (dd, J = 5.16, 1.40 Hz, 1H), 8.55 (dd, J = 2.64, 0.52 Hz, 1H), 8.50 (d, J = 8.92 Hz, 1H), 8.22 (d, J = 2.68 Hz, 1H), 8.08 (d, J = 8.92, 2.63 Hz, 1H), 7.94-7.90 (m, 2H), 7.61 (dd, J = 9.08, 2.68 Hz, 1H), 3.99 (s, 3H). The 1H of 2HCl was not observed.	DMSO	DMSO	J2
403				2 HCl	1H NMR (DMSO-d6) ppm 10.53 (s, 1H), 9.38 (s, 1H), 8.99 (d, J = 8.08 Hz, 1H), 8.88 (d, J = 4.60 Hz, 1H), 8.11 (brs, 1H), 8.01-7.95 (m, 2H), 7.90 (d, J = 8.60 Hz, 2H), 7.66 (dd, J = 9.12, 2.68 Hz, 1H), 7.50 (d, J = 8.60 Hz, 2H), 6.93 (s, 1H), 4.00 (s, 3H), 3.80 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	J2
404				HCl	1H NMR (DMSO-d6) ppm 11.81 (brs, 1H), 9.64 (s, 1H), 9.00 (d, J = 7.48 Hz, 1H), 8.85 (brs, 1H), 8.14 (d, J = 2.56 Hz, 1H), 8.00 (d, J = 8.60 Hz, 2H), 7.96 (d, J = 9.08 Hz, 1H), 7.86 (br, 1H), 7.66-7.63 (m, 3H), 7.25 (s, 1H), 4.00 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	J2

405			HCl	¹H NMR (DMSO-d₆) ppm 11.80 (brs, 1H), 9.65 (brs, 1H), 9.04 (d, J = 7.92 Hz, 1H), 8.88 (brs, 1H), 8.14 (d, J = 2.60 Hz, 1H), 8.05-8.02 (m, 2H), 7.96 (d, J = 9.12 Hz, 1H), 7.89 (m, 1H), 7.64 (dd, J = 9.12, 2.60 Hz, 1H), 7.42 (t, J = 8.84 Hz, 2H), 7.23 (s, 1H), 3.99 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	J2
406			HCl	¹H NMR (DMSO-d₆) ppm 10.31 (s, 1H), 9.37 (s, 1H), 8.88 (m, 2H), 8.04 (d, J = 2.60 Hz, 1H), 7.94 (d, J = 9.12 Hz, 1H), 7.91 (m, 1H), 7.64 (dd, J = 9.12, 2.63 Hz, 1H), 7.56 (d, J = 1.88 Hz, 1H), 6.40 (d, J = 1.88 Hz, 1H), 3.98 (s, 3H), 3.74 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	J2
407			2 HCl	¹H NMR (DMSO-d₆) ppm 12.89 (brs, 1H), 9.56 (s, 1H), 8.92 (d, J = 4.92 Hz, 2H), 8.04 (d, J = 8.84 Hz, 1H), 7.98 (d, J = 2.63 Hz, 1H), 7.84 (br, 1H), 7.71-7.66 (m, 3H), 7.45-7.39 (m, 2H), 4.00 (s, 3H), 3.94 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	J2
408			HCl	¹H NMR (DMSO-d₆) ppm 10.39 (s, 1H), 9.38 (s, 1H), 8.90 (d, J = 8.00 Hz, 1H), 8.84 (d, J = 4.76 Hz, 1H), 8.06 (d, J = 2.63 Hz, 1H), 7.95 (d, J = 9.16 Hz, 1H), 7.90-7.88 (m, 3H), 7.65 (dd, J = 9.16, 2.63 Hz, 1H), 7.44 (brt, J = 7.40 Hz, 2H), 7.35-7.31 (m, 1H), 6.88 (s, 1H), 3.99 (s, 3H), 3.80 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	J2 Temperature at 100 °C

409			HCl	¹H NMR (DMSO-d₆) ppm: 10.44 (brs, 1H), 9.01 (s, 1H), 8.68 (brs, 1H), 8.39 (brs, 1H), 8.23 (s, 1H), 8.07 (brs, 1H), 7.94 (d, J = 9.04 Hz, 1H), 7.70-7.54 (m, 2H), 7.44 (brd, J = 7.32 Hz, 2H), 7.15 (brt, J = 7.56 Hz, 2H), 7.00 (brt, 7.40 Hz, 1H), 3.97 (s, 3H), 3.93 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	J2 Temperature at 100 °C
410			HCl	¹H NMR (DMSO-d₆) ppm: 11.13 (s, 1H), 9.59 (d, J = 1.64 Hz, 1H), 9.03 (brs, 1H), 8.88 (brs, 1H), 8.63 (d, J = 2.52 Hz, 1H), 8.23 (d, J = 2.32 Hz, 1H), 7.93-7.82 (brm, 4H), 7.61-7.53 (m, 3H), 7.35-7.31 (brm, 2H), 4.00 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	J2 Temperature at 100 °C
411				¹H NMR (DMSO-d₆) ppm: 9.54 (s, 1H), 8.79 (s, 1H), 8.69 (d, J = 7.60 Hz, 1H), 7.89 (brd, J = 8.96 Hz, 2H), 7.68-7.58 (m, 4H), 7.33-7.30 (m, 2H), 3.98 (s, 3H). The 1H of NH- was not observed.	DMSO	>98	J2 Temperature at 100 °C
412			HCl	¹H NMR (DMSO-d₆) ppm: 9.70 (s, 1H), 9.05 (d, J = 7.88 Hz, 1H), 8.85 (brd, J = 4.44 Hz, 1H), 8.24 (brs, 1H), 8.04 (d, J = 8.44 Hz, 2H), 7.94 (d, J = 9.00 Hz, 1H), 7.84 (m, 1H), 7.56 (d, J = 8.44 Hz, 2H), 7.62 (dd, J = 9.00, 2.64 Hz, 1H), 3.99 (s, 3H). The 1H of HCl and NH- were not observed.	DMSO	>98	G12 Temperature at 100 °C

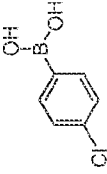
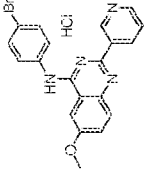
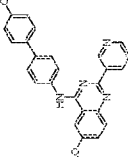
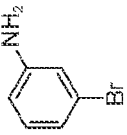
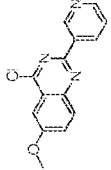
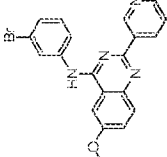
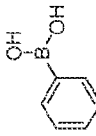
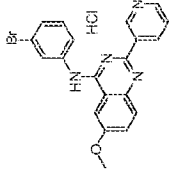
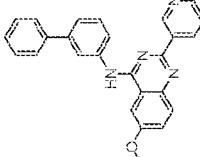
5		J_2 Temperature at 100 °C
10		>98
15		CDCl ₃
20		¹ H NMR (CDCl ₃) ppm 15.16 (brs, 1H), 9.57 (d, J = 2.32 Hz, 1H), 8.86 (dd, J = 4.80, 1.56 Hz, 1H), 8.59-8.56 (m, 1H), 7.78-7.73 (m, 4H), 7.55-7.52 (m, 1H), 7.43 (d, J = 8.56 Hz, 2H), 7.38 (dd, J = 8.88, 2.96 Hz, 1H), 7.34 (s, 1H), 3.97 (s, 3H), 3.30 (s, 3H), 3.11 (s, 3H).
25		
30		
35		
40		
45		
50		
55	413	

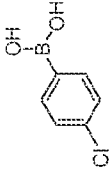
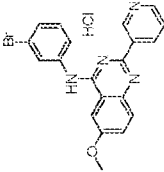
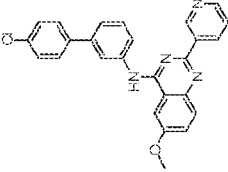
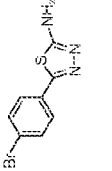
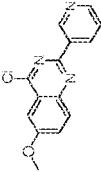
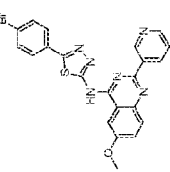
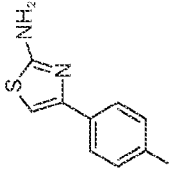
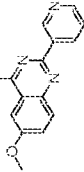
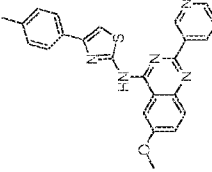
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	Retention Time	LCMS Method
414				2 HCl	¹ H NMR (DMSO-d ₆) ppm 9.72 (s, 1H), 9.16 (brs, 1H), 8.93 (d, J = 4.36 Hz, 1H), 8.14 (br, 1H), 7.96 (brm, 2H), 7.63 (dd, J = 9.08, 2.56 Hz, 1H), 7.00 (s, 1H), 3.98 (3H), 2.37 (s, 3H). The 1H of 2HCl and NH- were not observed.	DMSO	>98	G12 Temperature at 100°C			
415				2 HCl	¹ H NMR (DMSO-d ₆) ppm 12.60 (brs, 1H), 9.78 (s, 1H), 9.12 (d, J = 8.00 Hz, 1H), 8.86 (brs, 1H), 8.35 (brd, J = 2.60 Hz, 1H), 8.01 (d, J = 8.56 Hz, 2H), 7.96 (d, J = 9.08 Hz, 1H), 7.92 (s, 1H), 7.86 (m, 1H), 7.69 (d, J = 8.56 Hz, 2H), 7.63 (dd, J = 9.08, 2.60 Hz, 1H), 4.02 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G12 Temperature at 100°C			
416				HCl	¹ H NMR (DMSO-d ₆) ppm 12.80-12.40 (br, 1H), 9.79 (d, J = 1.64 Hz, 1H), 9.16 (d, J = 7.88 Hz, 1H), 8.88 (d, J = 3.88 Hz, 1H), 8.36 (brd, J = 2.64 Hz, 1H), 8.06-8.04 (m, 2H), 7.96 (d, J = 9.08 Hz, 1H), 7.90 (m, 1H), 7.85 (s, 1H), 7.63 (dd, J = 9.08, 2.64 Hz, 1H), 7.49 (m, 2H), 7.38 (m, 1H), 4.02 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G12 Temperature at 100°C			

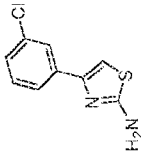
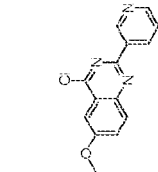
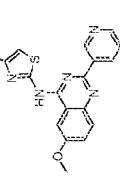
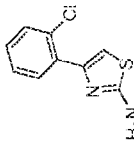
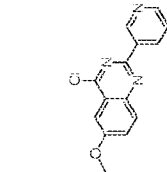
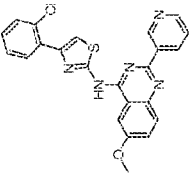
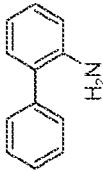
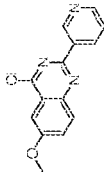
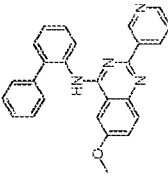
417			HCl	¹ H NMR (DMSO-d ₆) ppm 12.80 (s, 1H), 9.74 (d, J = 1.60 Hz, 1H), 9.01 (brs, 1H), 8.83 (brs, 1H), 8.31 (d, J = 2.64 Hz, 1H), 8.15 (d, J = 1.00 Hz, 1H), 7.96 (d, J = 9.12 Hz, 1H), 7.78 (br, 1H), 7.64 (dd, J = 9.12, 2.64 Hz, 1H), 3.99 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G12 Temperature at 100°C
418			HCl	¹ H NMR (DMSO-d ₆) ppm 12.60 (br, 1H), 9.79 (d, J = 1.72 Hz, 1H), 9.19 (d, J = 8.00 Hz, 1H), 8.93 (dd, J = 5.16, 1.44 Hz, 1H), 8.23 (brs, 1H), 8.10 (s, 1H), 7.99 (m, 1H), 7.95 (d, J = 9.08 Hz, 1H), 7.77-7.75 (br, 2H), 7.62 (dd, J = 9.08, 2.72 Hz, 1H), 7.50 (m, 2H), 7.37 (m, 1H), 3.99 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G12 Temperature at 100°C
419			HCl	¹ H NMR (DMSO-d ₆) ppm 12.56 (s, 1H), 9.78 (s, 1H), 9.09 (d, J = 8.20 Hz, 1H), 8.84 (d, J = 3.96 Hz, 1H), 8.35 (d, J = 2.60 Hz, 1H), 8.10-8.07 (m, 2H), 7.95 (d, J = 9.12 Hz, 1H), 7.83 (brt, J = 6.60 Hz, 2H), 7.63 (dd, J = 9.12, 2.60 Hz, 1H), 7.32 (m, 2H), 4.02 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G12 Temperature at 100°C
420			HCl	¹ H NMR (DMSO-d ₆) ppm 9.74 (s, 1H), 9.04 (d, J = 6.68 Hz, 1H), 8.86 (brs, 1H), 8.29 (brs, 1H), 8.05 (brs, J = 7.44 Hz, 2H), 7.97 (d, J = 9.08 Hz, 1H), 7.89 (br, 1H), 7.67-7.58 (m, 4H), 4.00 (s, 3H). The 1H of HCl and NH- were not observed.	DMSO	>98	G12 Temperature at 100°C

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424				HCl	1H NMR (DMSO-d6) ppm 10.30-10.15 (br, 1H), 9.54 (s, 1H), 8.91 (brs, 1H), 8.82 (brs, 1H), 8.10 (brs, 1H), 8.04 (d, J = 8.56 Hz, 2H), 7.92 (d, J = 9.6 Hz, 1H), 7.85 (d, J = 8.56 Hz, 2H), 7.80 (m, 1H), 7.79 (d, J = 8.56 Hz, 2H), 7.65-7.59 (brm, 1H), 7.55 (d, J = 8.56 Hz, 2H), 4.01 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	33		
425				HCl	1H NMR (DMSO-d6) ppm 9.91 (s, 1H), 9.53 (d, J = 1.84 Hz, 1H), 8.69 (d, J = 5.56 Hz, 2H), 8.31 (t, J = 1.88 Hz, 1H), 7.99 (d, J = 2.68 Hz, 1H), 7.98-7.94 (m, 1H), 7.87 (d, J = 9.08 Hz, 1H), 7.60-7.54 (m, 2H), 7.46 (t, J = 8.04 Hz, 1H), 7.39-7.35 (m, 1H), 3.99 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G1		
426				HCl	1H NMR (DMSO-d6) ppm 10.25 (br, 1H), 9.52 (s, 1H), 8.89 (brs, 1H), 8.88 (brs, 1H), 8.22 (brs, 1H), 8.11 (brs, 1H), 7.98-7.92 (brm, 2H), 7.80-7.70 (brm, 3H), 7.63-7.69 (brm, 2H), 7.55-7.50 (brm, 3H), 7.41 (m, 1H), 4.00 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13 using K ₂ CO ₃ instead of Cs ₂ CO ₃ at 120 °C		

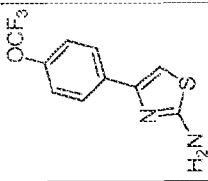
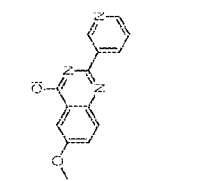
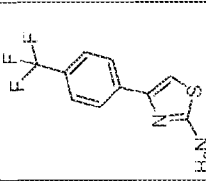
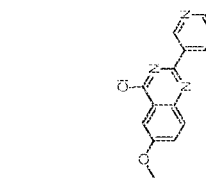
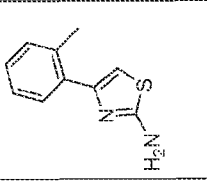
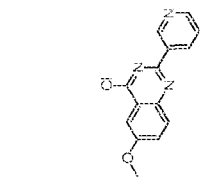
427				HCl	¹H NMR (DMSO-d₆) ppm 10.31 (br, 1H), 9.51 (brs, 1H), 8.92 (brd, J = 7.44 Hz, 1H), 8.84 (brs, 1H), 8.22 (brs, 1H), 8.12 (brd, J = 2.28 Hz, 1H), 8.00-7.93 (brm, 2H), 7.83-7.77 (brm, 3H), 7.64-7.54 (brm, 5H), 4.01 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	J3 using Na ₂ CO ₃ instead of K ₂ CO ₃		
428				4 HCl	¹H NMR (DMSO-d₆) ppm 9.73 (d, J = 1.72 Hz, 1H), 9.06 (d, J = 8.08 Hz, 1H), 8.88 (d, J = 3.72 Hz, 1H), 8.28 (brs, 1H), 8.01-7.96 (m, 3H), 7.88 (m, 1H), 7.81 (d, J = 8.60 Hz, 2H), 7.65 (dd, J = 9.08, 2.64 Hz, 1H), 4.00 (s, 3H). The 1H of 4HCl and NH- were not observed.	DMSO	>98	G13 using K ₂ CO ₃ instead of Cs ₂ CO ₃		
429				HCl	¹H NMR (DMSO-d₆) ppm 12.60 (br, 1H), 9.78 (d, J = 1.76 Hz, 1H), 9.28 (d, J = 8.04 Hz, 1H), 8.95 (dd, J = 5.20, 1.20 Hz, 1H), 8.36 (d, J = 2.28 Hz, 1H), 8.04-7.93 (m, 4H), 7.77 (s, 1H), 7.64 (dd, J = 9.12, 2.68 Hz, 1H), 7.30 (d, J = 8.04 Hz, 2H), 4.02 (s, 3H), 2.36 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13 using K ₂ CO ₃ instead of Cs ₂ CO ₃		

430				2 HCl	¹ H NMR (DMSO-d ₆) ppm 12.70-12.40 (br, 1H), 9.71 (s, 1H), 9.27 (d, J = 8.20 Hz, 1H), 8.94 (d, J = 4.24 Hz, 1H), 8.34 (d, J = 2.64 Hz, 1H), 8.13 (t, J = 1.80 Hz, 1H), 8.04-7.95 (m, 4H), 7.63 (dd, J = 9.08, 2.68 Hz, 1H), 7.52 (t, J = 7.88 Hz, 1H), 7.44-7.42 (m, 1H), 4.02 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13 using K ₂ CO ₃ instead of Cs ₂ CO ₃		
431				HCl	¹ H NMR (DMSO-d ₆) ppm 12.57 (br, 1H), 9.80 (brs, 1H), 9.04 (d, J = 7.36 Hz, 1H), 8.82 (brs, 1H), 8.34 (brs, 1H), 7.99 (brd, J = 7.28 Hz, 1H), 7.95 (d, J = 9.08 Hz, 1H), 7.82 (brs, 1H), 7.76 (brm, 1H), 7.62 (m, 2H), 7.50-7.41 (m, 2H), 4.00 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13 using K ₂ CO ₃ instead of Cs ₂ CO ₃		
432				HCl	¹ H NMR (DMSO-d ₆) ppm 10.40 (br, 1H), 9.23 (s, 1H), 8.80 (d, J = 4.68 Hz, 1H), 8.66 (brd, J = 8.20 Hz, 1H), 7.92 (brs, 1H), 7.87 (brd, J = 9.28 Hz, 1H), 7.78 (brm, 1H), 7.63-7.52 (m, 5H), 7.47-7.44 (m, 2H), 7.21-7.12 (m, 3H), 3.91 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13 using K ₂ CO ₃ instead of Cs ₂ CO ₃		

433			HCl	1H NMR (DMSO-d6) ppm 12.34 (br, 1H), 9.75 (s, 1H), 9.01 (d, J = 7.52 Hz, 1H), 8.81 (brs, 1H), 8.40-8.20 (br, 1H), 7.92 (d, J = 9.08 Hz, 1H), 7.77 (m, 1H), 7.60 (dd, J = 9.08, 2.72 Hz, 1H), 6.95 (br, 1H), 3.98 (s, 3H), 2.67 (m, 1H), 2.05 (m, 2H), 1.83-1.70 (m, 3H), 1.54-1.23 (m, 5H). The 1H of HCl was not observed.	DMSO	>98	G13 using K ₂ CO ₃ instead of Cs ₂ CO ₃		
434			2 HCl	1H NMR (DMSO-d6) ppm 12.58 (br, 1H), 9.79 (s, 1H), 9.19 (d, J = 7.32 Hz, 1H), 8.90 (d, J = 5.6 Hz, 1H), 8.36 (s, 1H), 7.96 (d, J = 9.08 Hz, 1H), 7.94 (m, 1H), 7.88 (s, 1H), 7.65-7.62 (m, 3H), 7.40 (t, J = 8.16 Hz, 1H), 6.96-6.93 (m, 1H), 4.02 (s, 3H), 3.84 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13 using K ₂ CO ₃ instead of Cs ₂ CO ₃		
435			2 HCl	1H NMR (DMSO-d6) ppm 12.80-12.40 (br, 1H), 9.79 (d, J = 1.44 Hz, 1H), 9.26 (d, J = 8.24 Hz, 1H), 8.93 (d, J = 4.08 Hz, 1H), 8.36 (d, J = 2.28 Hz, 1H), 8.02-7.96 (m, 4H), 7.68 (s, 1H), 7.64 (dd, J = 9.12, 2.68 Hz, 1H), 7.05 (d, J = 8.92 Hz, 2H), 4.02 (s, 3H), 3.82 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13 using K ₂ CO ₃ instead of Cs ₂ CO ₃		

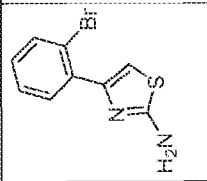
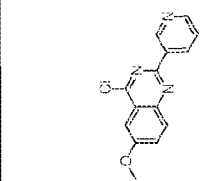
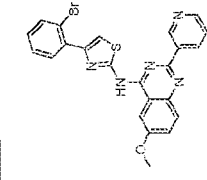
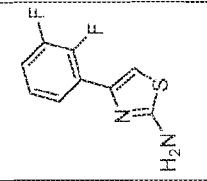
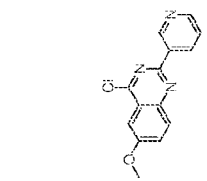
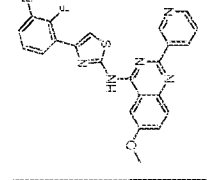
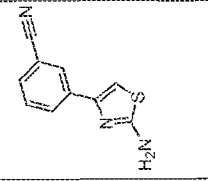
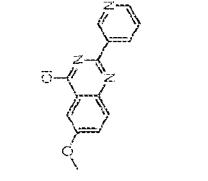
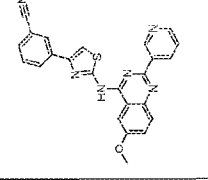
436				HCl	¹ H NMR (DMSO-d6) ppm 12.52 (br, 1H), 9.79 (s, 1H), 9.23 (d, J = 8.00 Hz, 1H), 8.91 (d, J = 4.12 Hz, 1H), 8.35 (brs, 1H), 8.25 (d, J = 7.6 Hz, 1H), 7.96 (d, J = 9.08 Hz, 1H), 7.93 (m, 1H), 7.85 (s, 1H), 7.64 (dd, J = 9.08, 2.68 Hz, 1H), 7.39-7.35 (m, 1H), 7.18 (d, J = 7.72 Hz, 1H), 7.11-7.07 (m, 1H), 4.02 (s, 3H), 3.96 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13 using K ₂ CO ₃ instead of Cs ₂ CO ₃		
437					¹ H NMR (DMSO-d6) ppm 9.94 (s, 1H), 9.27 (d, J = 1.92 Hz, 1H), 8.59 (dd, J = 4.72, 1.68 Hz, 1H), 8.45 (dt, J = 8.04, 1.8 Hz, 1H), 7.96 (d, J = 2.72 Hz, 1H), 7.84 (d, J = 8.28 Hz, 2H), 7.67 (d, J = 8.00 Hz, 1H), 7.55 (m, 2H), 7.46-7.43 (m, 1H), 7.33 (m, 1H), 3.96 (s, 3H).	DMSO	>98	G13 using K ₂ CO ₃ instead of Cs ₂ CO ₃ at 120 °C		
438				HCl	¹ H NMR (DMSO-d6) ppm 10.48 (br, 1H), 9.23 (s, 1H), 8.83 (brd, J = 3.68 Hz, 1H), 8.70 (brd, J = 7.36 Hz, 1H), 7.95 (brs, 1H), 7.91 (d, J = 9.12 Hz, 1H), 7.82 (m, 1H), 7.63-7.51 (m, 5H), 7.48 (d, J = 8.52 Hz, 2H), 7.26 (d, J = 8.52 Hz, 2H), 3.93 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	J3 using Na ₂ CO ₃ instead of K ₂ CO ₃		

439			HCl	¹H NMR (DMSO-d₆) ppm 12.60 (br, 1H), 9.78 (s, 1H), 9.17 (d, J = 7.96 Hz, 1H), 8.88 (d, J = 4.20 Hz, 1H), 8.35 (d, J = 2.60 Hz, 1H), 8.24-8.21 (m, 1H), 7.96 (d, J = 9.08 Hz, 1H), 7.91-7.88 (brm, 1H), 7.74 (d, J = 2.48 Hz, 1H), 7.63 (dd, J = 9.08, 2.60 Hz, 1H), 7.47-7.41 (m, 1H), 7.36 (m, 2H), 4.02 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13 at 100 °C		
440			HCl	¹H NMR (DMSO-d₆) ppm 12.60-12.45 (br, 1H), 9.79 (d, J = 1.52 Hz, 1H), 9.15 (d, J = 8.36 Hz, 1H), 8.88 (d, J = 5.04 Hz, 1H), 8.35 (d, J = 2.44 Hz, 1H), 7.96 (d, J = 9.12 Hz, 1H), 7.89 (brt, J = 7.08 Hz, 2H), 7.82 (brm, 1H), 7.81 (s, 1H), 7.63 (dd, J = 9.12, 2.44 Hz, 1H), 7.37 (t, J = 7.64 Hz, 1H), 7.19 (d, J = 7.48 Hz, 1H), 4.02 (s, 3H), 2.40 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13 at 100 °C		
441			HCl	¹H NMR (DMSO-d₆) ppm 12.60 (br, 1H), 9.77 (s, 1H), 9.25 (d, J = 8.08 Hz, 1H), 8.93 (d, J = 4.28 Hz, 1H), 8.34 (d, J = 2.60 Hz, 1H), 8.01-7.84 (m, 5H), 7.63 (dd, J = 9.12, 2.60 Hz, 1H), 7.56-7.51 (m, 1H), 7.21 (m, 1H). The 1H of HCl was not observed.	DMSO	>98	G13 at 100 °C		

442			2 HCl	1H NMR (DMSO-d6) ppm 12.66 (br, 1H), 9.78 (d, J = 1.68 Hz, 1H), 9.27 (d, J = 8.08 Hz, 1H), 8.94 (dd, J = 5.20, 1.40 Hz, 1H), 8.36 (d, J = 2.56 Hz, 1H), 8.17 (d, J = 8.84 Hz, 2H), 8.01 (m, 1H), 7.97 (d, J = 9.08 Hz, 1H), 7.93 (s, 1H), 7.64 (dd, J = 9.08, 2.56 Hz, 1H), 7.49 (brd, J = 8.84 Hz, 2H), 4.02 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13 at 100 °C		
443			2 HCl	1H NMR (DMSO-d6) ppm 12.64 (br, 1H), 9.79 (s, 1H), 9.13 (d, J = 7.56 Hz, 1H), 8.87 (d, J = 5.08 Hz, 1H), 8.36 (d, J = 2.60 Hz, 1H), 8.27 (d, J = 8.00 Hz, 2H), 8.08 (s, 1H), 7.96 (d, J = 9.12 Hz, 1H), 7.87 (m, 1H), 7.86 (d, J = 8.00 Hz, 2H), 7.64 (dd, J = 9.12, 2.60 Hz, 1H), 4.02 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13 at 100 °C		
444			2 HCl	1H NMR (MeOD-d4) ppm 9.79 (s, 1H), 9.37 (br, 1H), 8.49 (dd, J = 5.28, 1.32 Hz, 1H), 8.12 (m, 1H), 8.09 (d, J = 2.72 Hz, 1H), 7.98 (d, J = 9.16 Hz, 1H), 7.68 (dd, J = 9.16, 2.72 Hz, 1H), 7.58 (d, J = 7.12 Hz, 1H), 7.42-7.30 (m, 3H), 7.30 (s, 1H), 4.05 (s, 3H), 2.50 (s, 3H). The 1H of 2HCl was not observed.	MeOD	>96	G13		

445			2 HCl	¹H NMR (DMSO-d₆) ppm: 12.59 (br, 1H), 9.78 (d, J = 1.48 Hz, 1H), 9.19 (d, J = 8.00 Hz, 1H), 8.90 (dd, J = 5.16, 1.48 Hz, 1H), 8.34 (d, J = 2.60 Hz, 1H), 8.10-8.04 (m, 1H), 7.97-7.88 (m, 4H), 7.64 (dd, J = 2.60 Hz, 1H), 7.59-7.52 (m, 1H), 4.02 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13		
446			HCl	¹H NMR (DMSO-d₆) ppm: 12.61 (br, 1H), 9.78 (d, J = 1.72 Hz, 1H), 9.20 (d, J = 7.92 Hz, 1H), 8.90 (d, J = 4.00 Hz, 1H), 8.35 (d, J = 2.60 Hz, 1H), 8.27-8.21 (m, 1H), 7.96 (d, J = 9.12 Hz, 1H), 7.93 (m, 1H), 7.71 (d, J = 2.56 Hz, 1H), 7.64 (dd, J = 9.12, 2.60 Hz, 1H), 7.43 (m, 1H), 7.27 (m, 1H), 4.02 (s, 1H). The 1H of HCl was not observed.	DMSO	>98	G13		
447			2 HCl	¹H NMR (DMSO-d₆) ppm: 12.57 (br, 1H), 9.77 (d, J = 1.76 Hz, 1H), 9.25 (d, J = 8.08 Hz, 1H), 8.93 (dd, J = 5.16, 1.2 Hz, 1H), 8.34 (d, J = 2.60 Hz, 1H), 8.29 (t, J = 1.76 Hz, 1H), 8.05 (dd, J = 7.94, 1.12 Hz, 1H), 8.02-7.98 (m, 1H), 7.99 (s, 1H), 7.96 (d, J = 9.08 Hz, 1H), 7.64 (dd, J = 9.08, 2.60 Hz, 1H), 7.58-7.56 (m, 1H), 7.45 (t, J = 7.84 Hz, 1H), 4.02 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13		

448			HCl	¹H NMR (DMSO-d₆) ppm 12.62 (br, 1H), 9.77 (d, J = 1.52 Hz, 1H), 9.10 (d, J = 8.16 Hz, 1H), 8.85 (dd, J = 5.00, 1.56 Hz, 1H), 8.34 (d, J = 2.60 Hz, 1H), 8.24 (d, J = 8.60 Hz, 2H), 8.13 (s, 1H), 7.96 (d, J = 8.60 Hz, 2H), 7.96 (d, J = 9.08 Hz, 1H), 7.86-7.83 (brm, 1H), 7.63 (dd, J = 9.08, 2.60 Hz, 1H), 4.02 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13		
449			2 HCl	¹H NMR (DMSO-d₆) ppm 12.56 (brs, 1H), 9.79 (brs, 1H), 9.11 (d, J = 8.04 Hz, 1H), 8.87 (brs, 1H), 8.43 (brs, 1H), 8.34 (brs, 2H), 8.09 (s, 1H), 7.96 (d, J = 9.08 Hz, 1H), 7.87 (brs, 1H), 7.74 (brd, J = 5.00 Hz, 2H), 7.64 (dd, J = 9.08, 2.48 Hz, 1H), 4.03 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13 at 100 °C		
450			2 HCl	¹H NMR (DMSO-d₆) ppm 12.63 (br, 1H), 9.80 (d, J = 1.72 Hz, 1H), 9.27 (d, J = 8.16 Hz, 1H), 8.93 (dd, J = 5.20, 1.36 Hz, 1H), 8.33 (d, J = 2.24 Hz, 1H), 7.99 (m, 1H), 7.98 (d, J = 9.08 Hz, 1H), 7.88 (brd, J = 7.72 Hz, 1H), 7.71-7.75 (m, 2H), 7.70-7.63 (m, 2H), 7.45 (s, 1H), 3.98 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13 at 100 °C		

451				HCl	1H NMR (DMSO-d6) ppm 12.70 (br, 1H), 9.79 (d, J = 1.60 Hz, 1H), 9.30 (d, J = 7.56 Hz, 1H), 8.95 (d, J = 4.96 Hz, 1H), 8.34 (d, J = 2.20 Hz, 1H), 8.02 (m, 1H), 7.98 (d, J = 9.12 Hz, 1H), 7.85-7.77 (m, 2H), 7.73 (s, 1H), 7.64 (dd, J = 9.12, 2.58 Hz, 1H), 7.52 (m, 1H), 7.36 (m, 1H), 4.00 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13		
452				HCl	1H NMR (DMSO-d6) ppm 12.64 (br, 1H), 9.78 (d, J = 1.72 Hz, 1H), 9.19 (d, J = 7.92 Hz, 1H), 8.90 (d, J = 3.88 Hz, 1H), 8.36 (d, J = 2.64 Hz, 1H), 8.02-7.91 (m, 3H), 7.83 (d, J = 2.40 Hz, 1H), 7.64 (dd, J = 9.08, 2.64 Hz, 1H), 7.46 (m, 1H), 7.36 (m, 1H), 4.02 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13		
453				2 HCl	1H NMR (DMSO-d6) ppm 12.62 (br, 1H), 9.77 (d, J = 1.84 Hz, 1H), 9.30 (d, J = 8.00 Hz, 1H), 8.46 (dd, J = 5.36, 1.48 Hz, 1H), 8.48 (br, 1H), 8.39-8.33 (m, 2H), 8.08 (s, 1H), 8.05 (m, 1H), 7.97 (d, J = 9.12 Hz, 1H), 7.85-7.83 (m, 1H), 7.71 (t, J = 7.84 Hz, 1H), 7.64 (dd, J = 9.12, 2.60 Hz, 1H), 4.02 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13		

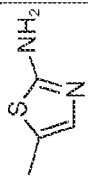
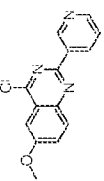
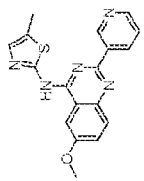
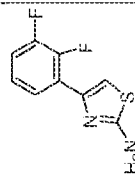
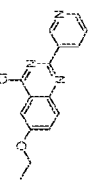
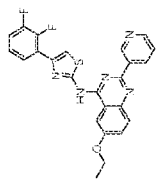
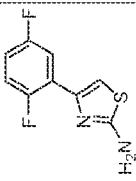
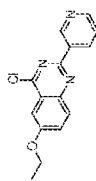
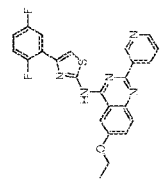
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	454			2 HCl	¹H NMR (DMSO-d6) ppm 12.65 (br, 1H), 9.79 (d, J = 1.72 Hz, 1H), 9.25 (d, J = 8.04 Hz, 1H), 8.92 (d, J = 5.36 Hz, 1H), 8.36 (d, J = 2.60 Hz, 1H), 8.24 (s, 1H), 7.97 (d, J = 9.08 Hz, 1H), 7.96 (m, 1H), 7.69 (s, 1H), 7.65 (dd, J = 9.08, 2.60 Hz, 1H), 7.55 (m, 3H), 4.02 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13	
	455			HCl	¹H NMR (DMSO-d6) ppm 12.52 (br, 1H), 9.78 (d, J = 1.48 Hz, 1H), 9.03 (d, J = 6.6 Hz, 1H), 8.81 (d, J = 4.72 Hz, 1H), 8.34 (d, J = 2.56 Hz, 1H), 8.09 (d, J = 7.89 Hz, 1H), 8.03 (brs, 2H), 7.96 (d, J = 9.12 Hz, 1H), 7.78 (br, 1H), 7.65 (m, 2H), 7.38 (m, 1H), 4.02 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13	
	456			2 HCl	¹H NMR (DMSO-d6) ppm 12.45 (br, 1H), 9.78 (d, J = 1.64 Hz, 1H), 9.15 (br, 1H), 8.89 (d, J = 4.52 Hz, 1H), 8.30 (brs, 1H), 7.94 (d, J = 9.08 Hz, 1H), 7.90 (brm, 1H), 7.76 (m, 2H), 7.62 (dd, J = 9.08, 2.64 Hz, 1H), 7.51 (m, 2H), 7.40 (m, 1H), 3.99 (s, 3H), 2.61 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13	

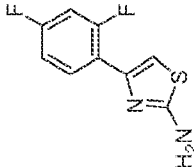
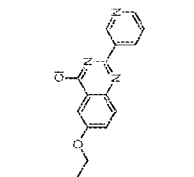
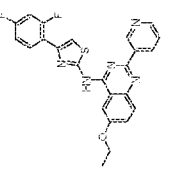
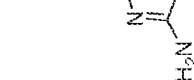
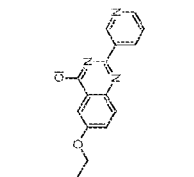
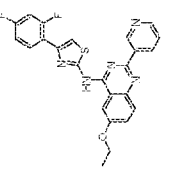
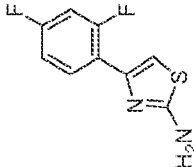
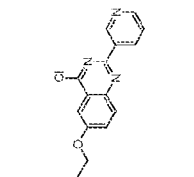
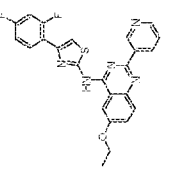
457			HCl	¹H NMR (DMSO-d₆) ppm: 12.55 (br, 1H), 9.78 (d, J = 1.48 Hz, 1H), 9.08 (d, J = 8.48 Hz, 1H), 8.84 (dd, J = 5.04, 1.56 Hz, 1H), 8.34 (d, J = 2.72 Hz, 1H), 8.09 (s, 1H), 7.96 (d, J = 9.12 Hz, 1H), 7.84 (m, 1H), 7.77 (m, 2H), 7.64 (dd, J = 9.12, 2.72 Hz, 1H), 7.25 (m, 1H), 4.02 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13		
458			HCl	¹H NMR (DMSO-d₆) ppm: 12.55 (br, 1H), 9.77 (d, J = 1.56 Hz, 1H), 9.14 (d, J = 8.00 Hz, 1H), 8.87 (dd, J = 5.08, 1.48 Hz, 1H), 8.34 (d, J = 2.60 Hz, 1H), 7.98 (m, 1H), 7.96 (d, J = 9.08 Hz, 1H), 7.89-7.86 (brm, 1H), 7.83 (d, J = 2.44 Hz, 1H), 7.64 (dd, J = 9.08, 2.60 Hz, 1H), 7.47-7.41 (m, 1H), 7.29 (m, 1H), 4.02 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13		
459			HCl	¹H NMR (DMSO-d₆) ppm: 12.66 (br, 1H), 9.79 (d, J = 1.60 Hz, 1H), 9.25 (d, J = 7.16 Hz, 1H), 8.92 (d, J = 5.20 Hz, 1H), 8.35 (d, J = 2.56 Hz, 1H), 7.98 (d, J = 9.12 Hz, 1H), 7.97 (m, 1H), 7.68 (s, 1H), 7.64 (dd, J = 9.12, 2.56 Hz, 1H), 7.55 (m, 1H), 7.27 (m, 2H), 3.99 (s, 3H). The 1H of HCl was not observed.	DMSO	>98	G13		

460			2 HCl	¹H NMR (DMSO-d₆) ppm 9.73 (s, 1H), 9.17 (brs, 1H), 8.92 (dd, J = 5.16, 1.4 Hz, 1H), 8.30-8.10 (br, 1H), 7.97 (d, J = 9.04 Hz, 1H), 7.96 (m, 1H), 7.63 (dd, J = 9.04, 2.72 Hz, 1H), 7.02 (s, 1H), 3.98 (s, 3H), 2.57 (d, J = 7.16 Hz, 2H), 2.08 (m, 1H), 0.94 (d, J = 6.60 Hz, 6H). The 1H of HCl and NH₂ were not observed.	DMSO	>98	G13		
461			2 HCl	¹H NMR (DMSO-d₆) ppm 12.80-12.20 (br, 1H), 9.78 (d, J = 1.76 Hz, 1H), 9.28 (d, J = 8.04 Hz, 1H), 8.96 (dd, J = 5.28, 1.32 Hz, 1H), 8.29 (brs, 1H), 8.07-8.03 (brm, 1H), 7.95 (d, J = 9.12 Hz, 1H), 7.79 (d, J = 8.60 Hz, 2H), 7.62 (dd, J = 9.12, 2.64 Hz, 1H), 7.57 (d, J = 8.60 Hz, 2H), 3.99 (s, 3H), 2.61 (brs, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13		
462			2 HCl	¹H NMR (DMSO-d₆) ppm 12.80-12.20 (br, 1H), 9.74 (s, 1H), 9.19 (d, J = 4.96 Hz, 1H), 8.91 (dd, J = 5.2, 1.4 Hz, 1H), 8.22 (brs, 1H), 7.96 (br, 1H), 7.95 (d, J = 9.08 Hz, 1H), 7.61 (dd, J = 9.08, 2.68 Hz, 1H), 7.35-7.31 (m, 4H), 7.23 (m, 1H), 7.09 (brs, 1H), 4.07 (brs, 2H), 3.96 (brs, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13		

463			2 HCl		1H NMR (DMSO-d6) ppm: 12.50 (br, 1H), 9.78 (d, J = 1.76 Hz, 1H), 9.25 (brs, 1H), 8.96 (brd, J = 4.68 Hz, 1H), 8.29 (brs, 1H), 8.04 (brs, 1H), 7.96 (d, J = 9.20 Hz, 1H), 7.70 (brm, 2H), 7.63 (dd, J = 9.20, 2.68 Hz, 1H), 7.51 (brm, 2H), 7.41 (brm, 1H), 3.99 (s, 3H), 3.02 (q, J = 7.40 Hz, 2H), 1.49 (t, J = 7.40 Hz, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13		
464			2 HCl		1H NMR (DMSO-d6) ppm: 12.53 (br, 1H), 9.78 (d, J = 1.76 Hz, 1H), 9.20 (d, J = 8.36 Hz, 1H), 8.94 (dd, J = 5.2, 1.4 Hz, 1H), 8.29 (brs, 1H), 8.00 (brm, 1H), 7.96 (d, J = 9.12 Hz, 1H), 7.66 - 7.61 (m, 3H), 7.52 (brm, 2H), 7.43 (t, J = 7.40 Hz, 1H), 3.98 (s, 3H), 1.43 (d, J = 6.8 Hz, 6H). The 1H of 2HCl was not observed. The CH proton of iPr group was not observed because of overlapping the H2O peak.	DMSO	>98	G13		
465			2 HCl		1H NMR (DMSO-d6) ppm: 12.59 (brs, 1H), 9.77 (d, J = 1.72 Hz, 1H), 9.25 (d, J = 7.76 Hz, 1H), 8.96 (d, J = 4.04 Hz, 1H), 8.29 (br, 1H), 8.04 (m, 1H), 7.96 (d, J = 9.12 Hz, 1H), 7.69 (brd, J = 7.12 Hz, 2H), 7.63 (dd, J = 9.12, 2.68 Hz, 1H), 7.51 (t, J = 7.36 Hz, 2H), 7.42 (t, J = 7.36 Hz, 1H), 3.99 (s, 3H), 2.96 (t, J = 7.52 Hz, 2H), 1.78 (m, 2H), 1.01 (t, J = 7.24 Hz, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13		

466			2 HCl	¹H NMR (DMSO-d₆) ppm 12.96 (brs, 1H), 9.74 (d, J = 2.04 Hz, 1H), 9.11 (d, J = 7.96 Hz, 1H), 8.90 (d, J = 4.96 Hz, 1H), 8.31 (brd, J = 2.64 Hz, 1H), 7.97 (d, J = 9.12 Hz, 1H), 7.87 (brm, 1H), 7.83-7.80 (m, 2H), 7.64 (dd, J = 9.12, 2.62 Hz, 1H), 7.48 (m, 3H), 4.25 (q, J = 7.04 Hz, 2H), 3.98 (s, 3H), 1.29 (t, J = 7.04 Hz, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13		
467			HCl	¹H NMR (DMSO-d₆) ppm 12.63 (br, 1H), 9.76 (d, J = 1.76 Hz, 1H), 9.29 (d, J = 8.16 Hz, 1H), 8.95 (dd, J = 5.28, 1.36 Hz, 1H), 8.32 (brd, J = 2.60 Hz, 1H), 8.07 (d, J = 8.52 Hz, 2H), 8.08-8.00 (m, 1H), 7.95 (d, J = 9.12 Hz, 1H), 7.90 (s, 1H), 7.61 (dd, J = 9.12, 2.60 Hz, 1H), 7.55 (d, J = 8.52 Hz, 2H), 4.24 (q, J = 6.88 Hz, 2H), 1.47 (t, J = 6.88 Hz, 3H). The 1H of HCl was not observed.	DMSO	>98	G13		
468			2 HCl	¹H NMR (DMSO-d₆) ppm 12.58 (brs, 1H), 9.76 (d, J = 1.80 Hz, 1H), 9.27 (d, J = 8.16 Hz, 1H), 8.94 (dd, J = 5.20, 1.32 Hz, 1H), 8.32 (d, J = 2.48 Hz, 1H), 8.09-8.01 (m, 2H), 7.96-7.89 (m, 3H), 7.62-7.52 (m, 2H), 4.29 (q, J = 6.88 Hz, 2H), 1.47 (t, J = 6.88 Hz, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13		

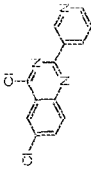
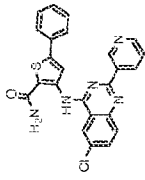
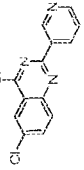
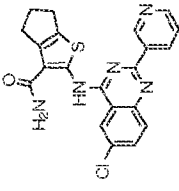
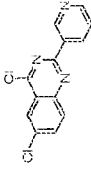
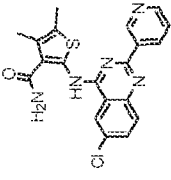
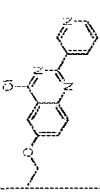
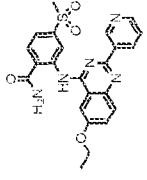
469				2 HCl	¹H NMR (DMSO-d₆) ppm 9.72 (s, 1H), 9.13 (brd, J = 5.52 Hz, 1H), 8.92 (d, J = 3.52 Hz, 1H), 8.16 (brs, 1H), 7.95 (brd, J = 9.08 Hz, 2H), 7.62 (dd, J = 2.4 Hz, 1H), 7.39 (brs, 1H), 3.97 (s, 3H), 2.47 (s, 3H). The 1H of 2HCl and NH- were not observed.	DMSO	>98	G13		
470				HCl	¹H NMR (DMSO-d₆) ppm 12.58 (brs, 1H), 9.76 (d, J = 1.68 Hz, 1H), 9.17 (d, J = 8.12 Hz, 1H), 8.89 (dd, J = 5.12, 1.48 Hz, 1H), 8.32 (d, J = 2.56 Hz, 1H), 8.01-8.00 (m, 1H), 7.94 (d, J = 9.08 Hz, 1H), 7.930-7.90 (m, 1H), 7.81 (d, J = 2.4 Hz, 1H), 7.60 (dd, J = 9.08, 2.40 Hz, 1H), 7.50-7.43 (m, 1H), 7.38-7.33 (m, 1H), 4.28 (q, J = 7.00 Hz, 2H), 1.47 (t, J = 7.00 Hz, 3H). The 1H of HCl was not observed.	DMSO	>98	G13		
471				2 HCl	¹H NMR (DMSO-d₆) ppm 12.53 (brs, 1H), 9.73 (d, J = 1.76 Hz, 1H), 9.29 (d, J = 8.16 Hz, 1H), 8.95 (dd, J = 5.32, 1.36 Hz, 1H), 8.29 (d, J = 2.56 Hz, 1H), 8.05-8.02 (brm, 1H), 7.97-7.92 (m, 1H), 7.93 (d, J = 9.08 Hz, 1H), 7.78 (d, J = 2.36 Hz, 1H), 7.60 (dd, J = 9.08, 2.6 Hz, 1H), 7.46-7.40 (m, 1H), 7.31-7.25 (m, 1H), 4.27 (q, J = 6.92 Hz, 2H), 1.47 (t, J = 6.92 Hz, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13		

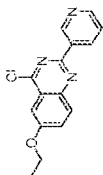
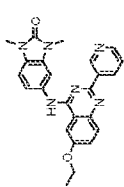
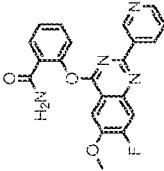
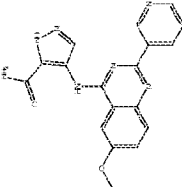
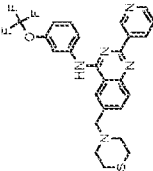
472				3 HCl	¹ H NMR (DMSO-d ₆) ppm 12.61 (brs, 1H), 9.76 (d, J = 1.8 Hz, 1H), 9.29 (d, J = 3.16 Hz, 1H), 8.95 (dd, J = 5.24, 1.2 Hz, 1H), 8.32 (d, J = 2.52 Hz, 1H), 8.24-8.20 (m, 1H), 8.05-8.01 (brm, 1H), 7.95 (d, J = 9.08 Hz, 1H), 7.68 (d, J = 2.52 Hz, 1H), 7.61 (dd, J = 9.08, 2.52 Hz, 1H), 7.46-7.40 (m, 1H), 7.28-7.24 (m, 1H), 4.28 (q, J = 6.92 Hz, 2H), 1.47 (t, J = 6.92 Hz, 3H). The 1H of 3HCl was not observed.	DMSO	>98	G13			
473				2 HCl	¹ H NMR (DMSO-d ₆) ppm 12.63 (brs, 1H), 9.78 (d, J = 1.68 Hz, 1H), 9.32 (d, J = 8.12 Hz, 1H), 8.97 (d, J = 5.24 Hz, 1H), 8.35 (d, J = 2.16 Hz, 1H), 8.08-8.04 (m, 3H), 7.96 (d, J = 9.08 Hz, 1H), 7.85 (s, 1H), 7.62 (dd, J = 9.08, 2.52 Hz, 1H), 7.49 (t, J = 7.44 Hz, 2H), 7.38 (t, J = 7.28 Hz, 1H), 4.30 (q, J = 6.96 Hz, 2H), 1.47 (t, J = 6.96 Hz, 3H). The 1H of 2HCl was not observed.	DMSO	>98	G13			
474					¹ H NMR (DMSO-d ₆) ppm 12.68 (s, 1H), 9.73 (d, J = 1.64 Hz, 1H), 8.87-8.84 (m, 1H), 8.75-8.73 (m, 1H), 8.33 (brs, 1H), 8.24 (brs, 1H), 7.93 (d, J = 9.16 Hz, 1H), 7.64-7.59 (m, 2H), 4.34 (q, J = 7.08 Hz, 2H), 3.98 (s, 3H), 1.34 (t, J = 7.08 Hz, 3H).	DMSO	>98	G13			

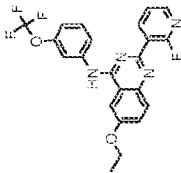
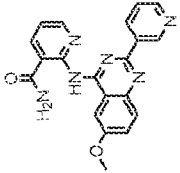
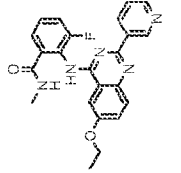
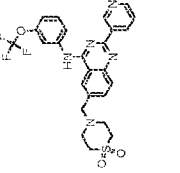
475			HCl	¹ H NMR (300 MHz, DMSO) δ 10.35 (s, 1H), 9.42 (s, 1H), 8.99 - 8.76 (m, 2H), 8.11 (d, J = 6.7 Hz, 1H), 7.99 - 7.75 (m, 3H), 7.67 - 7.43 (m, 2H), 7.23 (d, J = 8.2 Hz, 1H), 4.14 (s, 3H).	DMSO	98	G1	381 (M+1)	2.07	Method A (Formic acid)
476			HCl	¹ H NMR (300 MHz, DMSO) δ 13.78 (s, 1H), 9.62 (d, J = 1.7 Hz, 1H), 9.21 (d, J = 8.1 Hz, 1H), 8.98 (d, J = 4.2 Hz, 1H), 8.07 (dd, J = 7.9, 5.3 Hz, 1H), 7.83 (d, J = 9.1 Hz, 2H), 7.63 - 7.50 (m, 4H), 7.41 (d, J = 6.5 Hz, 2H), 7.14 (d, J = 2.5 Hz, 1H), 5.30 (s, 1H), 3.91 (s, 3H), 2.15 (s, 3H).	DMSO	95	Method G1	468.1 (M+1)		Method C
477			HCl	¹ H NMR (300 MHz, DMSO) δ 13.78 (s, 1H), 9.64 (d, J = 1.5 Hz, 1H), 9.11 (d, J = 8.2 Hz, 1H), 8.90 (d, J = 3.8 Hz, 1H), 7.95 (dd, J = 10.6, 2.6 Hz, 1H), 7.84 (d, J = 9.1 Hz, 2H), 7.60 - 7.48 (m, 4H), 7.38 (d, J = 6.5 Hz, 2H), 7.17 (d, J = 2.4 Hz, 1H), 5.28 (s, 1H), 4.14 (q, J = 6.9 Hz, 2H), 2.15 (s, 3H), 1.42 (t, J = 6.9 Hz, 3H).	DMSO	95	Method G1	482.1 (M+1)		Method C
478			HCl	¹ H NMR (300 MHz, DMSO) δ 12.59 (s, 1H), 9.56 (s, 1H), 8.70 (d, J = 5.8 Hz, 2H), 8.58 (d, J = 5.2 Hz, 1H), 7.95 (d, J = 5.2 Hz, 2H), 7.87 (d, J = 9.1 Hz, 2H), 7.64 - 7.49 (m, 2H), 7.40 (d, J = 2.4 Hz, 1H), 4.21 (q, J = 6.8 Hz, 2H), 1.44 (t, J = 6.9 Hz, 3H).	DMSO	99	Method G1	392.1 (M+1)		Method C

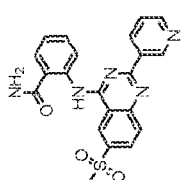
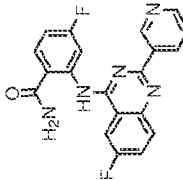
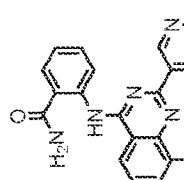
479			HCl	¹H NMR (300 MHz, DMSO) δ 13.75 (s, 1H), 9.60 (s, 1H), 8.98 (s, 1H), 8.82 (d, J = 3.6 Hz, 1H), 7.82 (d, J = 9.0 Hz, 3H), 7.52 (d, J = 9.3 Hz, 1H), 7.15 (s, 1H), 6.94 - 6.62 (s, 1H), 4.13 (d, J = 6.9 Hz, 2H), 2.90 (d, J = 21.9 Hz, 4H), 2.41 (s, 2H), 1.43 (t, J = 6.9 Hz, 3H).	DMSO	95	Method G1	432.1 (M+1)		Method C
480			HCl	¹H NMR (300 MHz, DMSO) δ 12.50 (s, 1H), 9.56 (s, 1H), 8.89 (s, 1H), 8.68 (d, J = 5.3 Hz, 2H), 8.00 - 7.79 (m, 3H), 7.73 (d, J = 7.2 Hz, 4H), 7.62 - 7.38 (m, 9H), 7.34 (d, J = 2.2 Hz, 2H), 4.18 (dd, J = 13.5, 6.5 Hz, 2H), 1.43 (t, J = 6.9 Hz, 3H).	DMSO	99	Method G1	468.1 (M+1)		Method C
481			HCl	¹H NMR (300 MHz, DMSO) δ 12.51 (s, 1H), 9.54 (s, 1H), 8.86 (s, 1H), 8.74 - 8.61 (m, 2H), 7.90 (s, J = 15.7 Hz, 2H), 7.83 (d, J = 9.1 Hz, 1H), 7.72 (d, J = 7.1 Hz, 2H), 7.63 - 7.40 (m, 6H), 7.33 (d, J = 2.3 Hz, 1H), 3.92 (s, 3H).	DMSO	99	Method G1	454.1 (M+1)		Method C
482			HCl	¹H NMR (300 MHz, DMSO) δ 13.70 (s, 1H), 9.70 (d, J = 2.1 Hz, 1H), 8.81 (d, J = 8.0 Hz, 1H), 8.70 (dd, J = 4.8, 1.7 Hz, 1H), 7.90 (d, J = 9.1 Hz, 1H), 7.66 - 7.53 (m, 2H), 7.29 (d, J = 2.2 Hz, 1H), 3.95 (s, 3H), 2.78 (d, J = 12.9 Hz, 4H), 1.78 (s, 4H).	DMSO	99	Method G1	432.1 (M+1)		Method C

483			HCl	¹H NMR (300 MHz, DMSO) δ 13.64 (s, 1H), 9.68 (s, 1H), 8.79 (d, J = 7.9 Hz, 1H), 8.69 (d, J = 4.7 Hz, 1H), 7.86 (d, J = 9.0 Hz, 1H), 7.63 - 7.50 (m, 2H), 7.24 (s, 1H), 4.18 (q, J = 6.7 Hz, 2H), 2.76 (d, J = 10.4 Hz, 4H), 1.78 (s, 4H), 1.43 (t, J = 6.9 Hz, 3H).	DMSO	99	Method G1	446.1 (M+1)		Method C
484			HCl	¹H NMR (300 MHz, DMSO) δ 12.82 (s, 1H), 9.67 (d, J = 2.1 Hz, 1H), 8.77 (dt, J = 8.1, 2.0 Hz, 1H), 8.71 (dd, J = 4.8, 1.7 Hz, 1H), 7.87 (d, J = 9.1 Hz, 1H), 7.83 (s, 1H), 7.60 - 7.53 (m, 3H), 7.48 (s, 1H), 7.20 (d, J = 2.6 Hz, 1H), 4.20 (q, J = 6.9 Hz, 2H), 2.90 (d, J = 4.7 Hz, 3H), 1.44 (t, J = 6.9 Hz, 3H).	DMSO	95	Method G1	422.1 (M+1)		Method C
485			HCl	¹H NMR (300 MHz, DMSO) δ 9.66 (s, 1H), 9.44 (d, J = 1.4 Hz, 1H), 8.64 - 8.55 (m, 2H), 7.90 (s, 1H), 7.81 (d, J = 9.1 Hz, 1H), 7.76 (d, J = 2.5 Hz, 1H), 7.69 (s, 2H), 7.54 - 7.45 (m, 2H), 4.19 (q, J = 7.0 Hz, 2H), 4.04 (s, 3H), 1.42 (t, J = 6.9 Hz, 3H).	DMSO	99	Method G1	390.1 (M+1)		Method C
486			2 HCl	¹H NMR (300 MHz, DMSO) δ 9.57 (d, J = 1.5 Hz, 1H), 9.07 (d, J = 4.6 Hz, 1H), 8.94 (dd, J = 5.2, 1.4 Hz, 1H), 8.04 (d, J = 9.1 Hz, 1H), 7.97 (dd, J = 6.7, 6.0 Hz, 1H), 7.67 (dd, J = 9.1, 2.8 Hz, 1H), 7.55 (d, J = 2.7 Hz, 1H), 7.36 (dd, J = 6.4, 2.3 Hz, 2H), 7.14 - 7.03 (m, 2H), 3.74 (s, 3H), 3.69 (s, 3H).	DMSO	99	Method G1	382.5 (M+1)		Method C

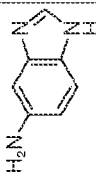
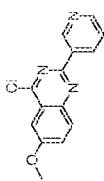
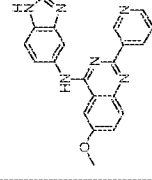
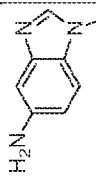
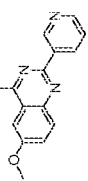
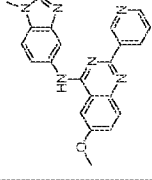
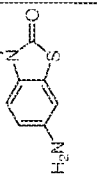
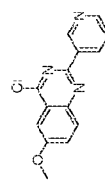
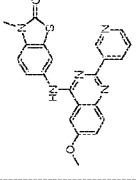
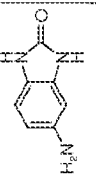
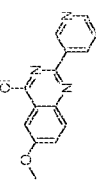
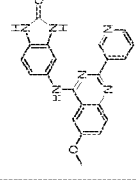
487			2 HCl	¹ H NMR (300 MHz, DMSO) δ 12.59 (s, 1H), 9.58 (d, J = 2.0 Hz, 1H), 8.94 (d, J = 8.2 Hz, 1H), 8.84 (d, J = 5.2 Hz, 1H), 8.73 (s, 1H), 8.09 (s, 1H), 7.95 (s, 3H), 7.82 (dd, J = 8.4, 5.4 Hz, 2H), 7.75 (d, J = 7.4 Hz, 2H), 7.59 - 7.42 (m, 3H).	DMSO	99	Method G1	458.4 (M+1)	Method C
488			2 HCl	¹ H NMR (300 MHz, DMSO) δ 13.79 (s, 1H), 9.49 (d, J = 1.5 Hz, 1H), 8.96 (d, J = 8.0 Hz, 1H), 8.86 (d, J = 3.7 Hz, 1H), 7.95 - 7.71 (m, 4H), 7.65 (d, J = 1.6 Hz, 1H), 6.76 (s, 1H), 2.85 (d, J = 22.5 Hz, 4H), 2.44 - 2.29 (m, 2H).	DMSO	93	Method G1	422.4 (M+1)	Method C
489			2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.53 (d, J = 1.8 Hz, 1H), 9.04 (d, J = 7.9 Hz, 1H), 8.93 (d, J = 3.9 Hz, 1H), 7.95 (dd, J = 8.0, 5.2 Hz, 1H), 7.90 - 7.78 (m, 2H), 7.70 (d, J = 1.7 Hz, 1H), 2.29 (s, 3H), 2.24 (s, 3H).	DMSO	94	Method G1	410.3 (M+1)	Method C
490				¹ H NMR (300 MHz, DMSO) δ 13.07 (s, 1H), 9.93 (d, J = 1.8 Hz, 1H), 9.64 (d, J = 1.6 Hz, 1H), 8.90 - 8.78 (m, 1H), 8.76 - 8.62 (m, 2H), 8.29 (s, 1H), 8.18 (d, J = 8.3 Hz, 1H), 7.89 (d, J = 9.1 Hz, 1H), 7.71 (dd, J = 8.3, 1.9 Hz, 1H), 7.65 (s, 1H), 7.57 (dd, J = 9.1, 2.5 Hz, 1H), 7.53 - 7.44 (m, 2H), 7.01 (s, 1H), 4.21 (q, J = 6.9 Hz, 2H), 3.34 (s, 3H), 1.44 (t, J = 6.9 Hz, 3H).	DMSO	99	Method G1	464.2 (M+1)	Method C

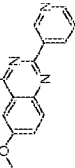
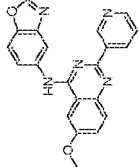
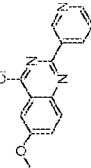
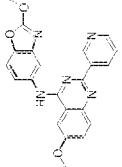
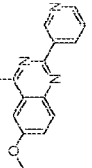
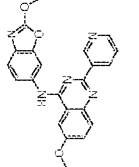
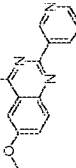
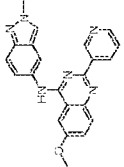
491				¹ H NMR (300 MHz, DMSO) δ 9.78 (s, 1H), 9.50 (d, J = 2.0 Hz, 1H), 8.68 - 8.57 (m, 2H), 7.97 (d, J = 2.5 Hz, 1H), 7.85 (d, J = 1.8 Hz, 1H), 7.79 (d, J = 9.1 Hz, 1H), 7.55 - 7.43 (m, 3H), 7.22 (d, J = 8.4 Hz, 1H), 4.22 (q, J = 6.9 Hz, 2H), 3.37 (d, J = 4.3 Hz, 6H), 1.43 (t, J = 6.9 Hz, 3H).	DMSO	99	Method G1	427.5 (M+1)		Method C
492				¹ H NMR (300 MHz, DMSO) δ 11.48 (s, 1H), 10.97 (s, 1H), 9.28 (d, J = 1.5 Hz, 1H), 8.64 (dd, J = 4.6, 1.7 Hz, 1H), 8.40 (d, J = 8.1 Hz, 1H), 7.94 - 7.80 (m, 3H), 7.53 - 7.35 (m, 2H), 7.02 (t, J = 7.5 Hz, 1H), 6.90 (d, J = 8.2 Hz, 1H), 4.04 (s, 1H).	DMSO	99	Method F5, G13	391.1 (M+1)		Method C
493			2 HCl	¹ H NMR (300 MHz, DMSO) δ 11.28 (s, 1H), 9.62 (d, J = 1.7 Hz, 1H), 9.17 (d, J = 7.8 Hz, 1H), 8.93 (dd, J = 5.2, 1.3 Hz, 1H), 8.60 (s, 1H), 8.05 (s, 1H), 8.03 - 7.94 (m, 2H), 7.77 (s, 1H), 7.64 (dd, J = 9.2, 2.5 Hz, 1H), 7.40 (d, J = 2.6 Hz, 1H), 3.97 (s, 3H).	DMSO	99	Method G14	362.4 (M+1)		Method C
494				¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.14 (s, 1H), 9.54 (s, 1H), 8.70 - 8.67 (m, 2H), 8.48 (s, 1H), 8.16 (s, 1H), 7.96 - 7.89 (m, 3H), 7.60 (t, J = 8.0 Hz, 1H), 7.55 - 7.52 (m, 1H), 7.17 (d, J = 8.8 Hz, 1H), 3.71 (s, 2H), 2.72 - 2.65 (m, 8H).	DMSO	95	Method G1	498.1 (M+1)	t = 2.128 min	Method B (NH ₄ HCO ₃)

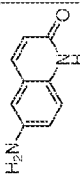
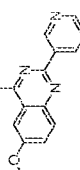
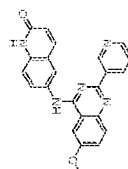
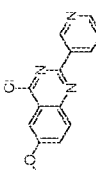
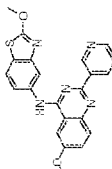
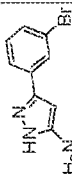
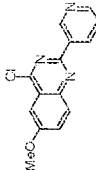
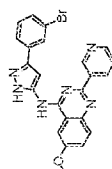
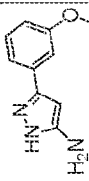
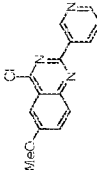
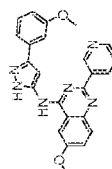
495					DMSO	95	Method G1	402.0 (M+1)	t = 1.746 min	Method B (NH ₄ HCO ₃)
496					DMSO	95	Method J1	373.0 (M+1)	t = 1.523 min	Method B (NH ₄ HCO ₃)
497		2HCl			DMSO	95	Method G1	418.2 (M+1)	1.782 min	Method B (NH ₄ HCO ₃)
498					DMSO	95	Method G1	530.0 (M+1)	1.956 min	Method B (NH ₄ HCO ₃)

499		2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.19 (s, 1H), 9.65 (s, 1H), 9.17 (d, J = 8.2 Hz, 1H), 8.97 (d, J = 4.5 Hz, 1H), 8.93 (s, 1H), 8.81 (d, J = 8.2 Hz, 1H), 8.45 (s, 1H), 8.40 (dd, J = 8.7, 1.7 Hz, 1H), 8.16 (d, J = 8.7 Hz, 1H), 7.99 (dd, J = 16.0, 10.4 Hz, 3H), 7.74 (t, J = 7.4 Hz, 1H), 7.32 (t, J = 7.6 Hz, 1H), 3.38 (s, 3H).	DMSO	95	Method G1	420.1 (M+1)	t = 1.478 min	Method B (NH ₄ HCO ₃)
501		HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.46 (s, 1H), 9.46 (s, 1H), 8.97 (d, J = 7.9 Hz, 1H), 8.93 (d, J = 4.4 Hz, 1H), 8.77 (d, J = 12.1 Hz, 1H), 8.55 (s, 1H), 8.16 ~ 7.91 (m, 4H), 7.83 (t, J = 7.5 Hz, 1H), 7.73 (d, J = 9.3 Hz, 1H), 7.05 (t, J = 6.9 Hz, 1H).	DMSO	95	Method G1	378.1 (M+1)	t = 1.873 min	Method B (NH ₄ HCO ₃)
502			¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.00 (s, 1H), 9.83 (s, 1H), 9.17 (d, J = 8.0 Hz, 1H), 9.00 (d, J = 8.0 Hz, 1H), 8.73 ~ 8.72 (m, 1H), 8.49 (s, 1H), 7.96 (d, J = 8.4 Hz, 1H), 7.92 (s, 1H), 7.76 (t, J = 7.2 Hz, 1H), 7.63 ~ 7.53 (m, 3H), 7.28 (d, J = 7.6 Hz, 1H), 7.22 (t, J = 7.6 Hz, 1H).	DMSO	95	Method G1	358.0 (M+1)	t = 1.644 min	Method B (NH ₄ HCO ₃)

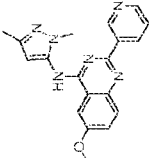
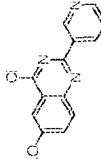
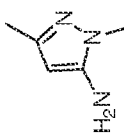
503			¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.34 (s, 1H), 9.50 (s, 1H), 8.69 (d, J = 3.6 Hz, 1H), 8.65 (d, J = 7.6 Hz, 1H), 8.40 (d, J = 8.4 Hz, 1H), 8.15 (s, 1H), 8.10 (s, 1H), 8.00 - 7.97 (m, 1H), 7.91 - 7.84 (m, 2H), 7.69 - 7.66 (m, 1H), 7.55 - 7.52 (m, 1H), 7.16 (t, J = 9.2 Hz, 1H).	DMSO	95	Method G1	378.0 (M+1)	t = 1.780 min	Method B (NH ₄ HCO ₃)
504		HCl	¹ H-NMR (300 MHz, DMSO): δ 10.48 (s, 1H), 9.57 (s, 1H), 9.13 (d, J = 7.6 Hz, 1H), 8.97 - 8.80 (m, 2H), 8.31 (s, 1H), 8.19 (s, 1H), 8.10 (d, J = 6.5 Hz, 1H), 7.97 (d, J = 9.1 Hz, 2H), 7.83 (d, J = 7.7 Hz, 1H), 7.66 (t, J = 7.9 Hz, 2H), 7.47 (d, J = 5.4 Hz, 1H), 4.02 (s, 3H).	DMSO	95	Method G1	396.1 (M+1)		Method C
505			¹ H NMR (400 MHz, DMSO) δ 15.71 (s, 1H), 10.01 (s, 1H), 9.59 - 9.51 (m, 1H), 8.73 - 8.62 (m, 2H), 8.60 - 8.54 (m, 1H), 8.13 - 7.96 (m, 2H), 7.93 - 7.82 (m, 2H), 7.62 - 7.49 (m, 2H), 4.01 (s, 3H).	DMSO	>98	G1			
506			¹ H NMR (400 MHz, DMSO) δ 10.77 (s, 1H), 9.75 (s, 1H), 9.55 - 9.46 (m, 1H), 8.73 - 8.60 (m, 2H), 7.96 (d, J = 2.7 Hz, 1H), 7.83 (d, J = 9.1 Hz, 1H), 7.63 (d, J = 2.1 Hz, 1H), 7.58 - 7.47 (m, 3H), 6.99 (d, J = 8.5 Hz, 1H), 4.64 (s, 2H), 3.97 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)			

507				¹ H NMR (400 MHz, DMSO) δ 12.54 (s, 1H), 9.85 (s, 1H), 9.51 (s, 1H), 8.70 – 8.60 (m, 2H), 8.27 – 8.16 (m, 2H), 8.03 (d, J = 2.5 Hz, 1H), 7.83 (d, J = 9.1 Hz, 1H), 7.72 – 7.59 (m, 2H), 7.56 – 7.47 (m, 2H), 3.99 (s, 3H).	DMSO	>98	G1		
508				¹ H NMR (400 MHz, DMSO) δ 9.87 (s, 1H), 9.50 – 9.47 (m, 1H), 8.66 – 8.61 (m, 2H), 8.21 (s, 1H), 8.16 (d, J = 1.6 Hz, 1H), 8.02 (d, J = 2.7 Hz, 1H), 7.83 (d, J = 9.1 Hz, 1H), 7.74 (dd, J = 8.7, 1.9 Hz, 1H), 7.67 (d, J = 8.6 Hz, 1H), 7.56 – 7.47 (m, 2H), 3.99 (s, 3H).	DMSO	>98	G1		
509				¹ H NMR (400 MHz, DMSO) δ 9.90 (s, 1H), 9.47 (dd, J = 2.1, 0.8 Hz, 1H), 8.67 – 8.60 (m, 2H), 8.15 (d, J = 2.1 Hz, 1H), 7.97 (d, J = 2.7 Hz, 1H), 7.88 – 7.81 (m, 2H), 7.57 – 7.49 (m, 2H), 7.44 (d, J = 8.7 Hz, 1H), 3.98 (s, 3H), 3.47 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)		
510				¹ H NMR (400 MHz, DMSO) δ 10.76 (s, 1H), 10.62 (s, 1H), 9.71 (s, 1H), 9.53 – 9.48 (m, 1H), 8.68 – 8.62 (m, 2H), 7.98 (d, J = 2.6 Hz, 1H), 7.81 (d, J = 9.1 Hz, 1H), 7.63 (d, J = 1.8 Hz, 1H), 7.55 – 7.49 (m, 2H), 7.37 (dd, J = 8.4, 2.0 Hz, 1H), 7.01 (d, J = 8.3 Hz, 1H), 3.97 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)		

511				¹ H NMR (400 MHz, DMSO) δ 9.96 (s, 1H), 9.49 (dd, J = 2.1, 0.8 Hz, 1H), 8.79 (s, 1H), 8.67 – 8.61 (m, 2H), 8.39 – 8.36 (m, 1H), 8.02 (d, J = 2.7 Hz, 1H), 7.90 – 7.84 (m, 3H), 7.56 (dd, J = 9.1, 2.7 Hz, 1H), 7.54 – 7.50 (m, 1H), 3.99 (s, 3H).	DMSO	>98	J2		
512				¹ H NMR (400 MHz, DMSO) δ 9.87 (s, 1H), 9.48 (dd, J = 2.1, 0.8 Hz, 1H), 8.67 – 8.61 (m, 2H), 8.10 – 8.08 (m, 1H), 8.00 (d, J = 2.7 Hz, 1H), 7.84 (d, J = 9.1 Hz, 1H), 7.65 – 7.63 (m, 2H), 7.57 – 7.49 (m, 2H), 4.22 (s, 3H), 3.99 (s, 3H).	DMSO	>98	G1		
513				¹ H NMR (400 MHz, DMSO) δ 9.92 (s, 1H), 9.49 (dd, J = 2.1, 0.8 Hz, 1H), 8.67 – 8.60 (m, 2H), 8.22 (d, J = 1.8 Hz, 1H), 8.00 (d, J = 2.7 Hz, 1H), 7.85 (d, J = 9.1 Hz, 1H), 7.70 – 7.64 (m, 1H), 7.60 (d, J = 8.5 Hz, 1H), 7.57 – 7.49 (m, 2H), 4.21 (s, 3H), 3.98 (s, 3H).	DMSO	>98	J2		
514			2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.97 (s, 1H), 9.43 (d, J = 1.6 Hz, 1H), 8.98 – 8.91 (m, 1H), 8.89 (dd, J = 5.2, 1.3 Hz, 1H), 8.43 (s, 1H), 8.27 (s, 1H), 8.13 – 8.04 (m, 2H), 7.92 – 7.85 (m, 1H), 7.73 (d, J = 9.2 Hz, 1H), 7.70 – 7.63 (m, 2H), 4.21 (s, 3H), 4.02 (s, 3H).	DMSO	>98	G1		

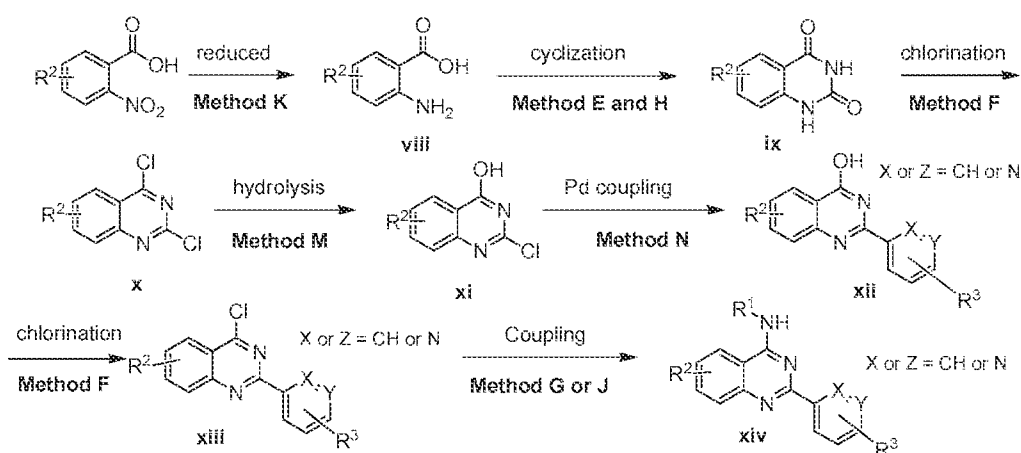
515				HCl	¹ H NMR (400 MHz, DMSO) δ 11.90 (s, 1H), 10.45 (s, 1H), 9.46 (d, J = 1.7 Hz, 1H), 8.97 – 8.89 (m, 1H), 8.85 (dd, J = 5.2, 1.5 Hz, 1H), 8.16 – 8.10 (m, 2H), 8.03 – 7.97 (m, 2H), 7.94 (d, J = 9.1 Hz, 1H), 7.90 – 7.82 (m, 1H), 7.63 (dd, J = 9.1, 2.6 Hz, 1H), 7.44 (d, J = 8.8 Hz, 1H), 6.58 (d, J = 9.4 Hz, 1H), 4.00 (s, 3H).	DMSO	>98	G1 (0.1 N HCl added)			
516					¹ H NMR (400 MHz, DMSO) δ 9.90 (s, 1H), 9.53 – 9.50 (m, 1H), 8.68 – 8.63 (m, 2H), 8.27 (d, J = 2.0 Hz, 1H), 8.01 (d, J = 2.7 Hz, 1H), 7.96 (d, J = 8.6 Hz, 1H), 7.85 (d, J = 9.1 Hz, 1H), 7.81 (dd, J = 8.7, 2.1 Hz, 1H), 7.58 – 7.50 (m, 2H), 4.20 (s, 3H), 3.99 (s, 3H).	DMSO	>98	G1			
517				HCl	¹ H NMR (300 MHz, DMSO) δ 11.35 (s, 1H), 9.58 (d, J = 1.6 Hz, 1H), 9.24 (d, J = 8.2 Hz, 1H), 9.05 – 8.97 (m, 1H), 9.05 – 8.95 (m, 1H), 8.21 (d, J = 2.4 Hz, 1H), 8.11 – 8.02 (m, 3H), 7.90 (d, J = 7.8 Hz, 1H), 7.58 (dd, J = 8.8, 2.0 Hz, 2H), 7.52 – 7.46 (m, 1H), 7.28 (s, 1H), 3.98 (d, J = 9.2 Hz, 3H).	DMSO	>98	G1			
518				2 HCl	¹ H NMR (300 MHz, DMSO) δ 11.32 (s, 1H), 9.60 (s, 1H), 9.23 (d, J = 8.0 Hz, 1H), 8.98 (d, J = 5.4 Hz, 1H), 8.26 (s, 1H), 8.06 (dd, J = 13.2, 7.3 Hz, 2H), 7.63 (d, J = 9.1 Hz, 1H), 7.50 – 7.36 (m, 4H), 7.32 (s, 1H), 7.06 – 6.90 (m, 1H), 3.99 (s, 3H), 3.86 (s, 3H).	DMSO	>98	G1			

519			2HCl	¹ H NMR (300 MHz, DMSO) δ 13.17 (s, 1H), 11.46 (s, 1H), 9.19 (d, J = 1.7 Hz, 1H), 8.95 – 8.78 (m, 2H), 8.31 (d, J = 2.6 Hz, 1H), 8.18 (d, J = 9.2 Hz, 1H), 7.86 (dd, J = 7.9, 5.4 Hz, 1H), 7.77 – 7.54 (m, 3H), 7.44 (dd, J = 11.4, 3.9 Hz, 1H), 7.20 – 7.02 (m, 1H), 4.01 (s, 3H).	DMSO	>98	G1		
520			HCl	¹ H NMR (300 MHz, DMSO) δ 11.10 (s, 1H), 9.59 (d, J = 1.8 Hz, 1H), 9.15 (d, J = 8.0 Hz, 1H), 8.93 (d, J = 3.9 Hz, 1H), 8.22 (d, J = 2.6 Hz, 1H), 7.99 (t, J = 7.4 Hz, 2H), 7.72 (dd, J = 8.4, 4.7 Hz, 2H), 7.63 – 7.50 (m, 2H), 7.35 (s, 1H), 7.27 – 7.18 (m, 1H), 3.99 (s, 3H).	DMSO	>98	G1		
521			HCl	¹ H NMR (300 MHz, DMSO) δ 11.51 (s, 1H), 9.63 (d, J = 1.9 Hz, 1H), 9.18 (d, J = 8.1 Hz, 1H), 8.98 (dd, J = 5.2, 1.2 Hz, 1H), 8.31 (d, J = 2.5 Hz, 1H), 8.17 (d, J = 9.1 Hz, 1H), 8.01 (dd, J = 8.0, 5.3 Hz, 1H), 7.84 (dd, J = 7.7, 1.5 Hz, 1H), 7.66 (dd, J = 9.1, 2.5 Hz, 1H), 7.48 – 7.35 (m, 2H), 7.21 (d, J = 8.2 Hz, 1H), 7.10 (t, J = 7.5 Hz, 1H), 3.99 (d, J = 6.3 Hz, 6H).	DMSO	>98	G1		
1758			3HCl	¹ H NMR (DMSO-d6) ppm 13.00-12.40 (br, 1H), 9.79 (d, J = 1.68 Hz, 1H), 9.30 (d, J = 8.20 Hz, 1H), 8.95 (dd, J = 5.32, 1.44 Hz, 1H), 8.71 (d, J = 4.24 Hz, 1H), 8.37 (brd, J = 2.56 Hz, 1H), 8.25-8.21 (m, 2H), 8.11-8.02 (m, 2H), 7.98 (d, J = 9.12 Hz, 1H), 7.66 (dd, J = 9.12, 2.68 Hz, 1H), 7.53-7.50 (m, 1H), 4.03 (s, 3H). The 1H of 3HCl was not observed.	DMSO	>98	G13		

			J2	>98	DMSO	¹ H NMR (DMSO-d ₆) ppm 10.35 (br, 1H), 9.37 (d, J = 1.6 Hz, 1H), 8.96 (d, J = 7.28 Hz, 1H), 8.90 (d, J = 4.2 Hz, 1H), 8.04 (s, 1H), 7.98 (m, 1H), 7.95 (d, J = 9.16 Hz, 1H), 7.64 (dd, J = 9.16, 2.56 Hz, 1H), 6.18 (s, 1H), 3.98 (s, 3H), 3.66 (s, 3H), 2.24 (s, 3H). The 1H of 3HCl was not observed.	3HCl				1759
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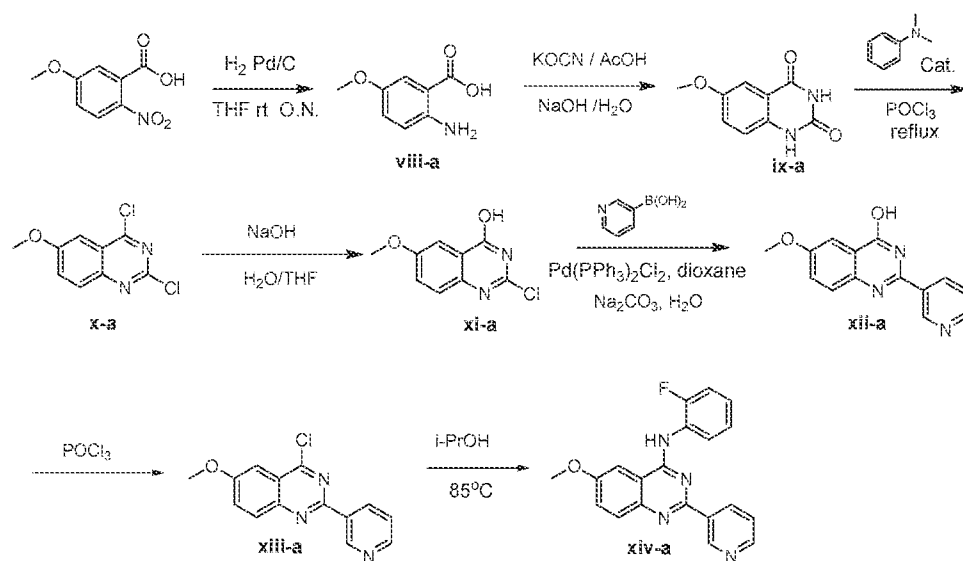
Scheme 6: General route for the synthesis of compounds with general formula xiv

[0301]



Scheme 7: Representative synthesis of pyridine of formula xiv-a (see Scheme 6)

[0302]



[0303] Method K: 2-Amino-5-methoxybenzoic acid (viii-a) 5-Methoxy-2-nitrobenzoic acid (30.0 g, 152.2 mmol) was hydrogenated over Pd/C (10%, 300 mg) in THF (250 mL) at room temperature under H₂ balloon. The mixture was stirred for 18 h. After the reaction was completed, the catalyst was removed by filtration over Celite and the filtrate was concentrated to afford 25.0 g of 2-amino-5-methoxybenzoic acid as a brown solid (98%). LCMS m/z = 168.1 (M+1), 150.1 (M-17) (Method B) (retention time = 0.53 min) ¹H NMR (400 MHz, DMSO-d₆): δ 7.31 (d, J = 2.8 Hz, 1H), 6.61 (dd, J = 8.8, 3.2 Hz, 1H), 6.44 (d, J = 8.8 Hz, 1H), 3.60 (s, 3H).

[0304] Method H: 6-Methoxyquinazolin-2,4(1H, 3H)-dione (ix-a) 2-Amino-5-methoxybenzoic acid (13.0 g, 77.8 mmol, 1.0 eq.) was suspended in water (200 mL) and glacial acetic acid (5.2 mL) at 35 °C. A freshly prepared solution of potassium cyanate (8.21 g, 101.4 mmol, 1.3 eq.) in water (86 mL) was added dropwise to the stirred mixture. After 4 h, NaOH (104.0 g, 2600 mL, 33.4 eq.) was added in portions, keeping the reaction temperature below 40 °C. A clear solution was obtained momentarily before a precipitate formed. After cooling, the precipitate was filtered off and dissolved in hot water which was acidified to pH 5. The precipitate was collected and washed with water, dried by lyophilization to afford 9.63 g of 6-methoxyquinazolin-2,4(1H, 3H)-dione as a white solid (65%). LCMS m/z = 193.1

(M+1) (Method B) (retention time = 1.22 min). ¹H-NMR (400 MHz, DMSO-d₆) : δ 11.26 (s, 1H), 11.01 (s, 1H), 7.31 - 7.25 (m, 2H), 7.10 (d, 1H, *J* = 8.8 Hz), 3.77 (s, 3H).

[0305] Method F1: 2,4-Dichloro-6-methoxyquinazoline (x-a) To a mixture of 6-methoxyquinazoline-2,4-(1*H*, 3*H*)-dione (9.63 g, 50.2 mmol) in POCl₃ (150 mL) was added *N,N*-dimethylaniline (0.5 mL). The resulting mixture was stirred at 120°C for 2 h. After the reaction was completed, POCl₃ was removed *in vacuo*, and the residue was added to ice-water slowly. The pH was adjusted to ~ 7 by slowly adding NaHCO₃ (sat.) at 0°C, then a precipitate formed. The solid was collected and dried *in vacuo* to give 11.2 g of 2,4-dichloro-6-methoxyquinazoline in a 98% yield as a brown solid. LCMS *m/z* = 229.1, 231.0 (M+1) (Method B) (retention time = 1.87 min). ¹H-NMR (400 MHz, DMSO-d₆) : δ 7.89 (d, *J* = 9.2 Hz, 1H), 7.71 (dd, *J* = 8.8, 2.4 Hz, 1H), 7.38 (d, *J* = 1.6 Hz, 1H), 3.91 (s, 3H).

[0306] Method M: 2-Chloro-6-methoxyquinazolin-4-ol (xi-a) A mixture of 2,4-dichloro-6-methoxyquinazoline (4.20 g, 18.5 mmol, 1.0 eq.) in THF (60 mL) and H₂O (60 mL) was treated with NaOH (4.00 g, 100 mmol, 5.4 eq.). The resulting mixture was stirred at 40°C for 2 h. The reaction color turned to dark green and then a precipitate formed. After the reaction was completed, the mixture was cooled to room temperature. The precipitate was filtered off and the filtrate was concentrated down to 60 mL. The pH was then adjusted to 6 by adding 2N HCl in water. The precipitate which formed was collected and dried *in vacuo* to give 4.00 g of 2-chloro-6-methoxyquinazolin-4-ol as a grey solid (98%). LCMS *m/z* = 211.1, 213.0 (M+1) (Method B) (retention time = 1.11 min).

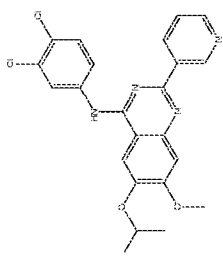
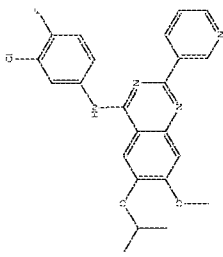
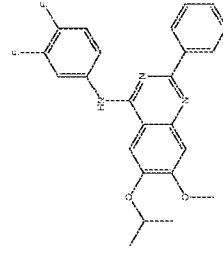
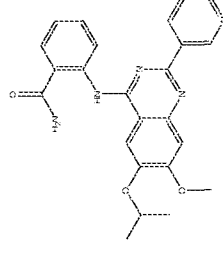
[0307] Method N: 6-Methoxy-2-(pyridin-3-yl)quinazolin-4-ol (xii-a) To a mixture of 2-chloro-6-methoxyquinazolin-4-ol (1.20 g, 5.7 mmol, 1.0 eq.), pyridin-3-ylboronic acid (1.27 g, 8.6 mmol, 1.5 eq.), K₂CO₃ (2.37 g, 17.1 mmol, 3.0 eq.) in dioxane (100 mL) and H₂O (10 mL) was added Pd(PPh₃)₂Cl₂ (230 mg, 0.29 mmol, 0.05 eq.) under N₂ atmosphere. The resulting mixture was stirred at 105 °C under N₂ atmosphere overnight. After reaction was completed, the mixture was cooled to room temperature, and the resultant precipitate was removed by filtration. The filtrate was concentrated *in vacuo* and the residue was partitioned between H₂O (30 mL) and ethyl acetate (100 mL x 3). The combined organic layers were washed with brine, dried over MgSO₄. After filtration and evaporation, the crude product was obtained, which was combined with the filter cake and dried *in vacuo* to give 1.35 g of 6-methoxy-2-(pyridin-3-yl)quinazolin-4-ol as a gray solid. LCMS *m/z* = 254.1 (M+1) (Method B) (retention time = 1.39 min). The crude product was used for the next step without further purification.

[0308] Method F1: 4-Chloro-6-methoxy-2-(pyridin-3-yl)quinazoline (xiii-a) To a mixture of 6-methoxy-2-(pyridin-3-yl)quinazolin-4-ol (600 mg, 2.37 mmol) in POCl₃ (5 mL) was added *N,N* dimethylaniline (1 drop). The resulting mixture was stirred at 120°C for 30 min. After the reaction was completed, POCl₃ was removed *in vacuo*, and the residue was added to ice-water slowly. The pH was adjusted to ~ 7 by slowly adding NaHCO₃ (sat.) at 0°C and then a precipitate formed. The solid was collected and was purified by chromatography on silica gel eluted with petroleum ether/ ethyl acetate (v/v = 4:1 to 1:1) to give 260 mg of 4-chloro-6-methoxy-2-(pyridin-3-yl)quinazoline as a pale yellow solid (40.4%). LCMS *m/z* = 272.1, 274.0 (M + 1) (Method B) (retention time = 1.90 min).

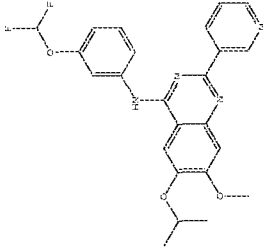
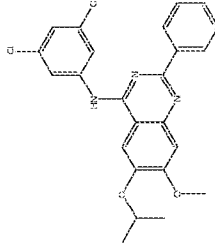
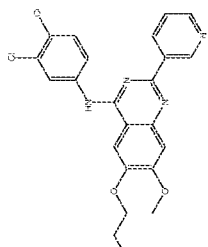
[0309] Method G1: N-(2-Fluorophenyl)-6-methoxy-2-(pyridin-3-yl)quinazolin-4-amine (xiv-a) A mixture of 4-chloro-6-methoxy-2-(pyridin-3-yl)quinazoline (80 mg, 0.293 mmol, 1.0 eq.) and 2-fluoroaniline (65 mg, 0.58 mmol, 2 eq.) in *i*-PrOH (5 mL) was stirred at 85 °C for 18 h. After the reaction was completed, the mixture was filtered, and the filter cake was washed with H₂O (10 mL) and diethyl ether (10 mL). After drying, the crude product was purified by PREP-HPLC (Condition C: Gradient: B = 5%-50%, Target Peak: at 7.2 min) to give 16.5 mg of N-(2-fluorophenyl)-6-methoxy-2-(pyridin-3-yl)quinazolin-4-amine as a yellow solid, yield 16.4%. LCMS *m/z* = 347.0 (M + 1) (Method A) (retention time = 1.31 min). ¹H-NMR (400 MHz, DMSO-d₆) : δ 10.08 (s, 1H), 9.33 (s, 1H), 8.72 (d, *J* = 4.4 Hz, 1H), 8.66 (d, *J* = 8.0 Hz, 1H), 8.80 (d, *J* = 1.2 Hz, 1H), 7.88 (d, *J* = 9.2 Hz, 1H), 7.70 - 7.65 (m, 2H), 7.58 (dd, *J* = 8.8, 1.2 Hz, 1H), 7.44 - 7.35 (m, 3H), 3.97 (s, 3H).

[0310] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 6** substituting with appropriate nitro benzoic acid, boronic acid and aniline

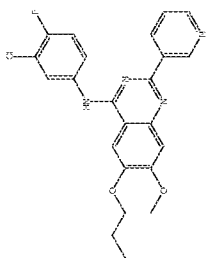
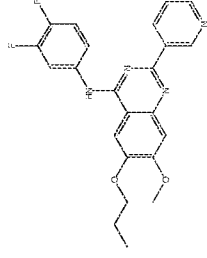
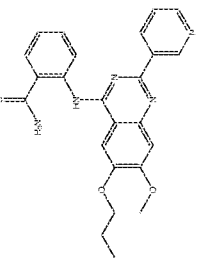
Table 2:

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
522			455.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.76 (s, 1H), 9.52 - 9.51 (m, 1H), 8.63 - 8.66 (m, 2H), 8.32 (d, J = 2.4 Hz, 1H), 7.94-7.91 (m, 2H), 7.72 (d, J = 8.8 Hz, 1H), 7.54 (q, J = 3.2 Hz, 1H), 7.32 (s, 1H), 4.87 - 4.90 (m, 1H), 3.96 (s, 3H), 1.38 (d, J = 6.0 Hz, 6H).	DMSO	454.9, 457.0 (M+1), 228.8 (M/2 +1)	Method A (TFA)	95	Method N, G1
523			438.88	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.68 (s, 1H), 9.49 (d, J = 2.0 Hz, 1H), 8.60-8.66 (m, 2H), 8.18 (dd, J = 6.4, 2.4 Hz, 1H), 7.84 - 7.88 (m, 2H), 7.50 - 7.54 (m, 2H), 7.31 (s, 1H), 4.85 - 4.88 (m, 1H), 3.96 (s, 3H), 1.38 (d, J = 6.4 Hz, 6H).	DMSO	439.1, 441.1 (M+1), 220.1, 220.8 (M/2 +1)	Method A (TFA)	95	Method N, G1
524			422.43	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.76 (s, 1H), 9.48 (s, 1H), 8.60 - 8.65 (m, 2H), 8.04 - 8.10 (m, 1H), 7.92 (s, 1H), 7.49-7.66 (m, 3H), 7.30 (s, 1H), 4.86 - 4.92 (m, 1H), 3.96 (s, 3H), 1.37 (d, J = 6.0 Hz, 6H).	DMSO	423.2 (M+1), 212.2 (M/2 +1)	Method A (TFA)	95	Method N, G1
525		HCl	429.47	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.96 (s, 1H), 9.60 (d, J = 8.0 Hz, 1H), 9.14 (d, J = 8.0 Hz, 1H), 8.68 - 8.74 (m, 2H), 8.46 (s, 1H), 7.95 - 7.96 (m, 2H), 7.74-7.70 (m, 1H), 7.59 - 7.54 (m, 2H), 7.36 (s, 1H), 7.19 - 7.15 (m, 1H), 4.76 - 4.82 (m, 1H), 3.99 (s, 3H), 1.44 (d, J = 6.0 Hz, 6H).	DMSO	430.2 (M+1), 215.7 (M/2 +1)	Method A (TFA)	95	Method N, G1

(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
526			452.45	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.70 (s, 1H), 9.52 - 9.53 (m, 1H), 8.64 - 8.67 (m, 2H), 7.89 - 7.94 (m, 2H), 7.73 - 7.74 (m, 1H), 7.48 - 7.53 (m, 3H), 7.32 (t, J = 83.6 Hz, 1H), 6.96 - 6.99 (m, 1H), 4.89-4.92 (m, 1H), 3.98 (s, 3H), 1.38 (d, J = 6.0 Hz, 6H).	DMSO	453.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
527			455.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.53 (s, 1H), 9.52 (d, J = 1.6 Hz, 1H), 8.64-8.68 (m, 2H), 8.12 - 8.13 (m, 2H), 7.90 (s, 1H), 7.54 - 7.58 (m, 1H), 7.34 - 7.36 (m, 2H), 4.86 - 4.92 (m, 1H), 3.98 (s, 3H), 1.40 (d, J = 6.0 Hz, 6H).	DMSO	455.0, 457.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
528			455.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.76 (s, 1H), 9.52 (d, J = 1.2 Hz, 1H), 8.64-8.67 (m, 2H), 8.32 (d, J = 2.8 Hz, 1H), 7.94 (dd, J = 8.8, 2.4 Hz, 1H), 7.86 (s, 1H), 7.72 (d, J = 8.8 Hz, 1H), 7.53 - 7.56 (m, 1H), 7.32 (s, 1H), 4.14 (t, J = 6.4 Hz, 2H), 3.99 (s, 3H), 1.85 - 1.92 (m, 2H), 1.06 (t, J = 7.2 Hz, 3H).	DMSO	455.1, 457.1 (M+1), 229.1 (M/2 +1)	Method A (TFA)	95	Method N, G1

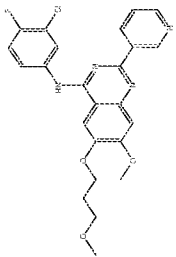
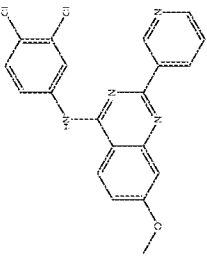
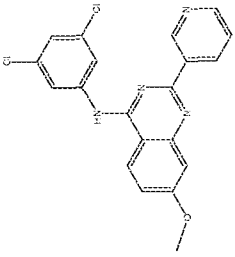
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
529			438.88	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.72 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 8.62-8.66 (m, 2H), 8.20 (dd, J = 6.4, 2.4 Hz, 1H), 7.85-7.89 (m, 2H), 7.50-7.55 (m, 2H), 7.31 (s, 1H), 4.13 (t, J = 6.4 Hz, 2H), 3.98 (s, 3H), 1.82-1.90 (m, 2H), 1.06 (t, J = 7.6 Hz, 3H).	DMSO	439.1, 441.1 (M+1) 220.1, 220.8 (M/2+1)	Method A (TFA)	95	Method N, G1
530			422.43	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.74-9.76 (m, 1H), 9.52 (d, J = 1.6 Hz, 1H), 8.64-8.69 (m, 2H), 8.08-8.14 (m, 1H), 7.90 (s, 1H), 7.68-7.70 (m, 1H), 7.52-7.60 (m, 2H), 7.34 (s, 1H), 4.16 (t, J = 6.4 Hz, 2H), 4.01 (s, 3H), 1.85-1.94 (m, 2H), 1.09 (t, J = 7.2 Hz, 3H).	DMSO	423.0 (M+1) 445.0 (M+23) 212.1 (M/2+1)	Method A (TFA)	95	Method N, G1
531		HCl	429.47	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.86 (s, 1H), 9.59 (d, J = 1.6 Hz, 1H), 9.05 (d, J = 8.4 Hz, 1H), 8.85 (d, J = 8.0 Hz, 1H), 8.76-8.78 (m, 1H), 8.44 (s, 1H), 7.96 (dd, J = 8.0, 1.6 Hz, 2H), 7.70-7.73 (m, 2H), 7.53 (s, 1H), 7.37 (s, 1H), 6.96-7.22 (m, 1H), 4.16 (t, J = 6.4 Hz, 2H), 4.01 (s, 3H), 1.86-1.90 (m, 2H), 1.07 (t, J = 7.2 Hz, 3H).	DMSO	430.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1

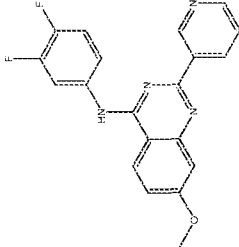
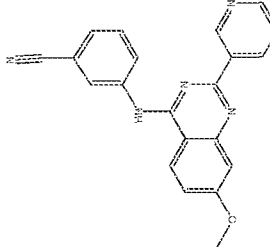
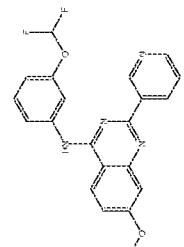
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
532			452.45	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.67 (s, 1H), 9.50 (d, J = 4.0 Hz, 1H), 8.62-8.65 (m, 2H), 7.89 - 7.92 (m, 3H), 7.44-7.54 (m, 1H), 7.29 - 7.31 (m, 3H), 7.26 (t, J = 74.0 Hz, 1H), 4.14 (t, J = 6.8 Hz, 2H), 3.98 (s, 3H), 1.84 - 1.90 (m, 2H), 1.06 (t, J = 7.2 Hz, 3H).	DMSO	453.1 227.2 (M/2+1)	Method A (TFA)	95	Method N, G1
533		HCl	454.88	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.73 (s, 1H), 9.50 (d, J = 2.4 Hz, 1H), 8.62-8.67 (m, 2H), 8.20 (dd, J = 7.2 Hz, 2.8 Hz, 1H), 7.85 - 7.89 (m, 2H), 7.51 - 7.57 (m, 2H), 7.33 (s, 1H), 4.31 (t, J = 4.0 Hz, 2H), 3.99 (s, 3H) 3.80 (t, J = 4.8 Hz, 2H), 3.37 (s, 3H).	DMSO	455.1, 457.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
534			445.47	¹ H-NMR (400 MHz, CD ₃ OD): δ 9.34 (s, 1H), 8.92 (d, J = 6.0 Hz, 1H), 8.73 (m, 2H), 7.84 (s, 1H), 7.76 (d, J = 8.0 Hz, 1H), 7.54 (t, J = 8.0 Hz, 1H), 7.32 (s, 1H), 7.15 (s, 2H), 4.21 (s, 2H), 3.92 (s, 3H), 3.78 (t, J = 4.4 Hz, 2H), 3.40 (s, 3H).	CD ₃ OD	446.2, 447.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
535		HCl	468.45	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.72 (s, 1H), 9.53 (s, 1H), 8.66 (d, J = 5.6 Hz, 2H), 7.91 (d, J = 6.4 Hz, 2H), 7.75 (d, J = 7.2 Hz, 1H), 7.51 (dd, J = 7.6 Hz, 2H), 7.34 (s, 1H), 7.30 (t, J = 74.0 Hz, 1H), 6.97 (dd, J = 8.0, 2.0 Hz, 1H), 4.32 (t, J = 4.4 Hz, 2H), 3.99 (s, 3H), 3.80 (t, J = 4.4 Hz, 2H), 3.37 (s, 3H).	DMSO	469.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1

(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
536			468.91	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.42 (s, 1H), 9.47 (s, 1H), 8.96 (d, J = 6.4 Hz, 1H), 8.90 (d, J = 5.2 Hz, 1H), 8.11 (dd, J = 6.8 Hz, 2.4 Hz, 1H), 8.05 (s, 2H), 7.85 - 7.94 (m, 2H), 7.55 (t, J = 9.2 Hz, 1H), 7.47 (s, 1H) 4.26 (t, J = 6.4 Hz, 2H), 4.01 (s, 3H), 3.55 (t, J = 6.0 Hz, 2H), 3.29 (s, 3H), 2.06-2.13 (m, 2H).	DMSO	469.0, 471.0 (M+1) 234.9, 235.6 (M/2+1)	Method A (TFA)	95	Method N, G1
537			397.26	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.08 (s, 1H), 9.54 (s, 1H), 8.76 (s, 2H), 8.49 (d, J = 7.0 Hz, 1H), 8.36 (s, 1H), 7.95 (d, J = 7.5 Hz, 1H), 7.66 - 7.23 (m, 2H), 7.33 (s, 2H), 3.98 (s, 3H).	DMSO	397.1, 399.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
538			397.26	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.00 (s, 1H), 8.54 (d, J = 1.3 Hz, 1H), 8.71-8.56 (m, 2H), 8.47 (d, J = 9.2 Hz, 1H), 8.17 (s, 1H), 8.16 (s, 1H), 7.59 - 7.56 (m, 1H), 7.35 - 7.30 (m, 3H), 3.98 (s, 3H).	DMSO	397.0, 399.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1

(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
539			364.35	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.94 (s, 1H), 9.53 (d, J = 1.6 Hz, 1H), 8.70-8.65 (m, 2H), 8.47 (d, J = 8.4 Hz, 1H), 8.12 (ddd, J = 13.2, 7.6, 2.8 Hz, 1H), 7.72 - 7.70 (m, 1H), 7.58 - 7.49 (m, 2H), 7.32 - 7.27 (m, 2H), 3.97 (s, 3H).	DMSO	365.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
540			353.38	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.06 (s, 1H), 9.53 (d, J = 1.6 Hz, 1H), 8.71-8.66 (m, 2H), 8.50 - 8.43 (m, 2H), 8.26-8.24 (m, 1H), 7.70 - 7.54 (m, 3H), 7.34-7.30 (m, 2H), 3.98 (s, 3H).	DMSO	354.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
541			394.37	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.93 (s, 1H), 9.54 (s, 1H), 8.68 (d, J = 5.2 Hz, 2H), 8.51 (d, J = 9.2 Hz, 1H), 7.96 (s, 1H), 7.79 (d, J = 7.6 Hz, 1H), 7.55 - 7.10 (m, 5H), 6.96 (dd, J = 8.0, 2.0 Hz, 1H), 3.96 (s, 3H).	DMSO	395.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1

(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
542			371.39	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.97 (s, 1H), 9.60 (s, 1H), 9.10 (d, J = 8.4 Hz, 1H), 8.75 - 8.71 (m, 2H), 8.48 (s, 1H), 8.07 (d, J = 9.6 Hz, 1H), 7.95 - 7.90 (m, 2H), 7.74 - 7.70 (m, 1H), 7.58 (dd, J = 8.0, 4.8 Hz, 1H), 7.35 - 7.33 (m, 2H), 7.19 (t, J = 7.2 Hz, 1H), 3.97 (s, 3H).	DMSO	372.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
543			431.7	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.03 (s, 1H), 9.49 - 9.48 (m, 1H), 8.69 - 8.67 (m, 1H), 8.63 - 8.59 (m, 1H), 8.31 (d, J = 2.4 Hz, 1H), 8.04 (s, 1H), 8.00 (s, 1H), 7.93 - 7.90 (m, 1H), 7.74 (d, J = 8.8 Hz, 1H), 7.56 - 7.53 (m, 1H), 4.07 (s, 3H).	DMSO	430.9, 432.9, 434.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
544			431.7	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.94 (s, 1H), 9.46 (d, J = 3.2 Hz, 1H), 8.68 (dd, J = 4.8, 2.0 Hz, 1H), 8.60 - 8.57 (m, 1H), 8.09 (s, 1H), 8.08 (s, 1H), 7.97 (s, 2H), 7.54 (dd, J = 8.8, 4.8 Hz, 1H), 7.38 (t, J = 2.0 Hz, 1H), 4.05 (s, 3H).	DMSO	430.9, 432.9, 434.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
545			415.25	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.02 (s, 1H), 9.47 (s, 1H), 8.70 - 8.62 (m, 2H), 8.19 - 8.17 (m, 1H), 8.04 (s, 1H), 8.00 (s, 1H), 7.84 - 7.88 (m, 1H), 7.59 - 7.53 (m, 2H), 4.07 (s, 3H).	DMSO	415.1, 417.0, 419.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1

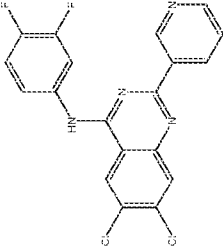
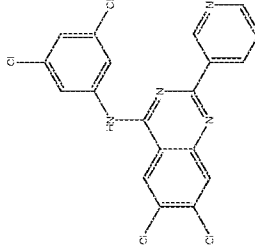
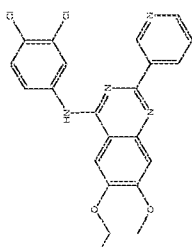
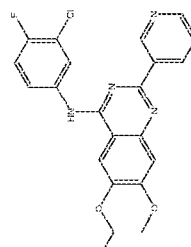
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
546			398.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.01 (s, 1H), 9.47 (d, J = 1.6 Hz, 1H), 8.69-8.67 (m, 1H), 8.62 - 8.59 (m, 1H), 8.08-8.04 (m, 2H), 8.00 (s, 1H), 7.65 - 7.53 (m, 3H), 4.07 (s, 3H).	DMSO	399.1, 401.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
547			428.82	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.32 (s, 1H), 9.45 (d, J = 1.2 Hz, 1H), 8.89 (d, J = 8.0 Hz, 1H), 8.83 (d, J = 4.0 Hz, 1H), 8.21 (s, 1H), 8.00 (s, 1H), 7.85 - 7.78 (m, 3H), 7.55 - 7.51 (m, 1H), 7.31 (t, J = 72.8 Hz, 1H), 7.05 - 7.03 (m, 1H), 4.07 (s, 3H).	DMSO	429.1, 431.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
548			387.82	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.62 (s, 1H), 9.38 (s, 1H), 8.97 - 8.90 (m, 2H), 8.32 - 8.26 (m, 3H), 7.97 - 7.94 (m, 2H), 7.69 - 7.65 (m, 2H), 4.08 (s, 3H).	DMSO	387.9, 390.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
549			405.84	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.98 (s, 1H), 9.36 (s, 1H), 8.94 - 8.77 (m, 3H), 8.48 (s, 1H), 7.98 - 7.83 (m, 4H), 7.59-7.14 (m, 2H), 7.17 (s, 1H), 3.96 (s, 3H).	DMSO	405.9, 408.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1

(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
550			436.12	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.24 (s, 1H), 9.39 (s, 1H), 8.87 (s, 1H), 8.81 (d, J = 4.4 Hz, 1H), 8.73 (d, J = 8.0 Hz, 1H), 8.24 (s, 1H), 8.01 (s, 1H), 7.88 (d, J = 8.8 Hz, 1H), 7.76 - 7.73 (m, 1H), 7.66 (d, J = 8.8 Hz, 1H).	DMSO	434.9, 436.9, 438.8 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
551			410.26	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.15 (s, 1H), 9.46 (s, 1H), 8.87 (d, J = 8.4 Hz, 1H), 8.74 (d, J = 3.6 Hz, 1H), 8.68 (d, J = 7.2 Hz, 1H), 8.49 (s, 1H), 8.23 (s, 1H), 8.02 (s, 2H), 7.94 (d, J = 8.0 Hz, 1H), 7.69 - 7.61 (m, 2H), 7.23 - 7.20 (m, 1H).	DMSO	410.0, 412.0, 413.9 (M+1), 432.0 (M+22)	Method A (TFA)	95	Method N, G1
552			433.24	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.30 (s, 1H), 9.53 (s, 1H), 9.02 (s, 1H), 8.83-8.80 (m, 2H), 8.20 (s, 1H), 7.91 (s, 1H), 7.81 - 7.79 (m, 1H), 7.74 - 7.71 (m, 1H), 7.56 - 7.52 (m, 1H), 7.30 (t, J = 73.6 Hz, 1H), 7.05 (d, J = 8.8 Hz, 1H).	DMSO	432.9, 434.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
553			419.67	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.43 (s, 1H), 9.40 (s, 1H), 8.97 (s, 1H), 8.89-8.85 (m, 2H), 8.15 - 8.14 (m, 1H), 8.05 (s, 1H), 7.89 - 7.87 (m, 2H), 7.51 - 7.47 (m, 1H).	DMSO	419.0, 421.0, 423.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1

(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
554			403.21	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.39 (s, 1H), 9.40 (s, 1H), 8.96 (s, 1H), 8.84-8.89 (m, 2H), 8.04 - 8.00 (m, 2H), 7.88 (s, 1H), 7.69 - 7.67 (m, 1H), 7.54 - 7.47 (m, 1H).	DMSO	402.9, 404.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
555			436.12	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.26 (s, 1H), 8.28 (s, 1H), 8.88 (d, J = 4.8 Hz, 1H), 8.85 (s, 1H), 8.76 (d, J = 8.0 Hz, 1H), 7.97 (s, 1H), 7.96 (s, 1H), 7.90-7.86 (m, 2H), 7.29 (s, 1H).	DMSO	434.9, 436.9, 438.9 (M+1)	Method A (TFA)	95	Method N, G1
556			441.31	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.74 (s, 1H), 9.50 (s, 1H), 8.66 - 8.64 (m, 2H), 8.31 (s, 1H), 7.90 (d, J = 8.8 Hz, 1H), 7.84 (s, 1H), 7.68 (d, J = 8.8 Hz, 1H), 7.52 (dd, J = 7.4, 5.0 Hz, 1H), 7.29 (s, 1H), 4.22 (q, J = 6.8 Hz, 2H), 3.97 (s, 3H), 1.45 (t, J = 6.8 Hz, 3H).	DMSO	441.0, 443.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
557			424.86	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.93 (s, 1H), 9.48 (s, 1H), 8.74 - 8.73 (m, 2H), 8.18 - 8.16 (m, 1H), 7.90 - 7.85 (m, 2H), 7.67 - 7.65 (m, 1H), 7.53 (t, J = 9 Hz, 1H), 7.33 (s, 1H), 4.24 (q, J = 7.2 Hz, 2H), 3.98 (s, 3H), 1.46 (t, J = 7.2 Hz, 3H).	DMSO	425.0, 427.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1

(continued)

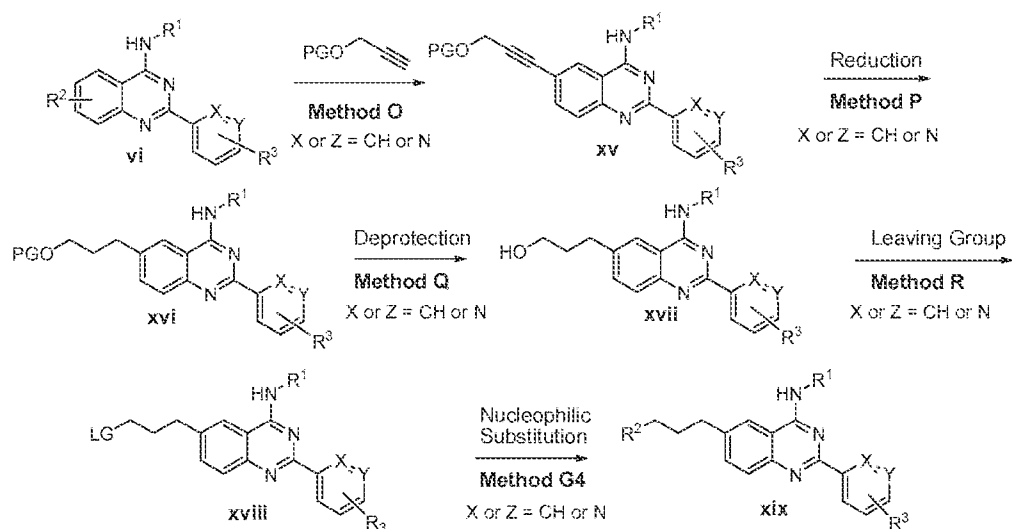
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
558			408.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.68 (s, 1H), 9.49 (s, 1H), 8.66–8.61 (m, 2H), 8.10–8.04 (m, 1H), 7.84 (s, 1H), 7.66–7.64 (m, 1H), 7.56–7.49 (m, 2H), 7.30 (s, 1H), 4.23 (q, J = 6.8 Hz, 2H), 3.98 (s, 3H), 1.46 (t, J = 6.8 Hz, 3H).	DMSO	409.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
559			397.43	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.27 (s, 1H), 9.43 (s, 1H), 8.82–8.78 (m, 2H), 8.32 (s, 1H), 8.23 (d, J = 7.6 Hz, 1H), 7.98 (s, 1H), 7.77–7.73 (m, 1H), 7.68–7.60 (m, 2H), 7.35 (s, 1H), 4.23 (q, J = 6.8 Hz, 2H), 3.96 (s, 3H), 1.44 (t, J = 6.8 Hz, 3H).	DMSO	398.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
560			410.83	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.89 (s, 1H), 9.48 (s, 1H), 8.74 (d, J = 6.0 Hz, 2H), 8.16 (dd, J = 6.8, 2.4 Hz, 1H), 7.85–7.88 (m, 2H), 7.67 (t, J = 6.4 Hz, 1H), 7.53 (t, J = 8.7 Hz, 1H), 7.32 (s, 1H), 3.98 (s, 6H).	DMSO	411.1, 413.1 (M+1)	Method A (TFA)	95	Method N, G1
561			394.37	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.92 (s, 1H), 9.49 (s, 1H), 8.77 (d, J = 6.4 Hz, 2H), 8.02–8.08 (m, 1H), 7.90 (s, 1H), 7.70–7.74 (m, 1H), 7.64–7.66 (m, 1H), 7.52–7.59 (m, 1H), 7.33 (s, 1H), 3.99 (s, 3H), 3.98 (s, 3H).	DMSO	395.0 (M+1), 198.1 (M/2+1)	Method A (TFA)	95	Method N, G1

(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
562			383.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.87 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 8.63-8.68 (m, 2H), 8.38 (s, 1H), 8.23 (d, J = 8.4 Hz, 1H), 7.89 (s, 1H), 7.69 (t, J = 7.8 Hz, 1H), 7.61 (d, J = 7.6 Hz, 1H), 7.52-7.56 (m, 1H), 7.33 (s, 1H), 4.00 (s, 3H), 3.96 (s, 3H).	DMSO	384.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
563			427.28	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.70 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 8.66 (d, J = 4.8 Hz, 1H), 8.62 (d, J = 8.0 Hz, 1H), 8.11 (d, J = 1.2 Hz, 2H), 7.79 (s, 1H), 7.53 (dd, J = 7.6, 4.8 Hz, 1H), 7.31 (s, 1H), 7.28 (s, 1H), 3.97 (s, 6H).	DMSO	427.1, 429.1, 431.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N, G1
564		2 HCl	389.3824	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.08 (s, 1H), 9.40 (s, 1H), 8.86 (d, J = 8.1 Hz, 1H), 8.78 (s, 1H), 8.53 (d, J = 10.0 Hz, 1H), 8.46 (s, 1H), 8.03 (d, J = 8.7 Hz, 1H), 8.00 - 7.88 (m, 2H), 7.73 (t, J = 7.8 Hz, 1H), 7.66 (d, J = 10.3 Hz, 2H), 7.28 (t, J = 7.6 Hz, 1H), 3.99 (s, 3H).	DMSO	390.0 (M+1)	Method B (NH ₄ CO ₃)	95	Method N, G1
565		2 HCl	401.4180	¹ H-NMR (400 MHz, CD ₃ OD): δ 9.20 (s, 1H), 8.85 (s, 1H), 8.78 (s, 1H), 8.72 (d, J = 8.2 Hz, 1H), 8.02 (d, J = 9.1 Hz, 1H), 7.95 (d, J = 7.7 Hz, 1H), 7.77 - 7.67 (m, 3H), 7.39 (t, J = 7.6 Hz, 1H), 4.18 (s, 3H), 4.08 (s, 3H).	MeOD	402.1 (M+1)	Method B (NH ₄ CO ₃)	95	Method N, G1

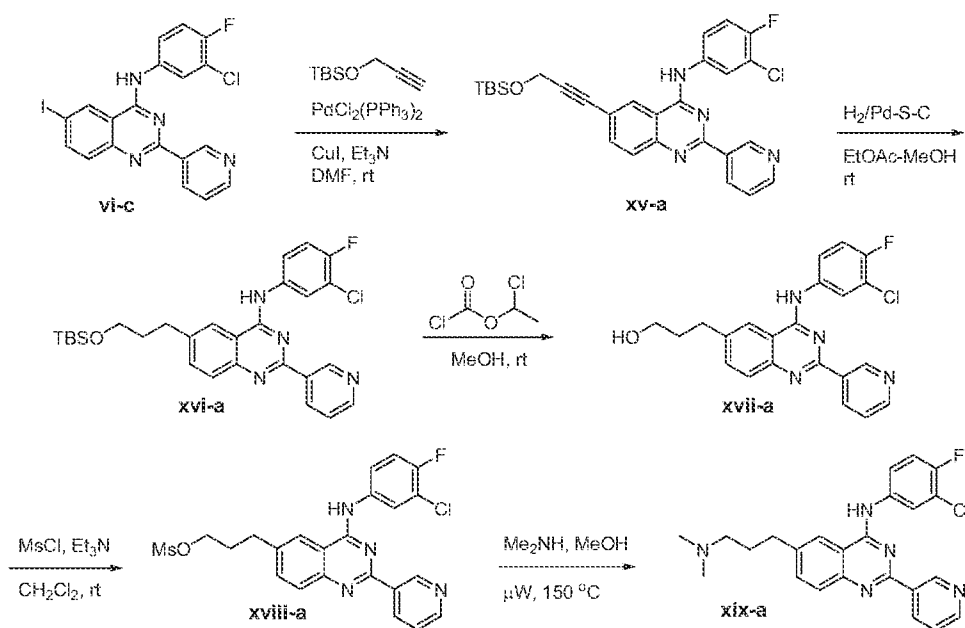
Scheme 8: General route for the synthesis of compounds with general formula xix

[0311]



Scheme 9: Representative synthesis of compounds of formula xix (see Scheme 8)

[0312]



[0313] Method

- O1: CuI/Pd(PPh₃)₂Cl₂/Et₃N/DMF/rt
 O2: Propargyl bromide/CuI/Pd(PPh₃)₂Cl₂/Nucleophile/DMF/rt
 O3: Pd(OAc)₂/PPh₃/TBAB/piperidine/THF-H₂O/rt

[0314] Method O1: 6-(3-(tert-butyldimethylsilyloxy)prop-1-ynyl)-N-(3-chloro-4-fluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine (xv-a) A suspension of N-(3-chloro-4-fluorophenyl)-6-iodo-2-(pyridin-3-yl)quinazolin-4-amine (synthesized as described in Scheme 1 and 4, substituting 5-iodo-2-nitrobenzoic acid for 2-nitro-5-propoxybenzoic acid and 3-chloro-4-fluoroaniline for 2-aminobenzamide) (1.00 g, 2.10 mmol), tert-butyldimethyl(2-propynyloxy)silane (0.85 ml, 4.20 mmol), copper(I) iodide (4.0 mg, 0.021 mmol), dichlorobis(triphenylphosphine) palladium

(II) (Pd(PPh₃)₂Cl₂) (29 mg, 0.042 mmol), and triethylamine (1.17 ml, 8.39 mmol) in DMF (15 mL) was stirred overnight at room temperature under argon atmosphere. Water (30 mL) and ethyl acetate (30 mL) were added to the mixture. The resultant precipitate was removed by filtration. The filtrate was extracted with EtOAc (2 x 50 mL). The combined organic layer was washed with water (1 x 100 mL) and brine (1 x 100 mL) and was dried over Na₂SO₄. After filtration and evaporation, the crude product was obtained, which was purified by column chromatography on silica gel (eluted with hexane/ethyl acetate 6:1 to 1:3) to give 0.73 g of 6-(3-(tert-butyldimethylsilyloxy)prop-1-ynyl)-N-(3-chloro-4-fluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine as light brown solid (67 %). LCMS m/z = 519 (M + 1) (Method D) (retention time = 3.22 min). ¹H NMR (300 MHz, DMSO) δ 9.99 (s, 1H), 9.35 (s, 1H), 8.62 - 8.44 (m, 3H), 8.11 (dd, J = 6.8, 2.6 Hz, 1H), 7.76 - 7.66 (m, 3H), 7.45 - 7.30 (m, 2H), 4.48 (s, 2H), 0.75 (s, 9H), 0.16 (s, 6H).

[0315] Method P: 6-(3-(tert-butyldimethylsilyloxy)propyl)-N-(3-chloro-4-fluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine (xvi-a) A suspension of 6-(3-(tert-butyldimethylsilyloxy)prop-1-ynyl)-N-(3-chloro-4-fluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine (0.20 g, 0.39 mmol) and 5 % Pd-S-C (40 mg) in EtOAc (5 mL) and MeOH (5 mL) was stirred overnight at room temperature under hydrogen atmosphere. The reaction mixture was filtered through Celite. The filtrate was concentrated *in vacuo*. The residue was purified by column chromatography on silica gel (eluted with hexane/ethyl acetate 6:1 to 2:3) to give 0.15 g of 6-(3-(tert-butyldimethylsilyloxy)propyl)-N-(3-chloro-4-fluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine as yellow solid (74 %). LCMS m/z = 523 (M + 1) (Method D) (retention time = 3.35 min). ¹H NMR (300 MHz, CDCl₃) δ 9.69 (dd, J = 2.2, 0.8 Hz, 1H), 8.79 - 8.65 (m, 2H), 8.07 (dd, J = 6.5, 2.7 Hz, 1H), 7.94 (d, J = 8.5 Hz, 1H), 7.76 - 7.61 (m, 3H), 7.47 - 7.37 (m, 2H), 7.31 - 7.18 (m, 1H), 3.68 (t, J = 6.1 Hz, 2H), 3.01 - 2.83 (m, 2H), 2.03 - 1.86 (m, 2H), 0.94 (s, 9H), 0.08 (s, 6H).

[0316] Method Q: 3-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)propan-1-ol (xvii-a) To a suspension of 6-(3-(tert-butyldimethylsilyloxy)propyl)-N-(3-chloro-4-fluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine (0.35 g, 0.67 mmol) in MeOH (10 mL) was added 1-chloroethyl chloroformate (7.2 μl, 0.067 mmol). The mixture was stirred overnight at room temperature. Methanol was removed *in vacuo*. Sat. NaHCO₃ aqueous (10 mL) and CH₂Cl₂ (10 mL) were added to the residue and stirred for a while. The resultant solid was collected by filtration and dried to give 0.25 g of 3-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)propan-1-ol as light yellow solid (91 %). MS m/z = 409 (M + 1) (Method D) (retention time = 1.71 min). ¹H NMR (300 MHz, DMSO) δ 10.01 (s, 1H), 9.51 (s, 1H), 8.66 (m, 2H), 8.37 (s, 1H), 8.29 - 8.20 (m, 1H), 7.94 - 7.67 (m, 3H), 7.63 - 7.43 (m, 2H), 4.60 (t, J = 5.0 Hz, 1H), 3.56 - 3.35 (m, 2H), 2.94 - 2.69 (m, 2H), 2.15 - 1.56 (m, 2H).

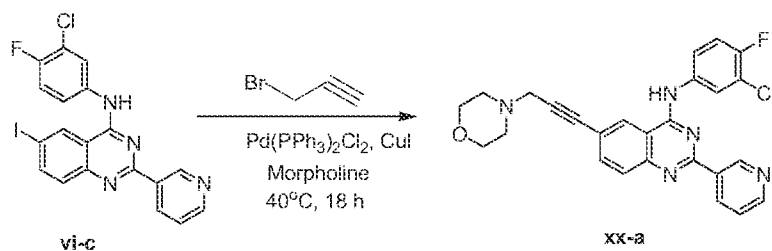
[0317] Method R: 3-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)propyl methanesulfonate (xviii-a) To a suspension of 3-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)propan-1-ol (0.25 g, 0.61 mmol) and triethylamine (0.17 ml, 1.22 mmol) in CH₂Cl₂ (10 mL) was added methanesulfonyl chloride (0.057 ml, 0.73 mmol). The mixture was stirred at room temperature for 1 h. Water (10 mL) was added to the mixture and stirred for a while. The resultant precipitate was collected by filtration and washed with CH₂Cl₂ and dried to give 0.27 g of 3-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)propyl methanesulfonate as pale yellow solid (91 %), which was used without further purification.

Method G4: N-(3-chloro-4-fluorophenyl)-6-(3-(dimethylamino)propyl)-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride (xviv-a) (This method is representative of method G4 and G5. These two methods can be implemented in a similar way except for substitution of the appropriate solvent and temperature)

[0318] A solution of 3-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)propyl methanesulfonate (40 mg, 82 mmol) and 40 % Me₂NH aqueous (1 mL) in methanol (2 mL) was placed in a microwave reaction vial. The mixture was heated under microwave irradiation conditions at 150°C for 30 minutes after which the solvent was removed *in vacuo*. The crude product was obtained, which was purified by column chromatography on basic silica gel (eluted with ethyl acetate/methanol 1:0→5:1). The HCl salt generated by 4 M HCl in dioxane was crystallized from 2-propanol to give 10 mg of N-(3-chloro-4-fluorophenyl)-6-(3-(dimethylamino)propyl)-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride as pale brown powder (24 %). LCMS m/z = 436 (M + 1) (Method C) (retention time = 1.68 min). ¹H NMR (300 MHz, DMSO) δ 10.59 (s, 1H), 10.18 (s, 1H), 9.51 (s, 1H), 8.98 - 8.76 (m, 2H), 8.65 (s, 1H), 8.26 (dd, J = 6.8, 2.6 Hz, 1H), 8.05 - 7.74 (m, 4H), 7.54 (t, J = 9.1 Hz, 1H), 3.19 - 3.03 (m, 2H), 2.97 - 2.83 (m, 2H), 2.78 (s, 3H), 2.77 (s, 3H), 2.16 (s, 2H).

Scheme 10: Method O2: Representative synthesis of compounds of formula xx

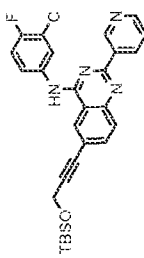
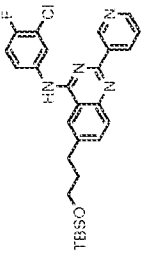
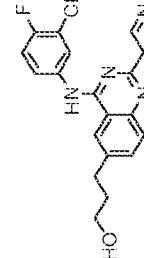
[0319]



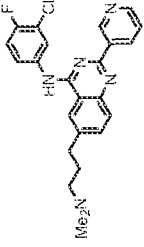
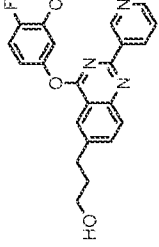
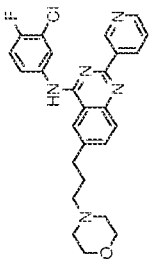
[0320] N-(3-chloro-4-fluorophenyl)-6-(3-morpholinoprop-1-ynyl)-2-(pyridin-3-yl) quinazolin-4-amine (xx-a) To a solution of N-(3-chloro-4-fluorophenyl)-6-iodo-2-(pyridin-3-yl)quinazolin-4-amine (synthesized as described in Scheme 1 and 4, substituting 5-iodo-2-nitrobenzoic acid for 2-nitro-5-propoxy-benzoic acid and 3-chloro-4-fluoroaniline for 2-aminobenzamide) (2.19 g, 4.6 mmol, 1 eq.), Pd(PPh₃)₂Cl₂ (161 mg, 0.23 mmol, 0.05 eq.), CuI (87 mg, 0.46 mmol, 0.1 eq.) in morpholine (15 mL) was added 3-bromoprop-1-yne (814 mg, 6.9 mmol, 1.5 eq.) at 0 °C under Ar atmosphere, following a procedure from Tetrahedron, 2007, 63, 10671-10683. The mixture was stirred at 40 °C overnight. After cooling, the mixture was filtered and methanol (60 mL) was added to the filtrate to form a precipitate. The precipitate was collected and re-crystallized from ethyl acetate twice to afford 1.60 g of xx-a as yellow solid (yield 74%). LCMS m/z = 474.1 (M + 1), 476.1 (M + 3) (Method C) (retention time = 1.97 min).

[0321] The compounds in the following table were prepared in a manner analogous to that described in Scheme 8 substituting with appropriate nucleophile.

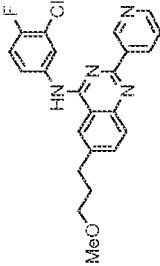
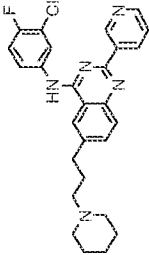
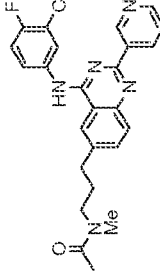
Table 3:

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (Min)	LCMS Protocol	Purity percent	Method for Coupling
566			519.085	¹ H NMR (300 MHz, DMSO) δ 9.99 (s, 1H), 9.35 (s, 1H), 8.62-8.44 (m, 3H), 8.11 (dd, J = 6.8, 2.6 Hz, 1H), 7.76-7.66 (m, 3H), 7.45-7.30 (m, 2H), 4.48 (s, 2H), 0.75 (s, 9H), 0.00 (s, 6H).	DMSO	519 (M+1)	3.22	Method D	96	Method O1
567			523.117	¹ H NMR (300 MHz, CDCl ₃) δ 9.69 (dd, J = 2.2, 0.8 Hz, 1H), 8.79-8.65 (m, 2H), 8.07 (dd, J = 6.5, 2.7 Hz, 1H), 7.94 (d, J = 8.5 Hz, 1H), 7.76-7.61 (m, 3H), 7.47-7.37 (m, 2H), 7.31-7.18 (m, 1H), 3.68 (t, J = 6.1 Hz, 2H), 3.01-2.83 (m, 2H), 2.03-1.	CDCl ₃	523 (M+1)	3.35	Method D	100	Methods O1, P
568			408.856	¹ H NMR (300 MHz, DMSO) δ 10.01 (s, 1H), 9.51 (s, 1H), 8.66 (m, 2H), 8.37 (s, 1H), 8.29-8.20 (m, 1H), 7.94-7.67 (m, 3H), 7.63-7.43 (m, 2H), 4.60 (t, J = 5.0 Hz, 1H), 3.56-3.35 (m, 2H), 2.94-2.69 (m, 2H), 2.15-1.56 (m, 2H).	DMSO	409 (M+1)	1.71	Method D	100	Methods O1, P, Q

(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (Min)	LCMS Protocol	Purity percent	Method for Coupling
569		2 HCl	508.846	¹ H NMR (300 MHz, DMSO) δ 10.59 (s, 1H), 10.18 (s, 1H), 9.51 (s, 1H), 8.98-8.76 (m, 2H), 8.65 (s, 1H), 8.26 (dd, J = 6.8, 2.6 Hz, 1H), 8.05 - 7.74 (m, 4H), 7.54 (t, J = 9.1 Hz, 1H), 3.19 - 3.03 (m, 2H), 2.97-2.83 (m, 2H), 2.78 (s, 3H), 2.77 (s, 3H), 2.	DMSO	436 (M+1)	1.68	Method C	100	Method G4
570			409.841	¹ H NMR (300 MHz, CDCl ₃) δ 9.49 (d, J = 1.4 Hz, 1H), 8.73 - 8.51 (m, 2H), 8.14 (s, 1H), 8.01 (d, J = 8.6 Hz, 1H), 7.81 (dd, J = 8.8, 1.9 Hz, 1H), 7.51 - 7.13 (m, 4H), 3.77 (dd, J = 11.5, 6.2 Hz, 2H), 3.11 - 2.86 (m, 2H), 2.16-1.93 (m, 2H), 1.44 (t, J = 5	CDCl ₃	410 (M+1)	1.95	Method D	100	Methods O1, P, Q
571		2 HCl	550.882	¹ H NMR (300 MHz, DMSO) δ 10.83-10.37 (m, 2H), 9.51 (s, 1H), 8.97 - 8.74 (m, 2H), 8.62 (s, 1H), 8.27 (d, J = 4.2 Hz, 1H), 8.05-7.72 (m, 4H), 7.54 (t, J = 9.0 Hz, 1H), 3.96 (d, J = 12.7 Hz, 2H), 3.76 (t, J = 12.1 Hz, 2H), 3.47 (d, J = 12.0 Hz, 2H), 3.24	DMSO	478 (M+1)	2.01	Method C	100	Method G4

(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (Min)	LCMS Protocol	Purity percent	Method for Coupling
572			422.882	¹ H NMR (300 MHz, DMSO) δ 9.99 (s, 1H), 9.51 (s, 1H), 8.75–8.60 (m, 2H), 8.37 (s, 1H), 8.26 (dd, J = 6.9, 2.6 Hz, 1H), 7.99–7.88 (m, 1H), 7.88–7.73 (m, 2H), 7.60–7.46 (m, 2H), 3.39 (dd, J = 8.4, 4.1 Hz, 2H), 3.27 (s, 3H), 2.92–2.78 (m, 2H), 2.05 (m, 2H), 2.05	DMSO	423 (M+1)	2.09	Method D	100	Method G4
573		HCl	548.909	¹ H NMR (300 MHz, DMSO) δ 10.69 (s, 1H), 10.12 (s, 1H), 9.51 (s, 1H), 9.02–8.80 (m, 2H), 8.69 (s, 1H), 8.27 (dd, J = 6.9, 2.7 Hz, 1H), 8.08–7.78 (m, 4H), 7.54 (t, J = 9.1 Hz, 1H), 3.46 (d, J = 10.2 Hz, 2H), 3.17–3.00 (m, 2H), 2.99–2.75 (m, 4H), 2.3	DMSO	476 (M+1)	1.79	Method C	100	Method G4
574		HCl	500.395	¹ H NMR (300 MHz, DMSO) δ 10.68 (s, 1H), 9.49 (s, 1H), 9.12–8.99 (m, 1H), 8.94 (d, J = 3.9 Hz, 1H), 8.58 (d, J = 11.5 Hz, 1H), 8.17 (dd, J = 6.7, 2.6 Hz, 1H), 8.07–7.82 (m, 5H), 7.55 (t, J = 9.1 Hz, 1H), 3.43–3.29 (m, 2H), 3.08–2.65 (m, 5H), 2.09–	DMSO	464 (M+1)	1.78	Method D	95	Method G4 followed by acylation with acetyl chloride/TEA/DCM, rt

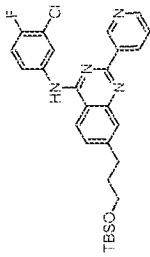
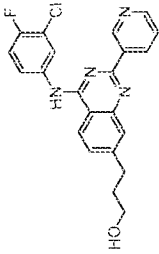
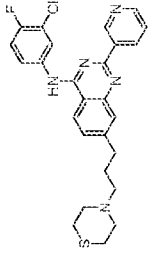
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (Min)	LCMS Protocol	Purity percent	Method for Coupling
575		HCl	600.39	¹ H NMR (300 MHz, DMSO) δ 11.79 (s, 1H), 10.81 (s, 1H), 9.50 (s, 1H), 8.98 (d, J = 8.2 Hz, 1H), 8.89 (d, J = 5.0 Hz, 1H), 8.75 (s, 1H), 8.27 (d, J = 4.8 Hz, 1H), 8.10-7.79 (m, 4H), 7.53 (t, J = 9.1 Hz, 1H), 5.18 - 3.09 (m, 10H), 3.01-2.69 (m, 5H), 2.37	DMSO	491 (M+1)	1.76	Method C	100	Method G4
576			473.93	¹ H NMR (300 MHz, DMSO) δ 10.17 (s, 1H), 9.52 (s, 1H), 8.70 (s, 1H), 8.66 (d, J = 8.1 Hz, 1H), 8.55 (d, J = 8.4 Hz, 1H), 8.27 (dd, J = 6.9, 2.6 Hz, 1H), 7.93 (s, 2H), 7.70 (d, J = 8.4 Hz, 1H), 7.61-7.51 (m, 2H), 3.67-3.61 (m, 6H), 2.61-2.55 (m, 4H).	DMSO	473.9 (M+1)	2.2	Method C	100	Method O1
577			435.92	¹ H NMR (300 MHz, DMSO) δ 10.06 (s, 1H), 9.54 (s, 1H), 8.77-8.63 (m, 2H), 8.52 (d, J = 8.4 Hz, 1H), 8.29 (d, J = 6.8 Hz, 1H), 7.93 (s, 1H), 7.79 (s, 1H), 7.64-7.51 (m, 3H), 3.00 (t, J = 8.2 Hz, 2H), 2.87 (t, J = 7.6 Hz, 2H), 2.72 (s, 6H), 2.14-1.99 (	DMSO	436.0 (M+1)	1.68	Method C	100	Methods O1, P

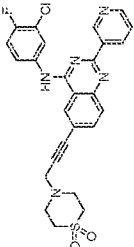
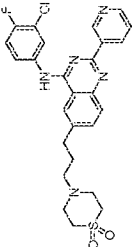
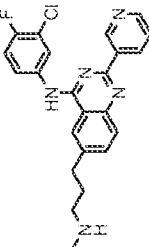
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (Min)	LCMS Protocol	Purity percent	Method for Coupling
578			519.09	¹ H NMR (300 MHz, DMSO) δ 10.17 (s, 1H), 9.67 - 9.43 (m, 1H), 8.68 (s, 2H), 8.55 (d, J = 8.4 Hz, 1H), 8.28 (s, 1H), 7.89 (s, 2H), 7.68 (s, 1H), 7.56 (d, J = 9.1 Hz, 2H), 4.64 (s, 2H), 0.92 (s, 9H), 0.18 (s, 6H).	DMSO	518.8 (M+1)	3.18	Method C	100	Method O1
579			404.82	¹ H NMR (300 MHz, DMSO) δ 10.19 (s, 1H), 9.51 (s, 1H), 8.65 (d, J = 8.1 Hz, 2H), 8.54 (d, J = 8.4 Hz, 1H), 8.26 (s, 1H), 7.85 (s, 2H), 7.67 - 7.60 (m, 1H), 7.52 (d, J = 7.0 Hz, 2H), 5.48 (s, 1H), 4.39 (d, J = 5.8 Hz, 2H).	DMSO	404.9 (M+1)	2.05	Method C	100	Methods O1, Q
580			477.96	¹ H NMR (300 MHz, DMSO) δ 9.99 (s, 1H), 9.51 (s, 1H), 8.73 - 8.61 (m, 2H), 8.45 (d, J = 8.6 Hz, 1H), 8.28 (dd, J = 6.8, 2.5 Hz, 1H), 7.97 - 7.86 (m, 1H), 7.71 (s, 1H), 7.60 - 7.47 (m, 3H), 3.63 - 3.51 (m, 4H), 2.82 (t, J = 7.5 Hz, 2H), 2.42 - 2.25 (m, 6H).	DMSO	477.9 (M+1)	1.99	Method C	100	Methods O1, P

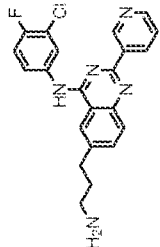
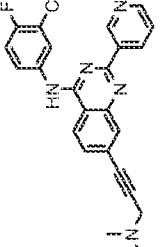
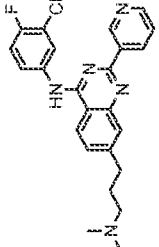
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (Min)	LCMS Protocol	Purity percent	Method for Coupling
581			523.12	¹ H NMR (300 MHz, CDCl ₃) δ 9.67 (d, J = 1.3 Hz, 1H), 8.76 - 8.66 (m, 2H), 8.03 (dd, J = 6.5, 2.7 Hz, 1H), 7.82 - 7.74 (m, 2H), 7.64 (ddd, J = 8.9, 4.0, 2.7 Hz, 1H), 7.46 (s, 1H), 7.44 - 7.37 (m, 2H), 7.20 (t, J = 8.7 Hz, 1H), 3.67 (t, J = 6.1 Hz, 2H), 2.95	CDCl ₃	523.1 (M+1)	3.35	Method C	100	Methods O1, P
582			408.86	¹ H NMR (300 MHz, CD ₃ OD) δ 9.50 (d, J = 1.3 Hz, 1H), 8.77 (d, J = 8.0 Hz, 1H), 8.62 (dd, J = 4.9, 1.6 Hz, 1H), 8.26 (d, J = 8.5 Hz, 1H), 8.18 (dd, J = 6.7, 2.6 Hz, 1H), 7.83 - 7.71 (m, 2H), 7.61 - 7.47 (m, 2H), 7.30 (t, J = 9.0 Hz, 1H), 3.65 (t, J = 6.4 Hz	CD ₃ OD	409.0 (M+1)	1.99	Method C	100	Methods O1, P, Q
583			494.03	¹ H NMR (300 MHz, CD ₃ OD) δ 9.50 - 9.46 (m, 1H), 8.77 - 8.70 (m, 1H), 8.61 (dd, J = 4.9, 1.6 Hz, 1H), 8.22 (d, J = 8.5 Hz, 1H), 8.17 (dd, J = 6.8, 2.6 Hz, 1H), 7.75 (ddd, J = 9.0, 4.2, 2.7 Hz, 1H), 7.70 (s, 1H), 7.54 (ddd, J = 8.0, 4.9, 0.8 Hz, 1H), 7.46 (d	CD ₃ OD	494.0 (M+1)	2.29	Method C	91	Method G4

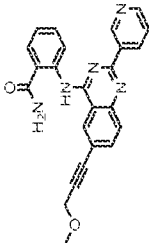
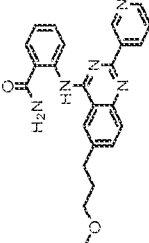
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (Min)	LCMS Protocol	Purity percent	Method for Coupling
584			522	¹ H NMR (300 MHz, DMSO) δ 10.15 (s, 1H), 8.74 - 8.61 (m, 2H), 8.25 (d, J = 3.5 Hz, 1H), 7.96 - 7.80 (m, 4H), 7.59 - 7.46 (m, 2H), 3.79 (s, 2H), 3.18 (s, 4H), 3.06 (s, 4H).	DMSO	522.0 (M+1)	2.18	Method C	100	Method 02
585			526.06	¹ H NMR (300 MHz, DMSO) δ 9.99 (s, 1H), 9.52 (d, J = 2.0 Hz, 1H), 8.72 - 8.62 (m, 2H), 8.37 (s, 1H), 8.27 (dd, J = 6.9, 2.6 Hz, 1H), 7.96 - 7.88 (m, 1H), 7.83 (d, J = 2.6 Hz, 2H), 7.61 - 7.50 (m, 2H), 3.10 (s, J = 17.4 Hz, 6H), 2.95-2.80 (m, 8H), 2.58 -	DMSO	526.1 (M+1)	2.11	Method C	91	Methods O2, P
586		HCl	494.82	¹ H NMR (300 MHz, DMSO) δ 10.49 (s, 1H), 9.50 (s, 1H), 8.96 - 8.53 (m, 5H), 8.25 (dd, J = 6.9, 2.5 Hz, 1H), 8.05 - 7.73 (m, 4H), 7.54 (t, J = 9.1 Hz, 1H), 3.02 - 2.78 (m, 4H), 2.64 - 2.37 (m, 3H), 2.15 - 1.99 (m, 2H).	DMSO	422 (M+1)	1.64	Method B (Ammonium formate)	100	Method 02

(continued)

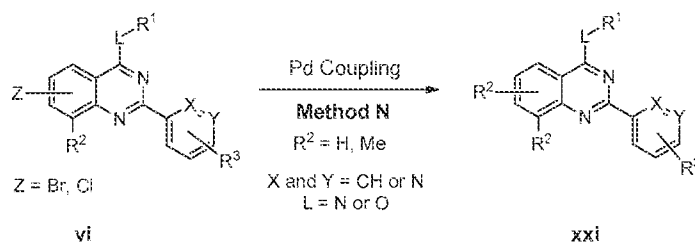
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (Min)	LCMS Protocol	Purity percent	Method for Coupling
587		HCl	480.79	¹ H NMR (300 MHz, DMSO) δ 10.61 (s, 1H), 9.50 (s, 1H), 9.00–8.78 (m, 2H), 8.66 (s, 1H), 8.25 (dd, J = 6.8, 2.6 Hz, 1H), 8.13–7.75 (m, 7H), 7.54 (t, J = 9.1 Hz, 1H), 3.02–2.73 (m, 4H), 2.14–1.94 (m, 2H).	DMSO	408 (M+1)	1.60	Method B (Ammonium formate)	100	Method O2
588			431.89	¹ H NMR (300 MHz, DMSO) δ 10.18 (s, 1H), 8.65 (d, J = 6.6 Hz, 1H), 8.56 (d, J = 8.7 Hz, 1H), 8.23 (dd, J = 6.9, 2.4 Hz, 1H), 8.01 (s, 1H), 7.91 (dd, J = 7.0, 3.5 Hz, 1H), 7.73 (d, J = 8.5 Hz, 1H), 7.68–7.39 (m, 3H), 4.07 (s, 2H), 2.67 (s, 6H).	DMSO	431.9 (M+1)		Method C	99	Method O1
589			435.92	¹ H NMR (300 MHz, DMSO) δ 10.06 (s, 1H), 9.53 (s, 1H), 8.76–8.62 (m, 2H), 8.51 (d, J = 8.4 Hz, 1H), 8.28 (dd, J = 6.9, 2.5 Hz, 1H), 7.92 (dd, J = 4.5, 1.8 Hz, 1H), 7.79 (s, 1H), 7.65–7.49 (m, 3H), 3.08–2.95 (m, 2H), 2.86 (t, J = 7.6 Hz, 2H), 2.71 (s, J = 7.2 Hz, 6H), 2.05 (dt, J = 13.8, 7.1 Hz, 2H).	DMSO	436.0 (M+1)		Method C	99	Method O1, P

(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (Min)	LCMS Protocol	Purity percent	Method for Coupling
590			409.44	¹ H-NMR (300 MHz, DMSO): δ 13.19 (s, 1H), 9.11 (s, 1H), 8.88 (s, 1H), 8.53 (s, 1H), 8.29 (s, 1H), 8.06 (s, 1H), 7.94 (d, J = 10.6 Hz, 4H), 7.78 (s, 1H), 7.25 (s, 1H), 4.44 (s, 2H), 3.37 (s, 3H).	DMSO	410.5 (M+1)		Method C	95	Method O1
591			413.47		DMSO	414.5 (M+1)		Method C	95	Method O1, P

Scheme 11: General route for the synthesis of compounds with general formula xxi:

[0322]



Method N1: Pd(APhos)₂Cl₂/ K₃PO₄ or Cs₂CO₃/ Boronic acid or ester/ Dioxane - H₂O, heat

Method N2: Pd(PPh₃)₂Cl₂/ K₂CO₃/ Boronic acid or ester/ DME-EtOH-H₂O / microwave, 120 °C

Method N3: Pd(OAc)₂/ Xphos/ Cs₂CO₃/ THF- H₂O, 80 °C

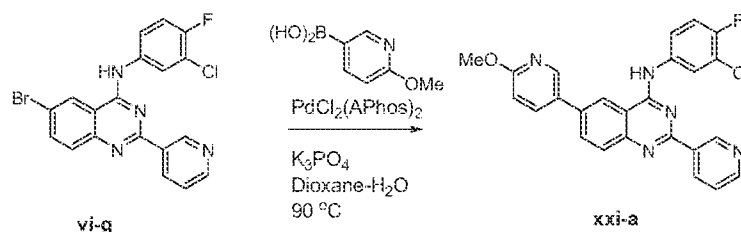
Method N4: Pd(OAc)₂/ Sphos/ K₂CO₃/ MeCN- H₂O microwave, 120 °C

Method N5: Pd(PPh₃)₄/K₃PO₄/ Dioxane - H₂O, heat

Method N6: Pd(dppf)Cl₂-CH₂Cl₂/K₃PO₄/ Dioxane - H₂O, heat

Scheme 12: Method N1: Representative synthesis of compounds of formula xxi (see Scheme 11).

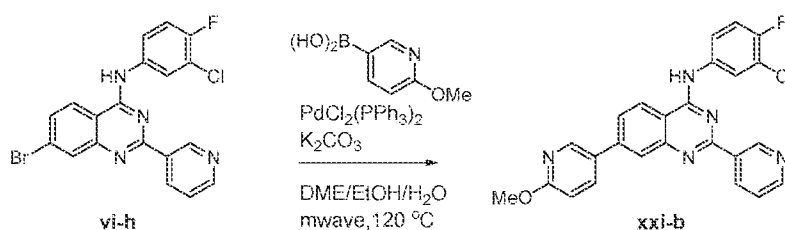
[0323]



[0324] N-(3-chloro-4-fluorophenyl)-6-(6-methoxypyridin-3-yl)-2-(pyridin-3-yl)quinazolin-4-amine (xxi-a) A 2.0 dram reaction vial was charged with 6-bromo-N-(3-chloro-4-fluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine (**synthesized as described in Scheme 1 and 4, substituting 5-bromo-2-nitrobenzoic acid for 2-nitro-5-propoxy-benzoic acid and 3-chloro-4-fluoroaniline for 2-aminobenzamide**) (100 mg, 0.233 mmol, 1.0 equiv), 6-methoxypyridin-3-ylboronic acid (44.5 mg, 0.291 mmol, 1.25 equiv), Pd(APhos)₂Cl₂ (6.6 mg, 0.0093 mmol, 4 mol%) and potassium phosphate monohydrate (69 mg, 0.70 mmol, 3.0 equiv). The mixture was suspended in dioxane/water (9:1, 4 mL), and the reaction was heated at 90 °C for 14 h. The reaction mixture was cooled to room temperature, diluted with water (15 mL) and the resultant precipitate was collected by filtration. The crude product was purified by stirring in methanol for 30 min at 60 °C to give the desired product as a pale yellow solid (58 mg, 54%) LCMS m/z = 458.1 (M+1) (Method C) (retention time = 2.56 min). ¹H NMR (300 MHz, DMSO) δ 10.12 (s, 1H), 9.53 (s, 1H), 8.82 (s, 1H), 8.70 (dd, J = 16.9, 5.4 Hz, 3H), 8.25 (d, J = 8.6 Hz, 3H), 8.01 - 7.87 (m, 2H), 7.57 (dd, J = 11.2, 6.0 Hz, 2H), 7.03 (d, J = 8.6 Hz, 1H), 3.94 (s, 3H).

Scheme 13: Method N2: Representative synthesis of compounds of formula xxi (see Scheme 11)

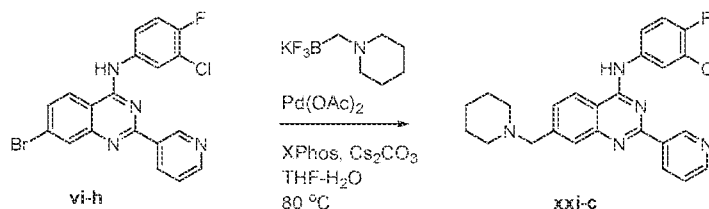
[0325]



[0326] N-(3-chloro-4-fluorophenyl)-7-(6-methoxypyridin-3-yl)-2-(pyridin-3-yl)quinazolin-4-amine (xxi-b) A microwave vial was charged with 7-bromo-N-(3-chloro-4-fluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine (**synthesized as described in Scheme 1 and 4, substituting 4-bromo-2-nitrobenzoic acid for 2-nitro-5-propoxybenzoic acid and 3-chloro-4-fluoroaniline for 2-aminobenzamide**) (100 mg, 0.233 mmol), 6-methoxypyridin-3-ylboronic acid (44.5 mg, 0.291 mmol, 1.25 equiv), Pd(PPh₃)₂Cl₂ (8.1 mg, 5 mol%) and potassium carbonate (160.1 mg, 1.16 mmol, 5.0 equiv). The mixture was suspended in DME/water/ethanol (7:3:2, 4 mL), and the reaction was heated under microwave irradiation conditions at 120°C for 10 minutes. The crude reaction mixture was diluted with water (10 mL) and then filtered. The solid residue was dissolved in methanol/THF (1:1, 5 mL) with heating and then filtered through Celite to remove the catalyst. The resulting filtrate was concentrated to afford the desired product as a tan solid (7.0 mg, 6.5%). LCMS m/z = 458.1 (M+1) (Method C) (retention time = 2.51 min). ¹H NMR (300 MHz, DMSO) δ 10.12 (s, 1H), 9.55 (s, 1H), 8.76 (d, J = 2.5 Hz, 1H), 8.72 - 8.65 (m, 2H), 8.62 (d, J = 8.6 Hz, 1H), 8.34 - 8.26 (m, 2H), 8.15 (d, J = 1.6 Hz, 1H), 8.02 (d, J = 8.7 Hz, 1H), 7.97 - 7.88 (m, 1H), 7.61 - 7.49 (m, 2H), 6.99 (d, J = 8.7 Hz, 1H), 3.94 (s, 3H).

Scheme 14: Method N3: Representative synthesis of compounds of formula xxi (see Scheme 11). (This method is representative of method N4 except for substitution of the appropriate ligand, solvent and the use of microwave instead of heating)

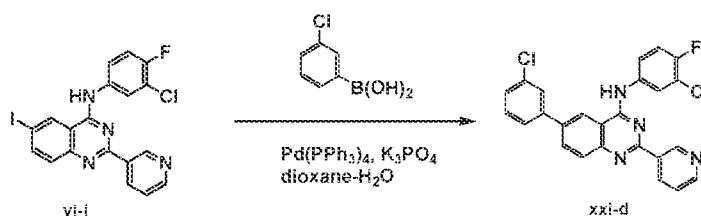
[0327]



[0328] N-(3-chloro-4-fluorophenyl)-7-(piperidin-1-ylmethyl)-2-(pyridin-3-yl)quinazolin-4-amine (xxi-c) A dry 15 mL sealed tube was charged with 7-bromo-N-(3-chloro-4-fluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine (**synthesized as described in Scheme 1 and 4, substituting 4-bromo-2-nitrobenzoic acid for 2-nitro-5-propoxybenzoic acid and 3-chloro-4-fluoroaniline for 2-aminobenzamide**) (100 mg, 0.233 mmol), potassium 1-trifluoroboratomethylpiperidine (52.5 mg, 0.256 mmol), cesium carbonate (227.5 mg, 0.698 mmol), Pd(OAc)₂ (1.6 mg, 3 mol%) and XPhos (6.7 mmol, 6 mol%) in THF/water (10:1, 3.3 mL). The reaction mixture was stirred at 80°C for 16 hours. The reaction mixture was filtered through a pad of Celite and solvent was removed *in vacuo*. The crude product was purified by ISCO (silica, 12 g column, 97% dichloromethane - 3% methanol - 0.1 % NH₄OH) to yield the desired compound as a yellow solid (23.3 mg, 22%). LCMS m/z = 448.0 (M+1) (Method C) (retention time = 1.95 min). ¹H NMR (300 MHz, CD₃OD) δ 9.40 (d, J = 1.4 Hz, 1H), 8.70 - 8.62 (m, 1H), 8.56 (dd, J = 4.9, 1.6 Hz, 1H), 8.18 - 8.09 (m, 2H), 7.69 (ddd, J = 6.8, 4.1, 2.1 Hz, 2H), 7.54 - 7.45 (m, 2H), 7.23 (t, J = 9.0 Hz, 1H), 3.61 (s, 2H), 2.47 (s, 4H), 1.64 (dd, J = 13.9, 8.9 Hz, 5H), 1.49 (d, J = 4.1 Hz, 2H).

Scheme 15: Method N5: Representative synthesis of compounds of formula xxi (see Scheme 11)

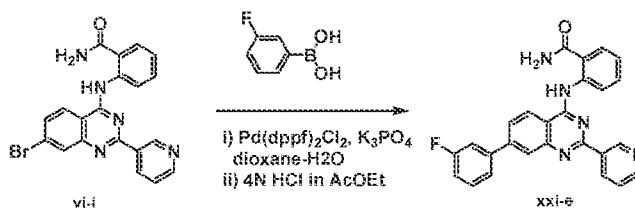
[0329]



[0330] N-(3-chloro-4-fluorophenyl)-6-(3-chlorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine (xxi-d) A mixture of N-(3-chloro-4-fluorophenyl)-6-iodo-2-(pyridin-3-yl)quinazolin-4-amine (1.0 g, 2.10 mmol), 3-chlorophenylboronic acid (0.49 g, 3.13 mmol), Pd(PPh₃)₄ (0.24 g, 0.210 mmol), K₃PO₄ (1.34 g, 6.31 mmol) in dioxane (20 mL) and water (2.0 mL) was stirred under reflux for 2 h. Ethyl acetate (20 mL) was added to the cooled mixture and filtered. The filtered solid was recrystallized from DMF and water to give the title compound (0.50 g, 51.6 %). ¹H NMR (400 MHz, DMSO) δ 10.20 (s, 1H), 9.54 (s, 1H), 8.87 (s, 1H), 8.74 - 8.65 (m, 2H), 8.31 - 8.23 (m, 2H), 8.04 - 7.85 (m, 4H), 7.64 - 7.49 (m, 4H).

Scheme 16: Method N6: Representative synthesis of compounds of formula xxi (see Scheme 11)

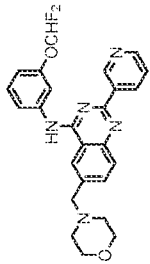
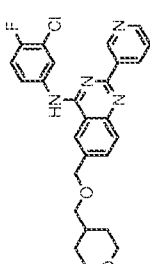
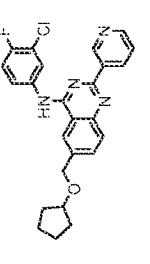
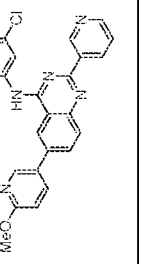
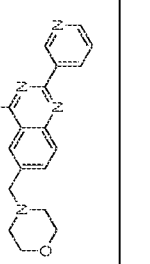
[0331]



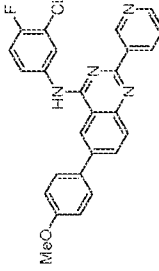
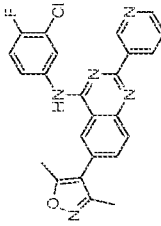
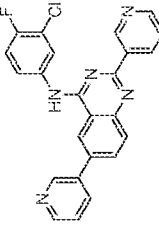
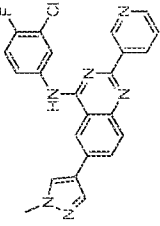
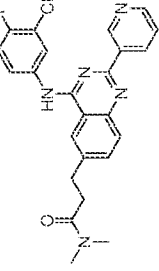
[0332] 2-(7-(3-fluorophenyl)-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide dihydrochloride (xxi-e) A mixture of 2-(7-bromo-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (0.40 g, 0.952 mmol), 3-fluorophenylboronic acid (0.20g 1.43 mmol), Pd(dppf)₂Cl₂ (77mg, 0.094mmol), K₃PO₄ (606 mg, 2.85 mmol) in dioxane (8 mL) and water (2 mL) was refluxed under argon atmosphere for 4 h and cooled. Ethyl acetate (10 mL) was added to the mixture and a precipitate formed and was filtered. The filtered solid was recrystallized from DMF and water to give the title compound as free form. The solid as free form was suspended in ethyl acetate (10 mL) and 4N HCl in ethyl acetate (0.71 mL) was added to the suspension. A precipitate formed and was subjected to sonication for 20 min, filtered and dried to give the title compound (0.24 g, 49.6 %). ¹H NMR (400 MHz, DMSO) δ 13.19 (s, 1H), 9.64 (s, 1H), 9.21 - 9.08 (m, 1H), 9.07 - 8.89 (m, 2H), 8.51 (s, 1H), 8.38 - 8.26 (m, 2H), 8.16 (d, J = 8.5 Hz, 1H), 8.07 - 7.87 (m, 3H), 7.83 - 7.70 (m, 3H), 7.62 (dd, J = 14.2, 7.7 Hz, 1H), 7.40 - 7.22 (m, 2H).

[0333] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 11** substituting with appropriate boronic acid/ester or boronate salt, catalyst and solvent

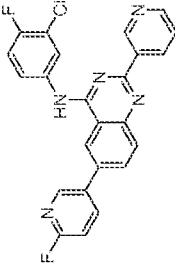
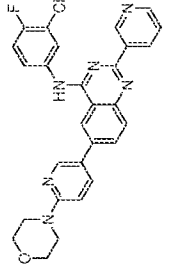
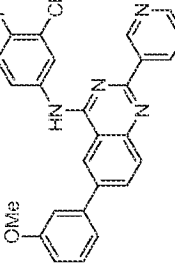
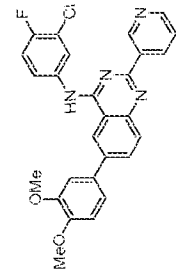
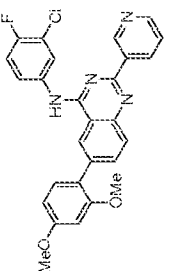
Table 4:

PRODUCT	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	Retention Time (min)	LCMS Protocol	Purity percent	Method Coupling
	2 HCl	536.401	¹ H NMR (300 MHz, DMSO) δ 11.34 (s, 1H), 10.53 (s, 1H), 9.56 (s, 1H), 9.17 - 8.95 (m, 2H), 8.89 (d, J = 4.5 Hz, 1H), 8.19 (d, J = 8.8 Hz, 1H), 8.08 - 7.81 (m, 4H), 7.62 - 6.98 (m, 3H), 4.53 (s, 2H), 4.05 - 3.74 (m, 4H), 3.48 - 3.07 (m, 4H).	DMSO	464 (M+1)	2.01	Method C	100	Method N3
	HCl	515.406	¹ H NMR (300 MHz, DMSO) δ 10.40 (s, 1H), 9.51 (s, 1H), 9.00 - 8.78 (m, 2H), 8.55 (s, 1H), 8.20 (d, J = 4.4 Hz, 1H), 8.03 - 7.75 (m, 4H), 7.54 (t, J = 9.1 Hz, 1H), 4.68 (s, 2H), 3.83 (d, J = 11.2 Hz, 2H), 3.44 - 3.18 (m, 4H), 1.98 - 1.78 (m, 1H), 1.62 (d, J = 1.62 Hz, 1H).	DMSO	479 (M+1)	2.08	Method D	97	Method N3
		448.920	¹ H NMR (300 MHz, DMSO) δ 10.10 (s, 1H), 9.51 (s, 1H), 8.75 - 8.60 (m, 2H), 8.47 (s, 1H), 8.32 - 8.19 (m, 1H), 7.97 - 7.77 (m, 3H), 7.63 - 7.44 (m, 2H), 4.62 (s, 2H), 4.16 - 3.97 (m, 1H), 1.88 - 1.42 (m, 8H).	DMSO	449 (M+1)	2.62	Method D	92	Method N3
		457.89	¹ H NMR (300 MHz, DMSO) δ 10.12 (s, 1H), 9.53 (s, 1H), 8.82 (s, 1H), 8.70 (dd, J = 16.9, 5.4 Hz, 3H), 8.25 (d, J = 8.6 Hz, 3H), 8.01 - 7.87 (m, 2H), 7.57 (dd, J = 11.2, 6.0 Hz, 2H), 7.03 (d, J = 8.6 Hz, 1H), 3.94 (s, 3H).	DMSO	458.2 (M+1)	2.56	Method C	100	Method N1
		449.91	¹ H NMR (300 MHz, DMSO) δ 10.13 (s, 1H), 9.52 (s, 1H), 8.67 (d, J = 9.8 Hz, 2H), 8.47 (s, 1H), 8.34 - 8.20 (m, 1H), 8.02 - 7.84 (m, 3H), 7.54 (t, J = 9.2 Hz, 2H), 3.64 (d, J = 17.6 Hz, 6H), 2.43 (s, 4H).	DMSO	449.9 (M+1)	2.13	Method C	100	Method N3

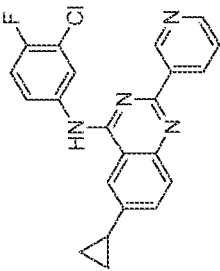
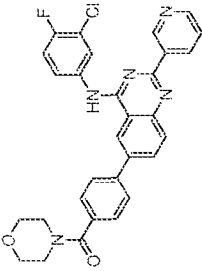
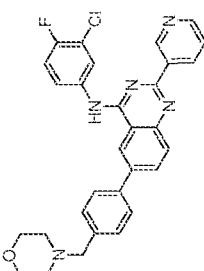
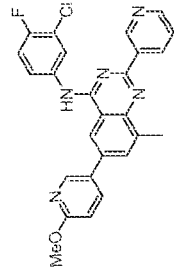
(continued)

PRODUCT	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	Retention Time (min)	LCMS Protocol	Purity percent	Method Coupling
		456.9		DMSO	456.0 (M+1)	2.64	Method C	100	Method N1
		445.88	¹ H NMR (300 MHz, DMSO) δ 10.04 (s, 1H), 9.52 (s, 1H), 8.68 (t, J = 6.5 Hz, 2H), 8.51 (s, 1H), 8.22 (dd, J = 6.7, 2.4 Hz, 1H), 8.03 - 7.83 (m, 3H), 7.54 (dd, J = 11.9, 6.6 Hz, 2H), 2.50 (s, 3H), 2.33 (s, 3H).	DMSO	446 (M+1)	2.35	Method C	100	Method N1
		427.86	¹ H NMR (300 MHz, DMSO) δ 10.10 (s, 1H), 9.51 (s, 1H), 9.13 (s, 1H), 8.86 (s, 1H), 8.66 (s, 3H), 8.24 (s, 3H), 7.94 (d, J = 8.7 Hz, 2H), 7.55 (dd, J = 16.7, 9.9 Hz, 3H).	DMSO	427.95 (M+1)	2.25	Method C	100	Method N1
		430.86	¹ H NMR (300 MHz, DMSO) δ 9.98 (s, 1H), 9.51 (d, J = 1.6 Hz, 1H), 8.66 (dd, J = 11.7, 2.2 Hz, 3H), 8.38 - 8.20 (m, 2H), 8.20 - 8.02 (m, 2H), 8.02 - 7.80 (m, 2H), 7.71 - 7.42 (m, 2H), 3.93 (s, 3H).	DMSO	430.9 (M+1)	2.17	Method C	100	Method N1
		449.91	¹ H NMR (300 MHz, DMSO) δ 9.99 (s, 1H), 9.52 (d, J = 1.3 Hz, 1H), 8.77 - 8.55 (m, 2H), 8.41 (s, 1H), 8.28 (dd, J = 6.9, 2.6 Hz, 1H), 8.03 - 7.77 (m, 3H), 7.65 - 7.44 (m, 2H), 3.34 (s, 8H), 3.15-2.90 (m, 5H), 2.90 - 2.64 (m, 5H).	DMSO	449.9 (M+1)	2.04	Method C	100	Method N1

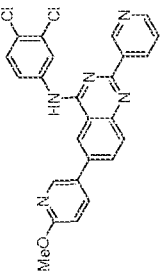
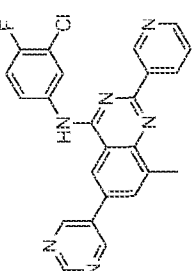
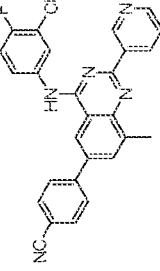
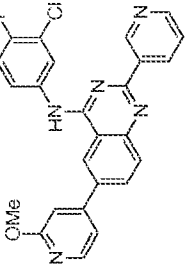
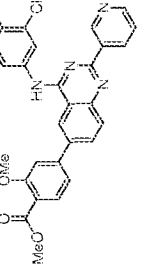
(continued)

PRODUCT	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	Retention Time (min)	LCMS Protocol	Purity percent	Method Coupling
		445.85	¹ H NMR (300 MHz, DMSO) δ 10.09 (s, 1H), 9.50 (s, 1H), 8.88 - 8.74 (m, 2H), 8.66 (dd, J = 14.1, 5.5 Hz, 2H), 8.56-8.42 (m, 1H), 8.23 (dd, J = 10.0, 5.7 Hz, 2H), 7.98 - 7.85 (m, 2H), 7.62 - 7.46 (m, 2H), 7.45 - 7.36 (m, 1H).	DMSO	445.9 (M+1)	2.43	Method C	99	Method N1
		512.97	¹ H NMR (300 MHz, DMSO) δ 10.05 (s, 1H), 9.51 (s, 1H), 8.70 (dd, J = 24.1, 7.0 Hz, 4H), 8.22 (dd, J = 14.3, 7.3 Hz, 2H), 8.10 (d, J = 7.9 Hz, 1H), 7.89 (d, J = 8.4 Hz, 2H), 7.53 (d, J = 7.4 Hz, 2H), 7.00 (d, J = 8.8 Hz, 1H), 3.74 (s, 4H), 3.55 (d, J = 4.7	DMSO	512.9 (M+1)	2.5	Method C	100	Method N1
		456.9	¹ H NMR (300 MHz, DMSO) δ 10.16 (s, 1H), 9.53 (s, 1H), 8.81 (s, 1H), 8.73 - 8.60 (m, 2H), 8.30 - 8.20 (m, 2H), 7.99 - 7.89 (m, 2H), 7.55 (dd, J = 8.2, 4.7 Hz, 2H), 7.46 (t, J = 7.6 Hz, 3H), 7.10 - 6.96 (m, 1H), 3.88 (s, 3H).	DMSO	458.9 (M+2)	2.65	Method C	100	Method N1
		486.92	¹ H NMR (300 MHz, DMSO) δ 10.16 (s, 1H), 9.54 (s, 1H), 8.79 (d, J = 9.7 Hz, 1H), 8.71 (s, 2H), 8.26 (d, J = 8.7 Hz, 1H), 7.95 (dd, J = 8.7, 3.7 Hz, 2H), 7.56 (dd, J = 11.7, 6.4 Hz, 1H), 7.43 (d, J = 15.2 Hz, 2H), 7.22 - 7.06 (m, 2H), 3.92 (s, 3H), 3.84 (s,	DMSO	486.9 (M+1)	2.48	Method C	100	Method N1
		486.92		DMSO	487.1 (M+1)	2.58	Method C	100	Method N1

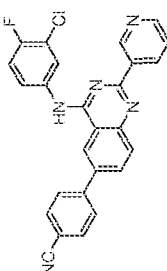
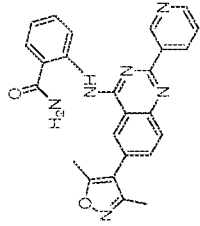
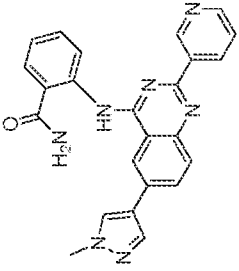
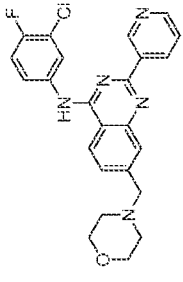
(continued)

PRODUCT	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	Retention Time (min)	LCMS Protocol	Purity percent	Method Coupling
		390.84		DMSO	390.9 (M+1)	2.48	Method C	100	Method N1
		539.99	¹ H NMR (300 MHz, DMSO) δ 10.15 (s, 1H), 9.51 (s, 1H), 8.84 (s, 1H), 8.66 (dd, J = 9.7, 6.5 Hz, 2H), 8.29 - 8.19 (m, 2H), 7.92 (ddd, J = 8.9, 7.5, 4.6 Hz, 4H), 7.65 - 7.48 (m, 4H), 3.64 (s, 6H), 3.35 (s, 2H)	DMSO	539.9 (M+1)	2.23	Method C	100	Method N1
		526			526.16 (M+1)	2.55	Method C	94	Method N1
		471.91	¹ H NMR (300 MHz, DMSO) δ 10.03 (s, 1H), 9.56 (d, J = 2.1 Hz, 1H), 8.77 - 8.66 (m, 3H), 8.64 (s, 1H), 8.24 (ddd, J = 8.7, 6.6, 2.6 Hz, 2H), 8.13 (s, 1H), 7.96 - 7.90 (m, 1H), 7.62 - 7.52 (m, 2H), 7.01 (d, J = 8.7 Hz, 1H), 3.94 (s, 3H), 2.78 (s, 3H)	CDCl ₃	471.9 (M+1)	2.93	Method C	100	Method N1

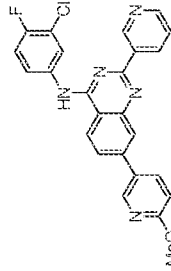
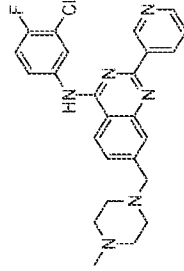
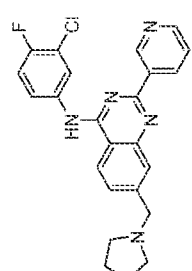
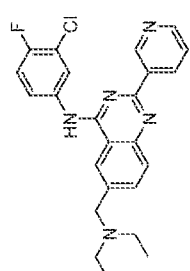
(continued)

PRODUCT	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	Retention Time (min)	LCMS Protocol	Purity percent	Method Coupling
		474.34	¹ H NMR (300 MHz, DMSO) δ 10.09 (s, 1H), 9.53 (s, 1H), 8.77 (s, 1H), 8.67 (dd, J = 12.3, 4.1 Hz, 3H), 8.35 (d, J = 2.3 Hz, 1H), 8.27 - 8.13 (m, 2H), 8.02 - 7.88 (m, 2H), 7.72 (d, J = 8.8 Hz, 1H), 7.55 (dd, J = 7.7, 4.6 Hz, 1H), 7.00 (d, J = 8.6 Hz, 1H), 3.	DMSO	474.02 (M+1)	2.72	Method C	100	Method N1
		442.88		DMSO	443.1 (M+1)	2.37	Method C	100	Method N1
		465.91	¹ H NMR (300 MHz, DMSO) δ 10.12 (s, 1H), 9.54 (s, 1H), 8.75 - 8.63 (m, 3H), 8.20 (dd, J = 8.2, 5.6 Hz, 2H), 8.09 (d, J = 8.5 Hz, 2H), 8.01 (d, J = 8.5 Hz, 2H), 7.88 (s, 1H), 7.55 (d, J = 9.1 Hz, 2H), 2.76 (s, 3H).	DMSO	466.1 (M+1)	2.78	Method C	100	Method N1
		457.89	¹ H NMR (300 MHz, DMSO) d 10.18 (s, 1H), 9.51 (s, 1H), 8.92 (s, 1H), 8.67 (dd, J = 12.7, 5.8 Hz, 2H), 8.39 - 8.16 (m, 3H), 7.91 (dd, J = 14.8, 5.7 Hz, 2H), 7.65 - 7.45 (m, 3H), 7.35 (s, 1H), 3.94 (s, 3H).	DMSO	458.1 (M+1)	2.62	Method C	100	Method N1
		514.93	¹ H NMR (300 MHz, DMSO) d 10.19 (s, 1H), 9.52 (s, 1H), 8.86 (s, 1H), 8.67 (dd, J = 10.1, 3.4 Hz, 2H), 8.28 (ddd, J = 9.4, 7.7, 2.1 Hz, 2H), 8.06 - 7.75 (m, 2H), 7.54 (dd, J = 9.0, 4.1 Hz, 4H), 3.99 (s, 3H), 3.82 (s, 3H).	DMSO	515.1 (M+1)	2.54	Method C	100	Method N1

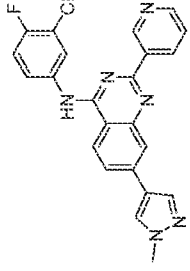
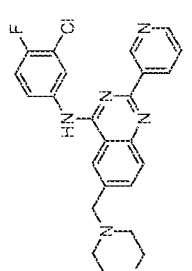
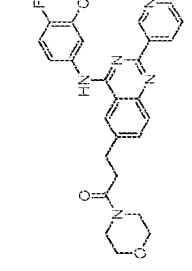
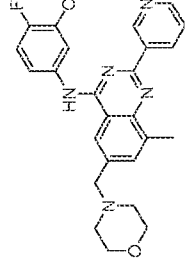
(continued)

PRODUCT	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	Retention Time (min)	LCMS Protocol	Purity percent	Method Coupling
		451.88	¹ H NMR (300 MHz, DMSO) δ 10.16 (s, 1H), 9.50 (s, 1H), 8.86 (s, 1H), 8.66 (dd, J = 15.2, 6.9 Hz, 2H), 8.35 - 8.16 (m, 2H), 8.15 - 7.82 (m, 6H), 7.55 (dd, J = 11.1, 6.0 Hz, 2H).	DMSO	452.0 (M+1)	2.61	Method C	96	Method N1
		436.47			437.2 (M+1)	2.08	Method C	95	Method N1
		421.45			422.2 (M+1)	1.8	Method C	100	Method N1
		449.91	¹ H NMR (300 MHz, DMSO) δ 10.06 (s, 1H), 9.52 (d, J = 1.3 Hz, 1H), 8.73 - 8.64 (m, 2H), 8.51 (d, J = 8.5 Hz, 1H), 8.27 J = 6.9, 2.6 Hz, 1H), 7.96 - 7.86 (m, 1H), 7.82 (s, 1H), 7.64 (d, J = 8.4 Hz, 1H), 7.57 (t, J = 3.9 Hz, 1H), 7.52 (t, J = 9.1 Hz, 1H), 3.69 (s, 2H), 3.62 (s, 4H), 2.44 (s, 4H).	(dd, DMSO)	449.9 (M+1)	2.13	Method C	100	Method N3

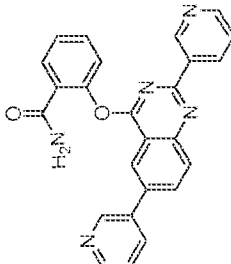
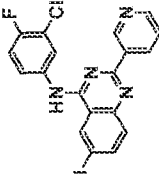
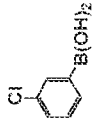
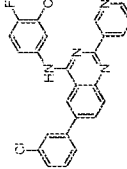
(continued)

PRODUCT	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	Retention Time (min)	LCMS Protocol	Purity percent	Method Coupling
		457.89	¹ H NMR (300 MHz, DMSO) δ 10.12 (s, 1H), 9.55 (s, 1H), 8.76 (d, J = 2.5 Hz, 1H), 8.72 - 8.65 (m, 2H), 8.62 (d, J = 8.6 Hz, 1H), 8.34 - 8.26 (m, 2H), 8.15 (d, J = 1.6 Hz, 1H), 8.02 (d, J = 8.7 Hz, 1H), 7.97 - 7.88 (m, 1H), 7.61 - 7.49 (m, 2H), 6.99 (d, J = 8.7 Hz, 1H), 3.94 (s, 3H).	DMSO	458.1 (M+1)	2.51	Method C	100	Method N2
		462.95	¹ H NMR (300 MHz, CD ₃ OD) δ 9.40 (s, 1H), 8.66 (d, J = 8.0 Hz, 1H), 8.56 (s, 1H), 8.19 - 8.09 (m, 2H), 7.74 (s, 1H), 7.72 - 7.65 (m, 1H), 7.54 - 7.45 (m, 2H), 7.24 (t, J = 9.0 Hz, 1H), 3.68 (d, J = 8.0 Hz, 2H), 2.75 - 2.41 (m, 8H), 2.30 (s, 3H).	CD ₃ OD	463.0 (M+1)	1.71	Method C	100	Method N3
		433.91			434.0 (M+1)	1.73	Method C	100	Method N3
		435.92			436.0 (M+1)	1.84	Method C	100	Method N3

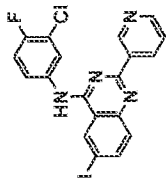
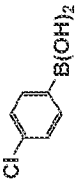
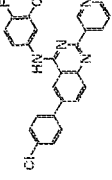
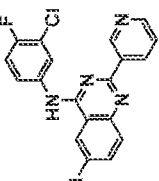
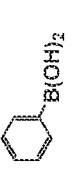
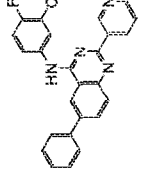
(continued)

PRODUCT	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	Retention Time (min)	LCMS Protocol	Purity percent	Method Coupling
		430.865	¹ H NMR (300 MHz, DMSO) δ 10.01 (s, 1H), 9.52 (d, J = 1.3 Hz, 1H), 8.72 - 8.60 (m, 2H), 8.49 (d, J = 8.6 Hz, 1H), 8.44 (s, 1H), 8.28 (dd, J = 6.9, 2.6 Hz, 1H), 8.15 (s, 1H), 8.03 (d, J = 1.5 Hz, 1H), 7.95-7.83 (m, 2H), 7.60-7.45 (m, 2H), 3.91 (s, 3H).	DMSO	431.0 (M+1)	2.12	Method C	100	Method N2
		465.97	¹ H NMR (300 MHz, CD ₃ OD) δ 9.56-9.50 (m, 1 H), 8.79 (dt, J = 8.1, 1.6 Hz, 1H), 8.63 (dd, J = 4.9, 1.7 Hz, 1H), 8.30 (s, 1H), 8.19 (dd, J = 6.7, 2.6 Hz, 1H), 7.91 (d, J = 1.1 Hz, 2H), 7.80 (ddd, J = 9.0, 4.2, 2.7 Hz, 1H), 7.58 (dd, J = 8.0, 4.9 Hz, 1H), 7.33 (t, J = 9.0 Hz, 1H), 3.75 (s, 2H), 2.84 - 2.66 (m, 8H).	CD ₃ OD	466.0 (M+1)	2.46	Method C	94	Method N3
		491.94	¹ H NMR (300 MHz, DMSO) δ 10.00 (s, 1H), 9.52 (s, 1H), 8.73 - 8.63 (m, 2H), 8.42 (s, 1H), 8.28 (dd, J = 6.8, 2.6 Hz, 1H), 7.92 (ddd, J = 6.8, 4.2, 2.0 Hz, 1H), 7.85 (s, 2H), 7.60 - 7.50 (m, 2H), 3.56-3.43 (m, 9H), 3.07 (t, J = 7.6 Hz, 2H), 2.79 (t, J = 7.7 Hz, 2H).	DMSO	491.9 (M+1)	2	Method C	93	Method N1
		463.93	¹ H NMR (300 MHz, DMSO) δ 9.99 (s, 1H), 9.54 (s, 1H), 8.67 (d, J = 5.3 Hz, 2H), 8.30 - 8.22 (m, 2H), 7.96 - 7.87 (m, 1H), 7.73 (s, 1H), 7.60 - 7.46 (m, 2H), 3.60 (s, 6H), 2.71 (s, 3H), 2.41 (s, 4H).	DMSO	464.0 (M+1)	2.52	Method C	100	Method N3

(continued)

PRODUCT	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solent	LCMS	Reten- tion Time (min)	LCMS Pro- tocol	Purity percent	Method Coupling
		419.43	¹ H NMR (300 MHz, DMSO) δ 9.41 (d, J = 1.4 Hz, 1H), 9.15 (d, J = 1.7 Hz, 1H), 8.74 - 8.64 (m, 3H), 8.55 (dt, J = 7.9, 1.8 Hz, 1H), 8.42 (dd, J = 8.7, 2.0 Hz, 1H), 8.36-8.29 (m, 1H), 8.15 (d, J = 8.7 Hz, 1H), 7.90 (dd, J = 7.8, 1.7 Hz, 1H), 7.61 (dd, J = 8.0, 4.8 Hz, 1H), 7.53 (dd, J = 7.9, 4.8 Hz, 1H), 7.43 (t, J = 7.6 Hz, 1H), 7.03 (t, J = 7.5 Hz, 1H), 6.93 (d, J = 7.7 Hz, 1H).	DMSO	420.1 (M+1)	2.12	Method C	99	Method N4
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Per- cent	Method Coupling	
629					¹ H NMR (400 MHz, DMSO) δ 10.20 (s, 1H), 9.54 (s, 1H), 8.87 (s, 1H), 8.74-8.65 (m, 2H), 8.31 - 8.23 (m, 2H), 8.04 - 7.85 (m, 4H), 7.64 - 7.49 (m, 4H).	DMSO	>98	N5	

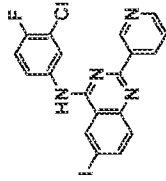
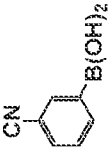
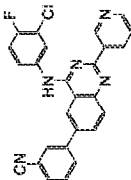
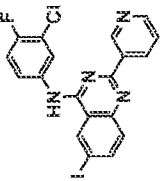
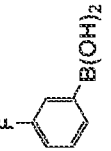
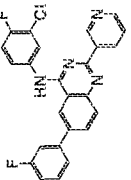
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
630					¹ H NMR (400 MHz, DMSO) δ 10.20 (s, 1H), 9.54 (d, J = 1.4 Hz, 1H), 8.86 (s, 1H), 8.76 - 8.64 (m, 2H), 8.26 (dd, J = 6.8, 2.3 Hz, 2H), 8.03 - 7.89 (m, 4H), 7.70 - 7.52 (m, 4H).	DMSO	>98	N5
631					¹ H NMR (400 MHz, DMSO) δ 10.20 (s, 1H), 9.54 (d, J = 1.4 Hz, 1H), 8.86 (s, 1H), 8.76 - 8.64 (m, 2H), 8.26 (dd, J = 6.8, 2.3 Hz, 2H), 8.03 - 7.89 (m, 4H), 7.70 - 7.52 (m, 4H).	DMSO	>98	N5

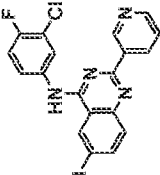
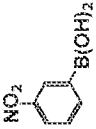
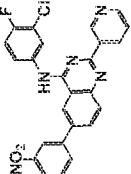
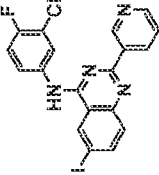
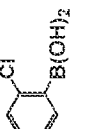
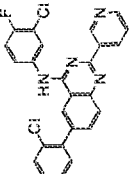
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
632					¹ H NMR (400 MHz, DMSO) δ 10.22 - 10.11 (m, 1H), 9.54 (s, 1H), 8.86 - 8.77 (m, 1H), 8.74 - 8.65 (m, 2H), 8.30 - 8.17 (m, 2H), 8.01 - 7.90 (m, 2H), 7.74 - 7.66 (m, 2H), 7.60 - 7.51 (m, 2H), 7.51 - 7.42 (m, 1H), 7.28 (d, J=7.5 Hz, 1H), 2.45 (s, 3H).	DMSO	>98	N5

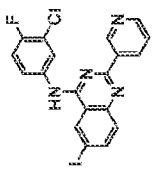
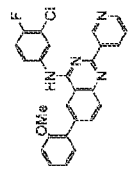
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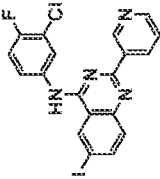
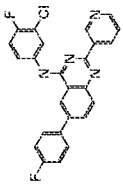
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Per-cent	Method Coupling
633					¹ H NMR (400 MHz, DMSO) δ 10.20 (s, 1H), 9.61 - 9.53 (m, 1H), 8.92 (s, 1H), 8.78 - 8.67 (m, 2H), 8.43 - 8.24 (m, 3H), 8.14 - 7.89 (m, 3H), 7.84 - 7.76 (m, 1H), 7.66 - 7.53 (m, 2H).	DMSO	>98	N5
634					¹ H NMR (400 MHz, DMSO) δ 10.20 (s, 1H), 9.55 (d, J = 1.2 Hz, 1H), 8.89 (s, 1H), 8.75 - 8.67 (m, 2H), 8.35 - 8.22 (m, 2H), 8.00 (d, J = 8.7 Hz, 1H), 7.96 - 7.90 (m, 1H), 7.84 - 7.75 (m, 2H), 7.66 - 7.52 (m, 3H), 7.34 - 7.27 (m, 1H).	DMSO	>98	N5

(continued)

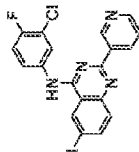
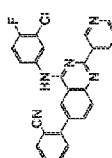
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
635					¹ H NMR (400 MHz, DMSO) δ 10.27 (s, 1H), 9.54 (d, J = 1.2 Hz, 1H), 8.94 (s, 1H), 8.77 - 8.66 (m, 3H), 8.40 - 8.21 (m, 4H), 7.99 (d, J = 8.9 Hz, 1H), 7.88 (t, J = 8.0 Hz, 2H), 7.61 - 7.50 (m, 2H).	DMSO	>98	N5
636					¹ H NMR (400 MHz, DMSO) δ 10.13 (s, 1H), 9.56 (s, 1H), 8.77 - 8.68 (m, 2H), 8.65 (s, 1H), 8.27 (dd, J = 6.9, 2.6 Hz, 1H), 8.04 - 7.97 (m, 2H), 7.96 - 7.88 (m, 1H), 7.70 - 7.43 (m, 6H).	DMSO	>98	N5

(continued)

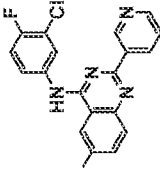
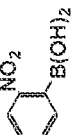
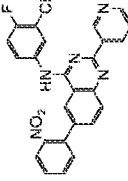
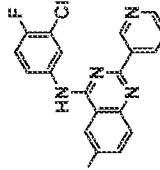
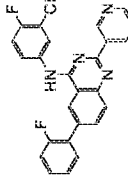
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
637					¹ H NMR (400 MHz, DMSO) δ 10.10 (s, 1H), 9.56 (s, 1H), 8.76 - 8.68 (m, 2H), 8.63 (s, 1H), 8.28 (dd, J = 6.7, 2.7 Hz, 1H), 8.05 (dd, J = 8.7, 1.8 Hz, 1H), 8.00 - 7.90 (m, 2H), 7.70 - 7.43 (m, 4H), 7.21 (d, J = 8.1 Hz, 1H), 7.14 (t, J = 7.4 Hz, 1H), 3.82 (s, 3H).	DMSO	>98	N5

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
638					¹ H NMR (400 MHz, DMSO) δ 10.16 (s, 1H), 9.54 (s, 1H), 8.81 (s, 1H), 8.74–1H), 8.74–8.64 (m, 2H), 8.30–8.17 (m, 2H), 8.01–7.87 (m, 4H), 7.61–7.51 (m, 2H), 7.42 (t, J = 8.8 Hz, 2H).	DMSO	>98	N5

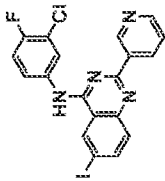
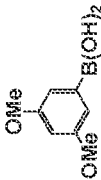
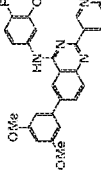
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
639					1H NMR (400 MHz, DMSO) δ 10.21 (s, 1H), 9.57 (s, 1H), 8.80 (d, J = 1.8 Hz, 1H), 8.76 - 8.69 (m, 2H), 8.27 (dd, J = 6.9, 2.6 Hz, 1H), 8.14 (dd, J = 8.6, 1.8 Hz, 1H), 8.10 - 8.02 (m, 2H), 7.97 - 7.88 (m, 2H), 7.83 (d, J = 6.8 Hz, 1H), 7.70 (td, J = 7.6, 1.2 Hz, 1H), 7.63 - 7.52 (m, 2H).	DMSO	>98	N5

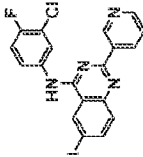
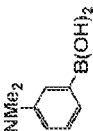
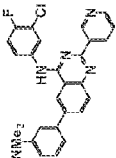
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
640					¹ H NMR (400 MHz, DMSO) δ 10.12 (s, 1H), 9.55 (s, 1H), 8.77 - 8.66 (m, 2H), 8.62 (s, 1H), 8.25 (dd, J = 6.8, 2.3 Hz, 1H), 8.16 (d, J = 8.3 Hz, 1H), 7.99 - 7.80 (m, 4H), 7.80 - 7.71 (m, 2H), 7.61 - 7.49 (m, 2H).	DMSO	>98	N5
641					¹ H NMR (400 MHz, DMSO) δ 9.57 (s, 1H), 8.85-8.55 (m, 3H), 8.20-7.95 (m, 3H), 7.85-7.20 (m, 7H).	DMSO	>98	N5

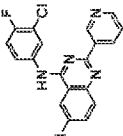
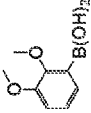
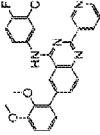
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
642					¹ H NMR (400 MHz, DMSO) δ 10.19 (s, 1H), 9.55 (d, J = 1.2 Hz, 1H), 8.83 (s, 1H), 8.74 - 8.66 (m, 2H), 8.32 - 8.23 (m, 2H), 8.02 - 7.89 (m, 2H), 7.61-7.53 (m, 2H), 7.05 (d, J = 2.2 Hz, 2H), 6.62 (t, J = 2.1 Hz, 1H), 3.88 (s, 6H).	DMSO	>98	N5

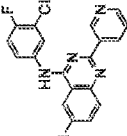
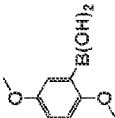
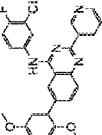
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
643					¹ H NMR (400 MHz, DMSO) δ 10.18 (s, 1H), 9.55 (d, J = 2.0 Hz, 1H), 8.81 (d, J = 1.7 Hz, 1H), 8.74 - 8.65 (m, 2H), 8.31 - 8.20 (m, 2H), 8.04 - 7.88 (m, 2H), 7.65 - 7.53 (m, 2H), 7.42 - 7.34 (m, 1H), 7.23 - 7.09 (m, 2H), 6.83 (dd, J = 8.3, 1.9 Hz, 1H), 3.02 (s, 6H).	DMSO	>98	N5

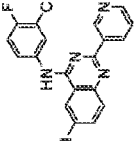
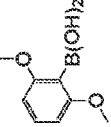
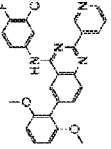
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
644					¹ H NMR (400 MHz, DMSO) δ 10.14 (s, 1H), 9.55 (s, 1H), 8.77 - 8.68 (m, 2H), 8.65 (d, J = 1.5 Hz, 1H), 8.28 (dd, J = 6.9, 2.6 Hz, 1H), 8.05 (dd, J = 8.6, 1.7 Hz, 1H), 8.01 - 7.90 (m, 2H), 7.60 - 7.51 (m, 2H), 7.29 - 7.21 (m, 1H), 7.18 (dd, J = 8.3, 1.6 Hz, 1H), 7.12 (dd, J = 7.6, 1.6 Hz, 1H), 3.89 (s, 3H), 3.58 (s, 3H).	DMSO	>98	N5

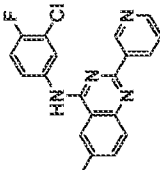
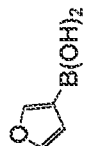
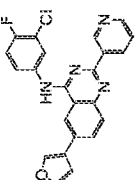
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
645					¹ H NMR (400 MHz, DMSO) δ 10.09 (s, 1H), 9.56 (s, 1H), 8.74 - 8.67 (m, 2H), 8.64 (s, 1H), 8.28 (dd, J = 6.9, 2.6 Hz, 1H), 8.06 (dd, J = 8.6, 1.7 Hz, 1H), 7.98 - 7.89 (m, 2H), 7.82 - 7.51 (m, 2H), 7.14 (d, J = 9.0 Hz, 1H), 7.09 (d, J = 3.1 Hz, 1H), 7.02 (dd, J = 8.9, 3.1 Hz, 1H), 3.81 (s, 3H), 3.76 (s, 3H).	DMSO	>98	N5

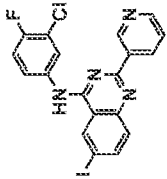
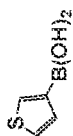
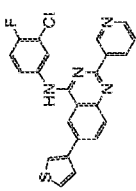
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
646					¹ H NMR (400 MHz, DMSO) δ 9.99 (s, 1H), 9.56 (d, J = 0.9 Hz, 1H), 8.71 (dd, J = 7.1, 1.6 Hz, 2H), 8.46 (d, J = 1.5 Hz, 1H), 8.29 (dd, J = 6.9, 2.6 Hz, 1H), 8.00 - 7.87 (m, 2H), 7.73 (dd, J = 8.5, 1.7 Hz, 1H), 7.63 - 7.47 (m, 2H), 7.41 (t, 3 = 8.4 Hz, 1H), 6.84 (d, J = 8.5 Hz, 2H), 3.77 - 3.66 (m, 6H).	DMSO	>98	N5

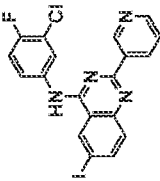
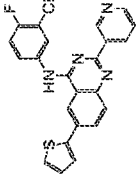
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
647					¹ H NMR (400 MHz, DMSO) δ 10.06 - 9.96 (m, 1H), 9.52 (s, 1H), 8.75 - 8.62 (m, 3H), 8.40 - 8.33 (m, 1H), 8.28 - 8.22 (m, 1H), 8.22 - 8.14 (m, 1H), 7.97 - 7.85 (m, 3H), 7.64 - 7.49 (m, 2H), 7.16 (s, 1H).	DMSO	>98	N5

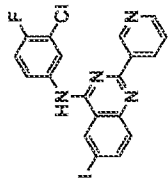
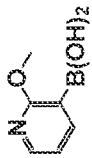
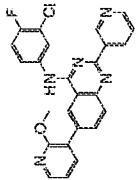
(continued)

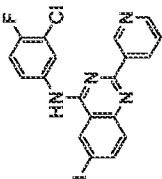
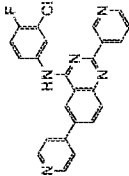
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
648					¹ H NMR (400 MHz, DMSO) δ 10.10 (s, 1H), 9.54 (s, 1H), 8.85 (s, 1H), 8.77 - 8.65 (m, 2H), 8.32 (d, J = 8.7 Hz, 1H), 8.26 (dd, J = 6.8, 2.6 Hz, 1H), 8.12 (s, 1H), 7.94 (d, J = 8.7 Hz, 2H), 7.86 - 7.76 (m, 2H), 7.64 - 7.54 (m, 2H).	DMSO	>98	N5

(continued)

5							
10							
15							
20							
25							
30							
35							
40							
45							
50							
55							
	649						
		Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent
						¹ H NMR (400 MHz, DMSO) δ 10.20 (s, 1H), 9.53 (s, 1H), 8.78 (s, 1H), 8.71 - 8.64 (m, 2H), 8.26 - 8.19 (m, 2H), 7.95 - 7.90 (m, 2H), 7.77 - 7.69 (m, 2H), 7.60 - 7.53 (m, 2H), 7.26 (t, J = 3.7 Hz, 1H).	DMSO
							>98
							N5

(continued)

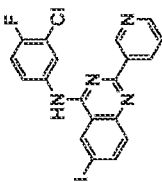
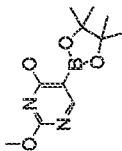
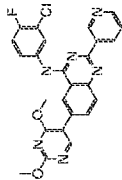
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
650					¹ H NMR (400 MHz, DMSO) δ 10.12 (s, 1H), 9.55 (d, J = 2.2 Hz, 1H), 8.75 - 8.67 (m, 3H), 8.33 - 8.23 (m, 2H), 8.13 (dd, J = 8.7, 1.8 Hz, 1H), 8.00 - 7.90 (m, 3H), 7.63 - 7.51 (m, 2H), 7.22 (dd, J = 7.3, 5.0 Hz, 1H), 3.95 (s, 3H).	DMSO	>98	N5

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
651					¹ H NMR (400 MHz, DMSO) δ 10.22 (s, 1H), 9.53 (s, 1H), 8.97 (s, 1H), 8.75 (d, J = 5.5 Hz, 2H), 8.73 - 8.64 (m, 2H), 8.38 - 8.30 (m, 1H), 8.24 (d, J = 6.8 Hz, 1H), 8.05 - 7.97 (m, 1H), 7.97 (m, 1H), 7.97 - 7.88 (m, 3H), 7.63 - 7.51 (m, 2H).	>98	N5

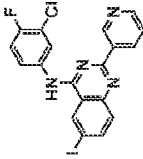
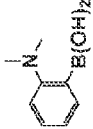
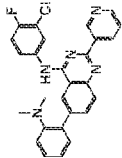
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
652					¹ H NMR (400 MHz, DMSO) δ 10.05 (s, 1H), 9.50 (s, 1H), 9.33 (s, 2H), 9.26 (s, 1H), 8.90 (s, 1H), 8.69 (d, J = 3.6 Hz, 1H), 8.64 (d, J = 8.0 Hz, 1H), 8.33 (d, J = 8.6 Hz, 1H), 8.22 (dd, J = 6.7, 2.0 Hz, 1H), 7.96 (d, J = 8.7 Hz, 1H), 7.93 - 7.86 (m, 1H), 7.61 - 7.48 (m, 2H).	DMSO	>98	N5

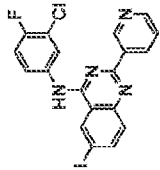
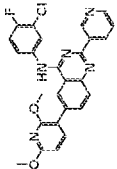
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Per-cent	Method Coupling
653					¹ H NMR (400 MHz, DMSO) δ 10.10 (s, 1H), 9.55 (d, J = 1.2 Hz, 1H), 8.77 - 8.67 (m, 2H), 8.61 (s, 1H), 8.26 (dd, J = 6.9, 2.6 Hz, 1H), 8.12 (dd, J = 8.7, 1.8 Hz, 1H), 8.01 - 7.89 (m, 3H), 7.62 - 7.51 (m, 2H), 4.02 (s, 3H), 4.00 (s, 3H).	DMSO	>98	N5

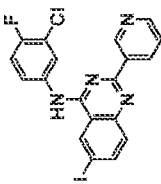
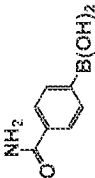
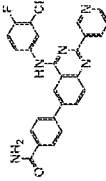
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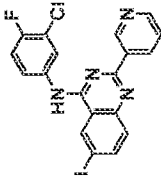
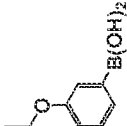
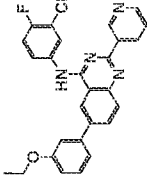
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
654					¹ H NMR (400 MHz, DMSO) δ 10.09 (s, 1H), 9.56 (s, 1H), 8.77 - 8.68 (m, 2H), 8.65 (d, J = 1.6 Hz, 1H), 8.30 (dd, J = 6.9, 2.6 Hz, 1H), 8.17 (dd, J = 8.7, 1.7 Hz, 1H), 7.99 - 7.88 (m, 2H), 7.61 - 7.50 (m, 2H), 7.42 - 7.33 (m, 2H), 7.21 - 7.08 (m, 2H), 2.53 (s, 6H).	DMSO	>98	N5

(continued)

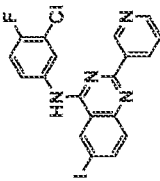
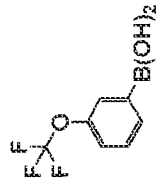
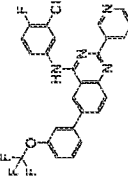
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
655					¹ H NMR (400 MHz, DMSO) δ 10.08 (s, 1H), 9.55 (s, 1H), 8.74 - 8.66 (m, 2H), 8.63 (s, 1H), 8.26 (dd, J = 6.8, 2.5 Hz, 1H), 8.12 (d, J = 8.7 Hz, 1H), 7.97 - 7.88 (m, 3H), 7.62 - 7.51 (m, 2H), 6.61 (d, J = 8.1 Hz, 1H), 3.97 (s, 3H), 3.96 (s, 3H).	DMSO	>98	N5

(continued)

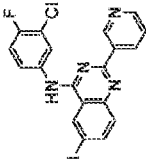
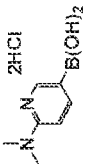
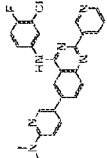
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
656					¹ H NMR (400 MHz, DMSO) δ 10.21 (s, 1H), 9.54 (d, J = 2.0 Hz, 1H), 8.91 (s, 1H), 8.76 - 8.65 (m, 2H), 8.31 (d, J = 8.8 Hz, 1H), 8.26 (dd, J = 6.8, 2.6 Hz, 1H), 8.15 - 7.89 (m, 7H), 7.61 - 7.52 (m, 2H), 7.46 (s, 1H).	>98	N5

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
657					¹ H NMR (400 MHz, DMSO) δ 10.21 (s, 1H), 9.54 (d, J = 2.0 Hz, 1H), 8.91 (s, 1H), 8.76 - 8.65 (m, 2H), 8.31 (d, J = 8.8 Hz, 1H), 8.26 (dd, J = 6.8, 2.6 Hz, 1H), 8.15 - 7.89 (m, 7H), 7.61 - 7.52 (m, 2H), 7.46 (s, 1H).	>98	N5

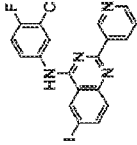
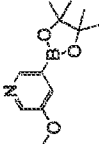
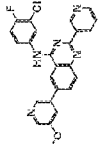
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
658					¹ H NMR (400 MHz, DMSO) δ 10.20 (s, 1H), 9.54 (s, 1H), 8.88 (s, 1H), 8.79 - 8.61 (m, 2H), 8.34 - 8.22 (m, 2H), 8.05 - 7.84 (m, 4H), 7.72 (t, J = 8.0 Hz, 1H), 7.65 - 7.53 (m, 2H), 7.47 (d, J = 8.4 Hz, 1H).	DMSO	>98	N5

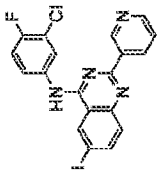
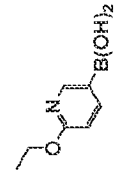
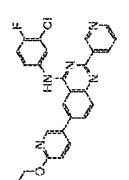
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
659					¹ H NMR (400 MHz, DMSO) δ 10.06 (s, 1H), 9.53 (s, 1H), 8.75 (s, 1H), 8.75 - 8.66 (m, 3H), 8.28 - 8.21 (m, 1H), 8.20 (dd, J = 7.7, 1.8 Hz, 1H), 8.06 (dd, J = 8.9, 2.5 Hz, 1H), 7.96 - 7.90 (m, 2H), 7.58 - 7.54 (m, 2H), 6.81 (d, J = 8.9 Hz, 1H), 3.11 (s, 6H).	DMSO	>98	N5

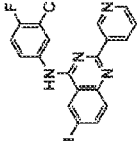
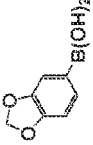
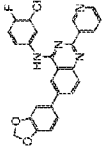
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Per-cent	Method Coupling
660					¹ H NMR (400 MHz, DMSO) δ 10.15 (s, 1H), 9.54 (s, 1H), 8.91 (s, 1H), 8.76 (s, 1H), 8.74 – 8.66 (m, 2H), 8.39 (d, J = 2.7 Hz, 1H), 8.37 – 8.30 (m, 1H), 8.26 (dd, J = 6.8, 2.6 Hz, 1H), 8.03 – 7.97 (m, 1H), 7.97 – 7.90 (m, 1H), 7.85 (d, J = 1.9 Hz, 1H), 7.62 – 7.52 (m, 2H), 3.98 (s, 3H).	DMSO	>98	N5

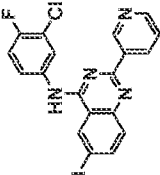
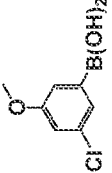
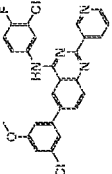
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
661					¹ H NMR (400 MHz, DMSO) δ 10.06 (s, 1H), 9.52 (d, $J = 1.3$ Hz, 1H), 8.76 (d, $J = 19.6$ Hz, 1H), 8.73 - 8.61 (m, 3H), 8.29 - 8.16 (m, 3H), 7.99 - 7.87 (m, 2H), 7.61 - 7.51 (m, 2H), 6.98 (d, $J = 8.6$ Hz, 1H), 4.39 (q, $J = 7.0$ Hz, 2H), 1.37 (t, $J = 7.0$ Hz, 3H).	DMSO	>98	N5

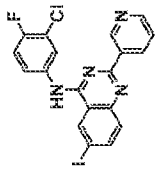
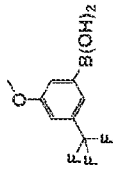
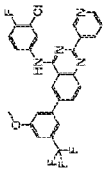
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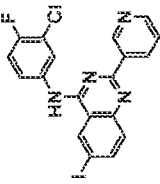
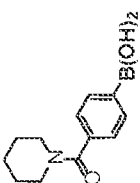
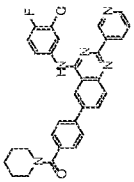
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Per-cent	Method Coupling
662					¹ H NMR (400 MHz, DMSO) δ 10.11 (s, 1H), 9.53 (dd, J=2.1, 0.7 Hz, 1H), 8.75 (d, J = 1.9 Hz, 1H), 8.73 - 8.64 (m, 2H), 8.25 (dd, J = 6.9, 2.6 Hz, 1H), 8.19 (dd, J = 8.7, 1.9 Hz, 1H), 7.97 - 7.89 (m, 2H), 7.61 - 7.50 (m, 3H), 7.41 (dd, J = 8.1, 1.9 Hz, 1H), 7.11 (d, J = 8.1 Hz, 1H), 6.13 (s, 2H).	DMSO	>98	N5

(continued)

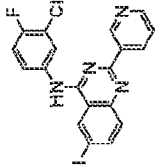
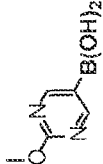
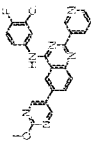
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
663					¹ H NMR (400 MHz, DMSO) δ 10.19 (s, 1H), 9.54 (s, 1H), 8.85 (s, 1H), 8.76 - 8.66 (m, 2H), 8.33 - 8.22 (m, 2H), 8.00 - 7.89 (m, 2H), 7.62 - 7.52 (m, 3H), 7.43 (s, 1H), 7.14 (s, 1H), 3.91 (s, 3H).	DMSO	>98	N5

(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
664					¹ H NMR (400 MHz, DMSO) δ 10.16 (s, 1H), 9.52 (d, J= 1.3 Hz, 1H), 8.89 - 8.81 (m, 1H), 8.73 - 8.64 (m, 2H), 8.37 - 8.28 (m, 1H), 8.28 - 8.21 (m, 1H), 8.01 - 7.86 (m, 2H), 7.81 - 7.69 (m, 2H), 7.60 - 7.51 (m, 2H), 7.33 (s, 1H), 3.97 (s, 3H).	DMSO	>98	N5

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
665					1H NMR (400 MHz, DMSO) δ 10.19 (s, 1H), 9.53 (d, J = 1.9 Hz, 1H), 8.89 (s, 1H), 8.73 - 8.61 (m, 2H), 8.57 (d, J = 4.5 Hz, 1H), 8.34 - 8.18 (m, 2H), 8.09 - 7.86 (m, 5H), 7.62 - 7.49 (m, 2H), 2.88 - 2.80 (m, 4H).	>98	N5

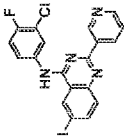
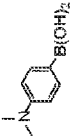
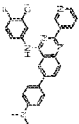
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
666				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.57 (s, 1H), 9.52 (d, J = 1.6 Hz, 1H), 9.21 (s, 2H), 9.08 - 9.02 (m, 1H), 9.02 - 8.96 (m, 1H), 8.89 (dd, J = 5.2, 1.5 Hz, 1H), 8.39 - 8.30 (m, 1H), 8.25 - 8.18 (m, 1H), 8.07 - 7.87 (m, 3H), 7.56 (t, J = 9.1 Hz, 1H).	DMSO	>98	N5

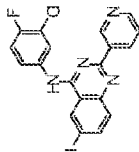
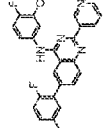
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
667				HCl	¹ H NMR (400 MHz, DMSO) δ 10.78 (s, 1H), 9.53 (s, 1H), 9.16 - 9.02 (m, 2H), 8.99 - 8.88 (m, 1H), 8.66 - 8.55 (m, 1H), 8.38 (d, J = 8.2 Hz, 1H), 8.20 (d, J = 5.0 Hz, 1H), 8.04 (dt, J = 27.3, 13.6 Hz, 7H), 7.57 (t, J = 8.8 Hz, 1H), 2.84 (d, J = 3.8 Hz, 3H).	DMSO	>98	N5

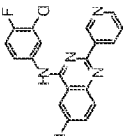
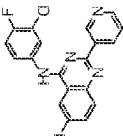
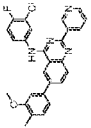
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
668				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.49 (s, 1H), 9.53 (d, J = 1.5 Hz, 1H), 8.99 – 8.83 (m, 3H), 8.34 – 8.25 (m, 1H), 8.21 (dd, J = 6.8, 2.6 Hz, 1H), 8.03 (d, J = 8.7 Hz, 1H), 7.98 – 7.91 (m, 1H), 7.86 (dd, J = 7.8, 5.2 Hz, 1H), 7.78 – 7.70 (m, 2H), 7.57 (t, J = 9.1 Hz, 1H), 7.47 (t, J = 7.6 Hz, 1H), 7.29 (d, J = 7.5 Hz, 1H), 2.45 (s, 6H).	DMSO	>98	N5

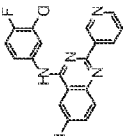
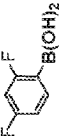
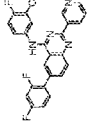
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
669				HCl	¹ H NMR (400 MHz, DMSO) δ 10.22 (s, 1H), 9.55 (d, $J = 1.4$ Hz, 1H), 8.81–8.70 (m, 3H), 8.25 (dd, $J = 6.8, 2.6$ Hz, 1H), 8.13–8.05 (m, 1H), 7.99 (d, $J = 8.7$ Hz, 1H), 7.97–7.89 (m, 1H), 7.64 (dd, $J = 7.9, 4.9$ Hz, 1H), 7.61–7.49 (m, 2H), 7.37–7.26 (m, 2H), 2.41 (s, 3H).	DMSO	>98	N5

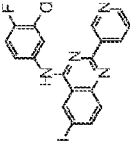
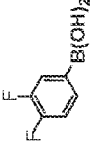
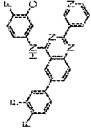
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
670				HCl	¹ H NMR (400 MHz, DMSO) δ 10.77 (s, 1H), 9.52 (d, J = 1.6 Hz, 1H), 9.09-8.95 (m, 2H), 8.90 (dd, J = 5.2, 1.5 Hz, 1H), 8.41 - 8.29 (m, 1H), 8.23 (dd, J = 6.8, 2.6 Hz, 1H), 8.05 (d, J = 8.7 Hz, 1H), 8.01 - 7.88 (m, 2H), 7.56 (t, J = 9.1 Hz, 1H), 7.50 - 7.38 (m, 2H), 7.32 (d, J = 8.2 Hz, 1H), 3.97 (s, 3H), 2.23 (s, 3H).	DMSO	>98	N5

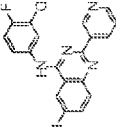
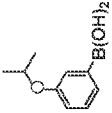
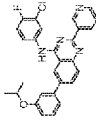
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
671				HCl	¹ H NMR (400 MHz, DMSO) δ 10.77 (s, 1H), 9.52 (d, J = 1.6 Hz, 1H), 9.09 - 8.95 (m, 2H), 8.90 (dd, J = 2.1, 1.5 Hz, 1H), 8.41 - 8.29 (m, 1H), 8.23 (dd, J = 6.8, 2.6 Hz, 1H), 8.05 (d, J = 8.7 Hz, 1H), 8.01 - 7.88 (m, 2H), 7.56 (t, J = 9.1 Hz, 1H), 7.50 - 7.38 (m, 2H), 7.32 (d, J = 8.2 Hz, 1H), 3.97 (s, 3H), 2.23 (s, 3H).	DMSO	>98	N5

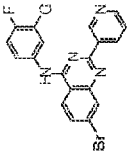
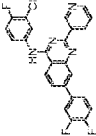
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Per-cent	Method Coupling
672				HCl	¹ H NMR (400 MHz, DMSO) δ 10.67 (s, 1H), 9.47 (d, J = 1.6 Hz, 1H), 9.04 – 8.94 (m, 2H), 8.89 (dd, J = 5.3, 1.5 Hz, 1H), 8.26 (dd, J = 8.8, 1.9 Hz, 1H), 8.18 (dd, J = 6.8, 2.6 Hz, 1H), 8.09 – 8.00 (m, 1H), 8.00 – 7.87 (m, 3H), 7.86 – 7.77 (m, 1H), 7.68 – 7.57 (m, 1H), 7.53 (t, J = 9.1 Hz, 1H).	DMSO	>98	N5

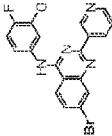
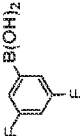
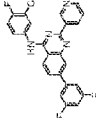
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Per-cent	Method Coupling
673				HCl	¹ H NMR (400 MHz, DMSO) δ 10.82 (s, 1H), 9.53 (s, 1H), 9.17 - 9.04 (m, 1H), 9.04 - 8.93 (m, 2H), 8.34 (dd, J = 8.7, 1.7 Hz, 1H), 8.18 (dd, J = 6.8, 2.6 Hz, 1H), 8.08 (d, J = 8.6 Hz, 1H), 8.05 - 7.89 (m, 2H), 7.57 (t, J = 9.0 Hz, 1H), 7.53 - 7.44 (m, 3H), 7.09 - 6.98 (m, 1H), 4.82 (dt, J = 11.7, 5.9 Hz, 1H), 1.33 (d, J = 6.0 Hz, 6H).	DMSO	>98	N5

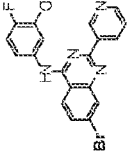
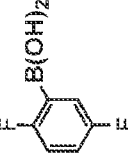
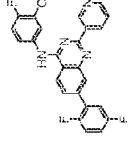
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Per-cent	Method Coupling
674				HCl	¹ H NMR (400 MHz, DMSO) δ 10.60 (s, 1H), 9.50 (d, J = 1.7 Hz, 1H), 9.11 - 9.02 (m, 1H), 8.99 - 8.91 (m, 1H), 8.76 (d, J = 8.8 Hz, 1H), 8.25 (d, J = 1.4 Hz, 1H), 8.19 (dd, J = 6.8, 2.6 Hz, 1H), 8.10 - 7.92 (m, 4H), 7.82 - 7.72 (m, 4H), 7.65 - 7.56 (m, 1H), 7.52 (t, J = 9.1 Hz, 1H).	DMSO	>98	N5

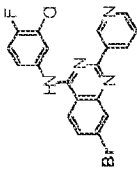
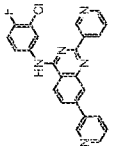
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
675				HCl	¹ H NMR (400 MHz, DMSO) δ 10.51 (s, 1H), 9.51 (d, J = 1.7 Hz, 1H), 9.07 - 9.00 (m, 1H), 8.93 (dd, J = 5.3, 1.4 Hz, 1H), 8.75 (d, J = 8.8 Hz, 1H), 8.28 (d, J = 1.7 Hz, 1H), 8.21 (dd, J = 6.8, 2.6 Hz, 1H), 8.10 (dd, J = 8.7, 1.9 Hz, 1H), 8.02 - 7.93 (m, 2H), 7.74 - 7.65 (m, 2H), 7.52 (t, J = 9.1 Hz, 1H), 7.40 - 7.31 (m, 1H).	DMSO	>98	N5

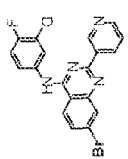
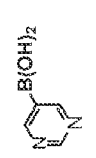
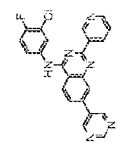
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
676				HCl	¹ H NMR (400 MHz, DMSO) δ 10.63 (s, 1H), 9.54 (d, J = 1.7 Hz, 1H), 9.10 (d, J = 8.1 Hz, 1H), 8.95 (dd, J = 5.3, 1.4 Hz, 1H), 8.81 (d, J = 8.7 Hz, 1H), 8.26 - 8.14 (m, 2H), 8.08 - 7.91 (m, 3H), 7.75 - 7.64 (m, 1H), 7.51 (ddd, J = 14.7, 14.2, 6.9 Hz, 2H), 7.44 - 7.34 (m, 1H).	DMSO	>98	N5

(continued)

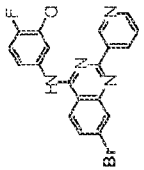
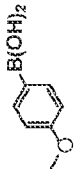
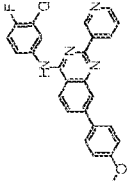
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
677				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.52 (s, 1H), 9.60 - 9.49 (m, 1H), 9.37 - 9.31 (m, 1H), 8.99 (1H), 8.95 - 8.81 (m, 3H), 8.75 (d, J = 8.3 Hz, 1H), 8.40 (1H), 8.30 - 8.18 (m, 2H), 8.04 - 7.87 (m, 3H), 7.56 (t, J = 9.1 Hz, 1H).	DMSO	>98	N5

(continued)

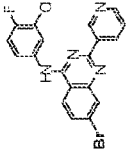
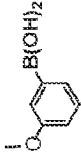
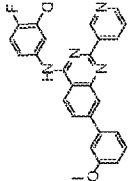
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
678				HCl	¹ H NMR (400 MHz, DMSO) δ 10.42 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 9.38 (s, 2H), 9.30 (s, 1H), 9.00 - 8.92 (m, 1H), 8.92 - 8.86 (m, 1H), 8.76 (d, J = 8.7 Hz, 1H), 8.34 (d, J = 1.7 Hz, 1H), 8.23 (dd, J = 6.8, 2.6 Hz, 1H), 8.15 (dd, J = 8.6, 1.8 Hz, 1H), 7.99 - 7.85 (m, 2H), 7.52 (t, J = 9.1 Hz, 1H).	DMSO	>98	N5

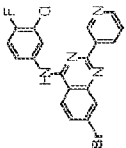
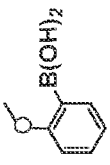
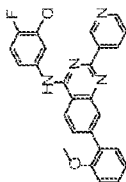
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
679				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.47 (s, 1H), 9.58 - 9.49 (m, 1H), 9.21 (s, 2H), 9.09 - 8.98 (m, 1H), 8.92 (d, J = 3.8 Hz, 1H), 8.77 (d, J = 8.8 Hz, 1H), 8.32 (d, J = 1.8 Hz, 1H), 8.27 - 8.20 (m, 1H), 8.15 (dd, J = 8.7, 1.8 Hz, 1H), 8.03 - 7.89 (m, 2H), 7.55 (t, J = 9.1 Hz, 1H), 4.02 (s, 3H).	>98	N5

(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
680					¹ H NMR (400 MHz, DMSO) δ 10.09 (s, 1H), 9.56 (d, J = 1.3 Hz, 1H), 8.75-8.67 (m, 2H), 8.60 (d, J = 8.7 Hz, 1H), 8.32 (dd, J = 6.9, 2.6 Hz, 1H), 8.10 (d, J = 1.6 Hz, 1H), 8.03 - 7.87 (m, 4H), 7.60 - 7.50 (m, 2H), 7.12 (d, J = 8.8 Hz, 2H), 3.85 (s, 3H).	DMSO	>98	N5

(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
681					¹ H NMR (400 MHz, DMSO) δ 10.05 (s, 1H), 9.55 - 9.46 (m, 1H), 8.69 - 8.60 (m, 2H), 8.56 (d, J = 8.7 Hz, 1H), 8.25 (dd, J = 6.9, 2.6 Hz, 1H), 8.10 (d, J = 1.8 Hz, 1H), 7.96 (dd, J = 8.7, 1.9 Hz, 1H), 7.91 - 7.85 (m, 1H), 7.55 - 7.35 (m, 5H), 7.03 - 6.95 (m, 1H), 3.83 (s, 3H).	DMSO	>98	N5

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
682					¹ H NMR (400 MHz, DMSO- <i>d</i> ₆) δ 10.09 (s, 1H), 9.55 (d, <i>J</i> = 1.9 Hz, 1H), 8.74–8.67 (m, 2H), 8.57 (d, <i>J</i> = 8.7 Hz, 1H), 8.32 (dd, <i>J</i> = 6.9, 2.6 Hz, 1H), 8.00–7.91 (m, 2H), 7.80 (dd, <i>J</i> = 8.6, 1.7 Hz, 1H), 7.60 – 7.41 (m, 4H), 7.21 (d, <i>J</i> = 7.8 Hz, 1H), 7.12 (td, <i>J</i> = 7.4, 0.9 Hz, 1H), 3.84 (s, 3H).	DMSO	>98	N5

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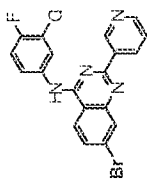
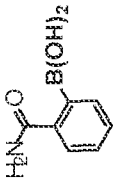
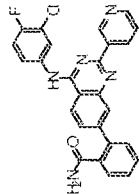
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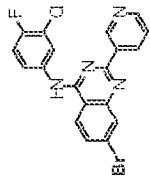
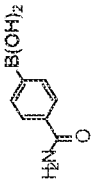
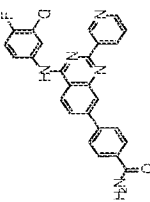
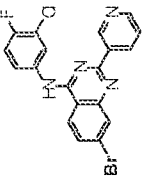
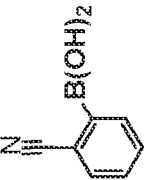
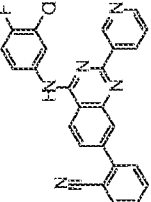
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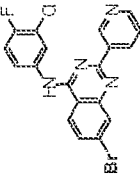
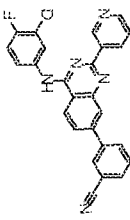
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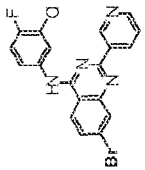
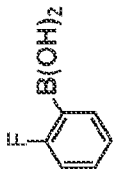
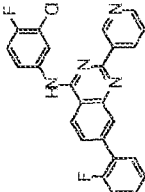
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
683					¹ H NMR (400 MHz, DMSO) δ 10.10 (s, 1H), 9.55 (s, 1H), 8.74 - 8.65 (m, 2H), 8.56 (d, J = 8.6 Hz, 1H), 8.35 - 8.26 (m, 1H), 7.99 - 7.89 (m, 2H), 7.81 (s, 1H), 7.70 (dd, J = 8.6, 1.6 Hz, 1H), 7.63 - 7.47 (m, 6H), 7.40 (s, 1H).	DMSO	>98	N5

(continued)

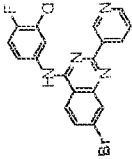
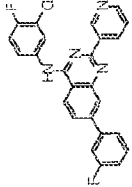
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
684					¹ H NMR (400 MHz, DMSO) δ 10.16 (s, 1H), 9.61 - 9.53 (m, 1H), 8.76 - 8.62 (m, 1H), 8.35 - 8.28 (m, 1H), 8.24 (d, J = 1.8 Hz, 1H), 8.16 - 7.99 (m, 6H), 7.99 - 7.92 (m, 1H), 7.63 - 7.51 (m, 2H), 7.45 (s, 1H).	DMSO	>98	N5
685					¹ H NMR (400 MHz, DMSO) δ 10.21 (s, 1H), 9.55 (d, J = 1.5 Hz, 1H), 8.78 - 8.62 (m, 3H), 8.31 (dd, J = 6.9, 2.6 Hz, 1H), 8.15 - 8.00 (m, 2H), 8.00 - 7.80 (m, 5H), 7.74 - 7.65 (m, 1H), 7.64 - 7.51 (m, 2H).	DMSO	>98	N5

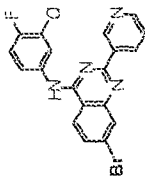
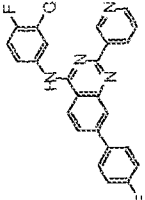
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
686					¹ H NMR (400 MHz, DMSO) δ 10.18 (s, 1H), 9.60 - 9.55 (m, 1H), 8.77 - 8.65 (m, 2H), 8.46 (s, 1H), 8.37 - 8.28 (m, 2H), 8.17 - 8.10 (m, 1H), 7.99 - 7.93 (m, 2H), 7.81 - 7.72 (m, 2H), 7.62 - 7.50 (m, 3H).	DMSO	>98	N5

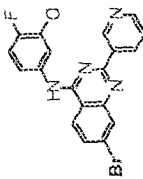
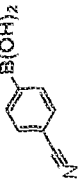
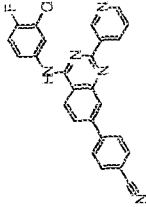
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	687	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
						¹ H NMR (400 MHz, DMSO) δ 10.18 (s, 1H), 9.56 (d, J = 2.2 Hz, 1H), 8.75-8.61 (m, 3H), 8.32 (dd, J = 6.8, 2.6 Hz, 1H), 8.07 (s, 1H), 7.99 - 7.91 (m, 1H), 7.91 - 7.85 (m, 1H), 7.82 - 7.75 (m, 1H), 7.61 - 7.50 (m, 3H), 7.47 - 7.36 (m, 2H).	DMSO	>98	N5

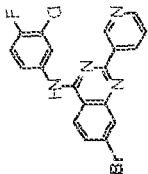
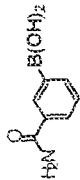
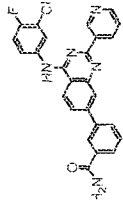
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
688					¹ H NMR (400 MHz, DMSO) δ 10.14 (s, 1H), 9.60 - 9.53 (m, 1H), 8.74 - 8.61 (m, 3H), 8.31 (dd, J = 6.9, 2.6 Hz, 1H), 8.20 (d, J = 1.8 Hz, 1H), 8.06 (dd, J = 8.7, 1.9 Hz, 1H), 7.99 - 7.90 (m, 1H), 7.86 - 7.77 (m, 2H), 7.65 - 7.48 (m, 3H), 7.37 - 7.26 (m, 1H).	DMSO	>98	N5

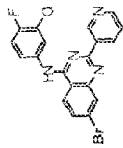
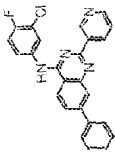
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
689					¹ H NMR (400 MHz, DMSO) δ 10.13 (s, 1H), 9.56 (d, J = 1.5 Hz, 1H), 8.75 - 8.60 (m, 3H), 8.31 (dd, J = 6.9, 2.6 Hz, 1H), 8.15 (d, J = 1.7 Hz, 1H), 8.07 - 7.88 (m, 4H), 7.65 - 7.49 (m, 2H), 7.39 (t, J = 8.8 Hz, 2H).	>98	N5

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
690					¹ H NMR (400 MHz, DMSO) δ 10.14 (s, 1H), 9.54 (d, J = 1.6 Hz, 1H), 8.76 - 8.60 (m, 3H), 8.34 - 8.25 (m, 1H), 8.21 (d, J = 1.7 MHz, 1H), 8.13 (d, J = 8.4 Hz, 2H), 8.10 - 7.97 (m, 3H), 7.97 - 7.87 (m, 1H), 7.62 - 7.45 (m, 2H).	>98	N5

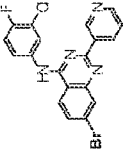
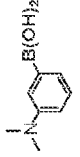
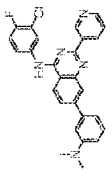
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
691				HCl	¹ H NMR (400 MHz, DMSO) δ 10.45 (s, 1H), 9.54 (d, J = 1.6 Hz, 1H), 8.99 – 8.91 (m, 1H), 8.91 – 8.83 (m, 1H), 8.75 (d, J = 8.8 Hz, 1H), 8.42 (s, 1H), 8.31 (t, J = 3.2 Hz, 1H), 8.29 – 8.22 (m, 2H), 8.13 (dd, J = 8.7, 1.8 Hz, 1H), 8.08 (d, J = 7.8 Hz, 1H), 8.03 – 7.92 (m, 2H), 7.86 (dd, J = 8.0, 5.2 Hz, 1H), 7.65 (t, J = 7.7 Hz, 1H), 7.59 – 7.46 (m, 2H).	DMSO	>98	N5

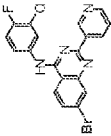
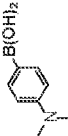
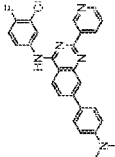
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
692				HCl	¹ H NMR (400 MHz, DMSO) δ 10.38 (s, 1H), 9.56 (d, J = 1.5 Hz, 1H), 8.93 (d, J = 8.2 Hz, 1H), 8.85 (dd, J = 5.1, 1.5 Hz, 1H), 8.71 (d, J = 8.8 Hz, 1H), 8.27 (dd, J = 6.8, 2.6 Hz, 1H), 8.21 (d, J = 1.7 Hz, 1H), 8.10 (dd, J = 8.7, 1.8 Hz, 1H), 8.01 - 7.90 (m, 3H), 7.83 (dd, J = 7.8, 5.0 Hz, 1H), 7.64 - 7.46 (m, 4H).	DMSO	>98	N5

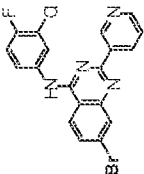
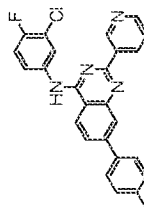
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
693				HCl	¹ H NMR (400 MHz, DMSO) δ 10.91 (s, 1H), 9.52 (d, J = 1.5 Hz, 1H), 9.14 (d, J = 8.1 Hz, 1H), 9.00 (dd, J = 5.4, 1.2 Hz, 1H), 8.91 (d, J = 8.8 Hz, 1H), 8.34 (d, J = 5.1 Hz, 1H), 8.20 (dd, J = 6.8, 2.6 Hz, 1H), 8.14 - 8.04 (m, 2H), 8.04 - 7.89 (m, 2H), 7.75 - 7.48 (m, 4H), 3.18 (s, 6H).	DMSO	>98	N5

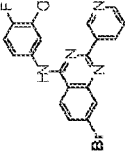
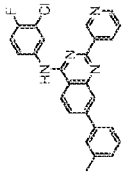
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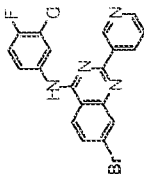
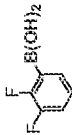
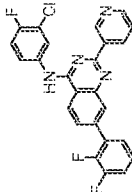
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
694				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.99 (s, 1H), 9.52 (d, J = 1.7 Hz, 1H), 9.08 (d, J = 8.0 Hz, 1H), 8.97 (dd, J = 5.3, 1.4 Hz, 1H), 8.80 (d, J = 8.8 Hz, 1H), 8.33 (d, J = 2.3 Hz, 1H), 8.18 (dd, J = 6.8, 2.6 Hz, 1H), 8.10 (dd, J = 8.8, 1.8 Hz, 1H), 8.04 - 7.91 (m, 2H), 7.85 (d, J = 8.7 Hz, 2H), 7.56 (t, J = 9.1 Hz, 1H), 7.09 (s, 2H), 3.05 (s, 6H).	DMSO	>98	N5

(continued)

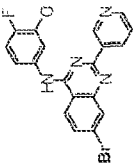
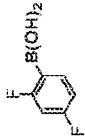
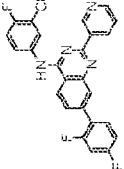
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
695				HCl	¹ H NMR (400 MHz, DMSO) δ 10.67 (s, 1H), 9.53 (d, J = 1.7 Hz, 1H), 9.07 (d, J = 9.07 Hz, 1H), 8.2 Hz, 1H), 8.95 (dd, J = 5.3, 1.4 Hz, 1H), 8.78 (d, J = 8.7 Hz, 1H), 8.26 (s, 1H), 8.21 (dd, J = 6.8, 2.6 Hz, 1H), 8.09 (dd, J = 8.7, 1.8 Hz, 1H), 8.04-7.92 (m, 2H), 7.79 - 7.67 (m, 2H), 7.55 (t, J = 9.1 Hz, 1H), 7.46 (t, J = 7.6 Hz, 1H), 7.32 (d, J = 7.5 Hz, 1H), 2.45 (s, 3H).	DMSO	>98	N5

(continued)

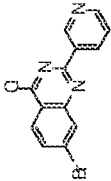
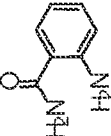
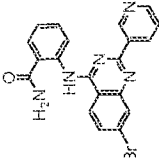
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
696				HCl	¹ H NMR (400 MHz, DMSO) δ 10.48 (s, 1H), 9.54 (d, J = 1.5 Hz, 1H), 8.97 (d, J = 8.1 Hz, 1H), 8.93 - 8.84 (m, 1H), 8.71 (d, J = 8.8 Hz, 1H), 8.30 - 8.16 (m, 2H), 8.08 (dd, J = 8.7, 1.8 Hz, 1H), 8.01 - 7.78 (m, 4H), 7.55 (t, J = 9.1 Hz, 1H), 7.39 (d, J = 7.9 Hz, 2H), 2.40 (s, 3H).	DMSO	>98	N5

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
697				HCl	¹ H NMR (400 MHz, DMSO) δ 10.68 (s, 1H), 9.52 (s, 1H), 9.08 (d, J = 8.1 Hz, 1H), 8.95 (d, J = 4.7 Hz, 1H), 8.82 (d, J = 8.7 Hz, 1H), 8.25 - 8.12 (m, 2H), 8.06 - 7.88 (m, 3H), 7.63 - 7.48 (m, 3H), 7.47 - 7.35 (m, 1H).	>98	N5

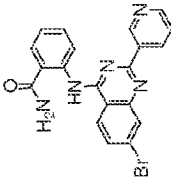
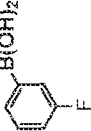
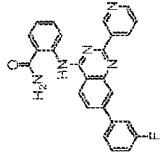
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
698				HCl	¹ H NMR (400 MHz, DMSO) δ 10.47 (s, 1H), 9.53 (d, J = 1.6 Hz, 1H), 9.01 - 8.92 (m, 1H), 8.87 (dd, J = 5.1, 1.4 Hz, 1H), 8.73 (d, J = 8.7 Hz, 1H), 8.24 (dd, J = 6.8, 2.6 Hz, 1H), 8.09 (s, 1H), 7.99 - 7.78 (m, 4H), 7.59 - 7.42 (m, 2H), 7.37 - 7.25 (m, 1H).	DMSO	>98	N5

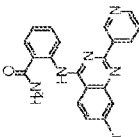
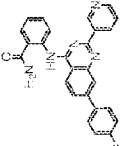
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
699				2 HCl	¹ H NMR (400 MHz, DMSO) δ 13.12 (s, 1H), 9.59 (s, 1H), 9.10 (d, J = 8.1 Hz, 1H), 9.00 - 8.86 (m, 2H), 8.50 (s, 1H), 8.22 - 8.07 (m, 2H), 8.03 - 7.86 (m, 4H), 7.72 (t, J = 7.9 Hz, 1H), 7.28 (t, J = 7.1 Hz, 1H).	DMSO	>98	Scheme 4 synthesis using method N6 coupling

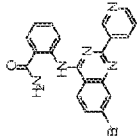
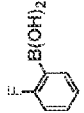
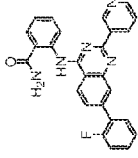
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
700				2 HCl	¹ H NMR (400 MHz, DMSO) δ 13.19 (s, 1H), 9.64 (s, 1H), 9.21 - 9.08 (m, 1H), 0.07 - 8.89 (m, 2H), 8.51 (s, 1H), 8.38 - 8.26 (m, 2H), 8.16 (d, J = 8.5 Hz, 1H), 8.07 - 7.87 (m, 3H), 7.83 - 7.70 (m, 3H), 7.62 (dd, J = 14.2, 7.7 Hz, 1H), 7.40 - 7.22 (m, 2H).	DMSO	>98	N6

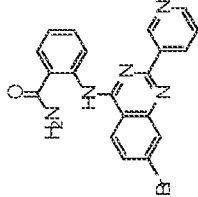
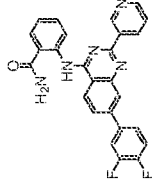
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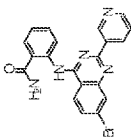
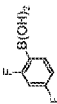
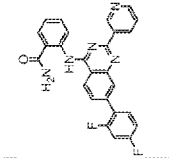
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Per-cent	Method Coupling
701				2 HCl	¹ H NMR (400 MHz, DMSO) δ 13.17 (s, 1H), 9.65 (s, 1H), 9.13 - 8.99 (m, 2H), 8.89 (d, J = 5.1 Hz, 1H), 8.50 (s, 1H), 8.31 (d, J = 8.7 Hz, 1H), 8.21 (d, J = 1.8 Hz, 1H), 8.11 (dd, J = 8.6, 1.9 Hz, 1H), 8.05 - 7.84 (m, 5H), 7.81 - 7.67 (m, 1H), 7.46 - 7.34 (m, 2H), 7.27 (dd, J = 11.8, 4.5 Hz, 1H).	DMSO	>98	N6

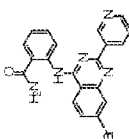
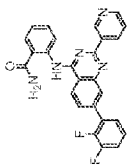
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
702				2 HCl	¹ H NMR (400 MHz, DMSO) δ 13.19 (s, 1H), 9.66 (d, J = 1.8 Hz, 1H), 9.20 (d, J = 7.5 Hz, 1H), 9.04 - 8.89 (m, 2H), 8.50 (s, 1H), 8.35 (d, J = 8.7 Hz, 1H), 8.17 (s, 1H), 8.11 - 7.86 (m, 4H), 7.86 - 7.68 (m, 2H), 7.65 - 7.52 (m, 1H), 7.52 - 7.39 (m, 2H), 7.39 - 7.23 (m, 1H).	DMSO	>98	N6

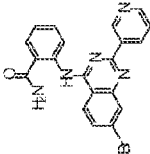
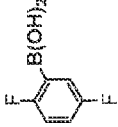
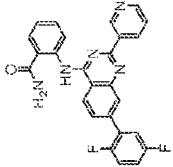
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	703	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Per-cent	Method Coupling
					2 HCl	¹ H NMR (400 MHz, DMSO) δ 13.17 (s, 1H), 9.62 (d, J = 1.6 Hz, 1H), 9.18 (d, J = 8.2 Hz, 1H), 9.03 - 8.91 (m, 2H), 8.51 (s, 1H), 8.34 - 8.20 (m, 2H), 8.16 - 7.86 (m, 5H), 7.84 - 7.68 (m, 2H), 7.68 - 7.54 (m, 1H), 7.35 - 7.19 (m, 1H).	DMSO	>98	N6

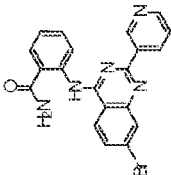
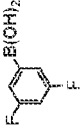
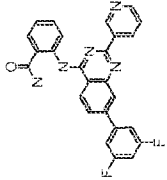
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
704				2 HCl	¹ H NMR (400 MHz, DMSO- <i>d</i> ₆) δ 13.19 (s, 1H), 9.61 (d, <i>J</i> = 1.8 Hz, 1H), 9.32 - 9.22 (m, 1H), 9.02 (dd, <i>J</i> = 5.5, 1.2 Hz, 1H), 8.88 (d, <i>J</i> = 7.8 Hz, 1H), 8.75 (s, 1H), 8.53 (s, 1H), 8.28 (d, <i>J</i> = 8.7 Hz, 1H), 8.16 - 8.07 (m, 2H), 8.01 - 7.88 (m, 3H), 7.85 - 7.76 (m, 1H), 7.76 - 7.65 (m, 1H), 7.54 - 7.42 (m, 1H), 7.37 - 7.19 (m, 2H).	>98	N6

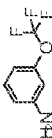
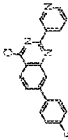
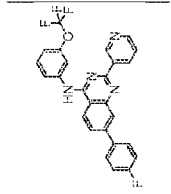
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
705				2 HCl	DMSO	>98	N6
					¹ H NMR (400 MHz, DMSO) δ 13.20 (s, 1H), 9.63 (s, 1H), 9.26 (d, J = 8.1 Hz, 1H), 9.01 (d, J = 5.0 Hz, 1H), 8.91 (d, J = 8.3 Hz, 1H), 8.52 (s, 1H), 8.33 (d, J = 8.7 Hz, 1H), 8.21 - 8.06 (m, 2H), 8.06 - 7.87 (m, 3H), 7.72 (t, J = 7.7 Hz, 1H), 7.66 - 7.52 (m, 2H), 7.49 - 7.35 (m, 1H), 7.29 (t, J = 7.5 Hz, 1H).		

(continued)

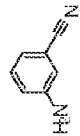
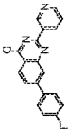
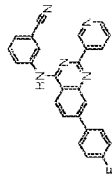
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
706				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.64 (d, J = 1.7 Hz, 1H), 9.25 (d, J = 8.2 Hz, 1H), 9.05 - 8.89 (m, 2H), 8.67 - 8.45 (m, 1H), 8.34 (d, J = 8.7 Hz, 1H), 8.19 (s, 1H), 8.09 (dd, J = 8.0, 5.5 Hz, 1H), 8.04 - 7.88 (m, 3H), 7.79 - 7.62 (m, 2H), 7.56 - 7.35 (m, 2H), 7.35 - 7.23 (m, 1H).	DMSO	>98	N6

(continued)

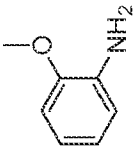
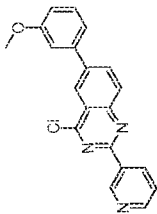
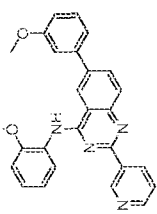
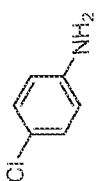
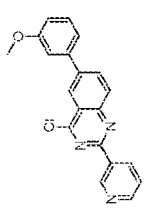
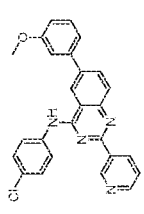
5							
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	707	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent Purity Percent Method Coupling
					2 HCl	¹ H NMR (400 MHz, DMSO) δ 13.18 (s, 1H), 9.62 (d, J = 1.6 Hz, 1H), 9.16 (d, J = 8.1 Hz, 1H), 9.08 - 8.88 (m, 2H), 8.51 (s, 1H), 8.37 - 8.23 (m, 2H), 8.19 - 8.09 (m, 1H), 8.09 - 7.88 (m, 3H), 7.83 - 7.63 (m, 3H), 7.43 - 7.32 (m, 1H), 7.32 - 7.21 (m, 1H).	DMSO >98 N6

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
708				HCl	¹ H NMR (400 MHz, DMSO) δ 10.52 (s, 1H), 9.56 (d, J = 1.6 Hz, 1H), 9.04 (d, J = 7.8 Hz, 1H), 8.92 (d, J = 4.8 Hz, 1H), 8.78 (d, J = 8.8 Hz, 1H), 8.24 (d, J = 1.7 Hz, 1H), 8.13 - 8.08 (m, 2H), 8.08 - 7.98 (m, 3H), 7.98 - 7.88 (m, 1H), 7.63 (t, J = 8.2 Hz, 1H), 7.42 (t, J = 8.8 Hz, 2H), 7.28 - 7.15 (m, 1H).	DMSO	>98	N6/F5 followed by G1

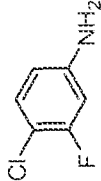
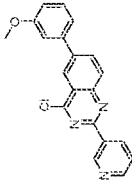
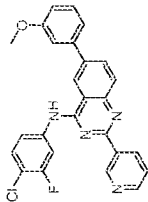
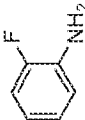
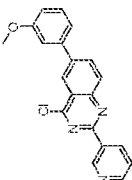
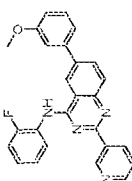
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
709				HCl	¹ H NMR (400 MHz, DMSO) δ 10.60 (s, 1H), 9.56 (s, 1H), 9.07 (d, J = 7.9 Hz, 1H), 8.93 (d, J = 4.3 Hz, 1H), 8.77 (d, J = 8.7 Hz, 1H), 8.41 (s, 1H), 8.37-8.27 (m, 1H), 8.24 (d, J = 1.7 Hz, 1H), 8.11 (dd, J = 8.7, 1.8 Hz, 1H), 8.08-7.94 (m, 3H), 7.79-7.65 (m, 2H), 7.41 (t, J = 8.8 Hz, 2H).	DMSO	>98	N6/F5 followed by G1

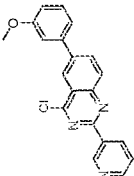
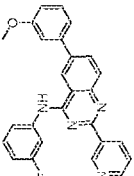
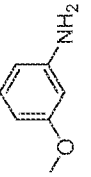
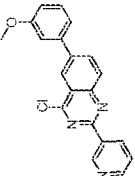
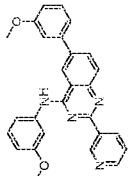
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
710					¹ H NMR (DMSO-d6) ppm 3.81(s, 3H), 3.88(s, 3H), 7.11-7.71(m, 9H), 7.91(d, 1H, J=8.7 Hz), 8.23(d, 1H, J=8.7 Hz), 8.53-8.64(m, 2H), 8.87(d, 1H, J=1.5 Hz), 9.37(d, 1H, J=1.5 Hz), 9.83(s, 1H)	DMSO	>98	N6/F5 followed by G1
711					¹ H NMR (DMSO-d6) ppm 3.89(s, 3H), 7.04(d, 1H, J=4.1 Hz), 7.48-7.57(m, 5H), 7.98(dd, 1H, J=7.1, 1.7 Hz), 8.26-8.29(m, 3H), 8.27(dd, 1H, J=7.1, 8.7 Hz), 8.78-9.53(m, 3H), 9.54(s, 1H), 10.37(s, 1H)	DMSO	>98	N6/F5 followed by G1

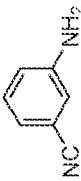
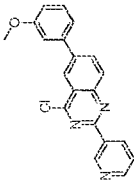
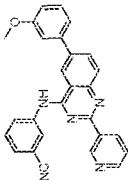
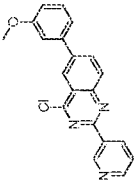
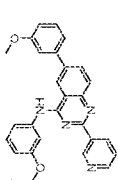
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
712					¹ H NMR (DMSO-d ₆) ppm 3.89 (s, 3H), 7.05 (d, 1H, J = 4.1 Hz), 7.48-7.57 (m, 4H), 7.88 (dd, 1H, J = 7.1, 1.7 Hz), 8.25-8.29 (m, 3H), 8.37 (dd, 1H, J = 7.1, 8.7 Hz), 8.78-9.53 (m, 3H), 9.54 (s, 1H), 10.37 (s, 1H)	DMSO	>98	N6/F5 followed by G1
713					¹ H NMR (DMSO-d ₆) ppm 3.88 (s, 3H), 7.05-7.51 (m, 9H), 7.95 (d, 1H, J = 8.7 Hz), 8.26 (d, 1H, J = 8.7 Hz), 8.54-8.88 (m, 3H), 9.38 (s, 1H), 10.19 (s, 1H)	DMSO	>98	N6/F5 followed by G1

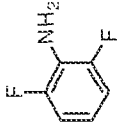
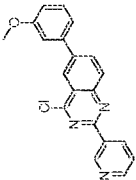
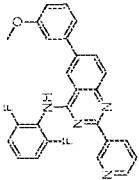
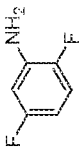
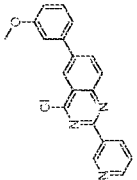
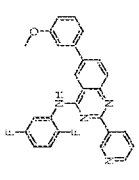
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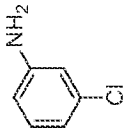
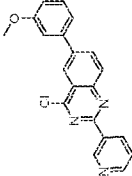
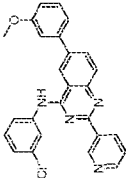
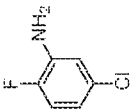
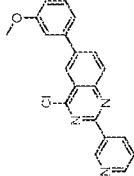
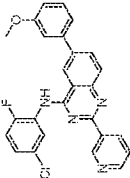
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
714					¹ H NMR (DMSO-d ₆) ppm 3.89 (s, 3H), 7.04-7.06 (m, 2H), 7.46-7.76 (m, 8H), 7.95 (d, 2H, J = 8.6 Hz), 8.24 (d, 1H, J = 1.7 Hz), 8.69-8.87 (m, 3H), 9.56 (s, 1H), 10.18 (s, 1H)	DMSO	>98	N6/F5 followed by G1
715					¹ H NMR (DMSO-d ₆) ppm 3.84 (s, 3H), 3.89 (s, 3H), 6.81-7.56 (m, 2H), 7.40-7.70 (m, 7H), 7.95 (d, 1H, J = 8.5 Hz), 8.23 (d, 1H, J = 1.5 Hz), 8.69-8.89 (m, 3H), 9.58 (s, 1H), 10.06 (s, 1H)	DMSO	>98	N6/F5 followed by G1

(continued)

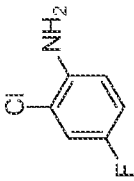
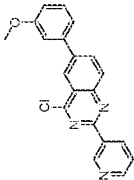
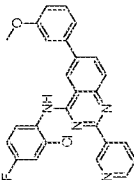
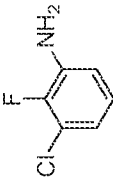
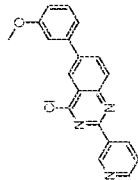
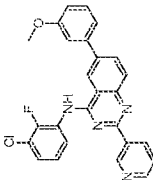
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
716					¹ H NMR (DMSO-d ₆) ppm 3.88 (s, 3H), 7.03-7.06 (m, 1H), 7.50-8.39 (m, 10H), 8.86-8.98 (m, 3H), 9.58 (s, 1H), 10.06 (s, 1H)	DMSO	>98	N6/F5 followed by G1
717					¹ H NMR (DMSO-d ₆) ppm 3.89 (s, 3H), 7.02-7.07 (m, 2H), 7.31 (s, 1H), 7.47-7.56 (m, 5H), 7.81-8.26 (m, 4H), 8.26-8.90 (m, 3H), 9.57 (s, 1H), 10.27 (s, 1H)	DMSO	>98	N6/F5 followed by G1

(continued)

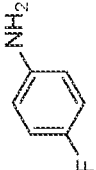
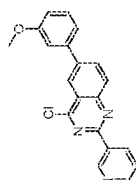
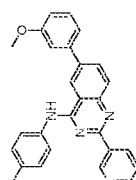
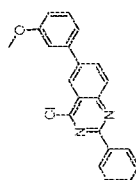
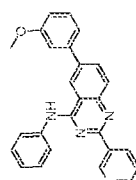
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
718					¹ H NMR (DMSO-d ₆) ppm 3.87 (s, 3H), 7.03-7.50 (m, 8H), 7.98 (d, 1H, J = 8.7 Hz), 8.30 (d, 1H, J = 8.7 Hz), 8.48-8.91 (m, 3H), 9.30 (s, 1H), 10.19 (s, 1H)	DMSO	>98	N6/F5 followed by G1
719					¹ H NMR (DMSO-d ₆) ppm 3.88 (s, 3H), 7.03-7.54 (m, 8H), 7.97 (d, 1H, J = 8.7 Hz), 8.28 (d, 1H, J = 8.7 Hz), 8.56-8.86 (m, 3H), 9.40 (s, 1H), 10.23 (s, 1H)	DMSO	>98	N6/F5 followed by G1

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
722					DMSO	>98	N6/F5 followed by G1
723					DMSO	>98	N6/F5 followed by G1

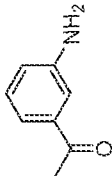
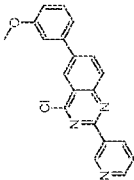
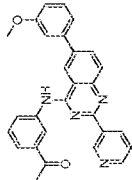
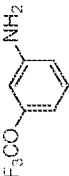
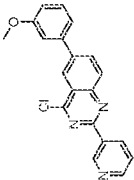
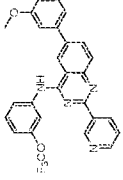
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
724					¹ H NMR (DMSO-d ₆) ppm 3.88 (s, 3H), 7.03-7.74 (m, 8H), 7.98 (d, 1H, J = 8.7 Hz), 8.27 (d, 1H, J = 8.7 Hz), 8.52-8.87 (m, 3H), 9.32 (s, 1H), 10.22 (s, 1H)	DMSO	>98	N6/F5 followed by G1
725					¹ H NMR (DMSO-d ₆) ppm 3.89 (s, 3H), 7.05-7.71 (m, 8H), 7.99 (d, 1H, J = 8.6 Hz), 8.30 (d, 1H, J = 8.7 Hz), 8.67-8.92 (m, 3H), 9.37 (s, 1H), 10.52 (s, 1H)	DMSO	>98	N6/F5 followed by G1

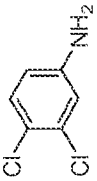
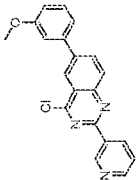
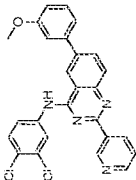
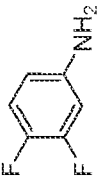
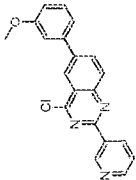
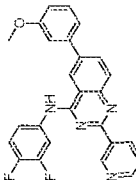
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
726					¹ H NMR (DMSO-d6) ppm 3.89 (s, 3H), 7.03-7.71 (m, 9H), 7.99 (d, 1H, J = 8.7 Hz), 8.29 (d, 1H, J = 8.7 Hz), 8.67-8.92 (m, 3H), 9.37 (s, 1H), 10.52 (s, 1H)	DMSO	>98	N6/F5 followed by G1
727					¹ H NMR (DMSO-d6 at 70 °C) ppm 3.89 (s, 3H), 7.02-7.52 (m, 8H), 7.90 (d, 1H, J = 8.5 Hz), 7.93 (s, 1H), 7.99 (d, 1H, J = 8.6 Hz), 8.22 (s, 1H), 8.74-8.91 (m, 3H), 9.52 (s, 1H), 10.23 (s, 1H)	DMSO	>98	N6/F5 followed by G1

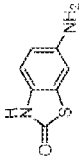
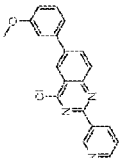
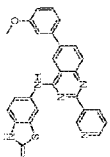
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
728					¹ H NMR (DMSO- <i>d</i> ₆) ppm 2.50 (s, 3H), 3.89 (s, 3H), 7.06 (s, 1H), 7.49-7.85 (m, 9H), 8.01-8.96 (m, 4H), 9.58 (s, 1H), 10.50 (s, 1H)	DMSO	>98	N6/F5 followed by G1
729					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.89 (s, 3H), 7.04 (br s, 1H), 7.22 (s, 1H), 7.49-8.30 (m, 9H), 8.77-8.91 (m, 3H), 9.55 (s, 1H), 10.37 (s, 1H)	DMSO	>98	N6/F5 followed by G1

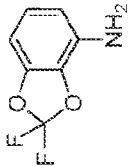
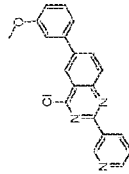
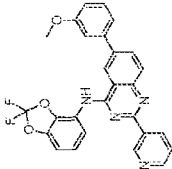
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
730					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.89 (s, 3H), 7.04 (s, 1H), 7.49-8.32 (m, 9H), 8.91-9.04 (m, 3H), 9.54 (s, 1H), 10.63 (s, 1H)	DMSO	>98	N6/F5 followed by G1
731					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.87 (s, 3H), 7.03 (s, 1H), 7.44-8.25 (m, 9H), 8.71-8.84 (m, 3H), 9.52 (s, 1H), 10.23 (s, 1H)	DMSO	>98	N6/F5 followed by G1

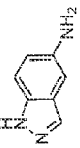
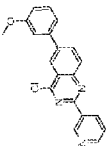
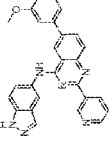
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
732					¹ H NMR (400 MHz, DMSO) δ 12.02 (s, 1H), 10.53 (s, 1H), 9.49 (s, 1H), 9.02 - 8.72 (m, 3H), 8.30 (dd, J = 8.7, 1.6 Hz, 1H), 8.05 (dd, J = 32.2, 5.3 Hz, 2H), 7.81 - 7.72 (m, 2H), 7.49 (dd, J = 8.0, 5.4 Hz, 3H), 7.25 (t, J = 9.2 Hz, 1H), 7.11 - 6.98 (m, 1H), 3.89 (s, 3H).	DMSO	>98	N6/F5 followed by G1

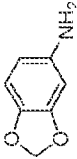
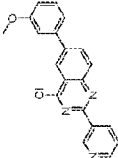
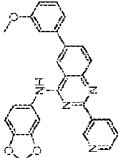
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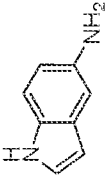
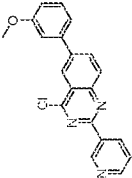
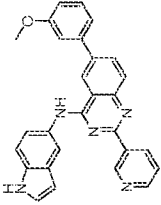
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
733					¹ H NMR (400 MHz, DMSO) δ 10.54 (s, 1H), 9.43 (s, 1H), 8.86 (d, J = 1.8 Hz, 1H), 8.64 (dd, J = 21.1, 5.9 Hz, 2H), 8.31 (dd, J = 8.7, 1.9 Hz, 1H), 8.01 (d, J = 8.7 Hz, 1H), 7.59 - 7.45 (m, 4H), 7.42 - 7.30 (m, 3H), 7.09 - 6.98 (m, 1H), 3.89 (s, 3H).	DMSO	>98	N6/F5 followed by G1

(continued)

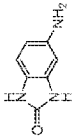
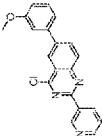
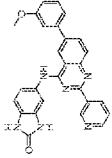
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
734					¹ H NMR (400 MHz, DMSO) δ 13.18 (s, 1H), 10.57 (s, 1H), 9.48 (s, 1H), 8.98 (d, J = 1.3 Hz, 1H), 8.78 (t, J = 7.1 Hz, 2H), 8.30 (dd, J = 8.7, 1.7 Hz, 1H), 8.18 (d, J = 8.4 Hz, 2H), 8.01 (d, J = 8.7 Hz, 1H), 7.83 (dd, J = 8.9, 1.8 Hz, 1H), 7.74 - 7.62 (m, 2H), 7.56 - 7.43 (m, 2H), 7.09 - 7.01 (m, 1H), 3.90 (s, 3H).	DMSO	>98	N6/F5 followed by G1

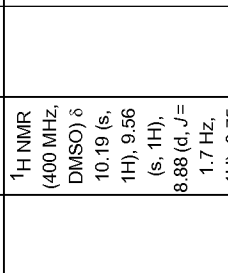
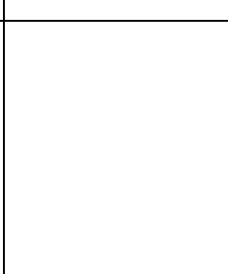
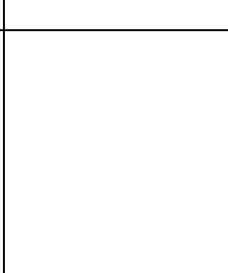
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Per-cent	Method Coupling
735					¹ H NMR (400 MHz, DMSO) δ 11.16 (s, 1H), 10.11 (s, 1H), 9.51 (d, J = 1.2 Hz, 1H), 8.91 (d, J = 1.7 Hz, 1H), 8.71-8.58 (m, 2H), 8.23 (dd, J = 8.7, 1.9 Hz, 1H), 7.95 (dd, J = 23.9, 5.2 Hz, 2H), 7.60 - 7.34 (m, 5H), 7.06 - 6.98 (m, 1H), 6.06 (s, 2H), 3.89 (s, 3H).	DMSO	>98	N6/F5 followed by G1

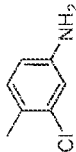
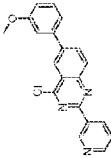
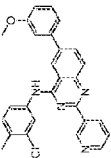
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
736					¹ H NMR (400 MHz, DMSO) δ 10.78 (s, 1H), 10.65 (s, 1H), 10.08 (s, 1H), 9.55 (d, J = 2.0 Hz, 1H), 8.88 (d, J = 1.8 Hz, 1H), 8.76 - 8.66 (m, 2H), 8.24 (dd, J = 8.7, 1.9 Hz, 1H), 7.94 (d, J = 8.7 Hz, 1H), 7.65 (d, J = 1.9 Hz, 1H), 7.58 (dd, J = 7.7, 5.1 Hz, 1H), 7.49 (dt, J = 4.7, 3.1 Hz, 3H), 7.41 (dd, J = 8.4, 2.0 Hz, 1H), 7.08 - 6.99 (m, 3H), 3.88 (s, 3H).	DMSO	>98	N6/F5 followed by G1

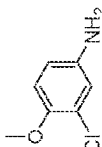
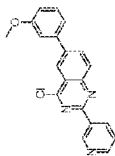
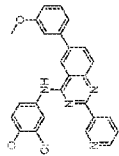
(continued)

5	737	Starting Material 1		Starting Material 2		Product		Salt type	^1H NMR	^1H NMR Solvent	Purity Percent	Method Coupling
10												
15												
20												
25												
30												
35												
40												
45												
50												
55												

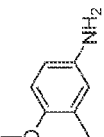
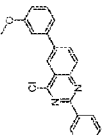
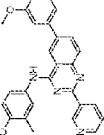
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Per- cent	Method Coupling
738					¹ H NMR (400 MHz, DMSO) δ 10.19 (s, 1H), 9.56 (s, 1H), 8.88 (d, <i>J</i> = 1.7 Hz, 1H), 8.75 (t, <i>J</i> = 7.2 Hz, 2H), 8.27 (dd, <i>J</i> = 8.6, 1.4 Hz, 1H), 8.14 (d, <i>J</i> = 1.6 Hz, 1H), 7.97 (d, <i>J</i> = 8.7 Hz, 1H), 7.84 (dd, <i>J</i> = 8.2, 2.0 Hz, 1H), 7.63 (dd, <i>J</i> = 7.7, 4.9 Hz, 1H), 7.48 (dd, <i>J</i> = 11.4, 4.0 Hz, 4H), 7.10 - 6.98 (m, 1H), 3.89 (s, 3H), 3.83 (s, 3H).	DMSO	>98	N6/F5 followed by G1

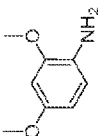
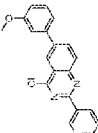
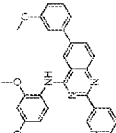
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
739					¹ H NMR (400 MHz, DMSO) δ 10.07 (s, 1H), 9.55 (s, 1H), 8.85 (s, 1H), 8.70 (d, J = 5.5 Hz, 2H), 8.24 (dd, J = 8.7, 1.5 Hz, 1H), 8.10 (d, J = 2.4 Hz, 1H), 7.99-7.83 (m, 2H), 7.60-7.40 (m, 4H), 7.30 (d, J = 9.0 Hz, 1H), 7.04 (dd, J = 7.1, 4.4 Hz, 1H), 3.90 (s, 3H), 2.29 (s, 3H).	DMSO	>98	N6/F5 followed by G1

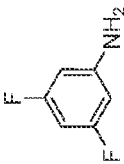
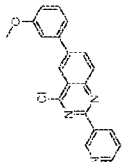
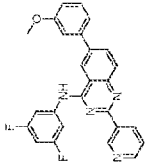
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
740					¹ H NMR (400 MHz, DMSO) δ 10.05 (s, 1H), 9.60 (s, 1H), 8.89 (d, J = 1.8 Hz, 1H), 8.71 (dd, J = 12.8, 6.3 Hz, 2H), 8.24 (dd, J = 8.7, 1.9 Hz, 1H), 7.95 (d, J = 8.7 Hz, 1H), 7.68 (d, J = 1.7 Hz, 1H), 7.61 - 7.45 (m, 5H), 7.44 - 7.37 (m, 1H), 7.24 (d, J = 8.0 Hz, 1H), 3.89 (s, 3H), 3.65 (s, 3H).	DMSO	>98	N6/F5 followed by G1

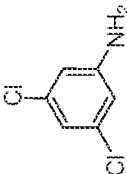
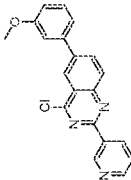
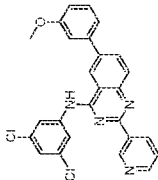
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50							
55							
	741	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	Method Coupling
						¹ H NMR (400 MHz, DMSO) δ 10.04 (s, 1H), 9.60 (s, 1H), 8.89 (d, J = 1.7 Hz, 1H), 8.71 (dd, J = 12.7, 6.3 Hz, 2H), 8.24 (dd, J = 8.7, 1.9 Hz, 1H), 7.95 (d, J = 8.7 Hz, 1H), 7.68 (d, J = 1.7 Hz, 1H), 7.59 - 7.45 (m, 4H), 7.44 - 7.37 (m, 1H), 7.25 (t, J = 6.7 Hz, 1H), 7.07 - 6.98 (m, 1H), 3.89 (s, 6H), 2.19 (s, 3H).	N6/F5 followed by G1
						DMSO	>98

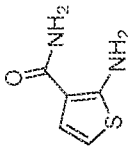
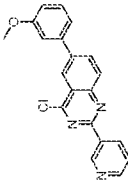
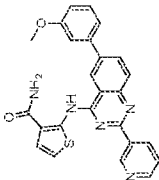
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
742					¹ H NMR (400 MHz, DMSO) δ 9.73 (s, 1H), 9.37 (s, 1H), 8.86 (d, J = 1.6 Hz, 1H), 8.63 (d, J = 3.4 Hz, 1H), 8.54 (d, J = 8.0 Hz, 1H), 8.23 (dd, J = 8.7, 1.8 Hz, 1H), 7.91 (d, J = 8.7 Hz, 1H), 7.55 - 7.42 (m, 5H), 7.05 - 6.98 (m, 1H), 6.77 (d, J = 2.6 Hz, 1H), 6.68 (dd, J = 8.6, 2.6 Hz, 1H), 3.89 (s, 3H), 3.85 (s, 3H), 3.79 (s, 3H).	DMSO	>98	N6/F5 followed by G1

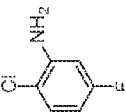
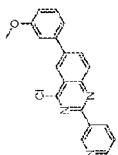
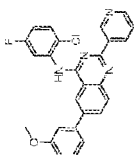
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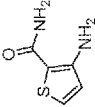
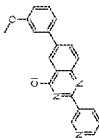
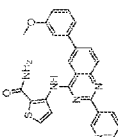
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
743					¹ H NMR (400 MHz, DMSO) δ 10.25 (s, 1H), 9.56 (s, 1H), 8.84 (d, J = 1.7 Hz, 1H), 8.71 (d, J = 9.1 Hz, 1H), 8.27 (dd, J = 8.7, 1.8 Hz, 1H), 7.99 (d, J = 8.7 Hz, 1H), 7.86 - 7.77 (m, 2H), 7.65 - 7.56 (m, 1H), 7.52 - 7.44 (m, 4H), 7.09 - 6.99 (m, 2H), 3.89 (s, 3H).	DMSO	>98	N6/F5 followed by G1

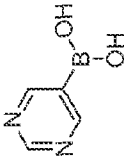
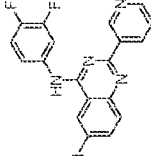
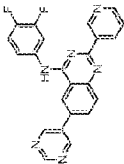
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
744					¹ H NMR (400 MHz, DMSO) δ 10.42 (s, 1H), 9.55 (s, 1H), 8.87 (d, J = 26.7 Hz, 3H), 8.31 (dd, J = 8.7, 1.7 Hz, 2H), 8.17 (d, J = 1.7 Hz, 2H), 8.02 (d, J = 8.7 Hz, 1H), 7.81 (s, 1H), 7.54 - 7.39 (m, 3H), 7.11 - 6.98 (m, 1H), 3.89 (s, 3H).	>98	N6/F5 followed by G1

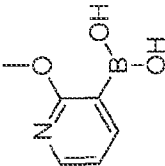
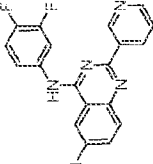
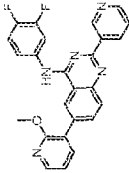
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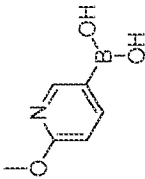
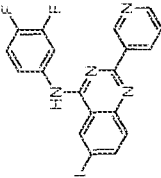
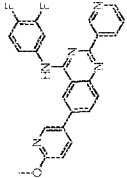
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
745					¹ H NMR (400 MHz, DMSO) δ 14.27 (s, 1H), 9.79 (s, 1H), 9.32 (d, J = 8.4 Hz, 1H), 8.98 (d, J = 5.2 Hz, 1H), 8.34 (dd, J = 8.7, 1.8 Hz, 1H), 8.29 - 8.19 (m, 2H), 8.07 (dd, J = 14.6, 8.3 Hz, 2H), 7.93 (s, 1H), 7.64 (d, J = 5.9 Hz, 1H), 7.52 (t, J = 7.9 Hz, 1H), 7.39 (dd, J = 8.9, 4.9 Hz, 2H), 7.23 (d, J = 5.8 Hz, 1H), 7.08 (dd, J = 8.2, 1.8 Hz, 1H), 3.86 (s, 3H).	DMSO	>98	N6/F5 followed by G1

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR Solvent	Purity Percent	Method Coupling
746				2 HCl	DMSO		N6/F5 followed by G1
					¹ H NMR (400 MHz, DMSO) δ 11.10 (s, 1H), 9.36 (s, 1H), 9.11 (s, 1H), 9.00 (dd, J = 23.2, 6.7 Hz, 2H), 8.41 (d, J = 9.1 Hz, 1H), 8.16 (d, J = 8.7 Hz, 1H), 8.05 (dd, J = 7.9, 5.5 Hz, 1H), 7.80 - 7.65 (m, 2H), 7.59 - 7.44 (m, 3H), 7.41 - 7.32 (m, 1H), 7.06 (d, J = 8.0 Hz, 1H), 3.90 (s, 3H).		

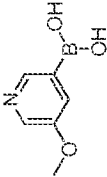
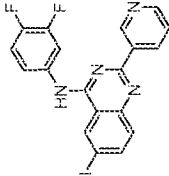
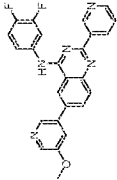
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity Percent	Method Coupling
747					¹ H NMR (400 MHz, DMSO) δ 12.92 (d, <i>J</i> = 17.9 Hz, 1H), 9.64 (s, 1H), 9.09 (d, <i>J</i> = 7.9 Hz, 1H), 8.92 (d, <i>J</i> = 4.9 Hz, 1H), 8.52 (d, <i>J</i> = 5.4 Hz, 1H), 8.38 (s, 1H), 8.30 (dd, <i>J</i> = 8.7, 1.7 Hz, 1H), 8.04 (d, <i>J</i> = 8.7 Hz, 1H), 7.98 - 7.84 (m, 3H), 7.50 (t, <i>J</i> = 7.9 Hz, 1H), 7.39 (dd, <i>J</i> = 8.1, 5.0 Hz, 2H), 7.06 (dd, <i>J</i> = 8.2, 1.8 Hz, 2H), 3.89 (s, 3H).	DMSO	>98	N6/F5 followed by G1

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
748				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.71 (s, 1H), 9.56 (s, 1H), 9.45 (s, 2H), 9.29 (s, 1H), 9.22 (s, 1H), 9.13 (d, J = 7.8 Hz, 1H), 8.96 (d, J = 8.96 (d, J = 4.3 Hz, 1H), 8.47 (d, J = 8.4 Hz, 1H), 8.16 - 8.08 (m, 2H), 8.07 - 7.99 (m, 1H), 7.75 (s, 1H), 7.59 (dd, J = 19.3, 9.0 Hz, 1H).	DMSO	>98	Method N6		

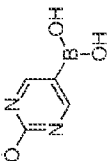
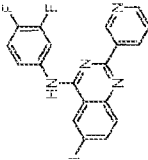
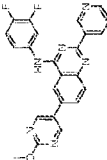
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
749					¹ H NMR (400 MHz, DMSO) δ 10.12 (s, 1H), 9.55 (d, J = 1.4 Hz, 1H), 8.74 - 8.66 (m, 3H), 8.28 (d, J = 4.9 Hz, 1H), 8.17 - 8.08 (m, 2H), 7.98 - 7.91 (m, 2H), 7.73 (d, J = 9.1 Hz, 1H), 7.61 - 7.51 (m, 2H), 7.22 (dd, J = 7.3, 5.0 Hz, 1H), 3.95 (s, 3H).	DMSO	>98	Method N6		

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
750				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.92 (s, 1H), 9.55 (d, J = 1.5 Hz, 1H), 9.33 (s, 1H), 9.10 - 8.99 (m, 2H), 8.93 (d, J = 5.2 Hz, 1H), 8.54 (d, J = 2.5 Hz, 1H), 8.46 (dd, J = 8.8, 1.8 Hz, 1H), 8.33 (s, 1H), 8.18 (ddd, J = 13.0, 7.4, 2.4 Hz, 1H), 8.08 (d, J = 8.7 Hz, 1H), 8.01 - 7.93 (m, 1H), 7.84 (d, J = 9.0 Hz, 1H), 7.57 (dd, J = 19.7, 9.1 Hz, 1H), 4.03 (s, 3H).	DMSO	>98	Method N6		

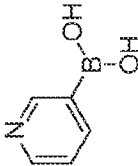
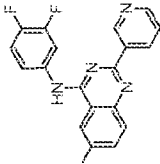
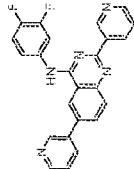
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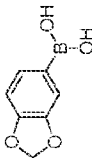
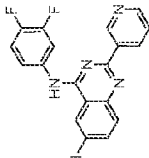
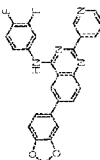
	Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
	751				HCl	¹ H NMR (400 MHz, DMSO) δ 10.57 (s, 1H), 9.53 (d, J = 1.7 Hz, 1H), 9.00 (dd, J = 14.9, 4.8 Hz, 2H), 8.90 (dd, J = 5.1, 1.4 Hz, 1H), 8.78 (d, J = 2.1 Hz, 1H), 8.31 (dt, J = 8.7, 2.2 Hz, 2H), 8.11 (ddd, J = 12.9, 7.5, 2.5 Hz, 1H), 8.03 (d, J = 8.7 Hz, 1H), 7.93 (dd, J = 8.0, 5.1 Hz, 1H), 7.75 (dd, J = 6.0, 2.9 Hz, 1H), 7.63 - 7.51 (m, 1H), 7.03 (d, J = 8.6 Hz, 1H), 3.94 (d, J = 8.4 Hz, 3H).	DMSO	>98	Method N6		

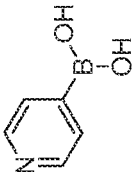
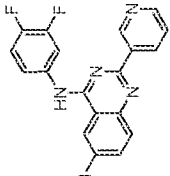
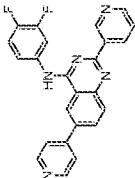
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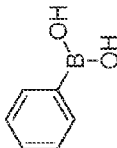
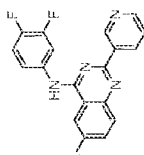
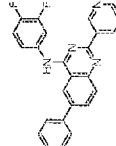
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
752					¹ H NMR (400 MHz, DMSO) δ 10.05 (s, 1H), 9.52 (s, 1H), 9.14 (d, J = 5.6 Hz, 2H), 8.84 (s, 1H), 8.68 (dd, J = 14.0, 6.4 Hz, 2H), 8.28 (dd, J = 8.7, 1.7 Hz, 1H), 8.17 - 8.06 (m, 1H), 7.96 (d, J = 8.7 Hz, 1H), 7.70 (d, J = 9.0 Hz, 1H), 7.62 - 7.50 (m, 2H), 4.02 (s, 3H).	DMSO	>98	Method N6		

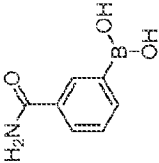
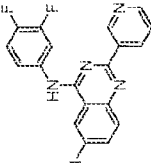
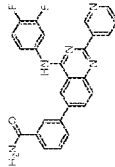
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
753					¹ H NMR (400 MHz, DMSO) δ 10.17 (s, 1H), 9.54 (d, J = 1.5 Hz, 1H), 9.16 (d, J = 1.9 Hz, 1H), 8.93 (d, J = 1.7 Hz, 1H), 8.74 - 8.60 (m, 3H), 8.35 - 8.26 (m, 2H), 8.19 - 8.07 (m, 1H), 8.01 (d, J = 8.7 Hz, 1H), 7.77 - 7.69 (m, 1H), 7.64 - 7.51 (m, 3H).	DMSO	>98	Method N6		

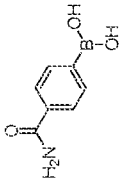
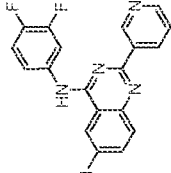
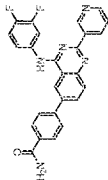
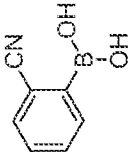
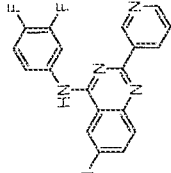
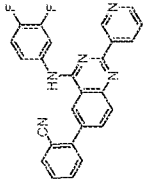
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
754					¹ H NMR (400 MHz, DMSO) δ 10.12 (s, 1H), 9.53 (d, <i>J</i> = 1.4 Hz, 1H), 8.79 - 8.62 (m, 3H), 8.24 - 8.06 (m, 2H), 7.94 (t, <i>J</i> = 7.7 Hz, 1H), 7.73 (dd, <i>J</i> = 6.2, 2.9 Hz, 1H), 7.56 (ddd, <i>J</i> = 15.3, 10.1, 4.8 Hz, 3H), 7.41 (dd, <i>J</i> = 8.1, 1.8 Hz, 1H), 7.11 (d, <i>J</i> = 8.1 Hz, 1H), 6.13 (s, 2H).	DMSO	>98	Method N6		

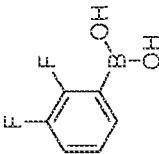
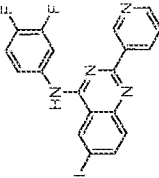
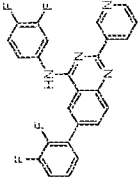
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
755				2 HCl	¹ H NMR (400 MHz, DMSO) δ 11.36 (s, 1H), 9.73 (d, J = 1.7 Hz, 1H), 9.52 (d, J = 1.8 Hz, 1H), 9.13 - 9.02 (m, 3H), 8.96 (dd, J = 5.4, 1.4 Hz, 1H), 8.79 (d, J = 6.9 Hz, 2H), 8.56 (dd, J = 8.9, 1.9 Hz, 1H), 8.19 (ddd, J = 13.0, 7.5, 2.5 Hz, 1H), 8.09 (d, J = 8.8 Hz, 1H), 8.02 (dd, J = 8.0, 5.4 Hz, 1H), 7.90 (dt, J = 9.0, 2.8 Hz, 1H), 7.62 - 7.50 (m, 1H).	DMSO	>98	Method N6		

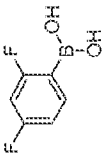
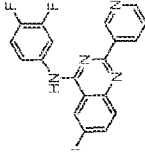
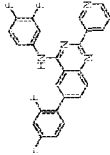
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
756					¹ H NMR (400 MHz, DMSO) δ 10.19 (s, 1H), 9.55 (d, J = 1.4 Hz, 1H), 8.86 (d, J = 1.7 Hz, 1H), 8.69 (dd, J = 8.8, 2.9 Hz, 2H), 8.25 (dd, J = 8.7, 1.8 Hz, 1H), 8.18 - 8.07 (m, 1H), 7.99 (d, J = 8.7 Hz, 1H), 7.92 (d, J = 7.2 Hz, 2H), 7.78 - 7.68 (m, 1H), 7.62 - 7.53 (m, 4H), 7.47 (t, J = 7.4 Hz, 1H).	DMSO	>98	Method N6		

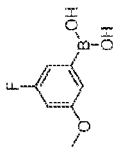
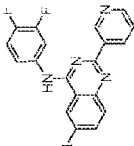
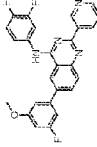
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
757				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.84 (s, 1H), 9.55 (s, 1H), 9.19 - 9.05 (m, 2H), 8.95 (d, J = 4.1 Hz, 1H), 8.53 (s, 1H), 8.41 (dd, J = 8.7, 1.7 Hz, 1H), 8.28 (s, 1H), 8.16 - 8.07 (m, 3H), 8.05 - 7.92 (m, 2H), 7.79 (d, J = 9.1 Hz, 1H), 7.70 - 7.46 (m, 3H).	DMSO	>98	Method N6		

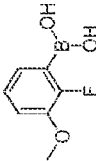
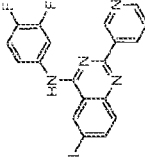
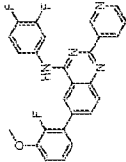
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
758				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.78 (s, 1H), 9.55 (d, J = 1.8 Hz, 1H), 9.12 (d, J = 8.6 Hz, 2H), 8.96 (dd, J = 5.4, 1.4 Hz, 1H), 8.40 (dd, J = 8.8, 1.8 Hz, 1H), 8.16 - 7.96 (m, 8H), 7.79 - 7.71 (m, 1H), 7.64 - 7.53 (m, 1H), 7.47 (s, 1H).	DMSO	>98	Method N6		
759				HCl	¹ H NMR (400 MHz, DMSO) δ 10.41 (s, 1H), 9.56 (s, 1H), 8.98 - 8.77 (m, 3H), 8.23 - 8.02 (m, 4H), 7.99 - 7.65 (m, 5H), 7.56 (dd, J = 19.1, 9.4 Hz, 1H).	DMSO	>98	Method N6		

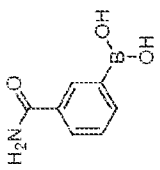
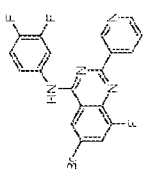
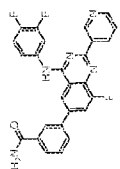
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
760				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.51 (s, 1H), 9.55 (d, J = 1.6 Hz, 1H), 9.04 (d, J = 8.1 Hz, 1H), 8.95 - 8.84 (m, 2H), 8.21-8.02 (m, 3H), 7.93 (dt, J = 10.6, 5.3 Hz, 1H), 7.77-7.68 (m, 1H), 7.63 - 7.51 (m, 3H), 7.49 - 7.38 (m, 1H).	DMSO	>98	Method N6		

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
761				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.51 (s, 1H), 9.55 (d, J = 1.5 Hz, 1H), 9.03 (d, J = 8.2 Hz, 1H), 8.91 (dd, J = 5.3, 1.5 Hz, 1H), 8.82 (s, 1H), 8.16 - 8.00 (m, 3H), 7.94 (dd, J = 8.0, 5.2 Hz, 1H), 7.87 - 7.78 (m, 1H), 7.75 - 7.66 (m, 1H), 7.61 - 7.44 (m, 2H), 7.39 - 7.28 (m, 1H).	DMSO	>98	Method N6		

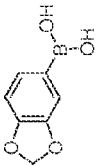
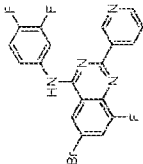
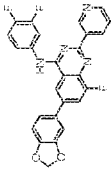
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
762				HCl	¹ H NMR (400 MHz, DMSO) δ 10.75 (s, 1H), 9.53 (d, J = 1.7 Hz, 1H), 9.07 (dd, J = 23.2, 4.9 Hz, 2H), 8.95 (dd, J = 5.3, 1.4 Hz, 1H), 8.36 (dd, J = 8.8, 1.9 Hz, 1H), 8.15 - 7.96 (m, 3H), 7.76 (dd, J = 9.0, 4.1 Hz, 1H), 7.64 - 7.50 (m, 1H), 7.41 (dt, J = 12.9, 1.6 Hz, 2H), 6.99-6.90 (m, 1H), 3.90 (s, 3H).	DMSO	>98	Method N6		

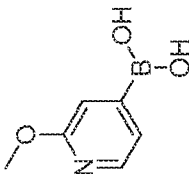
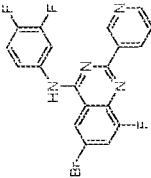
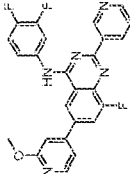
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
763				HCl	¹ H NMR (400 MHz, DMSO) δ 10.54 (s, 1H), 9.55 (s, 1H), 9.05 (d, J = 8.2 Hz, 1H), 8.92 (d, J = 4.6 Hz, 1H), 8.84 (s, 1H), 8.17 - 8.02 (m, 3H), 7.96 (dd, J = 7.9, 5.4 Hz, 1H), 7.72 (d, J = 9.0 Hz, 1H), 7.56 (dd, J = 19.8, 9.1 Hz, 1H), 7.39-7.21 (m, 3H), 3.89 (s, 3H).	DMSO	>98	Method N6		

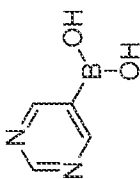
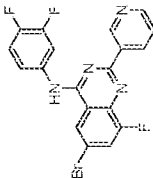
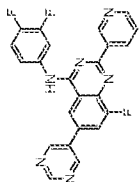
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Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
764				HCl	¹ H NMR (400 MHz, DMSO) δ 10.68 (s, 1H), 9.51 (d, J = 1.3 Hz, 1H), 9.04 (d, J = 8.1 Hz, 1H), 8.97 - 8.87 (m, 2H), 8.51 (s, 1H), 8.32 - 8.22 (m, 2H), 8.16 - 8.05 (m, 2H), 8.00 - 7.90 (m, 2H), 7.80 - 7.73 (m, 1H), 7.65 (t, J = 7.8 Hz, 1H), 7.60 - 7.47 (m, 2H).	DMSO	>98	Method N6		

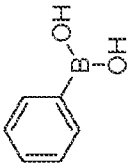
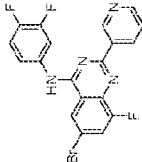
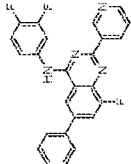
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
765				HCl	¹ H NMR (400 MHz, DMSO) δ 10.43 (s, 1H), 9.46 (d, J = 1.3 Hz, 1H), 9.01 (d, J = 8.1 Hz, 1H), 8.91 (d, 1H), 8.63 (s, 1H), 8.13 - 7.91 (m, 3H), 7.73 - 7.66 (m, 1H), 7.59 - 7.49 (m, 2H), 7.44 (dd, J = 8.2, 1.8 Hz, 1H), 7.07 (d, J = 8.1 Hz, 1H), 6.12 (s, 2H).	DMSO	>98	Method N6		

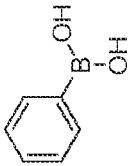
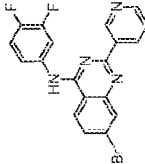
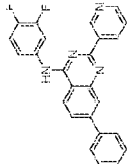
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
766					¹ H NMR (400 MHz, DMSO) δ 10.33 (s, 1H), 9.52 (s, 1H), 8.83 (s, 1H), 8.70 (1H), 8.70 (d, J = 6.3 Hz, 2H), 8.32 (d, J = 8.6 Hz, 2H), 8.13-8.03 (m, 1H), 7.70 (1H), 7.70 (d, J = 7.3 Hz, 1H), 7.58 (dd, J = 7.3, 5.7 Hz, 3H), 7.42 (s, 1H), 3.95 (s, 3H).	DMSO	>98	Method N6		

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
767				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.71 (s, 1H), 9.52 - 9.41 (m, 3H), 9.26 (s, 1H), 9.11 - 9.02 (m, 2H), 8.96 (d, J = 5.4 Hz, 1H), 8.37 (d, J = 11.6 Hz, 1H), 8.14 - 7.99 (m, 2H), 7.80 - 7.71 (m, 1H), 7.55 (d, J = 9.1 Hz, 1H).	DMSO	>98	Method N6		

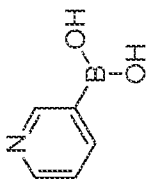
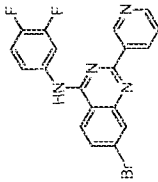
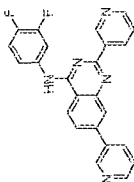
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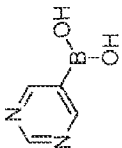
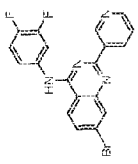
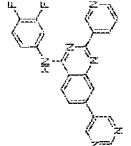
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
768				HCl	¹ H NMR (400 MHz, DMSO) δ 10.53 (s, 1H), 9.51 (d, J = 1.6 Hz, 1H), 9.07 - 9.00 (m, 1H), 8.91 (dd, J = 5.3, 1.5 Hz, 1H), 8.79 (s, 1H), 8.20 (dd, J = 11.9, 1.6 Hz, 1H), 8.13-8.03 (m, 1H), 8.00 - 7.91 (m, 3H), 7.76 - 7.68 (m, 1H), 7.62 - 7.44 (m, 4H).	DMSO	>98	Method N6		

(continued)

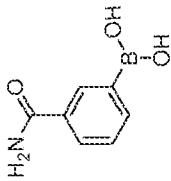
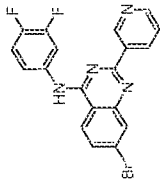
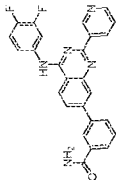
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
770				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.69 (s, 1H), 9.55 (d, J = 1.7 Hz, 1H), 9.12 (d, J = 8.2 Hz, 1H), 8.97 (dd, J = 5.4, 1.4 Hz, 1H), 8.81 (d, J = 8.7 Hz, 1H), 8.30 (d, J = 1.3 Hz, 1H), 8.15 - 7.99 (m, 3H), 7.96 - 7.88 (m, 2H), 7.81 - 7.72 (m, 1H), 7.63 - 7.47 (m, 4H).	DMSO	>98	Method N6		

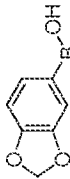
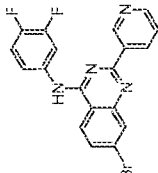
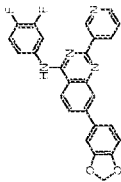
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Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
771				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.89 (s, 1H), 9.54 - 9.45 (m, 2H), 9.17 (t, J = 8.6 Hz, 1H), 9.10-8.93 (m, 4H), 8.48 (d, J = 1.5 Hz, 1H), 8.25 - 8.06 (m, 4H), 7.89 - 7.79 (m, 1H), 7.53 (dd, J = 19.6, 9.2 Hz, 1H).	DMSO	>98	Method N6		

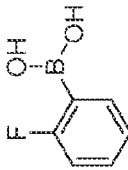
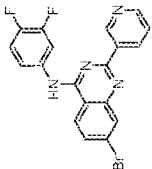
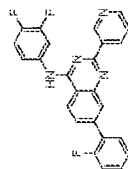
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
772				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.67 (s, 1H), 9.54 (d, J = 1.7 Hz, 1H), 9.40 (s, 2H), 9.31 (s, 1H), 9.20 (d, J = 8.2 Hz, 1H), 9.02 (d, J = 4.3 Hz, 1H), 8.89 (d, J = 8.7 Hz, 1H), 8.43 (s, 1H), 8.22 (dd, J = 8.7, 1.9 Hz, 1H), 8.17 - 8.06 (m, 2H), 7.82 - 7.74 (m, 1H), 7.56 (dd, J = 19.7, 9.2 Hz, 1H).	DMSO	>98	Method N6		

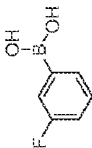
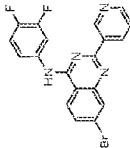
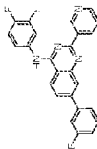
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
773				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.81 (s, 1H), 9.56 (s, 1H), 9.11 (d, J = 8.2 Hz, 1H), 8.94 (dd, J = 17.5, 6.7 Hz, 2H), 8.42 (d, J = 11.9 Hz, 2H), 8.28 (s, 1H), 8.22-7.94 (m, 5H), 7.81 (d, J = 8.8 Hz, 1H), 7.71-7.46 (m, 3H).	DMSO	>98	Method N6		

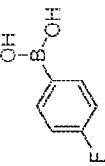
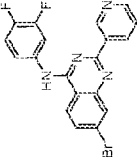
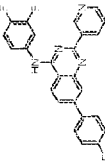
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
774				HCl	¹ H NMR (400 MHz, DMSO) δ 10.32 (s, 1 H), 9.54 (s, 1 H), 8.95 - 1 H), 8.95 - 8.78 (m, 2H), 8.65 (d, J = 8.8 Hz, 1H), 8.19 - 8.09 (m, 2H), 8.02 (d, J = 8.7, 1H), 7.85 - 7.69 (m, 2H), 7.62 - 7.37 (m, 3H), 7.09 (d, J = 7.0 Hz, 1H), 6.13 (s, 2H).	DMSO	>98	Method N6		

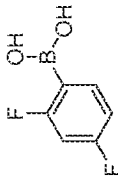
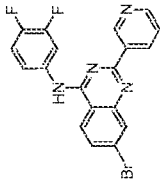
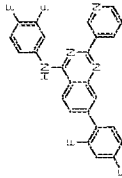
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Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
775				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.59 (s, 1H), 9.56 (d, J = 1.7 Hz, 1H), 9.11 (d, J = 8.1 Hz, 1H), 9.01 - 8.88 (m, 1H), 8.79 (d, J = 8.7 Hz, 1H), 8.18 - 8.06 (m, 2H), 8.05 - 7.91 (m, 2H), 7.82 - 7.72 (m, 2H), 7.61 - 7.52 (m, 2H), 7.48 - 7.37 (m, 2H).	DMSO	>98	Method N6		

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
776				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.46 (s, 1H), 9.53 (d, J = 1.6 Hz, 1H), 9.04 (dt, J = 8.1, 1.6 Hz, 1H), 8.91 (dd, J = 5.2, 1.4 Hz, 1H), 8.73 (d, J = 8.7 Hz, 1H), 8.24 (d, J = 1.8 Hz, 1H), 8.15 - 8.04 (m, 2H), 7.96 (dd, J = 8.0, 5.3 Hz, 1H), 7.84 - 7.70 (m, 2H), 7.65 - 7.48 (m, 2H), 7.39 - 7.25 (m, 1H).	DMSO	>98	Method N6		

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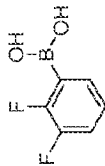
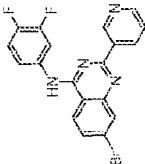
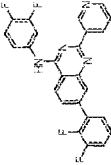
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
777				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.62 (s, 1H), 9.55 (d, J = 1.7 Hz, 1H), 9.12 (d, J = 8.1 Hz, 1H), 8.96 (dd, J = 5.3, 1.3 Hz, 1H), 8.78 (d, J = 8.7 Hz, 1H), 8.25 (d, J = 1.7 Hz, 1H), 8.16 - 7.91 (m, 6H), 7.81 - 7.69 (m, 1H), 7.56 (dd, J = 19.7, 9.1 Hz, 1H), 7.41 (t, J = 8.8 Hz, 1H).	DMSO	>98	Method N6		

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
778				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.65 (s, 1H), 9.55 (d, J = 1.7 Hz, 1H), 9.14 (d, J = 8.1 Hz, 1H), 8.97 (dd, J = 5.4, 1.3 Hz, 1H), 8.81 (d, J = 8.7 Hz, 1H), 8.20 - 8.00 (m, 3H), 7.96 - 7.71 (m, 3H), 7.63-7.42 (m, 2H), 7.38 - 7.25 (m, 1H).	DMSO	>98	Method N6		

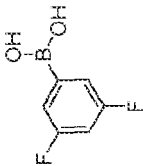
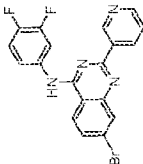
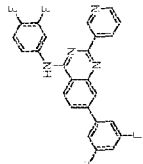
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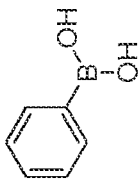
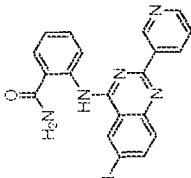
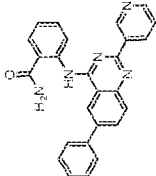
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
779				HCl	¹ H NMR (400 MHz, DMSO) δ 10.40 (s, 1H), 9.54 (d, J = 1.6 Hz, 1H), 9.00 (d, J = 8.1 Hz, 1H), 8.90 (dd, J = 5.2, 1.4 Hz, 1H), 8.71 (d, J = 8.7 Hz, 1H), 8.23 (d, J = 1.8 Hz, 1H), 8.17 - 8.01 (m, 3H), 7.93 (dd, J = 8.0, 5.2 Hz, 1H), 7.84 - 7.71 (m, 2H), 7.69-7.49 (m, 2H).	DMSO	>98	Method N6		

(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
780				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.78 (s, 1H), 9.54 (d, J = 1.6 Hz, 1H), 9.19 (d, J = 8.1 Hz, 1H), 9.06-8.96 (m, 1H), 8.87 (d, J = 8.7 Hz, 1H), 8.22-8.04 (m, 3H), 7.93 (t, J = 10.1 Hz, 1H), 7.78 (dd, J = 6.1, 2.9 Hz, 1H), 7.64-7.49 (m, 3H), 7.47-7.35 (m, 1H).	DMSO	>98	Method N6		

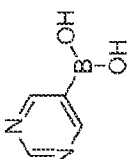
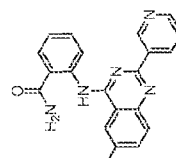
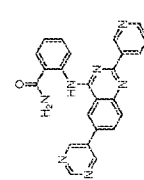
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
781				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.47 (s, 1H), 9.53 (d, $J = 1.7$ Hz, 1H), 9.08 (d, $J = 8.1$ Hz, 1H), 8.94 (dd, $J = 5.3, 1.3$ Hz, 1H), 8.74 (d, $J = 8.8$ Hz, 1H), 8.29 (d, $J = 1.8$ Hz, 1H), 8.17 - 7.95 (m, 3H), 7.80 - 7.65 (m, 3H), 7.61 - 7.48 (m, 1H), 7.42 - 7.30 (m, 1H).	DMSO	>98	Method N6		

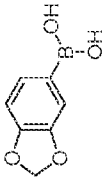
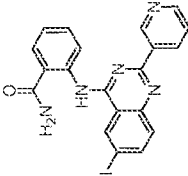
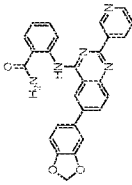
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
782				2 HCl	¹ H NMR (400 MHz, DMSO) δ 13.20 (s, 1H), 9.63 (s, 1H), 9.20 (d, J = 8.1 Hz, 1H), 8.97 (dd, J = 5.3, 1.3 Hz, 1H), 8.89 (d, J = 8.3 Hz, 1H), 8.56 (d, J = 1.6 Hz, 1H), 8.46 (s, 1H), 8.31 (dd, J = 8.7, 1.8 Hz, 1H), 8.13 - 7.92 (m, 4H), 7.90 - 7.83 (m, 2H), 7.77 - 7.70 (m, 1H), 7.59 (dd, J = 10.4, 4.8 Hz, 2H), 7.53-7.46 (m, 2H), 7.34 - 7.25 (m, 1H).	DMSO	>98	Method N6		

(continued)

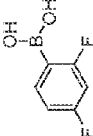
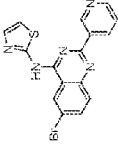
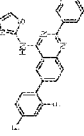
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
784				2 HCl	¹ H NMR (400 MHz, DMSO) δ 13.15 (s, 1H), 9.65 (s, 1H), 9.38 - 9.27 (m, 3H), 9.17 (d, J = 9.2 Hz, 1H), 8.92 (dd, J = 25.3, 6.7 Hz, 2H), 8.68 (s, 1H), 8.45 (d, J = 8.9 Hz, 2H), 8.12 (d, J = 9.8 Hz, 1H), 8.04 - 7.92 (m, 3H), 7.74 (t, J = 7.6 Hz, 1H), 7.30 (t, J = 8.1 Hz, 1H).	DMSO	>98	Method N6		

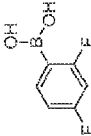
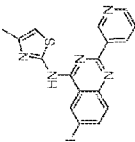
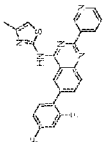
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
785				2 HCl	¹ H NMR (400 MHz, DMSO) δ 13.10 (s, 1H), 9.63 (s, 1H), 9.18 (d, J = 8.0 Hz, 1H), 8.96 (dd, J = 5.3, 1.3 Hz, 1H), 8.87 (d, J = 8.1 Hz, 1H), 8.46 (d, J = 21.1 Hz, 2H), 8.27 (dd, J = 8.7, 1.8 Hz, 1H), 8.09-7.89 (m, 4H), 7.74 (t, J = 7.1 Hz, 1H), 7.46 (d, J = 1.8 Hz, 1H), 7.38 (dd, J = 8.1, 1.9 Hz, 1H), 7.30 (t, J = 7.2 Hz, 1H), 7.12 (d, J = 8.1 Hz, 1H), 6.12 (s, 2H).	DMSO	>98	Method N6		

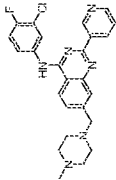
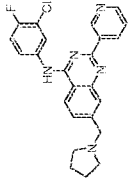
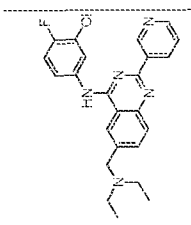
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
786				2 HCl	¹ H NMR (DMSO-d6) ppm 9.79 (d, J = 1.56 Hz, 1H), 9.26 (brd, J = 7.2 Hz, 1H), 9.00-8.90 (brm, 2H), 8.18 (d, J = 8.64 Hz, 1H), 8.11 (d, J = 8.64 Hz, 1H), 8.01 (brt, J = 6.28 Hz, 1H), 7.86-7.80 (brm, 1H), 7.72 (d, J = 3.96 Hz, 1H), 7.53-7.46 (m, 2H), 7.37-7.32 (brm, 1H). The ¹ H of 2HCl and NH-were not observed.	DMSO	>98	N6 using Na2CO3 instead of K3PO4		

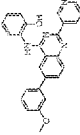
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
787					1H NMR (DMSO-d6) ppm 12.63 (brs, 1H), 9.80 (d, J = 1.64 Hz, 1H), 9.05-8.91 (brm, 2H), 8.76 (brd, J = 4.6 Hz, 1H), 8.10 (brd, J = 8.64 Hz, 1H), 8.02 (d, J = 8.64 Hz, 1H), 7.86-7.80 (brm, 1H), 7.66-7.62 (brm, 1H), 7.51-7.45 (brm, 1H), 7.34-7.30 (brm, 1H), 6.94 (brs, 1H), 2.35 (brs, 3H).	DMSO	>98	N6 using Na ₂ CO ₃ instead of K ₃ PO ₄		

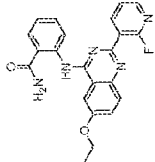
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
788					¹ H NMR (300 MHz, CD ₃ OD) δ 9.40 (s, 1H), 8.65 (d, J = 8.0 Hz, 1H), 8.56 (d, J = 3.2 Hz, 1H), 8.18 - 8.08 (m, 2H), 7.76 - 7.65 (m, 2H), 7.54 - 7.44 (m, 2H), 7.23 (t, J = 9.0 Hz, 1H), 3.66 (s, J = 8.0 Hz, 2H), 2.55 (s, J = 20.6 Hz, 8H), 2.30 (s, 3H).	CD ₃ OD	99	Method N3	463.0 (M+1)	Method C
789					-		99	Method N3	434.0 (M+1)	Method C
790					-		99	Method N3	436.0 (M+1)	Method C

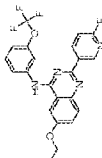
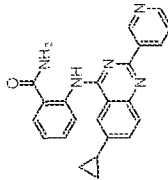
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
791					¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.20 (s, 1H), 9.64 (s, 1H), 9.58 (s, 1H), 8.73 (d, J = 5.5 Hz, 2H), 8.60 (s, 1H), 8.53 (dd, J = 7.2, 1.5 Hz, 1H), 8.28 - 8.25 (m, 1H), 7.99 (1H), 7.99 (d, J = 8.7 Hz, 1H), 7.61 - 7.58 (m, 1H), 7.50 (t, J = 4.5 Hz, 3H), 7.30 (d, J = 5.8 Hz, 1H), 7.06 (dd, J = 7.4, 4.1 Hz, 1H), 6.49 (t, J = 6.9 Hz, 1H), 3.91 (s, 3H).	DMSO	95	Method N1	422.2 (M+1)	Method B (NH ₄ HCO ₃)

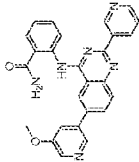
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
792				HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.42 (s, 1H), 8.99 (d, J = 8.0 Hz, 1H), 8.64 (td, J = 7.6, 2.0 Hz, 1H), 8.51 (s, 1H), 8.44 (d, J = 4.4 Hz, 1H), 8.01 - 7.95 (m, 3H), 7.68 - 7.58 (m, 4H), 7.24 (t, J = 7.8 Hz, 1H), 4.26 (q, J = 2.8 Hz, 2H), 1.47 (t, J = 2.8 Hz, 3H).	DMSO	95	Method N1	404.0 (M+1)	Method B (NH ₄ HCO ₃)

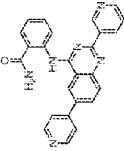
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Cou- pling	LCMS	LCMS Method	
793				HCl	¹ H-NMR (400 MHz, DMSO- d ₆): δ 10.31 (s, 1H), 9.13 (d, J = 2.0 Hz, 1H), 8.83 (td, J = 8.4, 2.4 Hz, 1H), 8.09 (d, J = 1.2 Hz, 2H), 7.93 (d, J = 8.4 Hz, 2H), 7.66 - 7.60 (m, 2H), 7.36 (dd, J = 8.8, 2.4 Hz, 1H), 7.22 (d, J = 8.0 Hz, 1H), 4.28 (q, J = 6.8 Hz, 2H), 1.47 (t, J = 6.8 Hz, 3H).	DMSO	95	Method N1	445.0 (M+1)	Method B (NH ₄ HCO ₃)	
794				2HCl			DMSO	95	Method N1	382.5 (M+1)	Method C

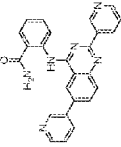
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
795				3HCl	¹ H-NMR (300 MHz, DMSO): δ 13.10 (s, 1H), 9.61 (s, 1H), 9.12 (d, J = 9.8 Hz, 1H), 8.93 (s, 1H), 8.83 (d, J = 15.2 Hz, 2H), 8.66 (s, 1H), 8.58–8.36 (m, 3H), 8.08 (d, J = 11.6 Hz, 2H), 7.96 (d, J = 9.2 Hz, 3H), 7.73 (s, 1H), 7.29 (t, J = 8.7 Hz, 1H), 4.02 (s, 3H).	DMSO	95	Method N1	449.4 (M+1)	Method C

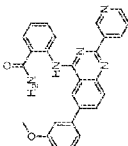
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
796				3HCl	¹ H-NMR (300 MHz, DMSO): δ 12.79 (s, 1H), 9.54 (d, J = 1.6 Hz, 1H), 9.18 (d, J = 8.2 Hz, 1H), 9.08 (d, J = 6.7 Hz, 2H), 9.05 - 8.96 (m, 2H), 8.63 - 8.46 (m, 4H), 8.39 (s, 1H), 8.08 (dd, J = 11.6, 8.5 Hz, 2H), 7.92 (d, J = 6.7 Hz, 1H), 7.77 (s, 1H), 7.68 (t, J = 7.1 Hz, 1H), 7.32 (t, J = 7.2 Hz, 1H).	DMSO	95	Method N1	419.1 (M+1)	Method C

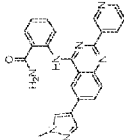
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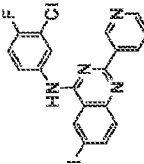
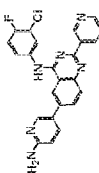
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
797				3HCl	¹ H-NMR (300 MHz, DMSO): δ 12.81 (s, 1H), 9.54 (s, 1H), 9.45 (s, 1H), 9.23 (d, J = 8.2 Hz, 1H), 9.08 - 8.93 (m, 3H), 8.83 (s, 1H), 8.52 (d, J = 8.2 Hz, 1H), 8.47 - 8.34 (m, 2H), 8.14 (ddd, J = 12.9, 8.1, 4.9 Hz, 3H), 7.92 (d, J = 8.6 Hz, 1H), 7.79 (s, 1H), 7.68 (t, J = 7.2 Hz, 1H), 7.32 (t, J = 7.2 Hz, 1H).	DMSO	95	Method N1	419.1 (M+1)	Method C

(continued)

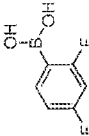
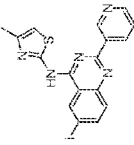
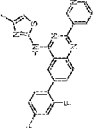
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
798				2HCl	1H-NMR (300 MHz, DMSO): δ 13.20 (s, 1H), 9.62 (s, 1H), 9.13 (d, J = 8.1 Hz, 1H), 8.94 (d, J = 3.5 Hz, 2H), 8.55 (s, 1H), 8.48 (s, 1H), 8.31 (d, J = 8.7 Hz, 1H), 8.06 (d, J = 8.7 Hz, 1H), 7.96 (s, 3H), 7.74 (s, 1H), 7.55 - 7.40 (m, 3H), 7.29 (s, 1H), 7.06 (d, J = 7.4 Hz, 1H), 3.90 (s, 3H).	DMSO	95	Method N1	448.1 (M+1)	Methods C

(continued)

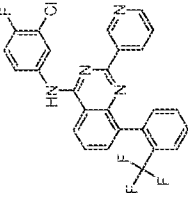
Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
799					¹ H-NMR (300 MHz, DMSO): δ 13.13 (s, 1H), 9.60 (d, J = 1.4 Hz, 1H), 9.12 (d, J = 7.8 Hz, 1H), 8.73 (dd, J = 9.7, 6.4 Hz, 2H), 8.52 (s, 1H), 8.35 (s, 1H), 8.28 (s, 1H), 8.14 (d, J = 8.7 Hz, 1H), 8.08 - 7.88 (m, 4H), 7.76 (d, J = 7.2 Hz, 1H), 7.59 (dd, J = 7.4, 4.8 Hz, 1H), 7.23 (t, J = 7.1 Hz, 1H), 3.94 (s, 3H).	DMSO	95	Method N1	422.2 (M+1)	Method C

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Cou-pling	LCMS	LCMS Method
800					¹ H NMR (400 MHz, DMSO) δ 10.08 (s, 1H), 9.59 - 9.49 (m, 1H), 8.79 - 8.63 (m, 3H), 8.55 (d, J = 2.1 Hz, 1H), 8.27 (dd, J = 6.9, 2.6 Hz, 1H), 8.18 (dd, J = 8.7, 1.9 Hz, 1H), 8.02 - 7.90 (m, 3H), 7.63 - 7.51 (m, 2H), 6.62 (d, J = 8.6 Hz, 1H), 6.24 (s, 2H).	DMSO	>98	N5		

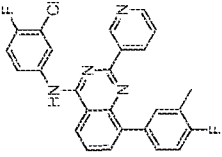
(continued)

Number	Starting Material 1	Starting Material 2	Produce	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
801				MsOH	¹ H NMR (DMSO-d6) ppm 9.78 (brs, 1H), 9.20-8.80 (br, 3H), 8.14 (brd, J = 8.64Hz, 1H), 8.04 (d, J = 8.64Hz, 1H), 7.83 (brm, 2H), 7.52-7.47 (m, 1H), 7.35-7.31 (m, 1H), 7.00 (brs, 1H), 2.36 (brs, 3H). The 1H of MsOH and NH were not observed.	DMSO	>98	N6 using Na2CO3 instead of K3PO4		

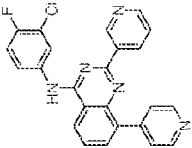
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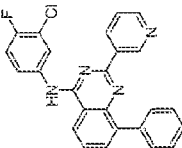
Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
802			494.87	¹ H-NMR (400 MHz, DMSO- _d ₆): δ 10.18 (s, 1H), 9.21 (d, J = 1.8 Hz, 1H), 8.65 - 8.60 (m, 2H), 8.34 - 8.24 (m, 2H), 7.96 - 7.90 (m, 2H), 7.85 - 7.68 (m, 4H), 7.56 (t, J = 9.0 Hz, 1H), 7.51 - 7.44 (m, 2H).	DMSO	495.0, 497.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method N1

(continued)

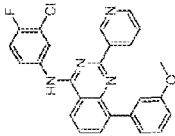
Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
803			458.89	¹ H-NMR (400 MHz, DMSO- _d ₆): δ 10.14 (s, 1H), 9.44 (s, 1H), 8.73-8.63 (m, 1H), 8.55 (t, J = 6.6 Hz, 2H), 8.27 (d, J = 4.6 Hz, 1H), 7.96 (d, J = 6.8 Hz, 2H), 7.76 - 7.64 (m, 3H), 7.58-7.52 (m, 2H), 7.31 (t, J = 9.2 Hz, 1H), 2.37 (s, 3H).	DMSO	459.0, 461.0 (M+1)	Method B (NH ₄ HCO ₄)	95	Method N1

(continued)

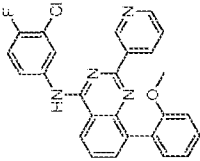
Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
804			427.86	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.19 (s, 1H), 9.42 (d, J = .5 Hz, 1H), 8.74 (d, J = 5.6 Hz, 2H), 8.69 - 8.63 (m, 2H), 8.52 (d, J = 8.0 Hz, 1H), 8.28 (dd, J = 6.8, 2.6 Hz, 1H), 8.05 (d, J = 7.2 Hz, 1H), 7.96 (m, 1H), 7.91 (m, 1H), 7.80 (dd, J = 14.6, 6.8 Hz, 3H), 7.59-7.52 (m, 2H).	DMSO	428.0, 430.0 (M+1) 214.6 (M/2+1)	Method A (TFA)	95	Method N1

Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
805			426.87	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.13 (s, 1H), 9.43 (d, J = 1.6 Hz, 1H), 8.66 (dd, J = 4.7, 1.6 Hz, 1H), 8.59 (dd, J = 8.56 Hz, 1H), 8.52 (td, J = 7.9, 1.8 Hz, 1H), 8.30 (dd, J = 6.84, 2.60 Hz, 1H), 7.99 (-7.92 (m, 2H), 7.82 - 7.73 (m, 3H), 7.58 - 7.51 (m, 4H), 7.46 (t, J = 7.4 Hz, 1H).	DMSO	427.1, 429.1 (M+1)	Method A (TFA)	95	Method N1

(continued)

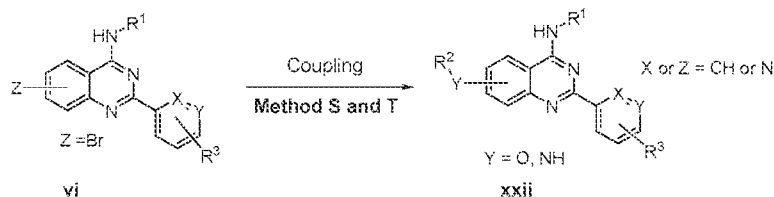
Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
806			456.9	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.13 (s, 1H), 9.45 (s, 1H), 8.67 (dd, J = 4.7, 1.7 Hz, 1H), 8.62 – 8.49 (m, 2H), 8.29 (dd, J = 6.8, 2.6 Hz, 1H), 8.03 – 7.88 (m, 2H), 7.81 – 7.68 (m, 1H), 7.62 – 7.49 (m, 2H), 7.48 – 7.30 (m, 3H), 7.04 (dd, J = 7.8, 2.1 Hz, 1H), 3.87 – 3.81 (m, 3H).	DMSO	457.1, 459.1 (M+1)	Method A (TFA)	95	Method N1

(continued)

Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
807			456.9	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.10 (s, 1H), 9.31 (d, J = 1.6 Hz, 1H), 8.63 (dd, J = 4.8, 1.7 Hz, 1H), 8.61–8.52 (m, 1H), 8.41 (td, J = 8.0, 1.9 Hz, 1H), 8.30 (dd, J = 6.9, 2.6 Hz, 1H), 7.97–7.92 (m, 1H), 7.83 (dd, J = 7.2, 1.2 Hz, 1H), 7.78–7.66 (m, 1H), 7.62–7.40 (m, 3H), 7.36 (dd, J = 7.4, 1.7 Hz, 1H), 7.19 (d, J = 8.1 Hz, 1H), 7.10 (t, J = 7.4 Hz, 1H), 3.66 (s, 3H).	DMSO	457.1, 459.1 (M+2)	Method B (NH ₄ HCO ₃)	95	Method N1

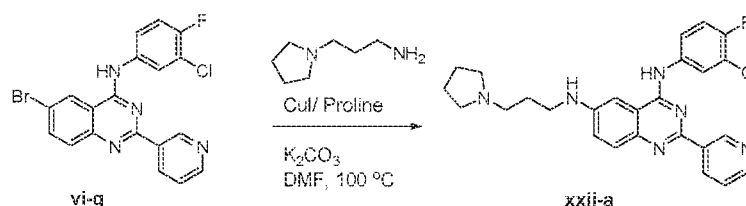
Scheme 20: General route for the synthesis of compounds with general formula xxii

[0334]



Scheme 21: Representative synthesis of compounds of formula xxii (see Scheme 20)

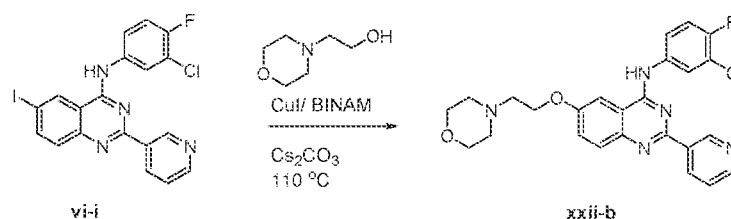
[0335]



[0336] Method S: N⁴-(3-chloro-4-fluorophenyl)-2-(pyridin-3-yl)-N⁶-(3-(pyrrolidin-1-yl)propyl)quinazoline-4,6-diamine (xxii-a) A 2.0 dram reaction vial was charged with 6-bromo-N-(3-chloro-4-fluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine (synthesized as described in Scheme 1 and 4, substituting **5-bromo-2-nitrobenzoic acid** for **2-nitro-5-propoxybenzoic acid** and 3-chloro-4-fluoroaniline for **2-aminobenzamide**) (100 mg, 0.233 mmol, 1.0 equiv), 3-(pyrrolidin-1-yl)propan-1-amine (40 mg, 0.345 mmol, 1.5 equiv), copper(I) iodide (4.4 mg, 0.023 mmol, 0.1 equiv), L-proline (5.3 mg, 0.046 mmol, 0.2 equiv), and potassium carbonate (96 mg, 0.69 mmol, 3.0 equiv) in DMF (3 mL). The reaction mixture was heated at 100 °C overnight. After cooling, water was added to the reaction mixture, and the resultant precipitate was collected by filtration. The crude product was purified via prep-TLC (silica, 2000 micron plate, 95% dichloromethane - 5% methanol - 0.1 % NH₄OH) to yield the desired compound as a brown solid (17.2 mg, 15%). LCMS m/z = 477.4 (M+1) (Method C) (retention time = 1.68 min). ¹H NMR (300 MHz, DMSO) δ 9.68 (s, 1H), 9.46 (s, 1H), 8.59 (d, J = 9.4 Hz, 2H), 8.25 (dd, J = 6.8, 2.6 Hz, 1H), 7.93 (dd, J = 8.3, 3.5 Hz, 1H), 7.69 - 7.59 (m, 1H), 7.52 (dd, J = 11.0, 7.1 Hz, 2H), 7.36 - 7.19 (m, 2H), 6.41 (s, 1H), 3.34 (m, 2H), 2.69 (m, 6H), 2.02 - 1.82 (m, 2H), 1.75 (bs, 4H).

Scheme 22: Representative synthesis of compounds of formula xxii (see Scheme 20)

[0337]



[0338] Method T: N-(3-chloro-4-fluorophenyl)-6-(2-morpholinoethoxy)-2-(pyridin-3-yl)quinazolin-4-amine (xxii-b) A 2.5 dram reaction vial was charged with N-(3-chloro-4-fluorophenyl)-6-iodo-2-(pyridin-3-yl)quinazolin-4-amine (0.250 g, 0.524 mmol), 2-morpholinoethanol (1 ml, 8.17 mmol) as solvent, copper(I) iodide (0.020 g, 0.105 mmol), racemic-2,2'-diamino-1,1'-binaphthyl (0.030 g, 0.105 mmol), and cesium carbonate (0.513 g, 1.573 mmol). The reaction mixture was heated at 110 °C overnight. After cooling, water was added to the reaction mixture, and the resultant precipitate was collected by filtration. The solid residue was purified via ISCO (silica, 12 g column, 95% dichloromethane - 5% methanol - 0.1 % NH₄OH) to yield the desired compound as a brown solid. The solid was further washed with a mixture of water and saturated NaHCO₃ solution and dried. The product off the column was further purified by prep TLC (silica gel, 1000 micron, 95% dichloromethane - 5% methanol - 0.1 % NH₄OH) to afford the desired product as a light

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brown solid (21.2 mg, 8%). LCMS m/z = 480.0 ($M+1$) (Method C) (retention time = 2.09 min). ^1H NMR (300 MHz, DMSO) δ 9.81 (s, 1H), 9.47 (d, J = 1.8 Hz, 1H), 8.66 - 8.57 (m, 2H), 8.21 (dd, J = 6.8, 2.6 Hz, 1H), 7.92 (d, J = 2.5 Hz, 1H), 7.87 (ddd, J = 8.9, 4.3, 2.7 Hz, 1H), 7.81 (d, J = 9.1 Hz, 1H), 7.57 - 7.47 (m, 3H), 4.25 (t, J = 5.8 Hz, 2H), 3.64 - 3.55 (m, 4H), 2.78 (t, J = 5.7 Hz, 2H), 2.55 - 2.49 (m, 4H).

5 **[0339]** The compounds in the following table were prepared in a manner analogous to that described in **Scheme 20** substituting with appropriate amine or alcohol

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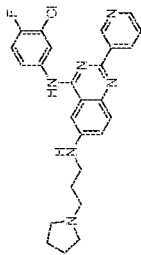
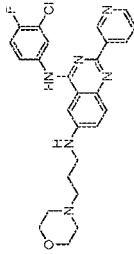
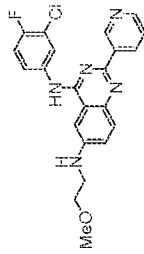
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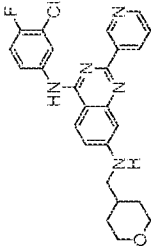
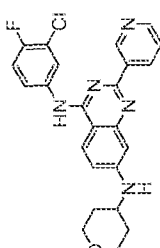
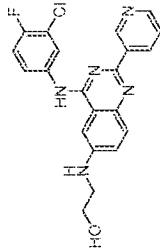
50

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Table 5:

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
808			476.98	¹ H NMR (300 MHz, DMSO) δ 9.68 (s, 1H), 9.46 (s, 1H), 8.59 (d, J = 9.4 Hz, 2H), 8.25 (dd, J = 6.8, 2.6 Hz, 1H), 7.93 (dd, J = 8.3, 3.5 Hz, 1H), 7.69 - 7.59 (m, 1H), 7.52 (dd, J = 11.0, 7.1 Hz, 2H), 7.36 - 7.19 (m, 2H), 6.41 (s, 1H), 3.34 (m, 2H), 2.69 (m, 2H).	DMSO	477.4 (M+1)	1.68	Method C	95	Method S
809			492.98	¹ H NMR (300 MHz, DMSO) δ 9.63 (s, 1H), 9.46 (s, 1H), 8.60 (td, J = 4.2, 2.4 Hz, 2H), 8.23 (dd, J = 6.8, 2.6 Hz, 1H), 7.90 (ddd, J = 8.9, 4.2, 2.6 Hz, 1H), 7.65 (d, J = 9.0 Hz, 1H), 7.57 - 7.45 (m, 2H), 7.32 (dd, J = 9.1, 2.2 Hz, 1H), 7.20 (s, 1H), 6.38 (s, 1H).	DMSO	493 (M+1)	1.97	Method C	100	Method S
810			423.87	¹ H NMR (300 MHz, DMSO) δ 9.60 (s, 1H), 9.46 (s, 1H), 8.60 (dd, J = 9.1, 3.1 Hz, 2H), 8.22 (dd, J = 6.9, 2.5 Hz, 1H), 7.89 (ddd, J = 9.0, 4.3, 2.7 Hz, 1H), 7.64 (d, J = 9.0 Hz, 1H), 7.58 - 7.46 (m, 2H), 7.37 (dd, J = 9.1, 2.2 Hz, 1H), 7.25 (s, 1H), 6.40 (t, 1H).	DMSO	423.9 (M+1)	2.16	Method C	100	Method S

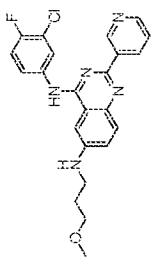
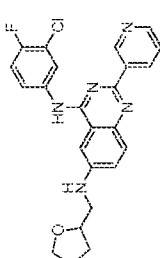
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
811			463.93	¹ H NMR (300 MHz, DMSO) δ 9.62 (s, 1H), 9.49 (d, J = 1.7 Hz, 1H), 8.70-8.59 (m, 2H), 8.27 (dd, J = 6.9, 2.5 Hz, 1H), 8.19 (dd, J = 9.0, 5.0 Hz, 1H), 7.91 - 7.82 (m, 1H), 7.57 - 7.43 (m, 2H), 7.01 (dd, J = 9.0, 2.1 Hz, 1H), 6.77 (t, J = 5.3 Hz, 1H), 6.72	DMSO	464.0 (M+1)	2.19	Method C	100	Method S
812			449.91	¹ H NMR (300 MHz, DMSO) δ 9.65 (s, 1H), 9.49 (d, J = 2.1 Hz, 1H), 8.69-8.59 (m, 2H), 8.27 (dd, J = 6.9, 2.6 Hz, 1H), 8.20 (d, J = 9.0 Hz, 1H), 7.87 (ddd, J = 9.0, 4.3, 2.7 Hz, 1H), 7.56-7.43 (m, 2H), 7.00 (dd, J = 9.1, 2.2 Hz, 1H), 6.79 (d, J = 2.1 Hz,	DMSO	449.9 (M+1)	2.12	Method C	100	Method S
813			409.84	¹ H NMR (300 MHz, DMSO) δ 9.71 (s, 1H), 9.46 (s, 1H), 8.66-8.55 (m, 2H), 8.24 (d, J = 4.3 Hz, 1H), 7.96-7.87 (m, 1H), 7.64 (d, J = 9.1 Hz, 1H), 7.58-7.46 (m, 3H), 7.36 (d, J = 9.0 Hz, 1H), 7.28 (s, 1H), 6.38 (s, 1H), 4.96 (s, 1H), 3.69 (s, 2H), 3.42	DMSO	409.9 (M+1)	1.86	Method C	100	Method S

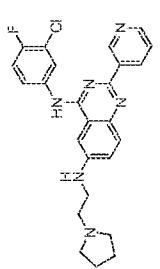
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
814			463.93	¹ H NMR (300 MHz, DMSO) δ 9.59 (s, 1H), 9.45 (d, J = 1.2 Hz, 1H), 8.60 (dd, J = 7.4, 5.7 Hz, 2H), 8.23 (dd, J = 2.5 Hz, 1H), 7.90 (dd, J = 9.0, 2.7 Hz, 1H), 7.64 (d, J = 9.0 Hz, 1H), 7.57-7.46 (m, 2H), 7.36 (d, J = 9.1 Hz, 1H), 7.19 (s, 1H), 6.38 (s, 1H).	DMSO	464.0 (M+1)	2.19	Method C	95	Method S
815			479.93	¹ H NMR (300 MHz, DMSO) δ 9.81 (s, 1H), 9.47 (d, J = 1.8 Hz, 1H), 8.66-8.57 (m, 2H), 8.21 (dd, J = 6.8, 2.6 Hz, 1H), 7.92 (d, J = 2.5 Hz, 1H), 7.87 (ddd, J = 8.9, 4.3, 2.7 Hz, 1H), 7.81 (d, J = 9.1 Hz, 1H), 7.57 - 7.47 (m, 3H), 4.25 (t, J = 5.8 Hz, 2H).	DMSO	480.0 (M+1)	2.09	Method C	91	Method T
816			423.87	¹ H NMR (300 MHz, DMSO) δ 9.60 (s, 1H), 9.47 (s, 1H), 8.67 - 8.55 (m, 2H), 8.23 (dd, J = 6.9, 2.5 Hz, 1H), 7.90 (ddd, J = 9.0, 4.3, 2.7 Hz, 1H), 7.65 (d, J = 9.0 Hz, 1H), 7.58 - 7.47 (m, 2H), 7.37 (dd, J = 9.1, 2.2 Hz, 1H), 7.25 (d, J = 1.3 Hz, 1H), 6.41 (t, J = 5.6 Hz, 1H), 3.63 (t, J = 5.7 Hz, 2H), 3.48-3.35 (m, 5H).	DMSO	423.9 (M+1)		Method C	99	Method S

(continued)

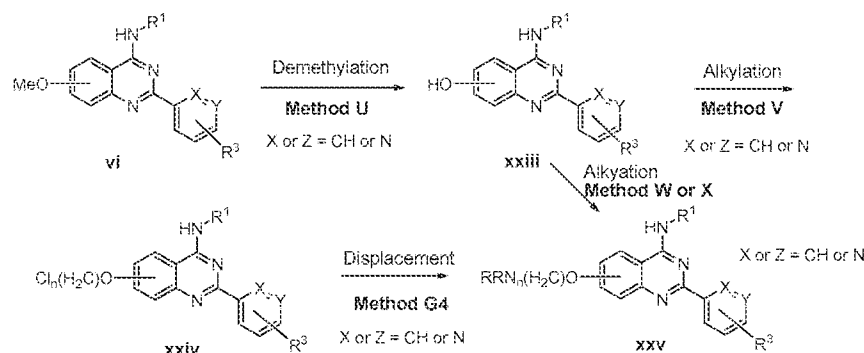
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
817			437.9	¹ H NMR (300 MHz, DMSO) δ 9.63 (s, 1H), 9.47 (s, 1H), 8.66 - 8.55 (m, 2H), 8.23 (dd, J = 6.9, 2.5 Hz, 1H), 7.90 (ddd, J = 8.8, 4.3, 2.5 Hz, 1H), 7.65 (d, J = 9.0 Hz, 1H), 7.58 - 7.45 (m, 2H), 7.32 (dd, J = 9.0, 1.8 Hz, 1H), 7.21 (s, 1H), 6.34 (t, J = 5.0 Hz, 1H), 3.50 (t, J = 6.1 Hz, 2H), 3.31 - 3.20 (m, 5H), 1.92 (p, J = 6.5 Hz, 2H).	DMSO	438.1 (M+1)		Method C	99	Method S
818			449.91	¹ H NMR (300 MHz, DMSO) δ 9.61 (s, 1H), 9.46 (d, J = 2.1 Hz, 1H), 8.65 - 8.56 (m, 2H), 8.23 (dd, J = 6.9, 2.6 Hz, 1H), 7.90 (ddd, J = 8.9, 4.2, 2.6 Hz, 1H), 7.64 (d, J = 9.0 Hz, 1H), 7.58 - 7.47 (m, 2H), 7.40 (dd, J = 8.9, 2.2 Hz, 1H), 7.26 (d, J = 1.7 Hz, 1H), 6.39 (t, J = 5.7 Hz, 1H), 4.12 (t, J = 6.2 Hz, 1H), 3.85 (dd, J = 14.2, 7.2 Hz, 1H), 3.70 (dd, J = 14.2, 7.7 Hz, 1H), 3.34 - 3.25 (m, 5H), 2.15 - 1.99 (m, 1H), 1.97 - 1.80 (m, 3H), 1.74 - 1.58 (m, 2H).	DMSO	449.9 (M+1)		Method C	94	Method S

(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
819				¹ H-NMR (300 MHz, DMSO): δ 10.04 (s, 1H), 9.48 (s, 1H), 8.67 (s, 2H), 8.43–8.31 (m, 1H), 8.14–8.01 (m, 1H), 7.72 (d, J = 9.0 Hz, 1H), 7.65 (s, 1H), 7.54 (dd, J = 9.7 Hz, 2H), 7.35 (d, J = 9.0 Hz, 1H), 6.77 (s, 1H), 3.67 (broad doublet, 4H), 3.41 (bs, 2H), 3.08 (d, J = 4.7 Hz, 2H), 2.03 (bs, 2H), 1.91 (bs, 2H).	DMSO	463.9 (M+1)		Method C	95	Method S

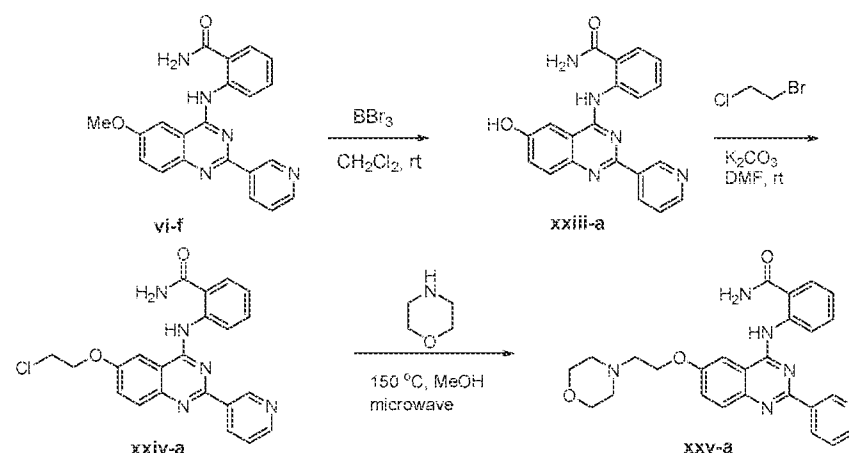
Scheme 23: General route for the synthesis of compounds with general formula xxv

[0340]



Scheme 24: Representative synthesis of compounds of formula xxv (see Scheme 23)

[0341]



[0342] Method U: 2-(6-hydroxy-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (xxiii-a) To a suspension of 2-(6-methoxy-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (synthesized as described in Scheme 1 and 4, substituting 5-methoxy-2-nitrobenzoic acid for 2-nitro-5-propoxy-benzoic acid) (371 mg, 1.0 mmol) in CH_2Cl_2 (4.5 mL) was slowly added boron tribromide, 1M solution in dichloromethane (4.5 mL, 4.5 mmol) at 0 °C. The reaction mixture was stirred overnight at room temperature after which it was carefully poured into a vigorously stirring mixture of ice and saturated solution of aqueous NaHCO_3 . The resultant solid was collected by filtration, dried and then stirred in a saturated solution of aqueous NH_4Cl for 1 h after which the suspension was filtered to give the desired product as a yellowish tan solid (262 mg, 73%). LCMS m/z = 357.9 (M + 1) (Method C) (retention time = 1.68 min). ^1H NMR (300 MHz, DMSO) δ 12.80 (s, 1H), 10.50 (s, 1H), 9.57 (s, 1H), 9.15 (d, J = 8.5 Hz, 1H), 8.70 (t, J = 7.3 Hz, 2H), 8.50 (s, 1H), 8.08 - 7.79 (m, 3H), 7.73 (t, J = 7.5 Hz, 1H), 7.57 (dd, J = 7.6, 4.8 Hz, 1H), 7.46 (d, J = 8.5 Hz, 2H), 7.19 (t, J = 7.6 Hz, 1H).

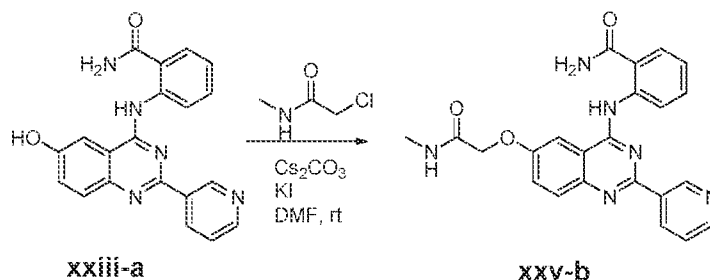
[0343] Method V: 2-(6-(2-chloroethoxy)-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (xxiv-a) The suspension of 2-(6-hydroxy-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (50 mg, 0.14 mmol), 1-bromo-2-chloroethane (0.015 mL, 0.15 mmol), and potassium carbonate (23 mg, 0.17 mmol) in DMF (5 mL) was stirred for 3 days at room temperature. Water (10 mL) was added to the mixture and extracted with ethyl acetate (2 x 10 mL). The combined organic layer was washed with water (1 x 20 mL) and brine (1 x 20 mL) and was dried over MgSO_4 . After filtration and evaporation, the crude product was obtained, which was washed with hexane and dried to give 48 mg of 2-(6-(2-chloroethoxy)-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide as light brown solid (79 %).

[0344] Method G4: 2-(6-(2-morpholinoethoxy)-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (xxv-a) 2-(6-(2-morpholinoethoxy)-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide was prepared from 2-(6-(2-chloroethoxy)-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide and morpholine in a manner analogous to that described for N-(3-chloro-4-fluorophenyl)-6-(3-(dimethylamino)propyl)-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride in Scheme 9 using Method G4

to give 50 mg of 2-(6-(2-morpholinoethoxy)-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide as light yellow solid. LCMS $m/z = 485$ ($M + 1$) (Method C) (retention time = 1.71 min). ^1H NMR (300 MHz, DMSO) δ 9.57 (s, 1H), 9.12 (d, $J = 8.5$ Hz, 1H), 8.78 - 8.63 (m, 2H), 8.48 (s, 1H), 8.02 - 7.93 (m, 2H), 7.88 (d, $J = 9.0$ Hz, 1H), 7.73 (t, $J = 8.0$ Hz, 1H), 7.63 - 7.49 (m, 3H), 7.20 (t, $J = 7.6$ Hz, 1H), 4.29 - 4.16 (m, 2H), 3.65 - 3.51 (m, 4H), 3.44 - 3.21 (m, 2H), 2.45 - 2.31 (m, 4H), 2.06 - 1.91 (m, 2H).

Scheme 25: Representative synthesis of compounds of formula xxv-b (see Scheme 23)

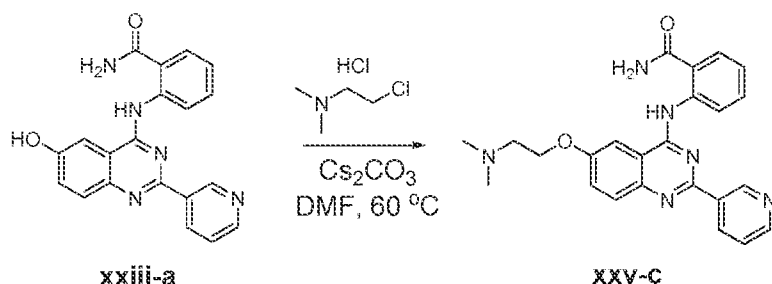
[0345]



[0346] Method W: 2-(6-(2-(methylamino)-2-oxoethoxy)-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide The suspension of 2-(6-hydroxy-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (synthesized as described in Scheme 24) (0.20 g, 0.56 mmol), 2-chloro-N-methylacetamide (90 mg, 0.80 mmol), cesium carbonate (0.37 g, 1.12 mmol) and potassium iodide (0.19 g, 1.12 mmol) in DMF (10 mL) was stirred for 4 days at room temperature. Water (20 mL) was added to the mixture. The resultant solid was collected by filtration. The obtained solid was washed with CH_2Cl_2 -THF (1:1) solution and dried to give 0.11 g of 2-(6-(2-(methylamino)-2-oxoethoxy)-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide as pale brown solid (46 %). LCMS $m/z = 429$ ($M + 1$) (Method C) (retention time = 1.65 min). ^1H NMR (300 MHz, DMSO) δ 9.58 (s, 1H), 9.12 (d, $J = 8.4$ Hz, 1H), 8.82 - 8.64 (m, 2H), 8.48 (s, 1H), 8.32 - 8.17 (m, 1H), 8.10 - 7.87 (m, 3H), 7.81 - 7.49 (m, 4H), 7.21 (t, $J = 7.5$ Hz, 1H), 4.66 (s, 2H), 2.71 (d, $J = 4.5$ Hz, 3H).

Scheme 26: Representative synthesis of compounds of formula xxv-c (see Scheme 23)

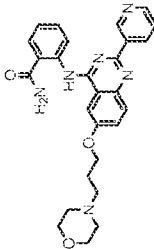
[0347]



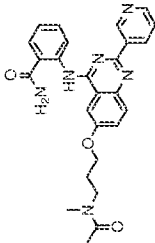
[0348] Method X: 2-(6-(2-(dimethylamino)ethoxy)-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide A suspension of 2-(6-hydroxy-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (synthesized as described in Scheme 24) (25 mg, 0.07 mmol), 2-chloro-N,N-dimethylethylamine hydrochloride (20 mg, 0.14 mmol) and cesium carbonate (68.4 mg, 0.21 mmol) in DMF (1 mL) was stirred at 60 °C overnight. The reaction mixture was cooled to room temperature, diluted with water (5 mL) and extracted with dichloromethane (3 X 5 mL). The combined organic extracts were dried over Na_2SO_4 and concentrated *in vacuo*. The crude product was purified by prep-TLC (silica, 2000 micron plate, 95% dichloromethane - 5% methanol - 0.1 % NH_4OH) to yield the desired compound as a brown solid (12.6 mg, 40%). LCMS $m/z = 428.9$ ($M + 1$) (Method C) (retention time = 1.41 min). ^1H NMR (300 MHz, DMSO) δ 12.99 (s, 1H), 9.59 (d, $J = 1.3$ Hz, 1H), 9.12 (d, $J = 8.4$ Hz, 1H), 8.81 - 8.66 (m, 2H), 8.52 (s, 1H), 8.06 - 7.88 (m, 3H), 7.80 - 7.53 (m, 4H), 7.22 (t, $J = 7.6$ Hz, 1H), 4.46 (s, 2H), 3.35 (s, 2H), 2.69 (s, 6H).

[0349] The compounds in the following table were prepared in a manner analogous to that described in Scheme 23 substituting with appropriate nucleophile.

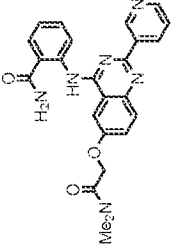
Table 6:

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
820			484.550	¹ H NMR (300 MHz, DMSO) δ 9.57 (s, 1H), 9.12 (d, J = 8.5 Hz, 1H), 8.78 - 8.63 (m, 2H), 8.48 (s, 1H), 8.02 - 7.93 (m, 2H), 7.88 (d, J = 9.0 Hz, 1H), 7.73 (t, J = 8.0 Hz, 1H), 7.63 - 7.49 (m, 3H), 7.20 (t, J = 7.6 Hz, 1H), 4.29 - 4.16 (m, 2H), 3.65-3.51 (m	DMSO	485 (M+1)	1.71	Method C	100	Methods V, G4

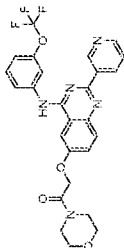
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
821			470.523	¹ H NMR (300 MHz, DMSO) δ 9.58 (s, 1H), 9.24 - 9.08 (m, 1H), 8.82 - 8.61 (m, 2H), 8.49 (s, 1H), 8.09 - 7.84 (m, 3H), 7.73 (t, J = 7.8 Hz, 1H), 7.67 - 7.47 (m, 3H), 7.19 (t, J = 7.5 Hz, 1H), 4.29 - 4.09 (m, 2H), 3.63 - 3.42 (m, 2H), 3.07 - 2.77 (m, 3H), 2.2	DMSO	471 (M+1)	1.75	Method C	100	Methods V, G4 followed by acylation with acetylchloride/TEA in DCM at rt.

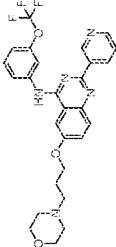
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
822			442.470	¹ H NMR (300 MHz, DMSO) δ 9.58 (s, 1H), 9.16 (d, J = 8.4 Hz, 1H), 8.82 - 8.60 (m, 2H), 8.49 (s, 1H), 8.05 - 7.80 (m, 3H), 7.79 - 7.68 (m, 1H), 7.67 - 7.51 (m, 2H), 7.43 (s, 1H), 7.29 - 7.12 (m, 1H), 5.03 (s, 2H), 3.10 (s, 3H), 2.89 (s, 3H).	DMSO	443 (M+1)	1.63	Method C	100	Method W

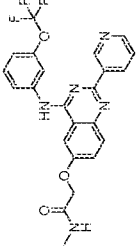
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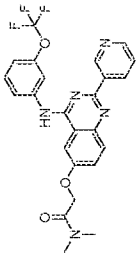


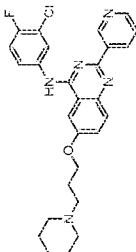
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
824		HCl	598.44	¹ H NMR (300 MHz, DMSO) δ 10.98 (s, 1H), 10.59 (s, 1H), 9.50 (s, 1H), 8.96 (s, 1H), 8.86 (d, J = 8.1 Hz, 1H), 8.86 (d, J = 5.1 Hz, 1H), 8.31 (s, 1H), 8.10 (s, 1H), 8.07 (s, 1H), 7.93 (m, 2H), 7.92 (m, 1H), 7.80 (m, 1H), 7.71 (m, 1H), 7.54 (m, 2H), 7.21 (d, J = 8.0 Hz, 1H), 4.47 (m, 2H), 4.23 (m, 2H).	DMSO	526 (M+1)	2.25	Method C	100	Methods V, G4

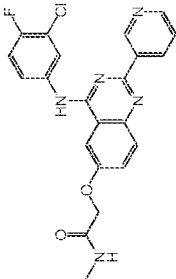
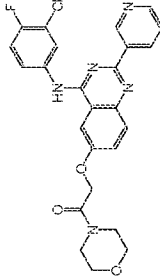
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
825			469.42	¹ H NMR (300 MHz, DMSO) δ 9.95 (s, 1H), 9.51 (s, 1H), 8.76 - 8.56 (m, 2H), 8.26 - 8.08 (m, 2H), 8.01 (s, 1H), 7.95 - 7.81 (m, 2H), 7.70 - 7.56 (m, 2H), 7.52 (dd, J = 7.9, 4.9 Hz, 1H), 7.17 (d, J = 8.3 Hz, 1H), 4.69 (s, 2H), 2.71 (d, J = 4.5 Hz, 3H).	DMSO	470 (M+1)	2.04	Method C	100	Method W

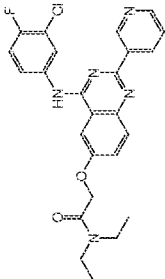
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
826			483.44	¹ H NMR (300 MHz, DMSO) δ 9.92 (s, 1H), 9.51 (s, 1H), 8.74 - 8.57 (m, 2H), 8.17 (s, 1H), 8.01 (d, J = 2.5 Hz, 1H), 7.96 - 7.83 (m, 2H), 7.70 - 7.46 (m, 3H), 7.16 (d, J = 7.0 Hz, 1H), 5.03 (s, 2H), 3.06 (s, 3H), 2.88 (s, 3H).	DMSO	484 (M+1)	1.71	Method D	100	Method W

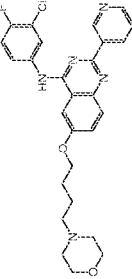
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
827		HCl	564.91	¹ H NMR (300 MHz, DMSO) δ 10.59 (s, 1H), 9.99 (s, 1H), 9.49 (s, 1H), 8.97 (d, 1H), 8.87 (d, 1H), 8.87 (d, 1H), 8.33 - 8.18 (m, 2H), 8.04 - 7.83 (m, 3H), 7.68 - 7.47 (m, 2H), 4.41 - 4.27 (m, 2H), 3.55 - 3.43 (m, 2H), 3.34 - 3.20 (m, 2H), 3.03 - 2.91 (m, 2H).	DMSO	492 (M+1)	1.81	Method C	100	Methods V, G4

(continued)

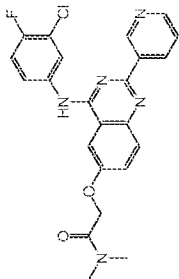
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
828			437.85	¹ H NMR (300 MHz, DMSO) δ 9.87 (s, 1H), 9.49 (s, 1H), 8.76 - 8.53 (m, 2H), 8.32 - 8.08 (m, 2H), 8.01 - 7.77 (m, 3H), 7.69 - 7.43 (m, 3H), 4.67 (s, 2H), 2.71 (d, J = 4.6 Hz, 3H).	DMSO	438 (M+1)	1.65	Method D	100	Method W
829			493.92	¹ H NMR (300 MHz, DMSO) δ 9.86 (s, 1H), 9.50 (s, 1H), 8.76 - 8.57 (m, 2H), 8.27 (d, J = 7.0 Hz, 1H), 8.05 - 7.80 (m, 3H), 7.69 - 7.45 (m, 3H), 5.04 (s, 2H), 3.78 - 3.41 (m, 8H).	DMSO	494 (M+1)	1.98	Method C	98	Method W

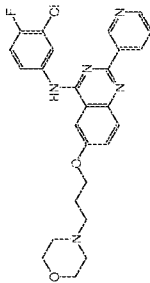
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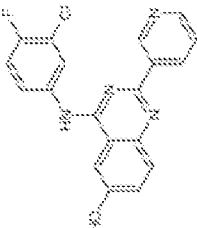
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
830			479.93	¹ H NMR (300 MHz, DMSO) δ 9.85 (s, 1H), 9.50 (s, 1H), 8.76 - 8.53 (m, 2H), 8.26 (dd, J = 6.9, 2.5 Hz, 1H), 8.08 - 7.76 (m, 3H), 7.71 - 7.41 (m, 3H), 4.98 (s, 2H), 3.52 - 3.09 (m, 4H), 1.37 - 0.88 (m, 6H).	DMSO	480 (M+1)	2.20	Method C	100	Method W

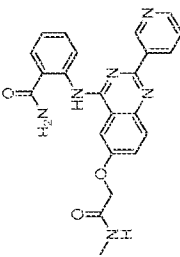
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
831		HCl	580.91	¹ H NMR (300 MHz, DMSO) δ 10.71 (s, 2H), 9.48 (s, 1H), 9.10 - 8.95 (m, 1H), 8.94 - 8.79 (m, 1H), 8.36 - 8.27 (m, 1H), 8.26 - 8.16 (m, 1H), 8.07 - 7.83 (m, 3H), 7.68 - 7.46 (m, 2H), 4.39 - 4.18 (m, 2H), 4.05 - 3.89 (m, 2H), 3.79 (t, J = 11.9 Hz, 2H), 3.53 -	DMSO	508 (M+1)	2.14	Method C	100	Methods V, G4

(continued)

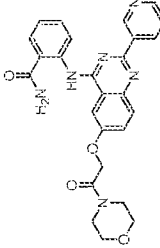
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
832			451.88	¹ H NMR (300 MHz, DMSO) δ 9.85 (s, 1H), 9.50 (s, 1H), 8.71 - 8.58 (m, 2H), 8.25 (dd, J = 6.8, 2.5 Hz, 1H), 8.01 - 7.80 (m, 3H), 7.66 - 7.45 (m, 3H), 5.02 (s, 2H), 3.06 (s, 3H), 2.88 (s, 3H).	DMSO	452 (M+1)	1.99	Method C	100	Method W

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
833		HCl	566.88	¹ H NMR (300 MHz, DMSO) δ 11.18 (s, 1H), 10.93 (s, 1H), 9.48 (s, 1H), 9.08 (s, 1H), 8.42 (d, J = 8.4 Hz, 1H), 8.94 (d, J = 5.0 Hz, 1H), 8.42 (s, 1H), 8.23 (dd, J = 6.8, 2.5 Hz, 1H), 8.12 - 7.89 (m, 3H), 7.63 (d, J = 9.1 Hz, 1H), 7.54 (t, J = 9.1 Hz, 1H), 4.45 - 4.26 (m,	DMSO	494 (M+1)	2.10	Method C	100	Methods V, G4

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
834			366.78	¹ H NMR (300 MHz, DMSO) δ 9.98 - 9.59 (m, 1H), 9.53 - 9.41 (m, 1H), 8.71 - 8.48 (m, 2H), 8.31 (dd, J = 6.9, 2.5 Hz, 1H), 8.01 - 7.85 (m, 1H), 7.83 - 7.66 (m, 2H), 7.59 - 7.35 (m, 3H).	DMSO	367 (M+1)	1.69	Method D	100	Method U

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
835			428.44	¹ H NMR (300 MHz, DMSO) δ 9.58 (s, 1H), 9.12 (d, J = 8.4 Hz, 1H), 8.82 - 8.64 (m, 2H), 8.48 (s, 1H), 8.32 - 8.17 (m, 1H), 8.10 - 7.87 (m, 3H), 7.81 - 7.49 (m, 4H), 7.21 (t, J = 7.5 Hz, 1H), 4.66 (s, 2H), 2.71 (d, J = 4.5 Hz, 3H).	DMSO	429 (M+1)	1.65	Method C	100	Method W

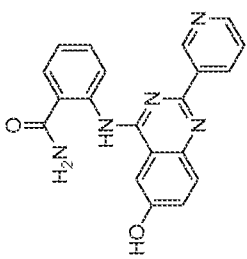
(continued)

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
836			484.51	¹ H NMR (300 MHz, DMSO) δ 9.58 (s, 1H), 9.13 (d, J = 8.3 Hz, 1H), 8.78 - 8.64 (m, 2H), 8.50 (s, 1H), 8.02 - 7.80 (m, 3H), 7.73 (t, J = 7.9 Hz, 1H), 7.68 - 7.53 (m, 2H), 7.49 (s, 1H), 7.20 (t, J = 7.5 Hz, 1H), 5.07 (s, 2H), 3.79 - 3.40 (m, 8H).	DMSO	485 (M+1)	1.72	Method C	91	Method W

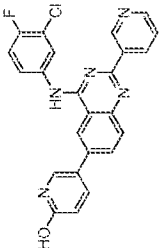
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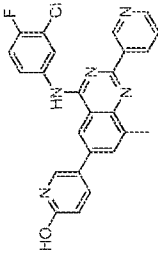
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
837			442.47	¹ H NMR (300 MHz, DMSO) δ 9.59 (s, 1H), 9.12 (d, J = 8.4 Hz, 1H), 8.78 - 8.63 (m, 1H), 8.49 (s, 1H), 8.29 (s, 1H), 8.11 (s, 1H), 7.88 (m, 2H), 7.79 - 7.47 (m, 4H), 7.26 - 7.14 (m, 1H), 4.58 (s, 2H), 3.27 - 3.08 (m, 2H), 1.18 - 0.93 (m, 3H).	DMSO	443 (M+1)	1.69	Method C	100	Method W

(continued)

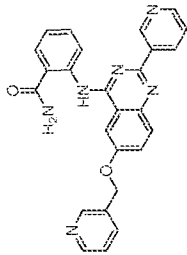
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
838			357.37	¹ H NMR (300 MHz, DMSO) δ 12.80 (s, 1H), 10.50 (s, 1H), 9.57 (s, 1H), 9.15 (s, 1H), 8.5 (d, J = 8.5 Hz, 1H), 8.70 (t, J = 7.3 Hz, 2H), 8.50 (s, 1H), 8.08 - 7.79 (m, 3H), 7.73 (t, J = 7.5 Hz, 1 H), 7.57 (dd, J = 7.6, 4.8 Hz, 1 H), 7.46 (d, J = 8.5 Hz, 2H), 7.19 (t, J = 7.6 Hz, 1H).	DMSO	357.9 (M+1)	1.68	Method C	100	Method U

(continued)

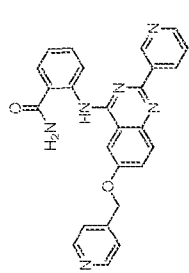
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
839		2HCl	443.86	¹ H NMR (300 MHz, DMSO) δ 10.57 (s, 1H), 9.49 (s, 1H), 8.97 (d, 1H), 8.89 (s, 1H), 8.81 (s, 1H), 8.16 (ddd, J = 12.2, 10.9, 5.2 Hz, 3H), 8.05 (d, J = 2.3 Hz, 1H), 8.00 - 7.86 (m, 3H), 7.55 (t, J = 9.1 Hz, 1H), 6.55 (d, J = 9.5 Hz, 1H).	DMSO	443.9 (M+1)	1.87	Method C	100	Method U

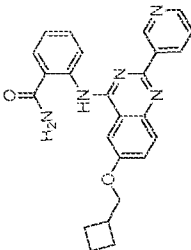
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
840		2HCl	457.89	¹ H NMR (300 MHz, DMSO) δ 10.27 (s, 1H), 9.47 (s, 1H), 9.09 (d, 1H), 8.91 (d, 1H), 8.56 (s, 1H), 8.18 - 8.08 (m, 2H), 8.01 (d, J = 6.3 Hz, 3H), 7.96 - 7.89 (m, 1H), 7.49 (t, J = 9.1 Hz, 1H), 6.52 (d, J = 9.6 Hz, 1H), 2.71 (s, 3H).	DMSO	458.0 (M+1)	2.12	Method D	100	Method U

(continued)

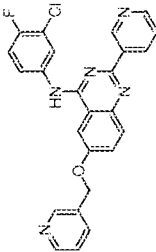
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
841			448.48	¹ H NMR (300 MHz, DMSO) δ 13.08 (s, 1H), 9.59 (s, 1H), 9.16 (d, J = 8.4 Hz, 1H), 8.88 - 8.44 (m, 5H), 8.12 - 7.85 (m, 4H), 7.73 (q, J = 8.7 Hz, 3H), 7.58 (dd, J = 7.9, 4.8 Hz, 1H), 7.49 (dd, J = 7.8, 4.8 Hz, 1H), 7.21 (t, J = 7.6 Hz, 1H), 5.36 (s, 2H).	DMSO	449.2 (M+1)	1.93	Method C	96	Method X

(continued)

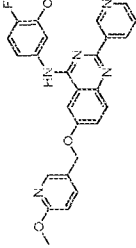
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
842			448.48	¹ H NMR (300 MHz, DMSO) δ 13.00 (s, 1H), 9.57 (s, 1H), 9.12 (d, 1H), 8.76 - 8.60 (m, 4H), 8.50 (s, 1H), 8.03 - 7.89 (m, 3H), 7.68 (dd, J = 15.6, 5.2 Hz, 3H), 7.57 (t, J = 4.6 Hz, 3H), 7.19 (t, J = 7.6 Hz, 1H), 5.37 (s, 2H).	DMSO	449.1 (M+1)	1.89	Method C	100	Method X

Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
843			425.48	¹ H NMR (300 MHz, DMSO) δ 12.95 (s, 1H), 9.56 (s, 1H), 9.10 (d, J = 8.4 Hz, 1H), 8.75 - 8.64 (m, 2H), 8.46 (s, 1H), 7.98 - 7.91 (m, 2H), 7.85 (dd, J = 9.0, 0.8 Hz, 1H), 7.71 (t, J = 7.9 Hz, 1H), 7.61 - 7.48 (m, 3H), 7.18 (t, J = 7.6 Hz, 1H), 4.12 (d, J = 6.6 Hz, 2H), 2.84 (dd, J = 14.3, 7.4 Hz, 1H), 2.18 - 2.05 (m, 2H), 1.97 - 1.84 (m, 5H).	DMSO	426.2 (M+1)	2.46	Method C	100	Method X

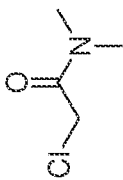
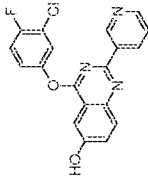
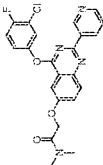
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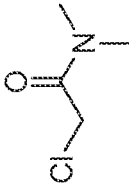
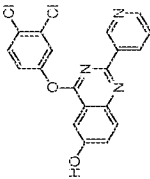
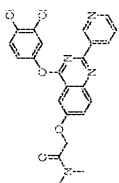
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
844			457.89	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.95 (s, 1H), 9.54 (d, J = 1.6 Hz, 1H), 8.82 (d, J = 1.7 Hz, 1H), 8.78 - 8.59 (m, 3H), 8.29 (dd, J = 6.8, 2.6 Hz, 1H), 8.16 (d, J = 2.5 Hz, 1H), 8.06 - 7.86 (m, 3H), 7.69 (dd, J = 9.1, 2.5 Hz, 1H), 7.64 - 7.47 (m, 3H), 5.37 (s, 2H).	DMSO	458.1, 460.1 (M+1), 229.6, 230.3 (M/2+1)		Method A	95	Method X

(continued)

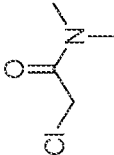
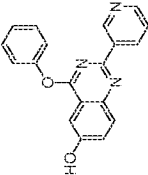
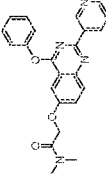
Number	Product	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time (min.)	LCMS Protocol	Purity Percent	Method for Coupling
845			487.91	¹ H-NMR (400 MHz, DMSO- <i>d</i> ₆): δ 9.74 (s, 1H), 9.34 (s, 1H), 8.57 - 8.41 (m, 2H), 8.22 (d, <i>J</i> = 2.2 Hz, 1H), 8.10 (dd, <i>J</i> = 6.8, 2.6 Hz, 1H), 7.95 (d, <i>J</i> = 2.5 Hz, 1H), 7.80 - 7.63 (m, 3H), 7.53 - 7.30 (m, 3H), 6.82 - 6.68 (m, 1H), 5.07 (s, 2H), 3.71 (s, 3H).	DMSO	488.1, 490.1 (M+1), 244.6, 245.4 (M/2+ 1)		Method A	95	Method X

(continued)

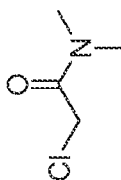
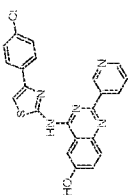
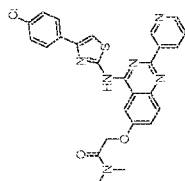
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
846				HCl	¹ H NMR (400 MHz, DMSO) δ 9.31 (d, J = 1.7 Hz, 1H), 8.83 - 8.76 (m, 1H), 8.76 - 8.67 (m, 1H), 8.07 (d, J = 9.2 Hz, 1H), 7.86 (dd, J = 6.3, 2.8 Hz, 1H), 7.83 - 7.73 (m, 2H), 7.70 (d, J = 2.8 Hz, 1H), 7.68 - 7.61 (m, 1H), 7.60 - 7.52 (m, 1H), 5.12 (s, 2H), 3.05 (s, 3H), 2.88 (s, 3H).	DMSO	>98	Method X using K ₂ CO ₃ instead of Cs ₂ CO ₃

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
847					¹ H NMR (400 MHz, DMSO) δ 9.28 (dd, J = 2.2, 0.8 Hz, 1H), 8.66 (dd, J = 4.8, 1.7 Hz, 1H), 8.50 - 8.44 (m, 1H), 8.04 (d, J = 9.2 Hz, 1H), 7.92 (d, J = 2.7 Hz, 1H), 7.85 (d, J = 8.8 Hz, 1H), 7.75 (dd, J = 9.2, 2.9 Hz, 1H), 7.67 (d, J = 2.8 Hz, 1H), 7.56 (dd, J = 8.8, 2.7 Hz, 1H), 7.52 (ddd, J = 8.0, 4.8, 0.8 Hz, 1H), 5.10 (s, 2H), 3.05 (s, 3H), 2.88 (s, 3H).	DMSO	>98	Method X using K ₂ CO ₃ instead of Cs ₂ CO ₃

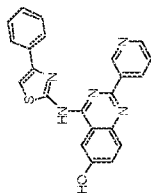
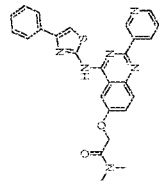
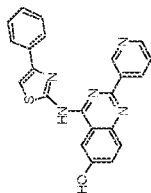
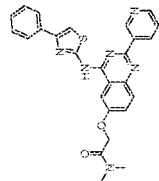
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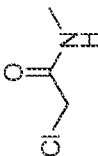
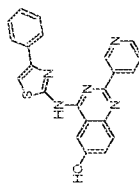
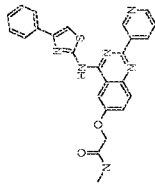
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
848					¹ H NMR (400 MHz, CDCl ₃) δ 9.45 (dd, J = 2.1, 0.8 Hz, 1H), 8.61 (dd, J = 4.8, 1.7 Hz, 1H), 8.59 - 8.53 (m, 1H), 8.01 (dd, J = 8.7, 0.8 Hz, 1H), 7.72 - 7.62 (m, 2H), 7.56 - 7.47 (m, 2H), 7.39 - 7.29 (m, 4H), 4.90 (s, 2H), 3.15 (s, 3H), 3.04 (s, 3H).	CDCl ₃	>98	Method X using K ₂ CO ₃ instead of Cs ₂ CO ₃

(continued)

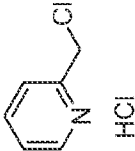
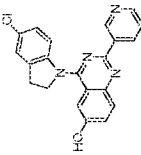
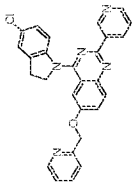
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
849				2HCl	1H NMR (300 MHz, DMSO) δ 13.01–11.72 (m, 1H), 9.70 (s, 1H), 9.40 - 9.30 (m, 1H), 8.99 (d, J = 4.5 Hz, 1H), 8.23 (d, J = 2.6 Hz, 1H), 8.12 (dd, J = 8.1, 5.5 Hz, 1H), 8.06 - 7.98 (m, 2H), 7.92 (d, J = 9.1 Hz, 1H), 7.84 (d, J = 4.9 Hz, 1H), 7.63 (dd, J = 9.1, 2.6 Hz, 1H), 7.56 - 7.49 (m, 2H), 5.05 (s, 3H), 3.11 (s, 3H), 2.92 (s, 3H).	DMSO	>98	W

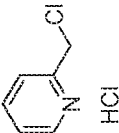
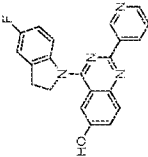
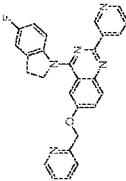
(continued)

5	850 			2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.74 (d, J = 1.8 Hz, 1H), 9.47 - 9.36 (m, 1H), 9.01 (dd, J = 5.5, 1.2 Hz, 1H), 8.28 (d, J = 2.6 Hz, 1H), 8.16 (dd, J = 8.2, 5.5 Hz, 1H), 8.06 - 7.93 (m, 3H), 7.81 (s, 1H), 7.66 (dd, J = 9.1, 2.6 Hz, 1H), 7.54 - 7.44 (m, 2H), 7.42 - 7.33 (m, 1H), 5.07 (s, 2H), 3.11 (s, 3H), 2.91 (s, 3H).	DMSO	>98	W	
10									
15									
20									
25									
30									
35									
40	850 			2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.74 (d, J = 1.8 Hz, 1H), 9.47 - 9.36 (m, 1H), 9.01 (dd, J = 5.5, 1.2 Hz, 1H), 8.28 (d, J = 2.6 Hz, 1H), 8.16 (dd, J = 8.2, 5.5 Hz, 1H), 8.06 - 7.93 (m, 3H), 7.81 (s, 1H), 7.66 (dd, J = 9.1, 2.6 Hz, 1H), 7.54 - 7.44 (m, 2H), 7.42 - 7.33 (m, 1H), 5.07 (s, 2H), 3.11 (s, 3H), 2.91 (s, 3H).	DMSO	>98	W	
45									
50									
55									

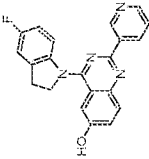
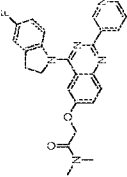
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
851				2HCl	¹ H NMR (300 MHz, DMSO) δ 12.67 (s, 1H), 9.31 - 9.25 (m, 1H), 8.73 (dd, J = 4.8, 1.6 Hz, 1H), 8.52 - 8.42 (m, 1H), 7.74 (d, J = 8.9 Hz, 1H), 7.66 (d, J = 2.9 Hz, 1H), 7.61 - 7.47 (m, 4H), 7.45 - 7.31 (m, 3H), 5.27 (s, 2H), 3.30 (s, 3H).	DMSO	>98	W

(continued)

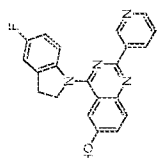
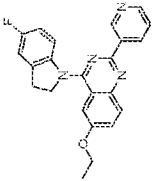
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
852				3HCl	¹ H NMR (300 MHz, DMSO) δ = 9.56 (d, J = 1.8 Hz, 1H), 9.25 (d, J = 8.2 Hz, 1H), 9.08 - 8.98 (m, 1H), 8.81 (d, J = 4.5 Hz, 1H), 8.31 (td, J = 7.8, 1.6 Hz, 1H), 8.20 - 8.08 (m, 1H), 7.95 (d, J = 7.9 Hz, 1H), 7.88 - 7.63 (m, 4H), 7.47 (d, J = 2.1 Hz, 1H), 7.30 (dd, J = 8.6, 2.2 Hz, 1H), 5.60 (s, 2H), 4.63 (t, J = 7.8 Hz, 2H), 3.25 (t, J = 7.7 Hz, 2H).	DMSO	>98	W

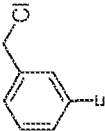
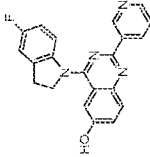
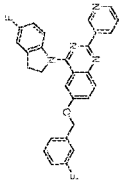
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
853				3HCl	¹ H NMR (300 MHz, DMSO) δ 9.56 (d, J = 1.8 Hz, 1H), 9.22 (d, J = 8.3 Hz, 1H), 9.00 (dd, J = 5.5, 1.3 Hz, 1H), 8.80 (d, J = 4.4 Hz, 1H), 8.29 (td, J = 7.8, 1.6 Hz, 1H), 8.21 - 8.05 (m, 2H), 8.00 - 7.66 (m, 5H), 7.29 (dd, J = 8.4, 2.6 Hz, 1H), 7.12 (td, J = 9.0, 2.8 Hz, 1H), 5.59 (s, 2H), 4.66 (t, J = 7.8 Hz, 2H), 3.25 (t, J = 7.6 Hz, 2H).	DMSO	>98	W

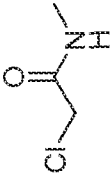
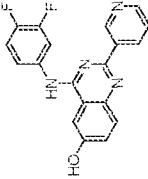
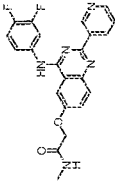
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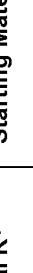
5	854	Starting Material R ¹		Starting Material R ³		Product		Salt Type	2HCl	1 ^H NMR	1 ^H NMR (300 MHz, DMSO) δ 9.55 (d, J = 1.8 Hz, 1H), 9.30 - 9.18 (m, 1H), 9.00 (dd, J = 5.5, 1.3 Hz, 1H), 8.18 - 8.02 (m, 2H), 7.87 (dd, J = 8.8, 4.8 Hz, 1H), 7.71 (dd, J = 9.2, 2.7 Hz, 1H), 7.49 (d, J = 2.7 Hz, 1H), 7.29 (dd, J = 8.4, 2.6 Hz, 1H), 7.13 (td, J = 9.1, 2.8 Hz, 1H), 5.05 (s, 2H), 4.63 (d, J = 7.7 Hz, 2H), 3.25 (t, J = 7.5 Hz, 2H), 3.02 (s, 3H), 2.86 (s, 3H).	¹ H NMR Solvent	DMSO	Purity percent	>98	Method of Coupling	W
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(continued)

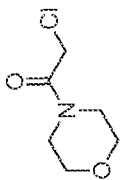
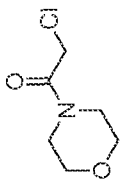
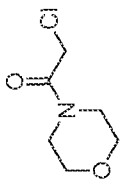
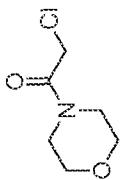
5	Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	1H NMR	1H NMR Solvent	Purity percent	Method of Coupling
10									
15	855				2HCl	1H NMR (300 MHz, DMSO) δ 9.55 (d, J = 1.7 Hz, 1H), 9.15 (d, J = 8.2 Hz, 1H), 8.99 - 8.90 (m, 1H), 8.10 - 7.98 (m, 2H), 7.77 (dd, J = 8.9, 4.7 Hz, 1H), 7.67 (dd, J = 9.1, 2.7 Hz, 1H), 7.51 (d, J = 2.6 Hz, 1H), 7.28 (dd, J = 8.4, 2.7 Hz, 1H), 7.12 (dd, J = 10.3, 7.7 Hz, 1H), 4.70 (t, J = 7.8 Hz, 2H), 4.23 - 4.14 (m, 2H), 3.26 (t, J = 7.9 Hz, 2H), 1.41 (t, J = 6.9 Hz, 3H).	DMSO	>98	W
20									
25									
30									
35									
40									
45									
50									
55									

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
856				2HCl	¹ H NMR (300 MHz, DMSO) δ = 9.54 (d, J = 1.7 Hz, 1H), 9.19 (d, J = 8.1 Hz, 1H), 8.98 (d, J = 4.3 Hz, 1H), 8.15 - 8.02 (m, 2H), 7.90 - 7.72 (m, 2H), 7.60 (d, J = 2.6 Hz, 1H), 7.53 - 7.41 (m, 1H), 7.40 - 7.05 (m, 5H), 5.34 (s, 2H), 4.59 (t, J = 7.8 Hz, 2H), 3.22 (t, J = 7.5 Hz, 2H).	DMSO	>98	W

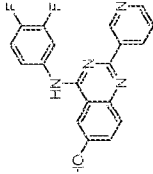
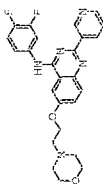
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
857				HCl	¹ H NMR (400 MHz, DMSO) δ 10.56 (s, 1H), 9.49 (d, J = 1.8 Hz, 1H), 9.11 (d, J = 8.0 Hz, 1H), 8.95 (dd, J = 5.4, 1.3 Hz, 1H), 8.26 (dd, J = 24.6, 3.5 Hz, 2H), 8.12 - 7.95 (m, 3H), 7.71 (dt, J = 9.1, 4.5 Hz, 2H), 7.56 (dd, J = 19.7, 9.1 Hz, 1H), 4.75 (s, 2H), 2.72 (d, J = 5.0 Hz, 3H).	DMSO	>98	Method W

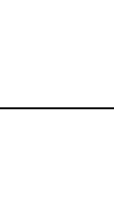
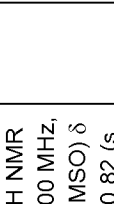
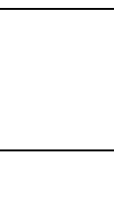
(continued)						
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent
858					¹ H NMR (400 MHz, DMSO) δ 9.77 (br s, J = 81.7 Hz, 1H), 9.50 (d, J = 1.5 Hz, 1H), 8.71 - 8.57 (m, 2H), 8.22 - 8.07 (m, 1H), 7.97 (t, J = 5.1 Hz, 1H), 7.87 (d, J = 9.1 Hz, 1H), 7.81 - 7.39 (m, 4H), 5.01 (s, 2H), 3.07 (s, 3H), 2.90 (s, 3H).	DMSO
					Purity percent	Method of Coupling
					>98	Method W

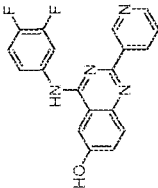
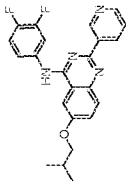
(continued)

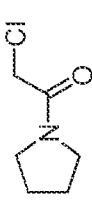
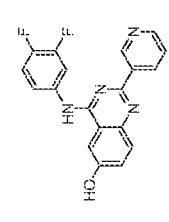
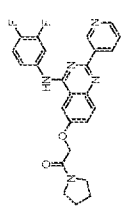
5	859 	Starting Material R¹ 	Starting Material R³ 	Product 	Salt Type HCl	¹H NMR ¹H NMR (400 MHz, DMSO) δ 10.62 (s, 1H), 9.50 (d, J = 1.7 Hz, 1H), 9.01 (d, J = 8.1 Hz, 1H), 8.90 (dd, J = 5.3, 1.5 Hz, 1H), 8.32 (d, J = 2.3 Hz, 1H), 8.20 - 8.09 (m, 1H), 7.95 (dd, J = 8.5, 4.7 Hz, 2H), 7.83 - 7.74 (m, 1H), 7.67 (dd, J = 9.1, 2.6 Hz, 1H), 7.55 (dd, J = 19.7, 9.1 Hz, 1H), 5.15 (s, 2H), 4.29 - 3.00 (m, 8H).	¹H NMR Solvent DMSO	Purity percent >98	Method of Coupling Method W
10									
15									
20									
25									
30									
35									
40									
45									
50									
55									

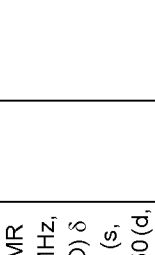
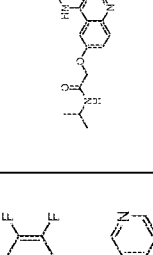
(continued)

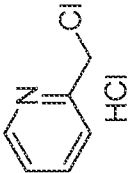
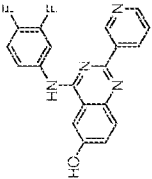
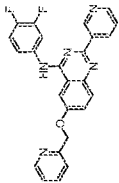
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
860				2HCl	¹ H NMR (400 MHz, DMSO) δ 10.39 (s, 1H), 9.51 (d, J = 1.6 Hz, 1H), 8.81 (dd, J = 19.3, 6.0 Hz, 2H), 8.29 (s, 1H), 8.16 (dd, J = 10.6, 7.5 Hz, 1H), 7.94 (d, J = 9.0 Hz, 1H), 7.77 (s, 2H), 7.69 – 7.51 (m, 2H), 4.67 (s, 2H), 3.99 (s, 2H), 3.81 (br s, 4H), 3.70 (br s, 4H).	DMSO	>98	Method W

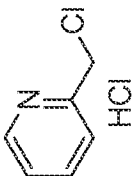
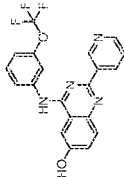
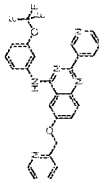
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
861				3HCl	¹ H NMR (400 MHz, DMSO) δ 10.82 (s, 1H), 9.50 (d, J = 1.3 Hz, 1H), 9.02 (d, J = 8.0 Hz, 1H), 8.90 (d, J = 4.4 Hz, 1H), 8.46 (s, 1H), 8.21 - 8.10 (m, 1H), 8.03 - 7.90 (m, 2H), 7.84 (d, J = 8.7 Hz, 1H), 7.68 (dd, J = 9.1, 2.3 Hz, 1H), 7.55 (dd, J = 19.6, 9.3 Hz, 1H), 4.67 (s, 2H), 3.23 - 3.05 (m, 6H), 2.10 - 1.78 (m, 4H).	DMSO	>98	Method W

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
862				2HCl	¹ H NMR (400 MHz, DMSO) δ 10.77 (s, 1H), 9.49 (d, J = 1.6 Hz, 1H), 9.12 (d, J = 8.1 Hz, 1H), 8.99 - 8.91 (m, 1H), 8.23 (d, J = 2.1 Hz, 1H), 8.12 - 7.99 (m, 3H), 7.78 - 7.71 (m, 1H), 7.66-7.54 (m, 2H), 4.02 (d, J = 6.5 Hz, 2H), 2.24 - 2.00 (m, 1H), 1.07 (d, J = 6.7 Hz, 6H).	DMSO	>98	Method W

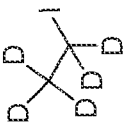
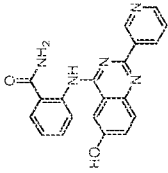
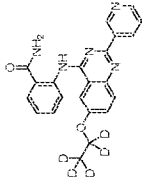
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
863				2HCl	¹ H NMR (400 MHz, CDCl ₃) δ 10.71 (s, 1H), 9.80 (s, 1H), 9.40 (d, 1H), 9.40 (d, 1H), 8.2 Hz, J = 8.2 Hz, 1H), 8.95 (d, 1H), 8.95 (d, 1H), 8.54 (d, 1H), 8.54 (d, 1H), 8.24 (d, 1H), 8.24 (d, 1H), 9.2 Hz, J = 9.2 Hz, 1H), 8.08 (dd, J = 8.0, 5.6 Hz, 1H), 7.86 - 7.76 (m, 1H), 7.69 (d, J = 8.7 Hz, 1H), 7.56 (d, J = 8.8, 2.2 Hz, 1H), 7.31 (s, 1H), 5.08 (s, 2H), 3.69 (t, J = 6.8 Hz, 2H), 3.55 (t, J = 6.9 Hz, 2H), 2.13 - 2.01 (m, 2H), 1.98 - 1.85 (m, 2H).	DMSO	>98	Method W

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	1H NMR	1H NMR Solvent	Purity percent	Method of Coupling
864				2HCl	1H NMR (400 MHz, DMSO) δ 10.62 (s, 1H), 9.50 (d, J = 1.9 Hz, 1H), 9.12 (d, J = 8.2 Hz, 1H), 8.96 (dd, J = 5.5, 1.4 Hz, 1H), 8.27 (d, J = 2.6 Hz, 1H), 8.20 (d, J = 8.0 Hz, 1H), 8.15 - 7.97 (m, 3H), 7.75 (ddd, J = 11.9, 7.4, 3.3 Hz, 2H), 7.56 (dt, J = 10.6, 9.1 Hz, 1H), 4.74 (s, 2H), 4.07 - 3.93 (m, 1H), 1.13 (d, J = 6.6 Hz, 6H)..	DMSO	>98	Method W

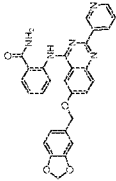
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
865				3HCl	¹ H NMR (400 MHz, DMSO) δ 10.83 (s, 1H), 9.51 (d, J = 1.8 Hz, 1H), 9.15 (d, J = 8.2 Hz, 1H), 8.97 (dd, J = 5.4, 1.3 Hz, 1H), 8.82 - 8.74 (m, 1H), 8.55 (d, J = 2.6 Hz, 1H), 8.28 - 8.19 (m, 1H), 8.16 - 8.01 (m, 3H), 7.94 (d, J = 7.9 Hz, 1H), 7.85 - 7.75 (m, 2H), 7.70 (dd, J = 7.0, 5.8 Hz, 1H), 7.61 - 7.49 (m, 1H), 5.63 (s, 2H).	DMSO	>98	Method W

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
866				4HCl	¹ H NMR (400 MHz, DMSO) δ 10.62 (s, 1H), 9.50 (d, J = 1.8 Hz, 1H), 9.11 (d, J = 8.1 Hz, 1H), 8.95 (d, J = 5.4 Hz, 1H), 8.76 (d, J = 5.0 Hz, 1H), 8.42 (s, 1H), 8.18 (t, J = 7.8 Hz, 1H), 8.08 – 7.96 (m, 4H), 7.89 (d, J = 7.8 Hz, 1H), 7.80 (dd, J = 9.1, 2.6 Hz, 1H), 7.64 (t, J = 8.2 Hz, 2H), 7.24 (d, J = 8.3 Hz, 1H), 5.57 (s, 2H).	DMSO	>98	Method W

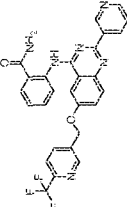
(continued)

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
867				HCl	¹ H NMR (300 MHz, DMSO) d 9.58 (s, 1H), 9.05 (d, J = 8.2 Hz, 1H), 8.98 (d, J = 8.1 Hz, 1H), 8.88 (d, J = 5.1 Hz, 1H), 8.47 (s, 1H), 8.05 - 7.85 (m, 4H), 7.71 (m, 1H), 7.61 (d, J = 8.1 Hz, 2H), 7.24 (dd, J = 7.6 Hz, 1H).	DMSO	94	Method X (K ₂ CO ₃ , DMF - THF (1 : 1), rt)

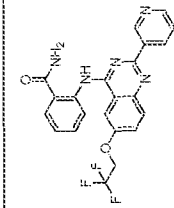
(continued)

5	868 	Starting Material R ¹	Starting Material R ³	Product	Salt Type	1H NMR (300 MHz, DMSO) δ 13.07 (s, 1H), 9.59 (d, J = 1.4 Hz, 1H), 9.13 (d, J = 8.1 Hz, 1H), 8.97 (d, J = 7.8 Hz, 1H), 8.92 (d, J = 4.1 Hz, 1H), 8.49 (s, 1H), 8.08 - 7.92 (m, 4H), 7.80 - 7.64 (m, 3H), 7.26 (td, J = 7.8, 1.1 Hz, 1H), 7.14 (d, J = 1.5 Hz, 1H), 7.09 (dd, J = 7.9, 1.6 Hz, 1H), 6.95 (d, J = 7.9 Hz, 1H), 6.03 (s, 2H), 5.19 (s, 2H).	DMSO	Purity percent	Method of Coupling
10									
15									
20									
25									
30									
35									
40									
45									
50									
55									

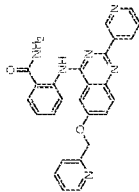
(continued)

5	869 	Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹H-NMR (400 MHz, DMSO-d₆): δ 9.67 (s, 1H), 9.40 (d, J = 7.6 Hz, 1H), 9.01 (d, J = 8.4 Hz, 2H), 8.93 (d, J = 8.4 Hz, 1H), 8.31 (d, J = 7.6 Hz, 1H), 8.22 (t, J = 7.2 Hz, 1H), 8.01 (d, J = 9.2 Hz, 1H), 7.97 - 7.92 (m, 2H), 7.79 - 7.71 (m, 3H), 7.33 (t, J = 8.0 Hz, 1H), 5.54 (s, 2H).	¹H NMR Solvent DMSO	Purity percent 98	Method of Coupling Method X
10										
15										
20										
25										
30										
35										
40										
45										
50										
55										

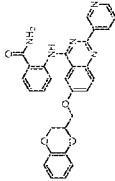
(continued)

5	Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	1H NMR	1H NMR Solvent	Purity percent	Method of Coupling
10									
15	870				2HCl	¹ H-NMR (400 MHz, CD3OD): δ 9.62 (d, J = 6.9 Hz, 1H), 9.40 (d, J = 8.2 Hz, 1H), 8.99 (d, J = 5.3 Hz, 1H), 8.82 (t, J = 8.2 Hz, 1H), 8.22 (dd, J = 8.1, 5.7 Hz, 1H), 8.02 - 7.88 (m, 2H), 7.76 - 7.60 (m, 3H), 7.31 (t, J = 7.6 Hz, 1H), 4.81 (q, J = 8.1 Hz, 2H).	MeOD	95	Method W
20									
25									
30									
35									
40									
45									
50									
55									

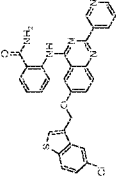
(continued)

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
871				4HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.96 (s, 1H), 9.59 (s, 1H), 9.24 (d, 1H), 8.1 Hz, J = 8.1 Hz, 1H), 9.00 (s, 1H), 8.80 (d, J = 5.7 Hz, 2H), 8.50 (s, 1H), 8.22 (t, J = 7.2 Hz, 1H), 8.09 (d, J = 9.2 Hz, 2H), 7.99 - 7.79 (m, 5H), 7.74 - 7.54 (m, 6H), 7.30 (t, J = 7.5 Hz, 1H), 5.55 (s, 2H).	DMSO-d ₆	95	Method X

(continued)

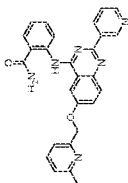
5	872 	Starting Material R ¹	Starting Material R ³	Product	Salt Type	1H-NMR (400 MHz, DMSO-d₆): δ 12.82 (s, 1H), 9.58 (s, 1H), 9.13 (d, 1H), 9.13 (d, 1H), 8.93 (s, 1H), 8.82 (d, 1H), 8.43 (s, 1H), 8.04 - 7.88 (m, 4H), 7.74 - 7.69 (m, 3H), 7.28 (t, J= 7.6 Hz, 1H), 7.01 - 6.91 (m, 2H), 6.87 - 6.85 (m, 2H), 4.74 - 4.70 (m, 1H), 4.55 - 4.53 (m, 1H), 4.50 - 4.47 (m, 1H), 4.29 - 4.24 (m, 1H).	DMSO	Purity percent	Method of Coupling
10									
15									
20									
25									
30									
35									
40									
45									
50									
55									

(continued)

5	873	Starting Material R ¹	Starting Material R ³	Product	Salt Type	1H NMR	1H NMR Solvent	Purity percent	Method of Coupling
10									
15					3HCl	1H-NMR (400 MHz, DMSO-d ₆): δ 13.07 (s, 1H), 9.61 (s, 1H), 9.05 - 9.00 (m, 2H), 8.88 (d, J = 4 Hz, 1H), 8.51 (s, 1H), 8.25 (s, 1H), 8.11 - 8.08 (m, 2H), 7.99 - 7.97 (m, 3H), 7.91 - 7.88 (m, 2H), 7.76 - 7.72 (m, 2H), 7.47 (dd, J = 2.0, 8.8 Hz, 1H), 7.27 (t, J = 8.8 Hz, 1H), 5.61 (s, 2H).	DMSO	95	Method X

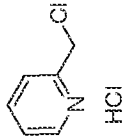
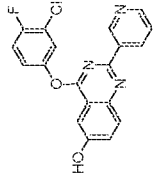
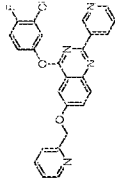
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
874			 <chem>Cc1ccc(cc1Oc2ccnc(c2)C(=O)N(C)C)[NH+]([O-])C3CCOC3.Cl</chem>	2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.89 (s, 1H), 10.72 (s, 1H), 9.83 (s, 1H), 9.38 (s, 1H), 9.38 (d, = 8.2 Hz, 1H), 9.22 (d, J = 5.6 Hz, 1H), 9.03 (d, J = 8.3 Hz, 1H), 8.51 (s, 1H), 8.37 - 8.27 (m, 1H), 8.02 - 7.86 (m, 3H), 7.71 (t, J = 7.6 Hz, 1H), 7.55 (d, J = 9.0 Hz, 1H), 7.51 (s, 1H), 7.25 (t, J = 7.5 Hz, 1H), 4.71 (d, J = 7.3 Hz, 2H), 1.56 (s, 1H), 0.81 - 0.64 (m, 4H).	DMSO	95	Method X

(continued)

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
875				3HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.88 (s, 1H), 9.57 (s, 1H), 9.24 (d, 1H), 9.24 (d, 1H), 8.1 Hz, J = 8.1 Hz, 1H), 9.01 (d, J = 5.3 Hz, 1H), 8.73 (d, J = 8.2 Hz, 1H), 8.49 (s, 1H), 8.37 (t, J = 7.8 Hz, 1H), 8.10 (dd, J = 11.3, 7.0 Hz, 2H), 7.97 (d, J = 7.8 Hz, 2H), 7.86 (s, 2H), 7.80 (d, J = 7.1 Hz, 2H), 7.71 (t, J = 7.8 Hz, 1H), 7.30 (t, J = 7.5 Hz, 1H), 5.65 (s, 2H), 2.78 (s, 3H).	DMSO	95	Method X

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
876				CF3COOH	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.93 (s, 1H), 10.74 (s, 1H), 9.75 (s, 1H), 9.40 (s, 1H), 9.40 (s, 1H), 9.05 (s, 1H), 8.0 (d, J = 8.0 Hz, 2H), 8.99 (d, J = 6.0 Hz, 1H), 7.99 (s, 1H), 7.98 - 7.96 (m, 1H), 7.89 - 7.86 (m, 2H), 7.72 - 7.51 (m, 2H), 7.24 (t, J = 6.0 Hz, 1H), 5.84 (s, 2H), 3.60 - 3.56 (m, 2H), 3.43 - 3.39 (m, 2H), 2.06 - 2.01 (m, 2H), 1.99 - 1.84 (m, 2H).	DMSO	95	Method W

(continued)

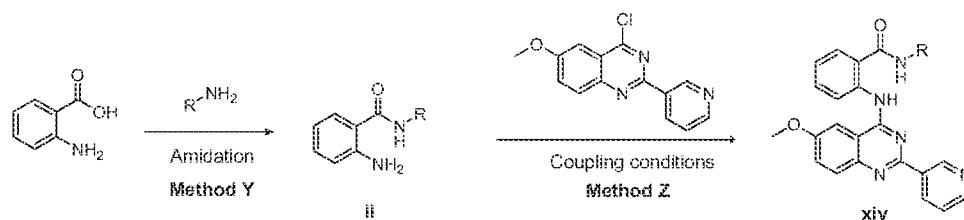
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1757	 <chem>ClCC1=CC=CC=N1</chem> HCl	 <chem>Oc1ccc2nc3cc(Oc4cc(F)c(Cl)cc4)nc(C5=CC=CC=N5)c2c1</chem>	 <chem>Oc1ccc2nc3cc(Oc4cc(F)c(Cl)cc4)nc(C5=CC=CC=N5)c2c1</chem>		¹ H NMR (400 MHz, DMSO) δ 9.32 - 9.27 (m, 1H), 8.66 (dd, J = 4.8, 1.7 Hz, 1H), 8.64 - 8.60 (m, 1H), 8.50 - 8.43 (m, 1H), 8.10 - 8.03 (m, 1H), 7.93 - 7.80 (m, 4H), 7.68 - 7.60 (m, 2H), 7.59 - 7.49 (m, 2H), 7.39 (ddd, J = 7.5, 4.8, 1.1 Hz, 1H), 5.43 (s, 2H).	DMSO	>98	Method W using K ₂ CO ₃ Instead of Cs ₂ CO ₃

Scheme 27: General route for the synthesis of compounds with general formula xiv

[0350]

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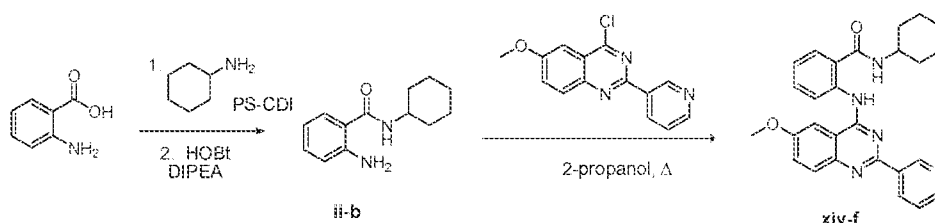
Scheme 28: Representative synthesis of compounds of formula xiv (see Scheme 27)

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[0351]

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[0352] **Method Y: 2-Amino-N-cyclohexyl-benzamide (ii-b)** To a dry reaction vial was added cyclohexylamine (40 μ mol, 1.0 eq) and PS-carbodiimide resin (72 μ mol, 1.8 eq). The solution of 2-amino-benzoic acid (44 μ mol, 1.1 eq), diisopropyl ethyl amine (10 μ L) and HOBT (44 μ mol, 1.1 eq) in THF (500 μ L) was added to the above vial. The reaction mixture was heated at 40°C for 6 h on a shaker. The resin was removed by filtration and washed with 10% MeOH/CH₂Cl₂. The solvent was removed *in vacuo* and the residue was applied to solid phase extraction cartridge (basic silica, 200 mg) and eluted with 50% EtOAc/CH₂Cl₂. After removal of the solvents, the crude 2-amino-N-cyclohexyl-benzamide was obtained and used for the following reaction.

[0353] **Method Z: N-Cyclohexyl-2-(6-methoxy-2-pyridin-3-yl-quinazolin-4-ylamino)-benzamide (xiv-f)** To the crude amide was added the solution of 4-chloro-6-methoxy-2-pyridin-3-yl-quinazoline (20 μ mol) in 2-propanol (200 μ L). The mixture was refluxed for 8 h. After evaporation, the residue was dissolved in 5% TFA/MeOH-DMF(1:1) and purified by **PREP-HPLC Condition D**. The target fraction was lyophilized to afford the titled compound whose structure was finally confirmed by LCMS using **LCMS Method E**.

[0354] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 27**, replacing cyclohexylamine with the appropriate amine.

Table 7:

Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
877		489	490	Methods Y, Z	884		467	468	Methods Y, Z
878		453	454	Methods Y, Z	885		441	442	Methods Y, Z
879		495	496	Methods Y, Z	886		529	530	Methods Y, Z
880		495	496	Methods Y, Z	887		543	544	Methods Y, Z
881		491	492	Methods Y, Z	888		543	544	Methods Y, Z

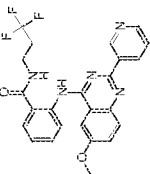
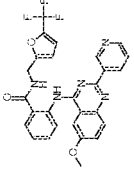
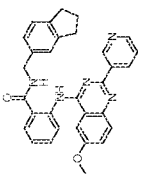
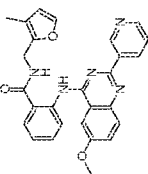
(continued)

Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
882		491	492	Methods Y, Z	889		543	544	Methods Y, Z
883		493	494	Methods Y, Z	890		487	488	Methods Y, Z
891		493	494	Methods	898		487	488	Methods
892		467	468	Methods Y, Z	899		467	468	Methods Y, Z
893		441	442	Methods Y, Z	900		501	502	Methods Y, Z

(continued)

Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
894		529	530	Methods Y, Z	901		465	466	Methods Y, Z
895		543	544	Methods Y, Z	902		435	436	Methods Y, Z
896		543	544	Methods Y, Z	903		519	520	Methods Y, Z
897		543	544	Methods Y, Z	904		487	488	Methods Y, Z
905		487	488	Methods Y, Z	909		435	436	Methods Y, Z

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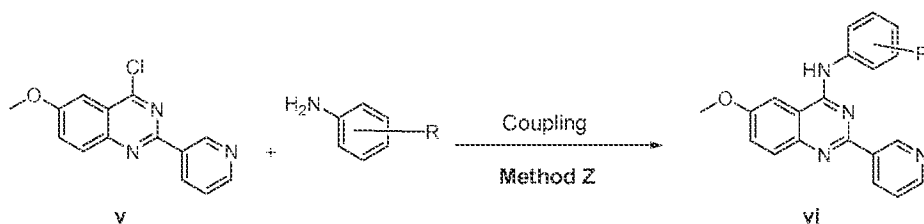
Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
906		467	468	Methods Y, Z	910		519	520	Methods Y, Z
907		501	502	Methods Y, Z					
908		465	466	Methods Y, Z					

Scheme 29: General route for the synthesis of compounds with general formula vi

[0355]

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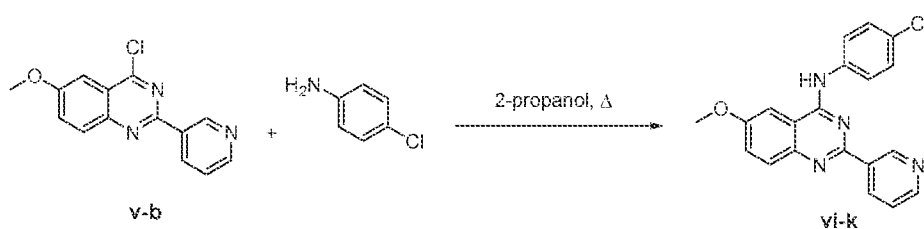
Scheme 30: Representative synthesis of compounds of formula vi-b (see Scheme 29)

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[0356]

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[0357] Method Z: Synthesis of **N-(4-chlorophenyl)-6-methoxy-2-(pyridin-3-yl)quinazolin-4-amine (vi-k)** To 4-chloroaniline (24 μ mol) was added the solution of 4-chloro-6-methoxy-2-pyridin-3-yl-quinazoline (20 μ mol) in 2-propanol (200 μ L). The mixture was refluxed for 8 h. After evaporation, the residue was dissolved in 5%TFA/MeOH-DMF(1:1) and purified by **PREP-HPLC** Condition **D**. The target fraction was lyophilized to afford the titled compound as the TFA salt whose structure was finally confirmed by LCMS using LCMS **Method E**.

[0358] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 29**, replacing 4-chloroaniline with the appropriate aniline.

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Table 8:

Number	Product	Exact Mass	LCMS (M+3)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
911		362	363	Method Z	918		342	343	Method Z
912		358	359	Method Z	919		353	354	Method Z
913		396	397	Method Z	920		416	417	Method Z
914		392	393	Method Z	921		396	397	Method Z
915		388	389	Method Z	922		396	397	Method Z
916		372	373	Method Z	923		356	357	Method Z

(continued)

Number	Product	Exact Mass	LCMS (M+3)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
917		388	389	Method Z	924		364	365	Method Z
925		372	373	Method Z	932		392	393	Method Z
926		392	393	Method Z	933		380	381	Method Z
927		376	377	Method Z	934		380	381	Method Z
928		426	427	Method Z	935		380	381	Method Z
929		407	408	Method Z	936		434	435	Method Z

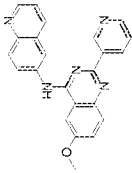
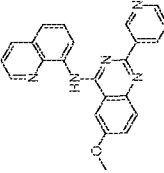
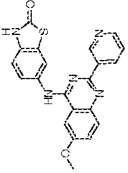
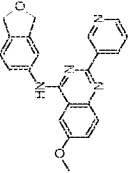
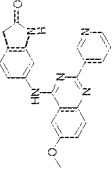
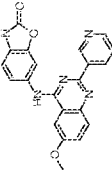
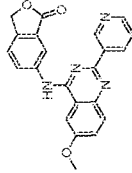
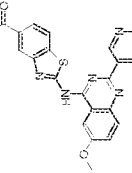
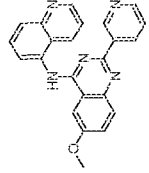
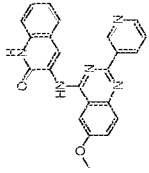
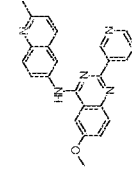
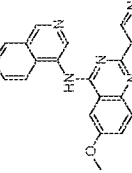
(continued)

Number	Product	Exact Mass	LCMS (M+3)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
930		430	431	Method Z	937		421	422	Method Z
931		440	441	Method Z	938		378	379	Method Z
939		376	377	Method Z	946		428	429	Method Z
940		387	388	Method Z	947		391	392	Method Z
941		414	415	Method Z	948		356	357	Method Z

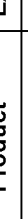
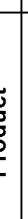
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Number	Product	Exact Mass	LCMS (M+3)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
942		392	393	Method Z	949		377	378	Method Z
943		380	381	Method Z	950		392	393	Method Z
944		390	391	Method Z	951		368	369	Method Z
945		428	429	Method Z	952		397	398	Method Z
953		441	442	Method Z	960		411	412	Method Z

(continued)

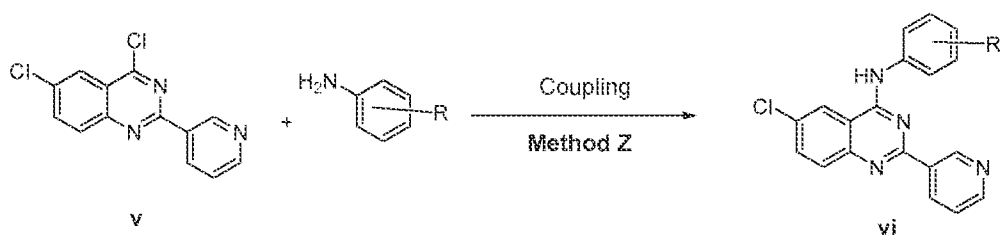
Number	Product	Exact Mass	LCMS (M+3)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
954		379	380	Method Z	961		379	380	Method Z
955		401	402	Method Z	962		370	371	Method Z
956		383	384	Method Z	963		385	386	Method Z
957		382	383	Method Z	964		427	428	Method Z
958		379	380	Method Z	965		395	396	Method Z
959		393	394	Method Z	966		379	380	Method Z

(continued)

Number	Product	Exact Mass	LCMS (M+3)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
967		402	403	Method Z	968		384	385	Method Z

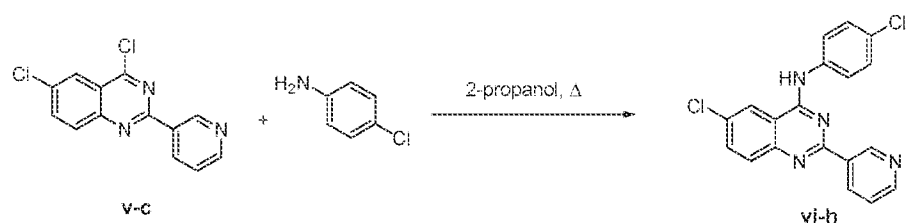
Scheme 31: General route for the synthesis of compounds with general formula vi

[0359]



Scheme 32: Representative synthesis of compounds of formula vi (see Scheme 31)

[0360]



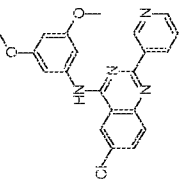
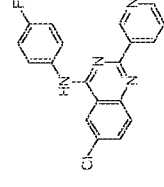
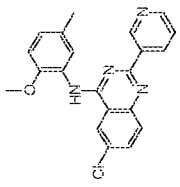
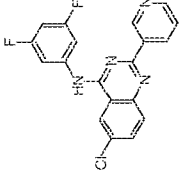
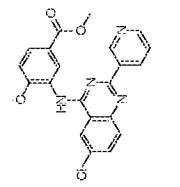
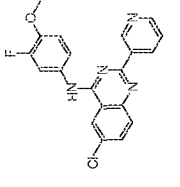
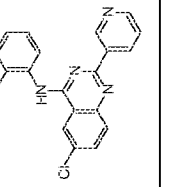
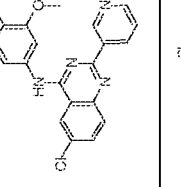
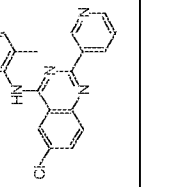
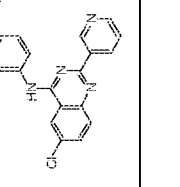
[0361] Method Z: Synthesis of 6-chloro-N-(4-chlorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine (vi-c): To 4-chloroaniline (24 μmol) was added the solution of 4,6-dichloro-2-(pyridin-3-yl)quinazoline (20 μmol) in 2-propanol (200 μL). The mixture was refluxed for 8 h. After evaporation, the residue was dissolved in 5% TFA/MeOH-DMF(1:1) and purified by **PREP-HPLC Condition D**. The target fraction was lyophilized to afford the titled compound as the TFA salt whose structure was finally confirmed by LCMS using LCMS **Method E**.

[0362] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 31**, replacing 4-chloroaniline with the appropriate aniline.

Table 9:

Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
969		366	367	Method Z	975		419	420	Method Z
970		362	363	Method Z	976		392	393	Method Z
971		366	367	Method Z	977		346	347	Method Z
972		346	347	Method Z	978		350	351	Method Z
973		392	393	Method Z	979		350	351	Method Z

(continued)

Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
974		392	393	Method Z	980		350	351	Method Z
981		376	377	Method Z	988		368	369	Method Z
982		420	421	Method Z	989		380	381	Method Z
983		360	361	Method Z	990		376	377	Method Z
984		360	361	Method Z	991		396	397	Method Z

(continued)

Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
985		360	361	Method Z	992		380	381	Method Z
986		360	361	Method Z	993		430	431	Method Z
987		368	369	Method Z	994		392	393	Method Z
995		396	397	Method Z	1002		425	426	Method Z
996		380	381	Method Z	1003		380	381	Method Z

(continued)

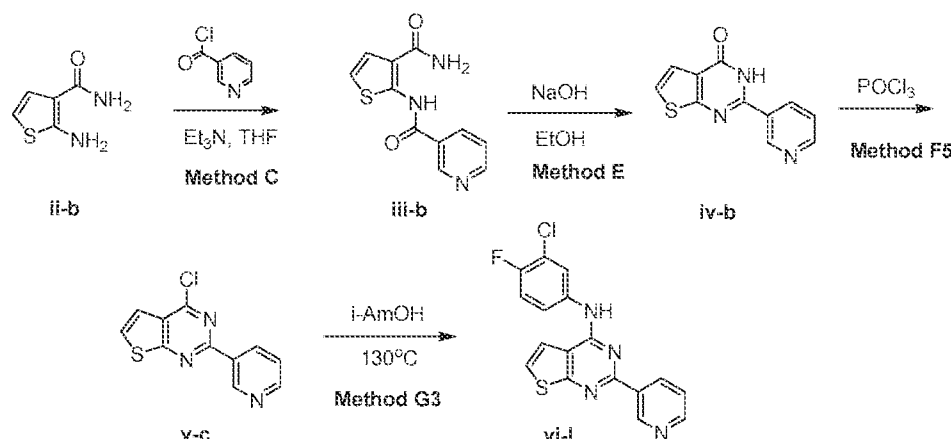
Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
997		382	383	Method Z	1004		434	435	Method Z
998		384	385	Method Z	1005		396	397	Method Z
999		465	466	Method Z	1006		384	385	Method Z
1000		357	358	Method Z	1007		446	447	Method Z
1001		396	397	Method Z	1008		415	416	Method Z
1009		381	382	Method Z	1011		396	397	Method Z

(continued)

Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling	Number	Product	Exact Mass	LCMS (M+1)	Method of Coupling
1010		381	382	Method Z					

Scheme 33: Representative synthesis of thiophene analogs of formula vi (see Scheme 4)

[0363]



[0364] N-(3-Carbamoylthiophen-2-yl)nicotinamide (iii-b) To a solution of 2-aminothiophene-3-carboxamide (800 mg, 5.63 mmol, 1.0 eq.) in THF (15 mL) and Et₃N (626 mg, 6.19 mmol, 1.1 eq.) was added nicotinoyl chloride (795 mg, 5.63 mmol, 1.0 eq.) in anhydrous THF (15 mL) dropwise. The resulted mixture was stirred at room temperature overnight. After the reaction was completed, the volatiles were evaporated. The residue was washed with CH₂Cl₂ (20 mL). The resulting solid was collected and dried *in vacuo* to give 1.50 g of N-(3-carbamoylthiophen-2-yl)nicotinamide as a brown solid (quantitative yield). LCMS *m/z* = 248.1(M+1) (Method B) (retention time = 1.34 min).

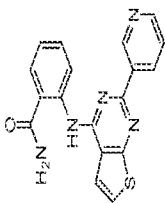
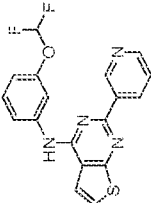
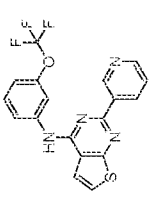
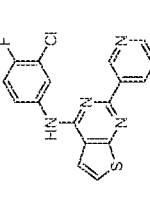
[0365] 2-(Pyridin-3-yl) thieno [2, 3-d] pyrimidin-4(3H)-one (iv-b) A mixture of N-(3-carbamoylthiophen-2-yl)nicotinamide (1.50 g salt, 6.07 mmol, 1.0 eq.) in EtOH (300 mL) was added NaOH (1.50 g, 37.5 mmol, 6.18 eq.). The resulting mixture was stirred at 80°C for 7 days. After the reaction was completed, the volatiles were removed *in vacuo*. Water (20 mL) was added to the residue and the pH was adjusted to around 2 by adding dilute HCl (2N in water). The solution was concentrated *in vacuo* to give 11.0 g of the HCl salt as a beige solid. LCMS *m/z* = 230.0 (M+1) (Method B) (retention time = 1.21 min). The crude product containing salts were used for the next step without further purification.

[0366] 4-Chloro-2-(pyridin-3-yl) thieno [2, 3-d] pyrimidine (v-c) The suspension of 2-(pyridin-3-yl)thieno[2,3-d]pyrimidin-4(3H)-one (6.0 g, containing salts) in POCl₃ (30 mL) was stirred at 120°C for 10 h. After the reaction was completed, the mixture was added to ice-water slowly. The pH was adjusted to ~7 by slowly adding NH₃·H₂O at 0°C, then a precipitate formed. The solid was collected and 540 mg of 4-chloro-2-(pyridin-3-yl) thieno [2, 3-d] pyrimidine was obtained as a brown solid. LCMS *m/z* = 247.9, 250.0(M+1) (Method B) (retention time = 1.85 min).

[0367] N-(3-Chloro-4-fluorophenyl)-2-(pyridin-3-yl)thieno[2,3-d]pyrimidin-4-amine (vi-1) A mixture of 4-chloro-2-(pyridin-3-yl)thieno[2,3-d]pyrimidine (80 mg, 0.32 mmol, 1.0 eq.) and 3-chloro-4-fluorobenzaniline (93 mg, 0.64 mmol, 2.0 eq.) in *i*-AmOH (8 mL) was stirred at 130°C overnight. A yellow precipitate formed and was collected and washed with MeOH (10 mL). The solid was suspended in H₂O (10 mL) and NH₃·H₂O (1 mL) was added. After filtration and drying *in vacuo*, 23.0 mg of the product was obtained as a yellow solid (20.0%). LCMS *m/z* = 357.0, 359.0 (M+1) (Method B) (retention time = 1.96 min). ¹H-NMR (400 MHz, DMSO-*d*₆): δ 9.99 (s, 1H), 9.48 (d, *J* = 1.6 Hz, 1H), 8.70 - 8.67 (m, 1H), 8.63 (d, *J* = 8.0 Hz, 1H), 8.27 (dd, *J* = 6.8, 2.6 Hz, 1H), 7.76 - 7.93 (m, 3H), 7.54 - 7.57 (m, 2H).

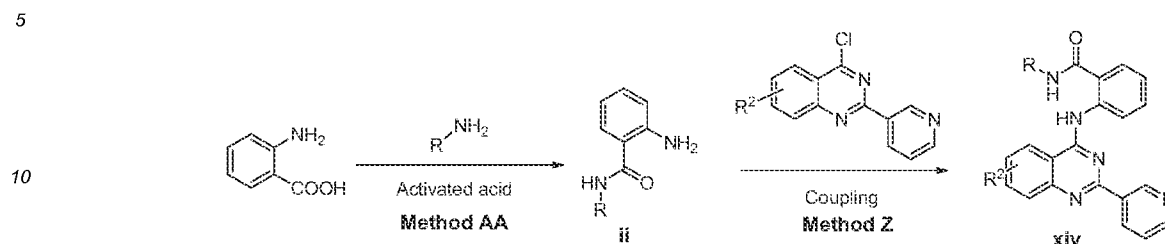
[0368] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 33**, replacing 4-fluoro-3-chloroaniline with the appropriate aniline.

Table 10:

Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Product	Purity percent	Method for Coupling
1012			347.39	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.69 (s, 1H), 9.55 (d, J = 1.3 Hz, 1H), 9.00 (d, J = 8.0 Hz, 1H), 8.44 (s, 1H), 8.60 - 8.78 (m, 2H), 7.82 - 8.00 (m, 3H), 7.42 - 7.79 (m, 3H), 7.20 (t, J = 7.2 Hz, 1H).	DMSO	348.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G3
1013			370.38	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.96 (s, 1H), 9.51 (d, J = 1.2 Hz, 1H), 8.60 - 8.77 (m, 2H), 7.98-7.94 (m, 2H), 7.71 - 7.86 (m, 2H), 7.56 - 7.50 (m, 1H), 7.30 (t, J = 74.0 Hz, 1H), 6.96 (dd, J = 8.8, 2.0 Hz, 1H).	DMSO	371.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G3
1014			388.37	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.04 (s, 1H), 9.50 (d, J = 1.6 Hz, 1H), 8.58 - 8.75 (m, 2H), 8.18 (s, 1H), 7.77 - 8.00 (m, 3H), 7.50-7.65 (m, 2H), 7.08 - 7.18 (m, 1H).	DMSO	389.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G3
1015			356.8	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.99 (s, 1H), 9.48 (d, J = 1.6 Hz, 1H), 8.70 - 8.67 (m, 1H), 8.63 (d, J = 8.0 Hz, 1H), 8.27 (dd, J = 6.8, 2.6 Hz, 1H), 7.76 - 7.93 (m, 3H), 7.54 - 7.57 (m, 2H).	DMSO	357.0, 359.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G3

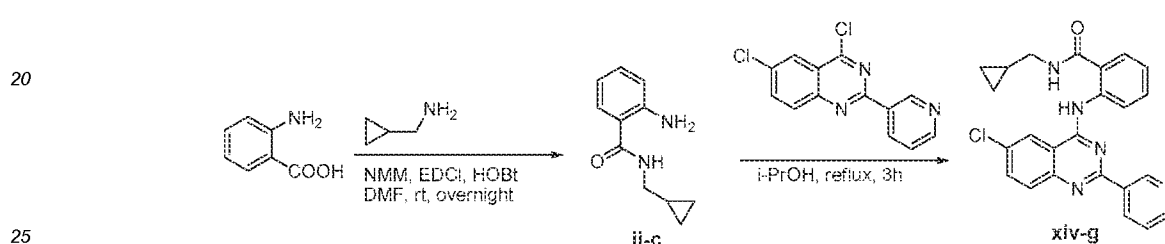
Scheme 34: General route for the synthesis of compounds with general formula xiv

[0369]



Scheme 35: Representative synthesis of compounds of formula xiv-g (see Scheme 34)

[0370]

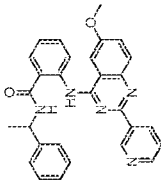
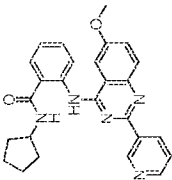
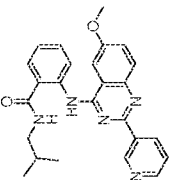
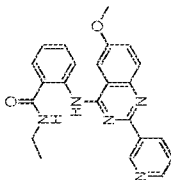


[0371] **Method AA: 2-Amino-N-(cyclopropylmethyl)benzamide (ii-c)** A 100 mL round bottom flask was charged with anthranilic acid (500 mg, 3.65 mmol), added DMF (15 mL) under nitrogen atmosphere and stirring. Then, added N-methylmorpholine (1 mL, 9.12 mmol), aminomethylcyclopropane (311 mg, 4.38 mmol), N-ethyl-N'-dimethylaminopropylcarbodiimide (EDCI) (840 mg, 4.38 mmol) and HOBT (670 mg, 4.38 mmol). The reaction mixture was stirred at room temperature overnight. The reaction mixture was quenched with water and extracted with diethyl ether (50 mL x 2). The organic extracts were combined, washed with brine, dried over MgSO₄, filtered and concentrated to afford amorphous colorless 2-amino-N-(cyclopropylmethyl)benzamide (500 mg, 72% yield), which was checked by NMR and used for next step without further purification. ¹H-NMR (Bruker 300 MHz, DMSO-d₆) δ 0.20-1.01 (m, 4H), 1.01-1.04 (m, 1H), 3.06-3.11 (m, 2H), 6.37-8.21 (m, 7H).

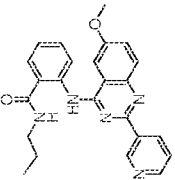
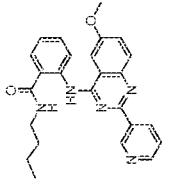
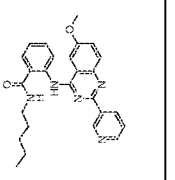
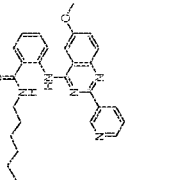
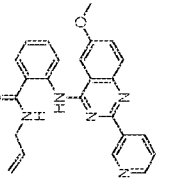
[0372] **Method Z: 2-(6-chloro-2-(pyridin-3-yl)quinazolin-4-ylamino)-N-(cyclopropylmethyl) benzamide (xiv-g):** A 100 mL round bottom flask was charged with 4,6-dichloro-2-(pyridin-3-yl)quinazoline (synthesized as described in Scheme 1 and 4, substituting 5-chloro-2-nitrobenzoic acid for 2-nitro-5-propoxy-benzoic acid) (150 mg, 0.54 mmol) and 2-amino-N-(cyclopropylmethyl)benzamide (310 mg, 1.63 mmol) and anhydrous *i*-PrOH (20 mL) was added and refluxed for 3h. The reaction mixture was cooled to room temperature and Et₃N (3eq) was added. The solvent was removed *in vacuo*. To the crude product was added a water-methanol mixture (5:1, 50 mL), and then sonicated for 5 min. The solidified compound was collected by filtration, and the solid was recrystallized from hot methanol and washed with water. The product was dried at 50 °C to give 2-(6-chloro-2-(pyridin-3-yl)quinazolin-4-ylamino)-N-(cyclopropylmethyl)benzamide as a colorless cotton (220 mg, 94%). The structure was confirmed by NMR and elemental analysis. ¹H NMR (Bruker 300 MHz, DMSO-d₆) ppm 0.20-0.98 (m, 5H), 3.07-3.21 (m, 2H), 7.24-8.26 (m, 7H), 8.68-9.55 (m, 5H), 12.22 (s, 1H). CHN Calcd. C: 67.05; H: 4.69; N: 16.29 Found C: 67.15; H: 4.89; N: 16.25

[0373] The compounds in the following table were prepared in a manner analogous to that described in Scheme 34, replacing aminomethylcyclopropane with the appropriate amine.

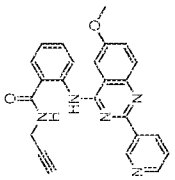
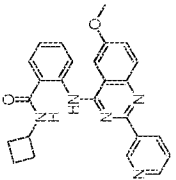
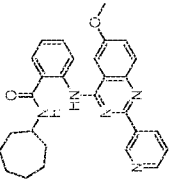
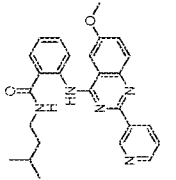
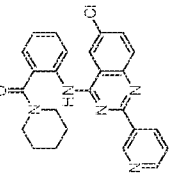
Table 11:

Number	Product	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1016		¹ H NMR (DMSO-d ₆) ppm 1.46 (d, 3H, J = 6.7 Hz), 3.90 (s, 3H), 5.17-5.22 (m, 1H), 7.19-7.98 (m, 12H), 8.67-8.71 (m, 2H), 8.90 (d, 1H, J = 8.2 Hz), 9.22 (d, 1H, J = 8.3 Hz), 9.55 (s, 1H), 12.18 (s, 1H).	DMSO	>98	Method Z
1017		¹ H NMR (DMSO-d ₆) ppm 1.49-1.62 (m, 8H), 3.99 (s, 3H), 4.24-4.26 (m, 1H), 7.24-7.71 (m, 7H), 8.65-9.55 (m, 5H), 12.19 (s, 1H).	DMSO	>98	Method Z
1018		¹ H NMR (DMSO-d ₆) ppm 0.88-1.99 (m, 7H), 3.14-3.16 (m, 2H), 3.02 (s, 3H), 7.57-7.91 (m, 7H), 8.68-9.55 (m, 5H), 12.19 (s, 1H).	DMSO	>98	Method Z
1019		¹ H NMR (DMSO-d ₆) ppm 1.07-1.12 (m, 3H), 3.44-3.52 (m, 2H), 4.01 (s, 3H), 7.24-7.86 (m, 8H), 8.68-9.55 (m, 5H), 12.19 (s, 1H).	DMSO	>98	Method Z

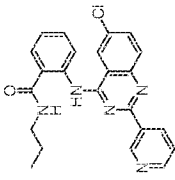
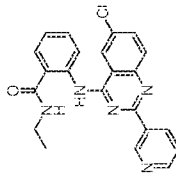
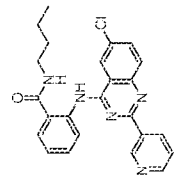
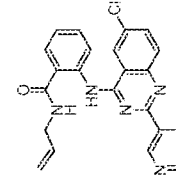
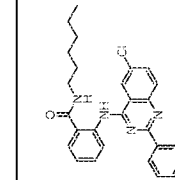
(continued)

Number	Product	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1020		¹ H NMR (DMSO-d ₆) ppm 0.87-0.92 (m, 3H), 1.53-1.62 (m, 2H), 3.30-3.32 (m, 2H), 3.98 (s, 3H), 7.21-7.89 (m, 7H), 8.67-9.55 (m, 5H), 12.38 (s, 1H).	DMSO	>98	Method Z
1021		¹ H NMR (DMSO-d ₆) ppm 0.83-1.51 (m, 7H), 3.30-3.32 (m, 2H), 3.99 (s, 3H), 7.24-7.90 (m, 7H), 8.67-9.55 (m, 5H), 12.38 (s, 1H).	DMSO	>98	Method Z
1022		¹ H NMR (DMSO-d ₆) ppm 0.79-1.52 (m, 9H), 3.25-3.30 (m, 2H), 3.98 (s, 3H), 7.21-7.88 (m, 7H), 8.68-9.55 (m, 5H), 12.30 (s, 1H).	DMSO	>98	Method Z
1023		¹ H NMR (DMSO-d ₆) ppm 0.80-1.49 (m, 11H), 3.26-3.30 (m, 2H), 3.99 (s, 3H), 7.24-7.90 (m, 7H), 8.67-9.56 (m, 5H), 12.30 (s, 1H).	DMSO	>98	Method Z
1024		¹ H NMR (DM80-d ₆) ppm 3.99-4.10 (m, 5H), 5.07-5.33 (m, 2H), 5.89-5.91 (m, 1H), 7.24-7.90 (m, 7H), 8.68-9.55 (m, 5H), 12.30 (s, 1H).	DMSO	>98	Method Z

(continued)

Number	Product	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1025		¹ H NMR (DMSO-d ₆) ppm 3.13 (s, 1H), 3.99-4.12 (m, 5H), 7.21-7.89 (m, 7H), 8.67-9.54 (m, 5H), 12.30 (s, 1H).	DMSO	>98	Method Z
1026		¹ H NMR (DMSO-d ₆) ppm 1.68-2.22 (m, 6H), 3.99 (s, 3H), 4.24-4.26 (m, 1H), 7.22-7.92 (m, 7H), 8.66-9.55 (m, 5H), 12.30 (s, 1H).	DMSO	>98	Method Z
1027		¹ H NMR (DMSO-d ₆) ppm 1.41-1.86 (m, 12H), 3.90 (br s, 4H), 7.23-7.89 (m, 7H), 8.63-9.55 (m, 5H), 12.30 (s, 1H).	DMSO	>98	Method Z
1028		¹ H NMR (DMSO-d ₆) ppm 0.85-1.60 (m, 9H), 3.27-3.33 (m, 2H), 3.99 (s, 3H), 7.21-7.91 (m, 7H), 8.68-9.55 (m, 5H), 12.19 (s, 1H).	DMSO	>98	Method Z
1029		¹ H NMR (DMSO-d ₆) ppm 1.03-1.34 (m, 6H), 3.17-3.31 (m, 4H), 7.40-7.88 (m, 7H), 8.55-10.03 (m, 5H), 12.32 (s, 1H).	DMSO	>98	Method Z

(continued)

Number	Product	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1030		¹ H NMR (DMSO-d ₆) ppm 0.83-1.52 (m, 5H), 3.20-3.29 (m, 2H), 7.27-8.21 (m, 7H), 8.67-9.53 (m, 5H), 12.33 (s, 1H).	DMSO	>98	Method Z
1031		¹ H NMR (DMSO-d ₆) ppm 1.02-1.07 (m, 3H), 3.24-3.32 (m, 2H), 7.27-8.22 (m, 7H), 8.67-9.53 (m, 5H), 12.34 (s, 1H).	DMSO	>98	Method Z
1032		¹ H NMR (DMSO-d ₆) ppm 0.75-1.37 (m, 7H), 3.18-3.20 (m, 2H), 7.21-8.19 (m, 7H), 8.64-9.48 (m, 5H), 12.15 (s, 1H).	DMSO	>98	Method Z
1033		¹ H NMR (DMSO-d ₆) ppm 3.90-3.92 (m, 2H), 5.02-5.17 (m, 2H), 5.81-5.86 (m, 1H), 7.27-8.23 (m, 7H), 8.68-9.55 (m, 5H), 12.25 (s, 1H).	DMSO	>98	Method Z
1034		¹ H NMR (DMSO-d ₆) ppm 0.78-1.42 (m, 11 H), 3.20-3.29 (m, 2H), 7.25-7.91 (m, 7H), 8.23-9.53 (m, 5H), 12.17 (s, 1H).	DMSO	>98	Method Z

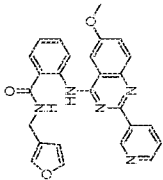
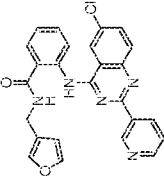
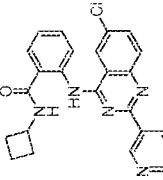
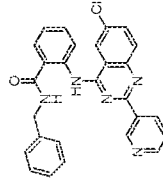
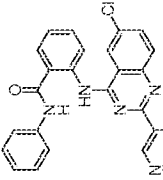
(continued)

Number	Product	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1035		¹ H NMR (DMSO-d ₆) ppm 3.06 (s, 1H), 4.05-4.08 (m, 2H), 7.28-8.22 (m, 7H), 8.65-9.51 (m, 5H), 12.12 (s, 1H).	DMSO	>98	Method Z
1036		¹ H NMR (DMSO-d ₆) ppm 0.88-1.92 (m, 7H), 3.05-3.16 (m, 2H), 7.23-7.94 (m, 7H), 8.22-9.54 (m, 5H), 12.21 (s, 1H).	DMSO	>98	Method Z
1037		¹ H NMR (DMSO-d ₆) ppm 1.37-1.79 (m, 12H), 3.90-3.96 (m, 1H), 7.25-8.24 (m, 7H), 8.55-9.52 (m, 5H), 12.07 (s, 1H).	DMSO	>98	Method Z
1038		¹ H NMR (DMSO-d ₆) ppm 1.49-1.85 (m, 8H), 4.24-4.26 (m, 1H), 7.23-7.69 (m, 6H), 7.85-9.53 (m, 6H), 12.06 (s, 1H).	DMSO	>98	Method Z
1039		¹ H NMR (DMSO-d ₆) ppm 1.12-1.14 (m, 6H), 4.11-4.18 (m, 1H), 7.23-7.94 (m, 8H), 8.33-9.53 (m, 4H), 12.30 (s, 1H).	DMSO	>98	Method Z

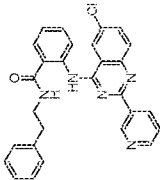
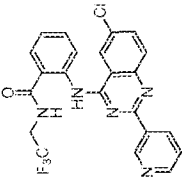
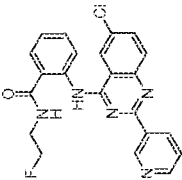
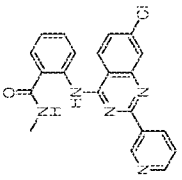
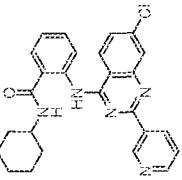
(continued)

Number	Product	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1040		¹ H NMR (DMSO-d ₆) ppm 0.84-1.16 (m, 9H), 3.28-3.34 (m, 2H), 7.23-7.91 (m, 6H), 8.25-9.53 (m, 6H), 12.16 (s, 1H).	DMSO	>98	Method Z
1041		¹ H NMR (DMSO-d ₆) ppm 0.79-1.52 (m, 9H), 3.25-3.32 (m, 2H), 7.24-7.91 (m, 6H), 8.23-9.53 (m, 6H), 12.18 (s, 1H).	DMSO	>98	Method Z
1042		¹ H NMR (DMSO-d ₆) ppm 1.04-1.81 (m, 10H), 3.77-3.82 (m, 1H), 7.21-8.05 (m, 8H), 8.65-9.52 (m, 4H), 11.96 (s, 1H).	DMSO	>98	Method Z
1043		¹ H NMR (DMSO-d ₆) ppm 0.47-0.67 (m, 4H), 2.85-2.88 (m, 1H), 7.26-8.26 (m, 7H), 8.66-9.52 (m, 5H), 12.26 (s, 1H).	DMSO	>98	Method Z
1044		¹ H NMR (DMSO-d ₆) ppm 1.14 (s, 9H), 7.30-7.93 (m, 7H), 8.21-9.50 (m, 5H), 12.22 (s, 1H).	DMSO	>98	Method Z

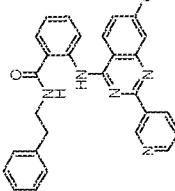
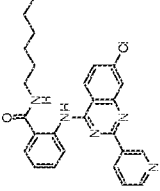
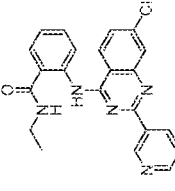
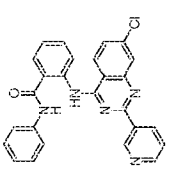
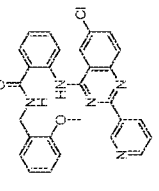
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Number	Product	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1045		¹ H NMR (DMSO-d ₆) ppm 3.98 (s, 3H), 4.36 (s, 2H), 6.47 (s, 1H), 7.21-7.91 (m, 9H), 8.68-9.56 (m, 5H), 12.22 (s, 1H).	DMSO	>98	Method Z
1046		¹ H NMR (DMSO-d ₆) ppm 4.30 (s, 2H), 6.42 (s, 1H), 7.28-8.24 (m, 9H), 8.70-9.53 (m, 5H), 12.22 (s, 1H).	DMSO	>98	Method Z
1047		¹ H NMR (DMSO-d ₆) ppm 1.69-2.55 (m, 6H), 4.41-4.44 (m, 1H), 7.23-8.24 (m, 7H), 8.65-9.54 (m, 5H), 12.06 (s, 1H).	DMSO	>98	Method Z
1048		¹ H NMR (DMSO-d ₆) ppm 4.55 (s, 2H), 7.17-8.21 (m, 12H), 8.65-9.54 (m, 5H), 12.06 (s, 1H).	DMSO	>98	Method Z
1049		¹ H NMR (DMSO-d ₆) ppm 7.02-8.67 (m, 15H), 9.47 (s, 1H), 10.04 (s, 1H), 12.06 (s, 1H).	DMSO	>98	Method Z

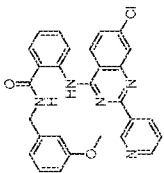
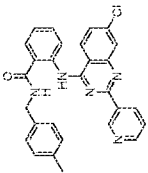
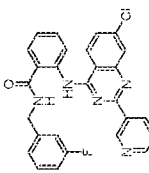
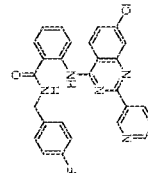
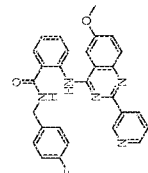
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Number	Product	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1050		¹ H NMR (DMSO-d ₆) ppm 2.75-2.80 (m, 2H), 3.48-3.54 (m, 2H), 7.08-8.20 (m, 12H), 8.69-9.56 (m, 5H), 12.16 (s, 1H).	DMSO	>98	Method Z
1051		¹ H NMR (DMSO-d ₆) ppm 4.01-4.10 (m, 2H), 7.28-8.27 (m, 7H), 8.59-9.51 (m, 5H), 11.45 (s, 1H).	DMSO	>98	Method Z
1052		¹ H NMR (DMSO-d ₆) ppm 4.38-4.57 (m, 4H), 7.27-8.24 (m, 7H), 8.69-9.56 (m, 5H), 12.10 (s, 1H).	DMSO	>98	Method Z
1053		¹ H NMR (DMSO-d ₆) ppm 2.85-2.92 (br s, 3H), 7.20-8.22 (m, 7H), 8.67-9.56 (m, 5H), 12.61 (s, 1H).	DMSO	>98	Method Z
1054		¹ H NMR (DMSO-d ₆) ppm 1.04-1.81 (m, 10H), 3.77-3.82 (m, 1H), 7.24-8.05 (m, 8H), 8.65-9.52 (m, 4H), 12.27 (s, 1H).	DMSO	>98	Method Z

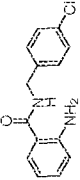
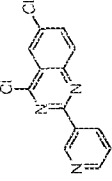
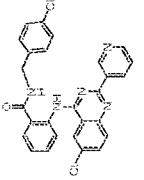
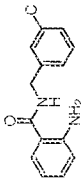
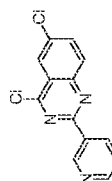
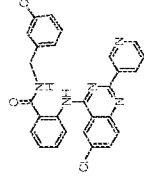
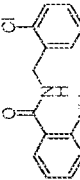
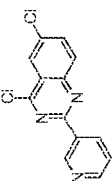
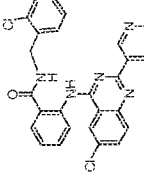
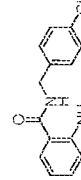
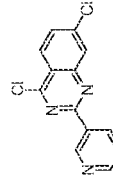
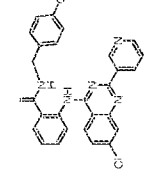
(continued)

Number	Product	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1055		¹ H NMR (DMSO-d ₆) ppm 2.49-2.51 (m, 2H), 3.57-3.59 (m, 2H), 7.21-8.16 (m, 12H), 8.69-9.56 (m, 5H), 12.16 (s, 1H).	DMSO	>98	Method Z
1056		¹ H NMR (DMSO-d ₆) ppm 0.79-1.44 (m, 11H), 3.26-3.30 (m, 2H), 7.24-7.92 (m, 7H), 8.67-9.52 (m, 5H), 12.45 (s, 1H).	DMSO	>98	Method Z
1057		¹ H NMR (DMSO-d ₆) ppm 1.04-1.16 (m, 3H), 3.33-3.38 (m, 2H), 7.20-8.22 (m, 7H), 8.60-9.53 (m, 5H), 12.34 (s, 1H).	DMSO	>98	Method Z
1058		¹ H NMR (DMSO-d ₆) ppm 7.09-8.68 (m, 15H), 9.51 (s, 1H), 10.34 (s, 1H), 12.06 (s, 1H).	DMSO	>98	Method Z
1059		¹ H NMR (DMSO-d ₆) ppm 3.78 (s, 3H), 4.43 (s, 2H), 6.77-8.23 (m, 11H), 8.68-9.54 (m, 5H), 12.06 (s, 1H).	DMSO	>98	Method Z

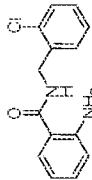
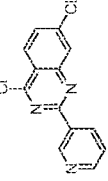
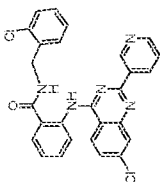
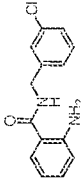
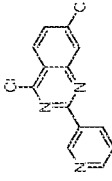
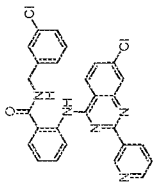
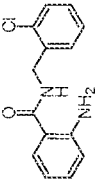
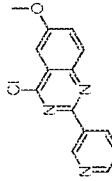
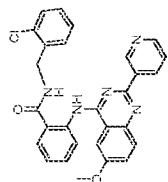
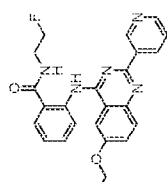
(continued)

Number	Product	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1060		¹ H NMR (DMSO-d ₆) ppm 3.69 (s, 3H), 4.48 (s, 2H), 6.89-8.15 (m, 11H), 8.68-9.54 (m, 5H), 12.44 (s, 1H).	DMSO	>98	Method Z
1061		¹ H NMR (DMSO-d ₆) ppm 2.24 (s, 3H), 4.46 (s, 2H), 7.03-8.13 (m, 11H), 8.67-9.54 (m, 5H), 12.21 (s, 1H).	DMSO	>98	Method Z
1062		¹ H NMR (DMSO-d ₆) ppm 4.51 (s, 2H), 7.11-8.14 (m, 11H), 8.67-9.54 (m, 5H), 12.34 (s, 1H).	DMSO	>98	Method Z
1063		¹ H NMR (DMSO-d ₆) ppm 4.47 (s, 2H), 7.05-8.14 (m, 11H), 8.67-9.54 (m, 5H), 12.37 (s, 1H).	DMSO	>98	Method Z
1064		¹ H NMR (DMSO-d ₆) ppm 3.95 (s, 3H), 4.47 (s, 2H), 7.05-8.14 (m, 11H), 8.67-9.54 (m, 5H), 12.32 (s, 1H).	DMSO	>98	Method Z

(continued)

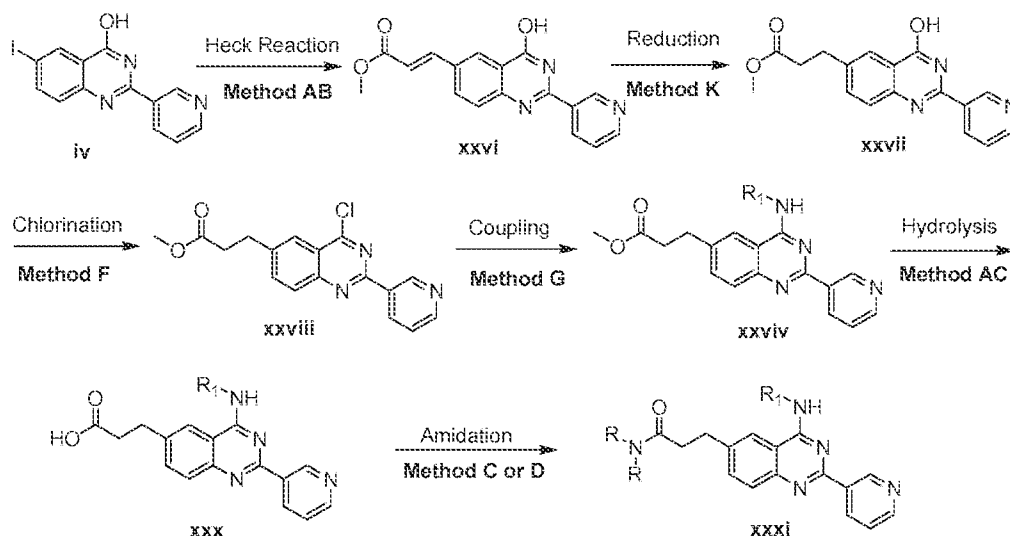
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1065					¹ H NMR (DMSO- <i>d</i> ₆) ppm 4.52 (d, 2H, <i>J</i> = 5.8 Hz), 7.21-7.30 (m, 5H), 7.50 (t, <i>J</i> = 7.9 Hz, 1H), 7.68 (t, <i>J</i> = 7.8 Hz, 1H), 7.89-7.92 (m, 3H), 8.21 (d, <i>J</i> = 1.8 Hz, 1H), 8.65-8.70 (m, 3H), 8.99 (s, 1H), 9.52 (s, 1H), 11.88 (s, 1H)	DMSO	>98	Method Y, Z
1066					¹ H NMR (DMSO- <i>d</i> ₆) ppm 4.49 (d, 2H, <i>J</i> = 5.6 Hz), 7.24-7.35 (m, 5H), 7.50 (t, <i>J</i> = 7.2 Hz, 1H), 7.70 (t, <i>J</i> = 7.8 Hz, 1H), 7.91-7.95 (m, 3H), 8.22 (d, <i>J</i> = 1.5 Hz, 1H), 8.67-8.73 (m, 3H), 9.19 (s, 1H), 9.53 (s, 1H), 11.98 (s, 1H)	DMSO	>98	Method Y, Z
1067					¹ H NMR (DMSO- <i>d</i> ₆) ppm 4.58 (d, 2H, <i>J</i> = 5.7 Hz), 7.16-7.36 (m, 5H), 7.50 (t, <i>J</i> = 7.5 Hz, 1H), 7.69 (t, <i>J</i> = 7.5 Hz, 1H), 7.88-7.96 (m, 3H), 8.22 (d, <i>J</i> = 1.5 Hz, 1H), 8.65-8.69 (m, 3H), 8.98 (s, 1H), 9.52 (s, 1H), 11.87 (s, 1H)	DMSO	>98	Method Y, Z
1068					¹ H NMR (DMSO- <i>d</i> ₆) ppm 4.47 (d, 2H, <i>J</i> = 5.6 Hz), 7.30 (br s, 5H), 7.54 (t, <i>J</i> = 7.7 Hz, 1H), 7.70 (t, <i>J</i> = 7.8 Hz, 1H), 7.88-7.96 (m, 3H), 8.15 (d, <i>J</i> = 1.5 Hz, 1H), 8.68-8.81 (m, 3H), 8.98 (s, 1H), 9.52 (s, 1H), 12.29 (s, 1H)	DMSO	>98	Method Y, Z

(continued)

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1069					¹ H NMR (DMSO- <i>d</i> ₆) ppm 4.57 (d, 2H, <i>J</i> = 5.6 Hz), 7.23-7.29 (m, 5H), 7.39 (t, <i>J</i> = 7.9 Hz, 1H), 7.71 (t, <i>J</i> = 7.9 Hz, 1H), 7.94-7.99 (m, 3H), 8.15 (d, <i>J</i> = 1.4 Hz, 1H), 8.68-8.84 (m, 3H), 9.24 (s, 1H), 9.55 (s, 1H), 12.23 (s, 1H)	DMSO	>98	Method Y, Z
1070					¹ H NMR (DMSO- <i>d</i> ₆) ppm 4.50 (d, 2H, <i>J</i> = 5.5 Hz), 7.25-7.53 (m, 5H), 7.69 (t, <i>J</i> = 7.1 Hz, 1H), 7.73 (t, <i>J</i> = 7.0 Hz, 1H), 7.94-8.15 (m, 3H), 8.18 (d, <i>J</i> = 1.4 Hz, 1H), 8.68-8.82 (m, 3H), 9.24 (s, 1H), 9.54 (s, 1H), 12.21 (s, 1H)	DMSO	>98	Method Y, Z
1071					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.90 (s, 3H), 4.56 (d, 2H, <i>J</i> = 5.6 Hz), 7.22-7.44 (m, 7H), 7.44 (t, <i>J</i> = 7.9 Hz, 1H), 7.74 (t, <i>J</i> = 7.9 Hz, 1H), 7.94-7.99 (m, 2H), 8.68-8.81 (m, 3H), 9.24 (s, 1H), 9.55 (s, 1H), 12.19 (s, 1H)	DMSO	>98	Method Y, Z
1072				HCl	¹ H-NMR (400 MHz, DMSO- <i>d</i> ₆): δ 12.17 (s, 1H), 9.52 (s, 1H), 9.08 (s, 1H), 8.95 (d, <i>J</i> = 7.6 Hz, 1H), 8.85 (s, 1H), 8.66 (d, <i>J</i> = 7.8 Hz, 1H), 7.91 (t, <i>J</i> = 9.4 Hz, 2H), 7.84 (s, 1H), 7.75-7.67 (m, 1H), 7.65 (s, 1H), 7.59 (d, <i>J</i> = 8.1 Hz, 1H), 7.30 (t, <i>J</i> = 7.3 Hz, 1H), 4.47 (d, <i>J</i> = 47.3 Hz, 2H), 4.26 (d, <i>J</i> = 6.8 Hz, 2H), 3.58-3.50 (m, 2H), 1.47 (t, <i>J</i> = 6.7 Hz, 3H).	DMSO	95	Method Y, Z

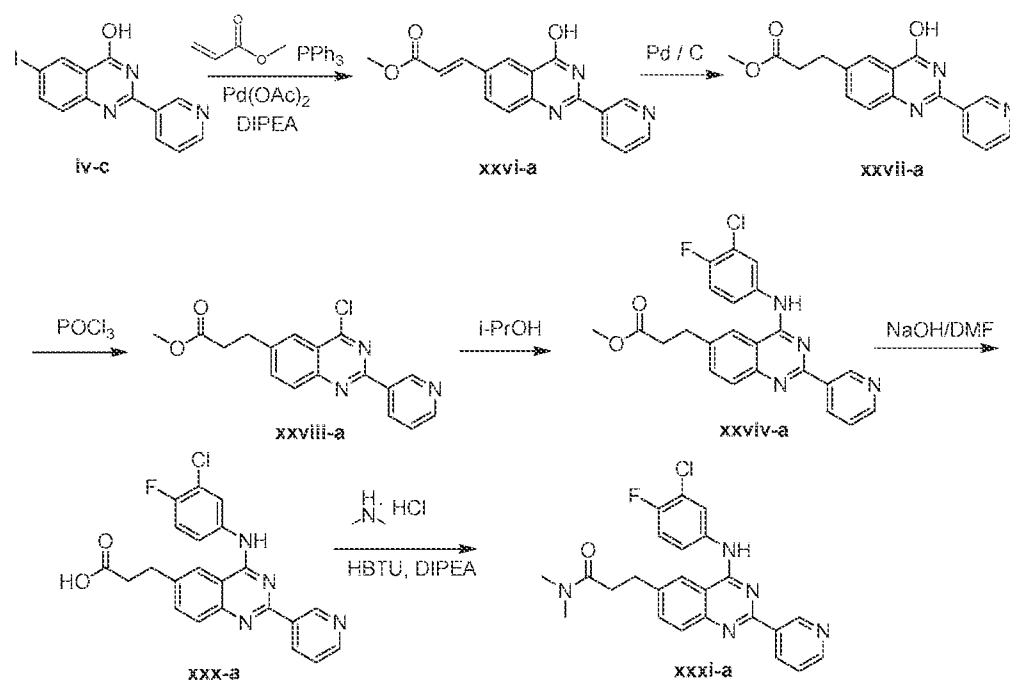
Scheme 36: General route for the synthesis of compounds with general formula xxxi

[0374]



Scheme 37: Representative synthesis of compounds of formula xxxi-a: (see Scheme 36)

[0375]



[0376] Method AB: Methyl 3-(4-hydroxy-2-(pyridin-3-yl)quinazolin-6-yl)acrylate (xxvi-a) To a solution of 6-iodo-2-(pyridin-3-yl)quinazolin-4-ol (synthesized as described in Scheme 1 and 4, substituting 5-iodo-2-nitrobenzoic acid for 2-nitro-5-propoxybenzoic acid) (3.00 g, 8.6 mmol, 1.0 eq.), Pd(OAc)₂ (48 mg, 0.21 mmol, 0.025 eq.) and PPh₃ (113 mg, 0.43 mmol, 0.05 eq.) in DMF (8 mL) was added methyl acrylate (3.22 g, 25.8 mmol, 3.0 eq.) and DIPEA (1.22 g, 9.46 mmol, 1.1 eq.) under Ar atmosphere. The mixture was stirred at 110 °C overnight. After cooling, the mixture was filtered and the solid was washed with ethyl acetate three times to afford 1.30 g of xxvi-a as green solid (yield 49%). LCMS m/z = 308.0 (M + 1) (Method B) (retention time = 1.41 min).

[0377] Methyl 3-(4-hydroxy-2-(pyridin-3-yl)quinazolin-6-yl)propanoate (xxvii-a) Methyl 3-(4-hydroxy-2-(pyridin-

3-yl) quinazolin-6-yl) propanoate was prepared in a manner analogous to that described for 2-amino-5-methoxybenzoic acid in **Method K**, replacing 5-methoxy-2-nitrobenzoic acid with methyl 3-(4-hydroxy-2-(pyridin-3-yl)quinazolin-6-yl)acrylate to afford 1.40 g of **xxvii-a** in quantitative yield as a yellow solid. LCMS m/z = 310.0 ($M + 1$) (Method B) (retention time = 1.42 min).

[0378] 3-(4-Chloro-2-pyridin-3-yl-quinazolin-6-yl)-propionic acid methyl ester (xxviii-a) 3-(4-Chloro-2-pyridin-3-yl-quinazolin-6-yl)-propionic acid methyl ester (prepared in a manner analogous to that described for 4-chloro-6-propoxy-2-pyridin-3-yl-quinazoline using Method F, replacing 6-propoxy-2-pyridin-3-yl-1H-quinazolin-4-one with methyl 3-(4-hydroxy-2-(pyridin-3-yl)quinazolin-6-yl)propanoate) was obtained in quantitative yield to give 1.50 g of **xxviii-a** as a red solid. LCMS m/z = 328.1, 330.0 ($M + 1$) (Method B) (retention time = 1.86 min).

[0379] 3-[4-(3-Chloro-4-fluoro-phenylamino)-2-pyridin-3-yl-quinazolin-6-yl]-propionic acid methyl ester (xxviv-a) 3-[4-(3-Chloro-4-fluoro-phenylamino)-2-pyridin-3-yl-quinazolin-6-yl]-propionic acid methyl ester (prepared in a manner analogous to that described for 2-(6-propoxy-2-pyridin-3-yl-quinazolin-4-ylamino)-benzamide in Method G1, replacing 4,7-dichloro-2-(4-chlorophenyl)quinazoline and 2-aminobenzamide with 3-(4-chloro-2-pyridin-3-yl-quinazolin-6-yl)-propionic acid methyl ester and 3-chloro-4-fluoro-phenylamine) was obtained in a 43% yield to give 1.04 g of **xxviv-a** as a yellow solid. LCMS m/z = 437.1, 439.1 ($M + 1$) (Method B) (retention time = 1.62 min).

[0380] Method AC: 3-(4-(3-Chloro-4-fluorophenylamino)-2-(pyridin-3-yl) quinazolin-6-yl)propanoic acid (xxx-a) To a solution of 3-[4-(3-chloro-4-fluoro-phenylamino)-2-pyridin-3-yl-quinazolin-6-yl]-propionic acid methyl ester (1.04 g, 2.39 mmol, 1.0 eq.) in DMF (20 mL) was added a solution of NaOH (0.57 g, 14.3 mmol, 6.0 eq.) in H₂O (8 mL). The mixture was stirred at room temperature for 2 h. 50 mL of water was added to the mixture. After filtration, the resulting filter cake was washed with water and dried *in vacuo* to give 933 mg of **xxx-a** as a yellow solid (yield 93%). LCMS m/z = 423.1, 425.1 ($M + 1$) (Method A) (retention time = 1.51 min).

[0381] 3-(4-(3-Chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)-N,N-dimethyl propanamide (xxxi-a) 3-(4-(3-Chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)-N,N-dimethyl propanamide (prepared in a manner analogous to that described for 2-benzamido-5-methoxy-3-methylbenzamide in Method D, replacing nicotinic acid and 2-amino-5-methoxy-3-methylbenzamide with 3-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)propanoic acid and dimethylamine hydrochloride) was obtained in 90% yield to give 891 mg of **xxxi-a** as a yellow solid. LCMS m/z = 450.0, 452.0 ($M + 1$) (Method B) (retention time = 1.834 min). ¹H-NMR (400 MHz, DMSO-*d*₆): δ 9.95 (s, 1H), 9.51 (d, J = 1.6 Hz, 1H), 8.69 - 8.63 (m, 2H), 8.38 (s, 1H), 8.28 (dd, J = 6.8, 2.4 Hz, 1H), 7.94 - 7.90 (m, 1H), 7.81 (s, 2H), 7.55 - 7.51 (m, 2H), 3.04 (t, J = 7.6 Hz, 2H), 2.99 (s, 3H), 2.85 (s, 3H), 2.77 (t, J = 8.0 Hz, 2H).

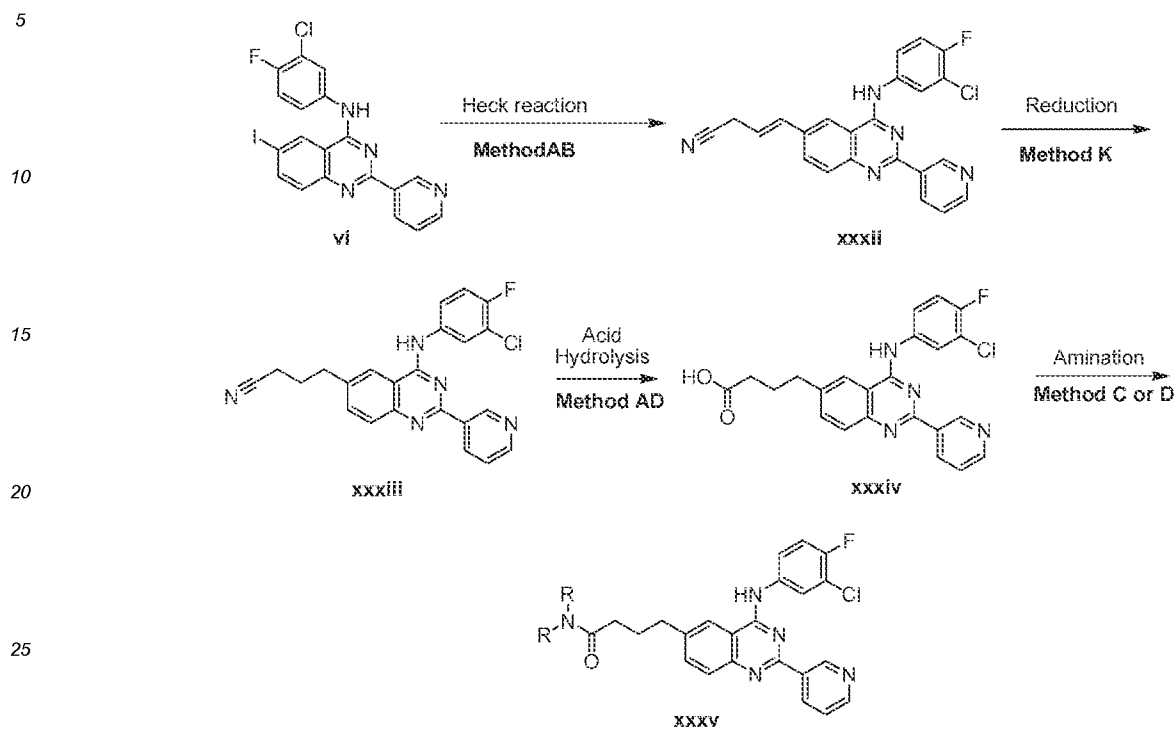
The compounds in the following table were prepared in a manner analogous to that described in **Scheme 36**, replacing dimethylamine with the appropriate amine and 4-fluoro, 3-chloro aniline with the appropriate aniline.

Table 12:

Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1073			421.85	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.01 (s, 1H), 9.52 (s, 1H), 8.65 - 8.69 (m, 2H), 8.39 (s, 1H), 8.28 (d, J = 5.2 Hz, 1H), 7.92 - 7.93 (m, 1H), 7.82 (q, J = 8.0 Hz, 2H), 7.52 - 7.57 (m, 2H), 7.35 (s, 1H), 6.84 (s, 1H), 3.05 (t, J = 7.2 Hz, 2H), 2.51 - 2.54 (m, 2H).	DMSO	422.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C
1074			435.43	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.01 (s, 1H), 9.55 (d, J = 2.4 Hz, 1H), 8.68 - 8.70 (m, 2H), 8.47 (s, 1H), 7.97 - 7.99 (m, 1H), 7.79 - 7.86 (m, 3H), 7.50 - 7.55 (m, 2H), 7.36 (s, 1H), 7.30 (t, J = 73.6 Hz, 1H), 6.99 (dd, J = 8.4, 2.4 Hz, 1H), 6.82 (s, 1H), 3.05 (t, J = 8.0 Hz, 2H), 2.52 (t, J = 8.4 Hz, 2H).	DMSO	436.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C
1075			435.88	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.00 (s, 1H), 9.52 (s, 1H), 8.65 - 8.69 (m, 2H), 8.39 (s, 1H), 8.27 - 8.29 (m, 1H), 7.79 (dd, J = 9.2, 2.4 Hz, 1H), 7.76 - 7.85 (m, 3H), 7.52 - 7.57 (m, 2H), 3.05 (t, J = 7.6 Hz, 2H), 2.57 (d, J = 4.4 Hz, 3H), 2.52 - 2.54 (m, 2H).	DMSO	436.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D
1076			449.91	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.95 (s, 1H), 9.51 (d, J = 1.6 Hz, 1H), 8.63 - 8.69 (m, 2H), 8.38 (s, 1H), 8.28 (dd, J = 6.8, 2.4 Hz, 1H), 7.90 - 7.94 (m, 1H), 7.81 (s, 2H), 7.51 - 7.55 (m, 2H), 3.04 (t, J = 7.6 Hz, 2H), 2.99 (s, 3H), 2.85 (s, 3H), 2.77 (t, J = 8.0 Hz, 2H).	DMSO	450.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D
1077			540.01	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.01 (s, 1H), 9.52 (d, J = 1.6 Hz, 1H), 8.61 - 8.74 (m, 2H), 8.41 (s, 1H), 8.28 (dd, J = 6.8, 2.6 Hz, 1H), 7.80 - 7.99 (m, 3H), 7.46 - 7.63 (m, 2H), 3.89 (s, 4H), 3.20 (s, 2H), 3.08 (t, J = 7.4 Hz, 4H), 2.90 (t, J = 7.6 Hz, 2H).	DMSO	540.2, 542.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D

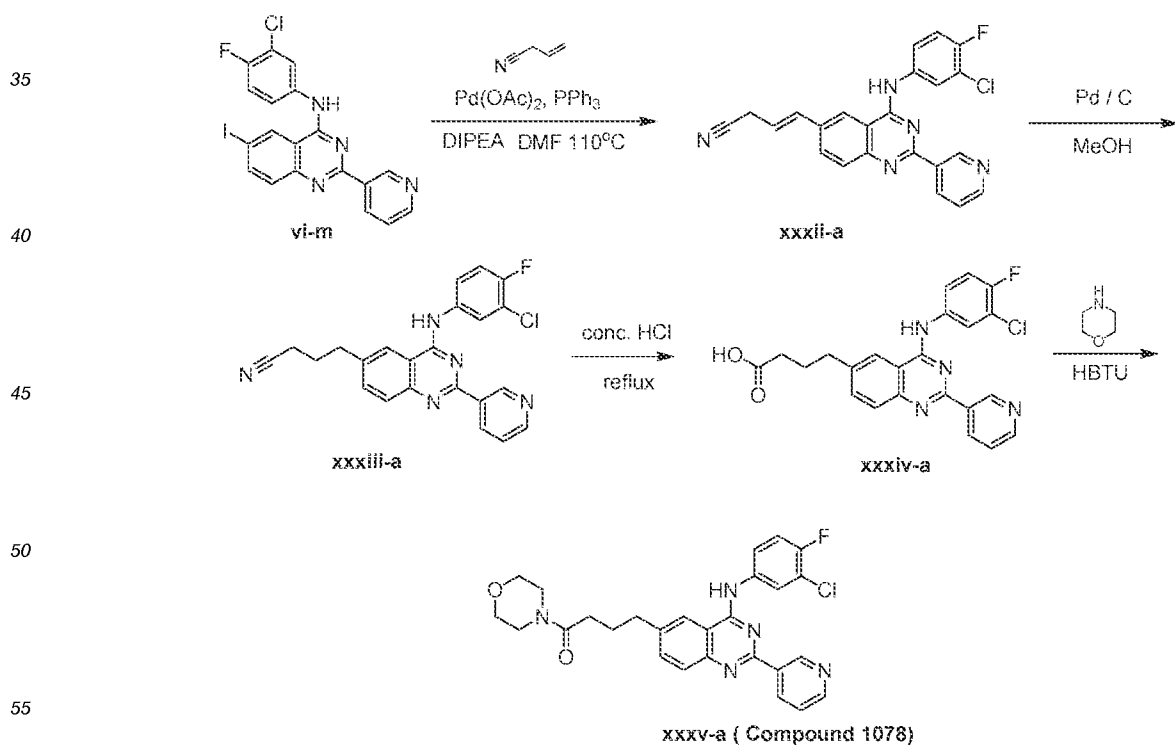
Scheme 38: General route for the synthesis of compounds with general formula xxxv:

[0382]



Scheme 39: Representative synthesis of compounds of formula xxxv-a: (see Scheme 38)

[0383]



[0384] (E)-4-[4-(3-Chloro-4-fluorophenylamino)-2-pyridin-3-yl-quinazolin-6-yl]-but-3-enenitrile (xxxii-a) (E)-

4-[4-(3-Chloro-4-fluoro-phenylamino)-2-pyridin-3-yl-quinazolin-6-yl]-but-3-enenitrile (prepared in a manner analogous to that described for (E)-methyl 3-(4-hydroxy-2-(pyridin-3-yl)quinazolin-6-yl)acrylate using Method AB, replacing 6-iodo-2-(pyridin-3-yl)quinazolin-4-ol and methyl acrylate with N-(3-chloro-4-fluorophenyl)-6-iodo-2-(pyridin-3-yl)quinazolin-4-amine and but-3-enenitrile) was obtained in a 48% yield to give 400 mg of **xxxii-a** as grey solid. LCMS $m/z = 416.0$ ($M + 1$) (Method B) (retention time = 1.99 min).

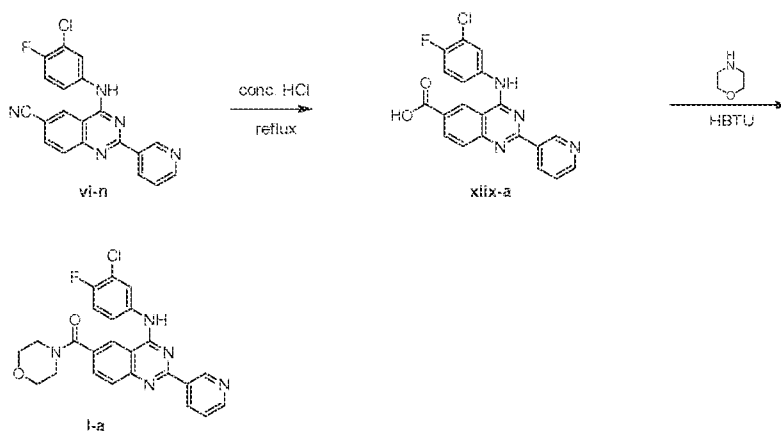
[0385] 4-[4-(3-Chloro-4-fluoro-phenylamino)-2-pyridin-3-yl-quinazolin-6-yl]-butyronitrile (**xxxiii-a**) 4-[4-(3-Chloro-4-fluoro-phenylamino)-2-pyridin-3-yl-quinazolin-6-yl]-butyronitrile (prepared in a manner analogous to that described for 2-amino-5-methoxybenzoic acid in Method K, replacing 5-methoxy-2-nitrobenzoic acid with (E)-4-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)but-3-enenitrile) was obtained in a 95% yield to give 190 mg of **xxxiii-a** as a brown solid. LCMS $m/z = 418.1$ ($M + 1$) (Method B) (retention time = 1.95 min).

[0386] Method AD: 4-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)butanoic acid (**xxxiv-a**) 4-[4-(3-Chloro-4-fluoro-phenylamino)-2-pyridin-3-yl-quinazolin-6-yl]-butyronitrile (190 mg, 046 mmol, 1.0 eq.) was treated with concentrated HCl (8 mL). The mixture was stirred at 100 °C for 2 days. The volatiles were removed *in vacuo*, and the residue was washed with water to afford 70 mg of **xxxiv-a** in a 35% yield as a yellow solid. LCMS $m/z = 437.1$, 439.1 ($M + 1$) (Method B) (retention time = 1.46 min).

[0387] 4-(4-(3-Chloro-4-fluorophenylamino)-2-(pyridin-3-yl) quinazolin-6-yl)-1-morpholino butan-1-one (**xxxv-a**) 4-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl) quinazolin-6-yl)-1-morpholinobutan-1-one was prepared in a manner analogous to that described for 2-benzamido-5-methoxy-3-methylbenzamide in Method D, replacing nicotinic acid and 2-amino-5-methoxy-3-methylbenzamide with 4-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl) butanoic acid and morpholine to give 32 mg of **xxxv-a** in a 40% yield as a beige solid. LCMS $m/z = 506.2$, 508.1 ($M + 1$) (Method B) (retention time = 1.85 min). ^1H NMR (400 MHz, DMSO- d_6) : δ 10.04 (s, 1H), 9.52 (d, $J = 1.6$ Hz, 1H), 8.69 - 8.65 (m, 2H), 8.39 - 8.38 (m, 1H), 8.27 (dd, $J = 6.8$, 2.8 Hz, 1H), 7.94 - 7.90 (m, 1H), 7.86 - 7.84 (m, 1H), 7.80 - 7.77 (m, 1H), 7.57 - 7.52 (m, 2H), 3.56 - 3.53 (m, 4H), 3.46 - 3.41 (m, 4H), 2.85 (t, $J = 8.0$ Hz, 2H), 2.40 (t, $J = 7.6$ Hz, 2H), 1.97 (t, $J = 7.6$ Hz, 2H).

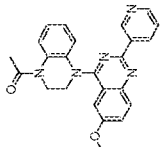
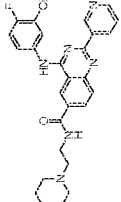
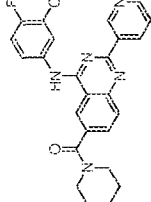
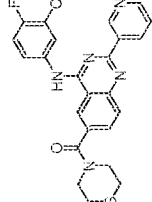
Scheme 40: Representative synthesis of compounds of formula I-a

[0388]



[0389] The compounds in the following table were prepared in a manner analogous to that described in Scheme 38 in the synthesis of 4-(4-(3-Chloro-4-fluorophenyl amino)-2-(pyridin-3-yl) quanazolin-6-yl)-1-morpholino butan-1-one, replacing (E)-4-[4-(3-Chloro-4-fluoro-phenylamino)-2-pyridin-3-yl-quinacolin-6-yl]-but-3-enenitrile with 4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazoline-6-carbonitrile.

Table 13:

Number	PRODUCT	Salt type	Molecular	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1078			411.46	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.64 (d, J = 1.6 Hz, 1H), 8.78 (td, J = 7.9, 1.8 Hz, 1H), 8.71 (dd, J = 4.7, 1.6 Hz, 1H), 7.94 (d, J = 9.2 Hz, 1H), 7.59 (brs, 1H), 7.58 (dd, J = 7.6, 4.8 Hz, 1H), 7.52 (dd, J = 9.2, 2.8 Hz, 1H), 7.16 (t, J = 7.6 Hz, 1H), 7.08 (d, J = 6.4 Hz, 1H), 6.84 (dd, J = 8.0, 1.1 Hz, 1H), 6.69 (s, 1H), 4.36 (t, J = 6.5 Hz, 2H), 4.09 (t, J = 6.4 Hz, 2H), 3.46 (s, 3H), 2.30 (s, 3H).	DMSO	412.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G
1079			506.96	¹ H NMR (400 MHz, DMSO-d ₆): δ 10.42 (s, 1H), 9.54 (d, J = 1.6 Hz, 1H), 9.08 (d, J = 1.2 Hz, 1H), 8.73 - 8.65 (m, 3H), 8.28 (dd, J = 6.8, 2.4 Hz, 1H), 7.97 - 7.93 (m, 2H), 7.60 - 7.53 (m, 2H), 3.60 - 3.58 (m, 4H), 3.51 - 3.46 (m, 2H), 2.55 - 2.53 (m, 2H), 2.41 - 2.48 (m, 4H).	DMSO	508.1 (M+1) 254.1 254.9 (M/2+1)	Method B (NH ₄ HCO ₃)	95	Method C, G, C
1080			461.92	¹ H NMR (400 MHz, DMSO-d ₆): δ 10.21 (s, 1H), 9.54 (d, J = 1.6 Hz, 1H), 8.72 - 8.67 (m, 2H), 8.64 (d, J = 1.2 Hz, 1H), 8.28 (dd, J = 7.0, 2.6 Hz, 1H), 7.95 - 7.86 (m, 3H), 7.60 - 7.53 (m, 2H), 3.68 (d, J = 2.4 Hz, 2H), 3.37 - 3.34 (m, 2H), 1.65 - 1.51 (m, 6H).	DMSO	462.0 464.0 (M+1) 232.3 (M/2+1)	Method A (TFA)	95	Method C, G, C
1081			479.96	¹ H NMR (400 MHz, DMSO-d ₆): δ 10.20 (s, 1H), 9.53 (d, J = 1.6 Hz, 1H), 8.72 - 8.67 (m, 2H), 8.64 (d, J = 1.2 Hz, 1H), 8.28 (dd, J = 6.8, 2.8 Hz, 1H), 7.97 - 7.90 (m, 3H), 7.60 - 7.54 (m, 2H), 3.97 - 3.94 (m, 2H), 3.66 - 3.60 (m, 2H), 2.77 - 2.67 (m, 4H).	DMSO	480.0 482.0 (M+1) 240.6 (M/2+1)	Method A (TFA)	95	Method C, G, C

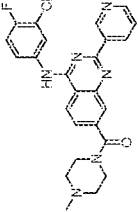
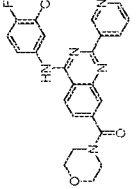
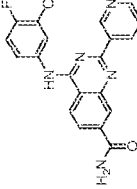
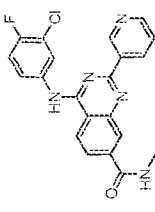
(continued)

Number	PRODUCT	Salt type	Molecular	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1082			421.85	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.20 (s, 1H), 9.55 (d, J = 1.6 Hz, 1H), 8.79 - 8.64 (m, 3H), 8.29 (dd, J = 6.8, 2.6 Hz, 1H), 8.01 - 7.87 (m, 3H), 7.53 - 7.60 (m, 2H), 3.15 - 3.05 (m, 3H), 3.02 (s, 3H).	DMSO	422.1, 424.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G, D
1083			463.89	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.22 (s, 1H), 9.55 (d, J = 1.4 Hz, 1H), 8.77 - 8.65 (m, 3H), 8.02 - 7.87 (m, 3H), 7.60 - 7.54 (m, 2H), 3.56 - 3.39 (m, 2H), 3.82 - 3.55 (m, 6H).	DMSO	464.1, 466.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G, D
1084			407.83	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.41 (s, 1H), 9.54 (d, J = 1.5 Hz, 1H), 9.08 (d, J = 1.3 Hz, 1H), 8.76 - 8.63 (m, 3H), 8.32 - 8.20 (m, 2H), 7.96 - 7.93 (m, 2H), 7.63 - 7.50 (m, 2H), 2.88 (d, J = 4.5 Hz, 3H).	DMSO	408.0, 410.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G, D
1085			447.89	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.23 (s, 1H), 9.55 (d, J = 1.5 Hz, 1H), 8.68 - 8.77 (m, 3H), 8.28 (dd, J = 6.9, 2.6 Hz, 1H), 8.02 (dd, J = 8.6, 1.6 Hz, 1H), 7.99 - 7.90 (m, 2H), 7.53 - 7.58 (m, 2H), 3.57 (t, J = 6.8 Hz, 2H), 3.51 (t, J = 6.4 Hz, 2H), 1.85 - 1.95 (m, 4H).	DMSO	448.1, 420.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G, D

(continued)

Number	PRODUCT	Salt type	Molecular	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1086			476.93	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.22 (s, 1H), 9.54 (d, J = 1.5 Hz, 1H), 8.75 - 8.63 (m, 3H), 8.29 (dd, J = 6.8, 2.6 Hz, 1H), 8.01 - 7.84 (m, 3H), 7.60 - 7.53 (m, 2H), 3.72 (d, J = 2.2 Hz, 2H), 3.51 - 3.39 (m, 2H), 2.48 - 2.40 (m, 2H), 2.33 (d, J = 1.3 Hz, 2H), 2.24 (s, 3H).	DMSO	477.1, 479.1 (M+1) 239.1, 239.9 (M/2+1)	Method B (NH ₄ HCO ₃)	95	Method C, G, D
1087			511.96	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.19 (s, 1H), 9.53 (d, J = 1.6 Hz, 1H), 8.59 - 8.82 (m, 3H), 8.26 (dd, J = 6.8, 2.6 Hz, 1H), 7.81 - 8.14 (m, 3H), 7.57 (m, 2H), 3.83 - 4.10 (m, 4H), 3.33 - 3.35 (m, 4H).	DMSO	512.1, 514.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G, D
1088			481.91	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.47 (s, 1H), 9.54 (d, J = 1.6 Hz, 1H), 9.23 (d, J = 1.2 Hz, 1H), 8.72 - 8.74 (m, 3H), 8.22 - 8.42 (m, 2H), 7.85 - 8.12 (m, 2H), 7.55 - 7.59 (m, 2H), 4.61 (t, J = 5.4 Hz, 1H), 3.47 (t, J = 5.2 Hz, 2H), 3.37 (s, 6H).	DMSO	482.1, 484.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G, D
1089			421.85	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.23 (s, 1H), 9.59 (d, J = 1.2 Hz, 1H), 8.72 - 8.77 (m, 2H), 8.68 (d, J = 8.4 Hz, 1H), 8.34 (dd, J = 7.2, 2.0 Hz, 1H), 7.96 - 8.00 (m, 1H), 7.92 (m, 1H), 7.72 - 7.74 (m, 1H), 7.58 - 7.65 (m, 2H), 3.13 (s, 3H), 3.03 (s, 3H).	DMSO	422.0 (M+1)	Method A (TFA)	95	Method C, G, D
1090			394.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.25 (s, 1H), 9.54 (s, 1H), 8.68 - 8.70 (m, 2H), 8.60 (d, J = 8.4 Hz, 1H), 8.37 (s, 1H), 8.32 (dd, J = 7.2, 2.8 Hz, 1H), 8.11 (d, J = 7.2 Hz, 1H), 7.93 - 7.97 (m, 1H), 7.51 - 7.58 (m, 2H).	DMSO	395.0 (M+1)	Method A (TFA)	95	Method C, G, D

(continued)

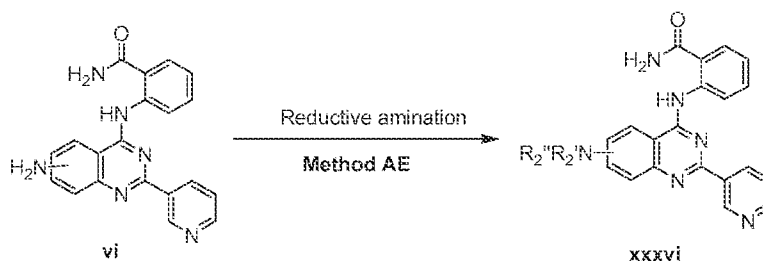
Number	PRODUCT	Salt type	Molecular	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1091			476.93	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.19 (s, 1H), 9.53 (s, 1H), 8.62 - 8.72 (m, 3H), 8.29 (dd, J = 6.8, 2.4 Hz, 1H), 7.91 - 7.94 (m, 1H), 7.83 (s, 1H), 7.66 (d, J = 8.8 Hz, 1H), 7.53 - 7.59 (m, 2H), 3.69 - 3.71 (s, 2H), 3.36 - 3.38 (s, 2H), 2.42 - 2.44 (s, 2H), 2.30 - 2.32 (s, 2H), 2.23 (s, 3H).	DMSO	477.2 (M+1) 239.1 (1/2M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G, D
1092			463.89	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.18 (s, 1H), 9.52 (s, 1H), 8.62 - 8.71 (m, 3H), 8.28 (d, J = 4.8 Hz, 1H), 7.87 - 7.96 (m, 2H), 7.68 (d, J = 8.0 Hz, 1H), 7.52 - 7.58 (m, 2H), 3.61 - 3.71 (m, 6H), 3.40 (m, 2H).	DMSO	464.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method G, D
1093			393.8	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.20 (s, 1H), 9.54 (d, J = 1.6 Hz, 1H), 8.62 - 8.72 (m, 3H), 8.42 (d, J = 1.2 Hz, 1H), 8.37 (s, 1H), 8.30 (dd, J = 7.2, 2.8 Hz, 1H), 8.08 (dd, J = 8.4, 1.2 Hz, 1H), 7.91 - 7.95 (m, 1H), 7.71 (s, 1H), 7.53 - 7.60 (m, 2H).	DMSO	394.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method G, D
1094			407.83	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.19 (s, 1H), 9.53 (s, 1H), 8.84 (d, J = 4.0 Hz, 1H), 8.60 - 8.72 (m, 3H), 8.35 (s, 1H), 8.29 (dd, J = 6.8, 2.8 Hz, 1H), 8.05 (d, J = 8.4 Hz, 1H), 7.90 - 7.93 (m, 1H), 7.52 - 7.59 (m, 2H), 2.87 (d, J = 4.8 Hz, 3H).	DMSO	408.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method G, D

(continued)

Number	PRODUCT	Salt type	Molecular	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1095		HCl	506.96	¹ H-NMR (400 MHz, DMSO-d ₆): 610.20 (s, 1H), 9.53 (d, J = 1.6 Hz, 1H), 8.85 (t, J = 5.6 Hz, 1H), 8.61 - 8.72 (m, 3H), 8.36 (d, J = 1.2 Hz, 1H), 7.98 (dd, J = 6.4, 2.4 Hz, 1H), 8.03 (dd, J = 8.8, 1.6 Hz, 1H), 7.91 - 7.95 (m, 1H), 7.52 - 7.59 (m, 2H), 3.60 (t, J = 4.4 Hz, 4H), 3.46 - 3.48 (q, J = 6.4 Hz, 2H), 2.51 - 2.54 (m, 2H), 2.42 - 2.48 (m, 4H).	DMSO	507.1 (M+1) 254.1 (M/2+1)	Method B (NH ₄ HCO ₃)	95	Method G, C
1096			511.96	¹ H-NMR (400 MHz, DMSO-d ₆): 610.22 (s, 1H), 9.56 (d, J = 1.8 Hz, 1H), 8.61 - 8.79 (m, 3H), 8.31 (dd, J = 6.8, 2.6 Hz, 1H), 7.88 - 8.11 (m, 2H), 7.78 (dd, J = 8.4, 1.4 Hz, 1H), 7.51 - 7.66 (m, 2H), 4.12 (brs, 2H), 3.76 (brs, 2H), 3.30 - 3.33 (m, 4H).	DMSO	512.1, 514.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method G, D
1097			474	¹ H-NMR (400 MHz, DMSO-d ₆): 8 10.64 (brs, 1H), 9.53 (s, 1H), 9.44 (t, J = 6.4 Hz, 1H), 9.35 (s, 1H), 8.90 (d, J = 8.0 Hz, 1H), 8.85 (d, J = 4.8 Hz, 1H), 8.35 (dd, J = 8.4, 1.2 Hz, 1H), 8.27 (dd, J = 6.8, 2.4 Hz, 1H), 8.00 - 7.97 (m, 2H), 7.82 (dt, J = 13.2, 5.2 Hz, 1H), 7.55 (t, J = 8.8 Hz, 1H), 4.22 - 4.18 (m, 2H).	DMSO	475.7, 476.6 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G, D

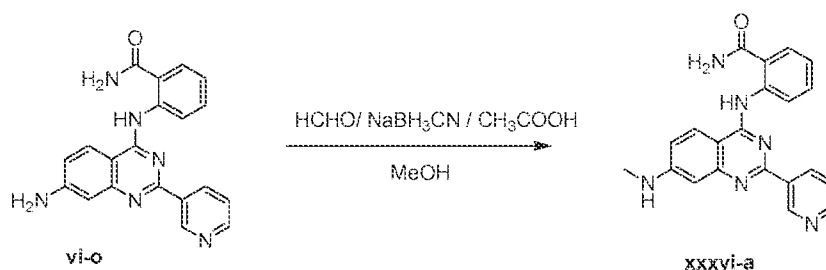
Scheme 41: General route for the synthesis of compounds with general formula xxxvi

[0390]



Scheme 42: Representative synthesis of compounds of formula xxxvi-a: (see Scheme 41)

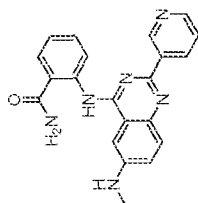
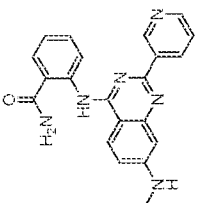
[0391]



[0392] Method AE: 2-(7-(Methylamino)-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (xxxvi-a): A mixture of 2-(7-amino-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (160 mg, 0.449 mmol, 1.0 eq.) and $HCHO$ (40%, 37 mg, 0.494 mmol, 1.1 eq.), acetic acid (2 drops) were stirred at room temperature for 0.5 h. $NaBH_3CN$ (34 mg, 0.449 mmol, 1.0 eq.) was added and the mixture was stirred at room temperature overnight. After filtration, the filtrate was concentrated to give the crude product, which was purified by reverse phase chromatography ($MeOH / H_2O = 3:7$) to afford 28 mg of **xxxvi-a** as a white solid (17%). LCMS $m/z = 371.1 (M+1)$, $372.1 (M+2)$ (Method B) (retention time = 1.60 min). 1H -NMR (400 MHz, $DMSO-d_6$): δ 12.71 (s, 1H), 9.58 (d, $J = 1.6$ Hz, 1H), 9.14 - 9.12 (m, 1H), 8.74 - 8.69 (m, 2H), 8.45 (s, 1H), 7.93 - 7.91 (m, 1H), 7.85 (d, $J = 9.2$ Hz, 2H), 7.67 - 7.71 (m, 1H), 7.59 - 7.55 (m, 1H), 7.12 - 7.16 (m, 1H), 7.06 - 7.03 (m, 1H), 6.82 - 6.81 (m, 1H), 6.70 (d, $J = 2.4$ Hz, 1H), 2.84 (d, $J = 4.8$ Hz, 3H).

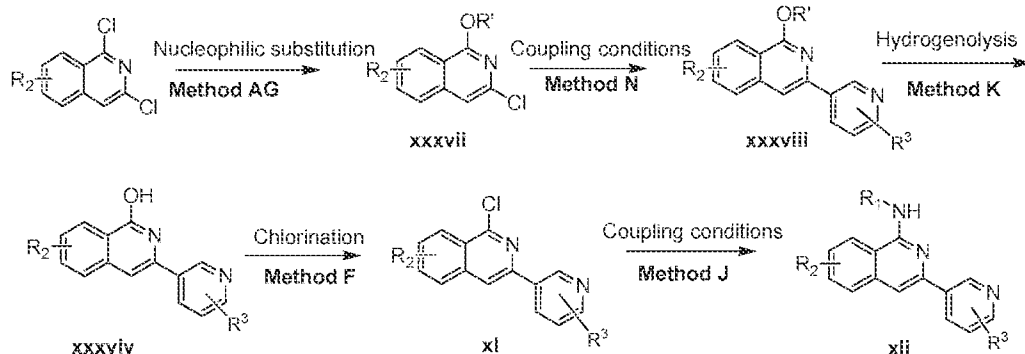
[0393] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 41**, replacing formaldehyde with the appropriate aldehyde and 2-aminobenzamide with the appropriate aniline.

Table 14:

Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1096			370.41	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.87 (s, 1H), 9.57 (d, J = 1.2 Hz, 1H), 9.25 (d, J = 8.8 Hz, 1H), 8.65 - 8.71 (m, 2H), 8.47 (s, 1H), 7.95 - 7.99 (m, 2H), 7.69 - 7.74 (m, 2H), 7.56 (dd, J = 7.7, 5.0 Hz, 1H), 7.33 (dd, J = 9.1, 1.9 Hz, 1H), 7.16 (t, J = 7.2 Hz, 1H), 6.90 (d, J = 1.2 Hz, 1H), 6.62 (d, J = 4.8 Hz, 1H), 2.88 (d, J = 4.8 Hz, 3H).	DMSO	371.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method AE
1099			370.41	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.71 (s, 1H), 9.58 (d, J = 1.4 Hz, 1H), 9.13 (d, J = 7.7 Hz, 1H), 8.80 - 8.64 (m, 2H), 8.45 (s, 1H), 7.92 (dd, J = 7.9, 1.4 Hz, 1H), 7.85 (d, J = 9.2 Hz, 2H), 7.76 - 7.62 (m, 1H), 7.57 (dd, J = 7.9, 4.8 Hz, 1H), 7.15 (dd, J = 11.2, 4.0 Hz, 1H), 7.04 (dd, J = 9.0, 2.3 Hz, 1H), 6.81 (q, J = 4.8 Hz, 1H), 6.70 (d, J = 2.2 Hz, 1H), 2.85 (d, J = 4.9 Hz, 3H).	DMSO	371.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method AE

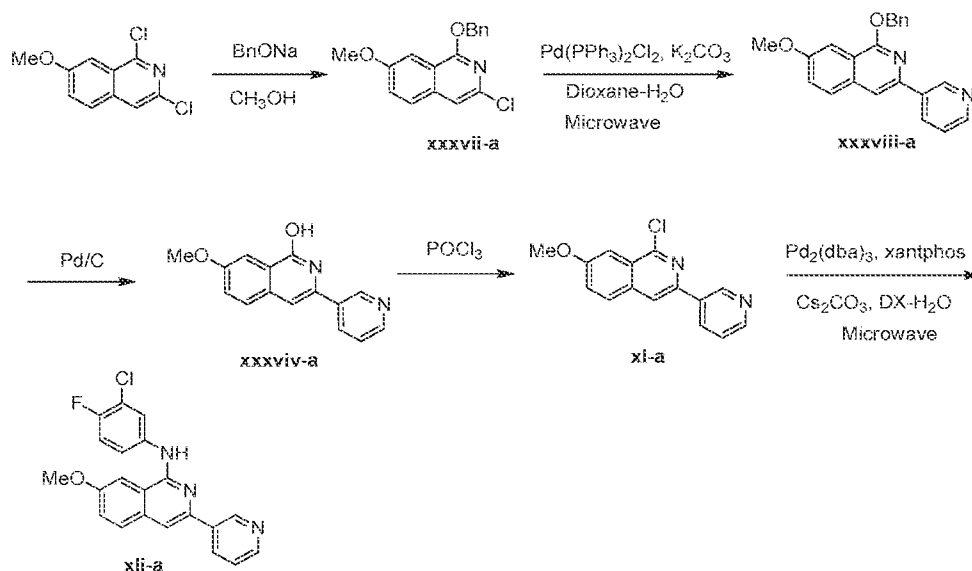
Scheme 43: General route for the synthesis of compounds with general formula xxxxi :

[0394]



Scheme 44: Representative synthesis of compounds of formula xli-a: (see Scheme 43)

[0395]



[0396] Method AG: 1-(Benzyloxy)-3-chloro-7-methoxyisoquinoline (xxxvii-a) 1,3-dichloro-7-methoxyisoquinoline (2.00 g, 8.8 mmol, 1.0 eq.) was dissolved in 30 mL anhydrous toluene and sodium benzyloxide (2.30 g, 17.6 mmol, 2.0 eq.) was added. The mixture was heated to 80°C for 18h. TLC indicated the reaction was complete.

[0397] The mixture was concentrated to give the crude product, which was purified by chromatography on silica gel (eluted with petroleum ether) to give 2.20 g of xxxvii-a as a white solid (84.6%). LCMS m/z = 300.1, 302.0 (M+1) (Method B) (retention time = 2.23 min).

[0398] 1-(Benzyloxy)-7-methoxy-3-(pyridin-3-yl)isoquinoline (xxxviii-a) 1-(Benzyloxy)-3-chloro-7-methoxyisoquinoline (93 mg, 0.5 mmol, 1.0 eq.), potassium carbonate (357mg, 2.5 mmol, 5.0 eq.), and Pd(PPh₃)₂Cl₂ (18 mg, 0.026 mmol, 0.05 eq.) was dissolved in the mixed solvent of 1,4-dioxane (3 mL) and water (1 mL). The resulting mixture was stirred at 120°C for 30 min under the microwave condition. The solid was filtrated off and the filtrate was concentrated. The residue was partitioned between ethyl acetate and water. The combined organic layers were washed with brine, dried over anhydrous sodium sulfate. After filtration and evaporation, the residue was purified by silica gel chromatography (petroleum ether/ethyl acetate = 1:10) to afford 100 mg of xxxviii-a as a yellow solid (84.7%).

[0399] 7-Methoxy-3-(pyridin-3-yl)isoquinolin-1-ol (xxxv-a): 7-Methoxy-3-(pyridin-3-yl)isoquinolin-1-ol (prepared in a manner analogous to that described for 2-amino-5-methoxybenzoic acid in Method K, replacing 5-methoxy-2-nitrobenzoic acid with 1-(benzyloxy)-7-methoxy-3-(pyridin-3-yl) isoquinoline) was obtained in a 63.6 % yield to

give 280 mg of xxxviv-a as a yellow solid. This was carried on without further purification.

[0400] 1-Chloro-7-methoxy-3-(pyridin-3-yl)isoquinoline (xl-a) 1-Chloro-7-methoxy-3-(pyridin-3-yl)isoquinoline (prepared in a manner analogous to that described for 4-chloro-6-propoxy-2-pyridin-3-yl-quinazoline in Method F1, replacing 6-propoxy-2-pyridin-3-yl-1H-quinazolin-4-one with 3-(pyridin-3-yl)isoquinolin-1-ol) was obtained in a 73.0% yield to give 60 mg of **xl-a** as a brown solid. MS $m/z = 241.1$ ($M+1$) (Method A) (retention time = 1.34 min).

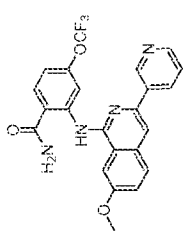
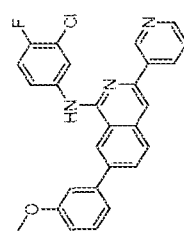
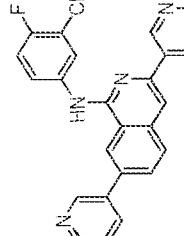
[0401] N-(3-Chloro-4-fluorophenyl)-7-methoxy-3-(pyridin-3-yl)isoquinolin-1-amine (xli-a) N-(N-(3-chloro-4-fluorophenyl)-7-methoxy-3-(pyridin-3-yl)isoquinolin-1-amine was prepared using **Method J**. A mixture of 1-chloro-7-methoxy-3-(pyridin-3-yl)isoquinoline (40 mg, 0.17 mmol, 1.0 eq.), 3-chloro-4-fluoro aniline (30 mg, 0.21 mmol, 1.2 eq.), $\text{Pd}_2(\text{dba})_3$ (10 mg, 0.011 mmol, 0.06 eq.), Xantphos (15 mg, 0.026 mmol, 0.15 eq.), cesium carbonate (167 mg, 0.52 mmol, 3.0 eq.) was suspended in the mixed solvent of 1,4-dioxane (4 mL) and water (1 mL). The resulting mixture was stirred at 120°C for 30 min under the microwave conditions. After cooling, the resulting mixture was partitioned between water and ethyl acetate. The combined organic layers were washed with water and brine,, dried over anhydrous sodium sulfate. After filtration and evaporation, the residue was purified by prep-HPLC to afford 3 mg of **xli-a** as a yellow solid (yield 5.2%). LCMS $m/z = 380.1$ ($M+1$) (Method B). ^1H NMR (400 MHz, DMSO-d_6): δ 9.35 (s, 1H), 9.30 (d, $J = 2.0$ Hz, 1H), 8.57 (dd, $J = 4.7, 1.5$ Hz, 1H), 8.41 (td, $J = 8.0, 1.8$ Hz, 1H), 8.30 (dd, $J = 6.9, 2.6$ Hz, 1H), 7.94 - 7.87 (m, 4H), 7.43 - 7.49 (m, 3H), 3.99 (s, 3H).

[0402] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 43**, replacing with the appropriate isoquinoline and aniline.

Table 15:

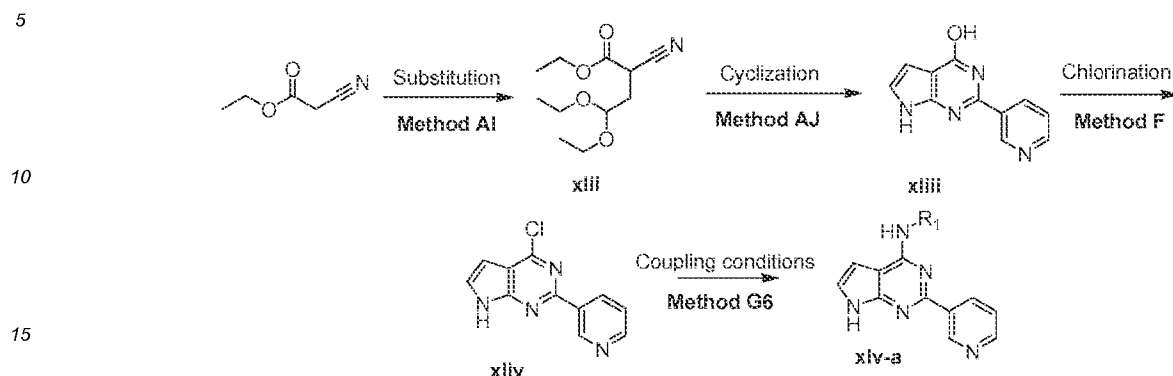
Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Parity percent	Method J
1100			379.81	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.35 (s, 1H), 9.30 (d, J = 2.0 Hz, 1H), 8.57 (dd, J = 4.7, 1.5 Hz, 1H), 8.41 (td, J = 8.0, 1.8 Hz, 1H), 8.30 (dd, J = 6.9, 2.6 Hz, 1H), 7.94 - 7.87 (m, 4H), 7.43 - 7.49 (m, 3H), 3.99 (s, 3H).	DMSO	380.1, 382.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method J
1101			411.38	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.46 (s, 1H), 9.31 (d, J = 1.75 Hz, 1H), 8.57 (dd, J = 4.6, 1.3 Hz, 1H), 8.45-8.41 (m, 1H), 8.19 (s, 1H), 7.99-7.94 (m, 2H), 7.91-7.86 (m, 2H), 7.54-7.44 (m, 3H), 7.01 (d, J = 8.1 Hz, 1H), 4.00 (s, 3H).	DMSO	412.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method J
1102			388.39	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.35 (s, 1H), 9.34 (d, J = 1.9 Hz, 1H), 8.80 (d, J = 8.4 Hz, 1H), 8.59 (dd, J = 4.7, 1.48 Hz, 1H), 8.49-8.45 (m, 1H), 8.26 (s, 1H), 8.13 (s, 1H), 8.02 (s, 1H), 7.97-7.92 (m, 1H), 7.61 (td, J = 14.9, 7.5 Hz, 1H), 7.56-7.48 (m, 3H), 6.95 (dd, J = 10.3, 8.3 Hz, 1H), 3.98 (s, 3H).	DMSO	389.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method J
1103			370.4	¹ H-NMR (400 MHz, CDCl ₃): δ 11.89 (s, 1H), 9.39 (s, 1H), 9.29 (s, 1H), 8.63-8.59 (m, 1H), 8.42-8.37 (m, 1H), 7.75 (d, J = 8.8 Hz, 1H), 7.60-7.62 (m, 4H), 7.44-7.33 (m, 3H), 7.05-7.00 (m, 1H), 4.06 (s, 3H).	CDCl ₃	371.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method J
1104			407.37	¹ H-NMR (400 MHz, CDCl ₃): δ 9.28 (s, 1H), 8.60 (d, J = 4.6 Hz, 1H), 8.34 (d, J = 7.8 Hz, 1H), 7.85 (s, 1H), 7.79 (d, J = 8.9 Hz, 1H), 7.63 (s, 1H), 7.43-7.35 (m, 2H), 7.22 (d, J = 8.6 Hz, 1H), 7.17 (s, 1H), 7.09-7.03 (m, 2H), 4.00 (s, 3H).	CDCl ₃	408.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method J

(continued)

Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Parity percent	Method J
1105			454.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.00 (s, 1H), 9.43 (d, J = 1.2 Hz, 1H), 9.35 (s, 1H), 8.63 (d, J = 3.7 Hz, 1H), 8.53 (s, 1H), 8.48 - 8.44 (m, 1H), 8.03 - 8.09 (m, 3H), 7.97 (d, J = 8.9 Hz, 1H), 7.65 (d, J = 1.8 Hz, 1H), 7.56 - 7.50 (m, 2H), 7.05 (dd, J = 8.6, 1.93 Hz, 1H), 4.01 (s, 3H).	DMSO	455.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method J
1106			455.91	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.65 (s, 1H), 9.37 (d, J = 1.7 Hz, 1H), 8.83 (s, 1H), 8.63 (dd, J = 4.7, 1.3 Hz, 1H), 8.54 - 8.43 (m, 1H), 8.35 (dd, J = 6.9, 2.6 Hz, 1H), 8.15 (dd, J = 8.5, 1.4 Hz, 1H), 8.03 (t, J = 4.3 Hz, 2H), 7.91 - 7.95 (m, 1H), 7.46 - 7.55 (m, 5H), 7.11 - 7.01 (m, 1H), 3.91 (s, 3H).	DMSO	456.1, 458.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method J
1107				¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.63 (s, 1H), 9.36 (d, J = 1.9 Hz, 1H), 9.19 (d, J = 1.9 Hz, 1H), 8.91 (s, 1H), 8.67 (dd, J = 4.8, 1.5 Hz, 1H), 8.62 (dd, J = 4.7, 1.5 Hz, 1H), 8.53 - 8.44 (m, 1H), 8.35 - 8.32 (m, 2H), 8.19 (dd, J = 8.5, 1.5 Hz, 1H), 8.13 - 8.01 (m, 2H), 7.92 (ddd, J = 9.1, 4.3, 2.7 Hz, 1H), 7.61 (dd, J = 7.9, 4.8 Hz, 1H), 7.55 (dd, J = 8.0, 4.8 Hz, 1H), 7.50 (t, J = 9.1 Hz, 1H).	DMSO	426.7, 428.7 (M+1)	Method B (NH ₄ HCO ₃)	95	Method J

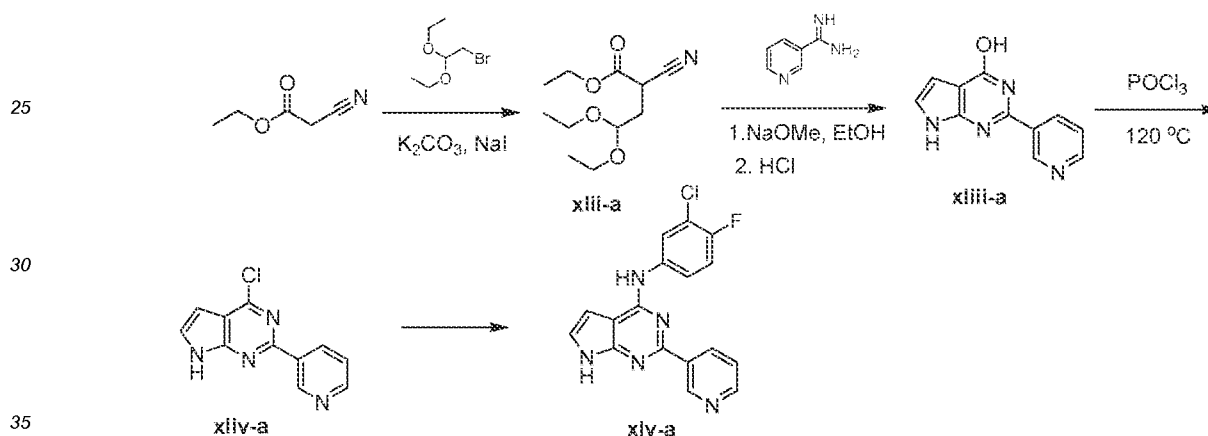
Scheme 45: General route for the synthesis of compounds with general formula xxxv-a :

[0403]



Scheme 46: Representative synthesis of compounds of formula xxxv-a: (see Scheme 45)

[0404]



[0405] **Method A1: Ethyl 2-cyano-4,4-diethoxybutanoate (xlii-a)** 2-Bromo-1,1-diethoxyethane (4 g, 20 mmol, 1.0 eq.) was added to a mixture of ethyl 2-cyanoacetate (11.4 g, 101 mmol, 5.0 eq.), K_2CO_3 (2.8 g, 20 mmol, 1.0 eq.) and NaI (200 mg, 1.3 mmol, 0.06 eq.), as described in *J. Chem. Soc.*, **1960**, 131-138. The reaction mixture was refluxed for 4 h at 145 °C. After cooling, the reaction mixture was purified by chromatography on silica gel (eluted with petroleum ether/ethyl acetate (80:1→40: 1→10:1) to give 3.57 g of **xlii-a** as a colorless oil (78%). 1H NMR (400 MHz, $CDCl_3$): δ 4.70 (t, J = 5.6 Hz, 1H), 4.26 (q, J = 7.2 Hz, 2H), 3.78 - 3.64 (m, 3H), 3.62 - 3.45 (m, 2H), 2.35 - 2.14 (m, 2H), 1.34 (q, J = 7.2 Hz, 3H), 1.25 - 1.16 (m, 6H).

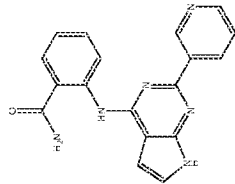
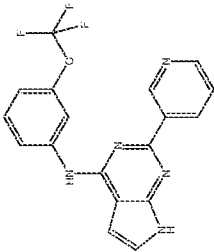
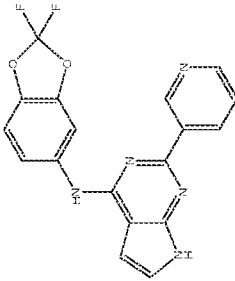
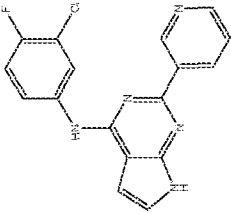
[0406] **Method AJ: 2-(Pyridin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-ol (xliii)** To a solution of nicotinimidamide (5.03 g, 41.6 mmol, 1.2 eq.) in EtOH (100 mL), was added NaOMe (4.8 g, 88.8 mmol, 2.5 eq.). The mixture was stirred at room temperature for 4 h. The reaction mixture was added to ethyl 2-cyano-4,4-diethoxybutanoate (8.00 g, 34.9 mmol, 1 eq.). This mixture was stirred at 105 °C overnight. After cooling, the reaction mixture was acidified with conc. HCl and stirred at room temperature for 2 h. A precipitate formed and was collected and washed with H_2O (20 mL X 2). After lyophilization, 3.10 g of product was obtained as a gray yellow solid (yield 41.8 %). LCMS m/z = 213.1 ($M + 1$) (Method B) (retention time = 1.07 min). 1H NMR (400 MHz, $DMSO-d_6$): δ 12.26 (s, 1H), 12.04 (s, 1H), 9.23 (d, J = 1.6 Hz, 1H), 8.70 (dd, J = 4.8, 1.2 Hz, 1H), 8.43 - 8.40 (m, 1H), 7.55 (dd, J = 8.0, 4.8 Hz, 1H), 7.12 (d, J = 1.6 Hz, 1H), 6.51 (d, J = 2.8 Hz, 1H).

[0407] **4-Chloro-2-(pyridin-3-yl)-7H-pyrrolo[2,3-d]pyrimidine (xliv-a)** 4-Chloro-2-(pyridin-3-yl)-7H-pyrrolo[2,3-d]pyrimidine (prepared in a manner analogous to that described for 4-chloro-6-propoxy-2-pyridin-3-yl-quinazolin-4-one in **Method F1**, replacing 6-propoxy-2-pyridin-3-yl-1H-quinazolin-4-one with 2-(pyridin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-ol) to give 450 mg of **xliv-a** in a 69.0 % yield as a brown solid. LCMS m/z = 231.0, 233.0 ($M + 1$) (Method B) (retention time = 1.60 min).

[0408] **N-(3-chloro-4-fluorophenyl)-2-(pyridin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine (xlv-a)** N-(3-chloro-4-

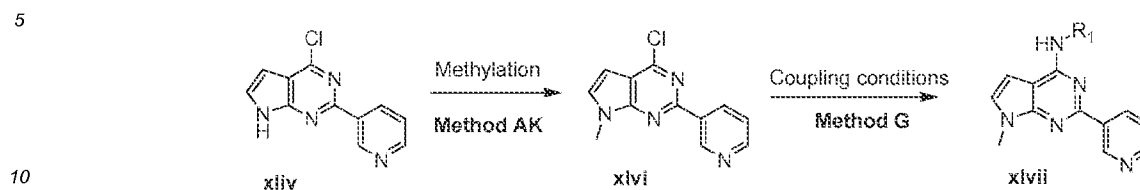
fluorophenyl)-2-(pyridin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine (prepared in a manner analogous to that described for N-(3-chloro-4-fluorophenyl)-6-(3-(dimethylamino)propyl)-2-(pyridin-3-yl)quinazolin-4-amine using Method G6, replacing 3-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)propyl methanesulfonate with 4-chloro-2-(pyridin-3-yl)-7H-pyrrolo[2,3-d]pyrimidine) to give a 11.3 % yield, 10 mg of **xlv-a** as brown solid. LCMS $m/z = 340.1, 342.0$ ($M + 1$) (Method B) (retention time = 1.814 min). ^1H NMR (400 MHz, DMSO- d_6): δ 9.48 (d, $J = 1.6$ Hz, 1H), 8.62 (dd, $J = 4.8, 1.6$ Hz, 1H), 8.60 - 8.56 (m, 1H), 8.27 (dd, $J = 6.4, 2.4$ Hz, 1H), 8.06 (ddd, $J = 8.8, 4.0, 2.8$ Hz, 1H), 7.71 (d, $J = 3.6$ Hz, 1H), 7.66 (t, $J = 9.2$ Hz, 1H), 7.50 (dd, $J = 8.0, 4.8$ Hz, 1H), 7.41 (br s, 2H), 6.85 (d, $J = 3.6$ Hz, 1H). **[0409]** The compounds in the following table were prepared in a manner analogous to that described in **Scheme 45**, replacing with the appropriate aniline.

Table 16:

Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for coupling
1108			330.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.29 (s, 1H), 12.09 (s, 1H), 9.54 (d, J = 1.2 Hz, 1H), 9.11 (d, J = 8.4 Hz, 1H), 8.65 - 8.68 (m, 1H), 8.35 (s, 1H), 7.89 (dd, J = 8.0, 1.2 Hz, 1H), 7.79 (s, 1H), 7.67 (td, J = 8.8, 1.2 Hz, 1H), 7.55 (dd, J = 7.2, 4.8 Hz, 1H), 7.43 - 7.41 (m, 1H), 7.10 (td, J = 8.0, 1.2 Hz, 1H), 6.49 (d, J = 3.6 Hz, 1H).	DMSO	331.1 (M + 1)	Method B (NH ₄ HCO ₃)	95	Method AJ, F, G6
1109			371.32	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.02 (d, J = 1.3 Hz, 1H), 9.75 (s, 1H), 9.50 (d, J = 1.5 Hz, 1H), 8.68 - 8.53 (m, 2H), 8.29 (s, 1H), 7.94 - 7.75 (m, 1H), 7.61 - 7.44 (m, 2H), 7.41 - 7.29 (m, 1H), 7.04 (s, 1H), 6.88 (dd, J = 3.4, 1.9 Hz, 1H).	DMSO	372.0 (M + 1)	Method A (TFA)	95	Method AJ, F, G6
1110			366.32	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.98 (s, 1H), 9.69 (s, 1H), 9.48 (d, J = 1.8 Hz, 1H), 8.64 - 8.59 (m, 2H), 8.19 (d, J = 2.0 Hz, 1H), 7.63 (dd, J = 8.8, 2.0 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 7.45 (d, J = 8.8 Hz, 1H), 7.34 (d, J = 3.0 Hz, 1H), 6.	DMSO	368.1 (M + 1)	Method A (TFA)	95	Method AJ, F, G6
1111			339.75	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.51 (s, 1H), 8.78 (d, J = 8.0 Hz, 2H), 8.17 (d, J = 9.2 Hz, 1H), 7.80 (dd, J = 9.2, 2.8 Hz, 1H), 7.70 - 7.64 (m, 1H), 7.46 (d, J = 2.8 Hz, 1H), 7.41 (d, J = 8.0 Hz, 1H), 7.20 (d, J = 4.0 Hz, 2H), 7.16 - 7.08 (m, 1H).	DMSO	340.1, 342.0 (M + 1)	Method B (NH ₄ HCO ₃)	95	Method AJ, F, G6

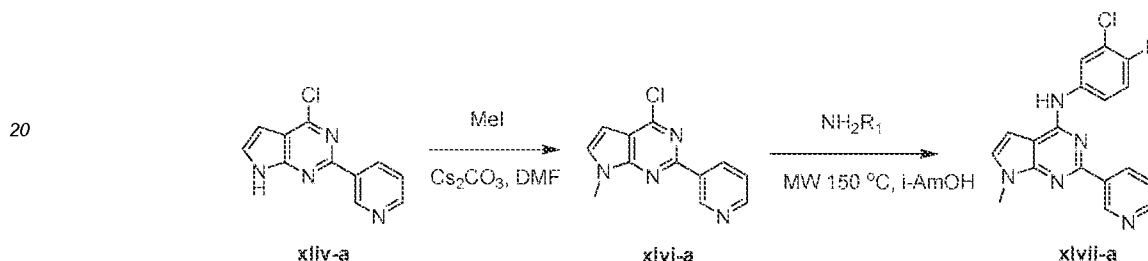
Scheme 47: General route for the synthesis of compounds with general formula xxxvii :

[0410]



Scheme 48: Representative synthesis of compounds of formula xxxvii-a: (see Scheme 47)

[0411]

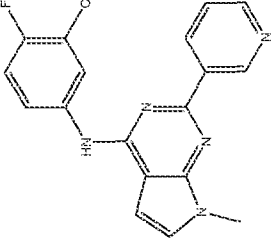
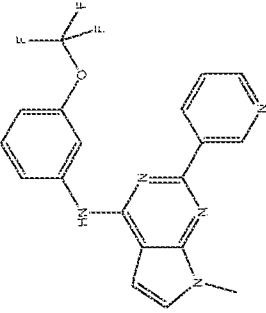
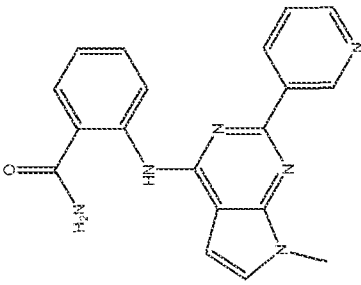


[0412] Method AK: 4-Chloro-7-methyl-2-(pyridin-3-yl)-7H-pyrrolo[2,3-d]pyrimidine (xliiv-a) To a solution of 4-chloro-2-(pyridin-3-yl)-7H-pyrrolo[2,3-d]pyrimidine (80 mg, 0.34 mmol, 1.0 eq.) in dry DMF (20 mL) was added Cs_2CO_3 (221 mg, 0.68 mmol, 2.0 eq.) and iodomethane (54.3 mg, 0.38 mmol, 1.1 eq.) at 0 °C. The reaction mixture was warmed to room temperature and stirred for 2.5 h. The reaction mixture was poured into ice water and extracted with EtOAc (20 mL x 3). The combined organic layers were dried over anhydrous Na_2SO_4 . After filtration and concentration, the residue was purified by chromatography on silica gel (petroleum ether/ethyl acetate 8:1) to give 65 mg of **xlvi-a** as brown solid (56.7%).

[0413] N-(3-chloro-4-fluorophenyl)-7-methyl-2-(pyridin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine (xlvi-a) N-(3-chloro-4-fluorophenyl)-7-methyl-2-(pyridin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine (prepared in a manner analogous to that described for N-(3-chloro-4-fluorophenyl)-6-(3-(dimethylamino)propyl)-2-(pyridin-3-yl)quinazolin-4-amine using Method G6, replacing 3-(4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazolin-6-yl)propyl methanesulfonate with 4-chloro-7-methyl-2-(pyridin-3-yl)-7H-pyrrolo[2,3-d]pyrimidine) to give a 70.0 % yield, 30 mg of **xlvi-a** as a brown solid. LCMS m/z = 354.1, 356.1 (M + 1) (Method B) (retention time = 1.94 min). ^1H NMR (400 MHz, DMSO-d_6): δ 9.97 (s, 1H), 9.53 (s, 1H), 9.15 (d, J = 8.0 Hz, 1H), 8.91 (d, J = 5.2 Hz, 1H), 8.25 (dd, J = 6.8, 2.4 Hz, 1H), 8.04 (dd, J = 7.6, 5.6 Hz, 1H), 7.94 - 7.90 (m, 1H), 7.48 - 7.43 (m, 2H), 6.95 (d, J = 3.2 Hz, 1H), 3.88 (s, 3H).

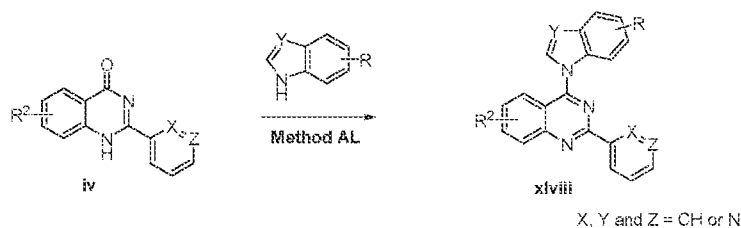
[0414] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 46**, replacing with the appropriate aniline.

Table 17:

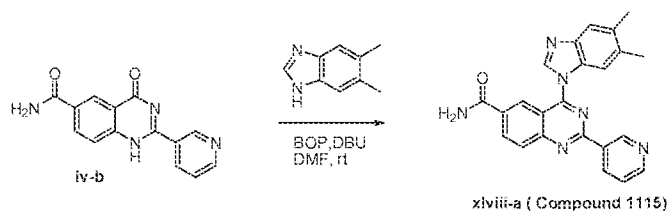
Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for coupling
1112		HCl	353.78	¹ H NMR (400 MHz, DMSO-d ₆): δ 9.97 (s, 1H), 9.53 (s, 1H), 9.15 (d, J = 8.0 Hz, 1H), 8.91 (d, J = 5.2 Hz, 1H), 8.25 (dd, J = 6.8, 2.4 Hz, 1H), 8.04 (dd, J = 7.6, 5.6 Hz, 1H), 7.94 - 7.90 (m, 1H), 7.48 - 7.43 (m, 2H), 6.95 (d, J = 3.2 Hz, 1H), 3.88 (s, 3H).	DMSO	354.1, 356.1 (M + 1)	Method B (NH ₄ HCO ₃)	95	Method AK, G6
1113			385.34	¹ H NMR (400 MHz, DMSO-d ₆): δ 9.81 (s, 1H), 9.56 (d, J = 1.6 Hz, 1H), 8.69 - 8.64 (m, 2H), 8.29 (s, 1H), 7.87 (d, J = 8.8 Hz, 1H), 7.55 - 7.51 (m, 2H), 7.41 (d, J = 3.6 Hz, 1H), 7.04 (d, J = 8.0 Hz, 1H), 6.90 (d, J = 3.6 Hz, 1H), 3.87 (s, 3H).	DMSO	386.1 (M + 1)	Method B (NH ₄ HCO ₃)	95	Method AK, G6
1114				¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.32 (s, 1H), 9.61 (s, 1H), 9.11 (d, J = 7.6 Hz, 1H), 9.11 (d, J = 7.6 Hz, 1H), 8.74 (dt, J = 8.0, 1.9 Hz, 1H), 8.67 (d, J = 3.6 Hz, 1H), 8.38 (s, 1H), 8.38 (s, 1H), 7.90 (dd, J = 7.9, 1.4 Hz, 1H), 7.82 (s, 1H), 7.68 (dd	DMSO	344.9 (M + 1)	Method B (NH ₄ HCO ₃)	95	Method AK, G6

Scheme 49: General route for the synthesis of compounds with general formula **xlvi**

[0415]

Scheme 50: Representative synthesis of compounds of formula **xlvi-a** (See Scheme 49)

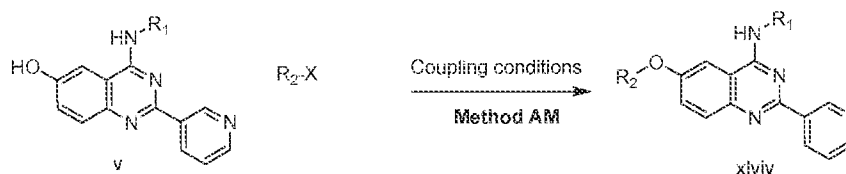
[0416]



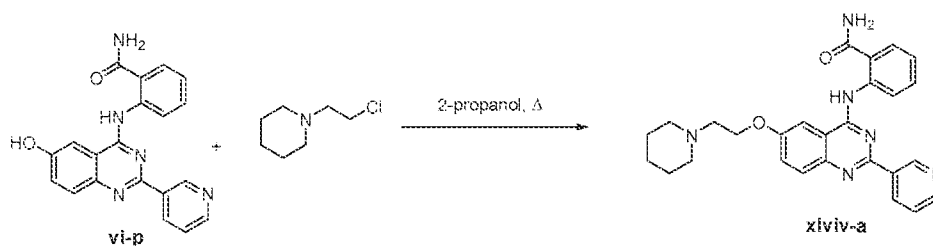
[0417] Method AL: 4-(5,6-dimethyl-1H-benzo[d]imidazol-1-yl)-2-(pyridin-3-yl)quinazoline-6-carboxamide (xlvi-a) To a suspension of 4-oxo-2-(pyridin-3-yl)-1,4-dihydroquinazolin-6-carboxamide (prepared as described in **scheme 4**) (100 mg, 0.376 mmol, 1.0 eq.) in dry DMF (20 mL) was added benzotriazole-1-yl-oxy-tris-(dimethylamino)-phosphonium hexafluorophosphate (216 mg, 0.488 mmol, 1.3 eq.) and diaza(1,3)bicyclo[5.4.0]undecene (114.5 mg, 0.752 mmol, 2.0 eq.) following a procedure outlined in J. Org. Chem., 2007, 72, 10194-10210. To the clear solution was then added 5,6-dimethyl-1H-benzo[d]imidazole (165.2 mg, 1.13 mmol, 3.0 eq.) and the mixture was stirred overnight at room temperature. The resultant precipitate was then collected by filtration and washed with dichloromethane, water, and ether. The product was dried *in vacuo* to give 26.3 mg of the desired product (**xlvi-a**) as an off-white solid (6.7 %). LCMS *m/z* = 395.1 (*M* + 1) (Method C) (retention time = 1.68 min). ¹H NMR (300 MHz, DMSO) δ 9.69 (s, 1H), 8.90 (s, 1H), 8.82 (d, *J* = 15.2 Hz, 2H), 8.66 (s, 1H), 8.54 (d, *J* = 8.6 Hz, 1H), 8.42 (s, 1H), 8.30 (d, *J* = 8.5 Hz, 1H), 7.79 (s, 1H), 7.68 (s, 3H), 2.39 (s, 3H), 2.36 (s, 3H).

Scheme 51: General route for the synthesis of compounds with general formula **xlvi**

[0418]

Scheme 52: Representative synthesis of compounds of formula **xlvi** (see Scheme 51)

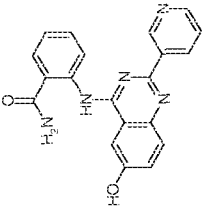
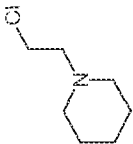
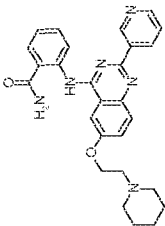
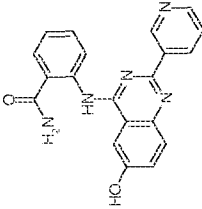
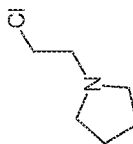
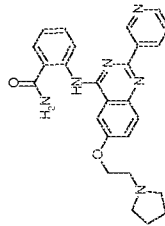
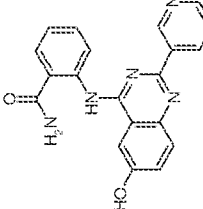
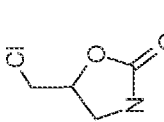
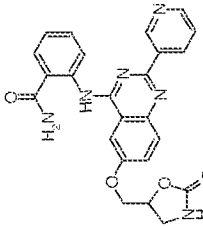
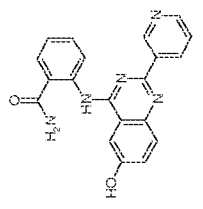
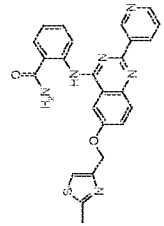
[0419]



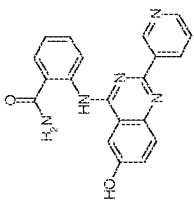
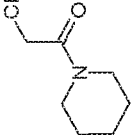
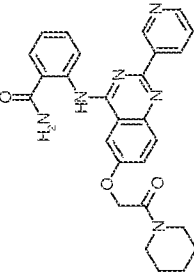
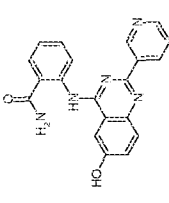
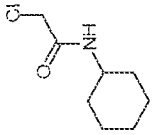
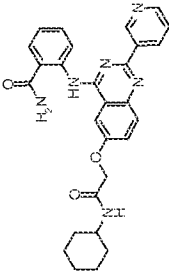
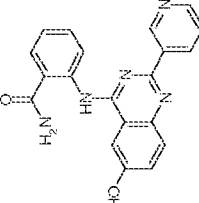
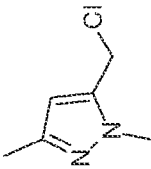
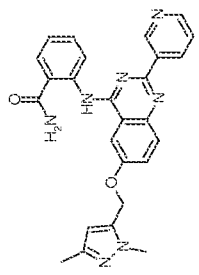
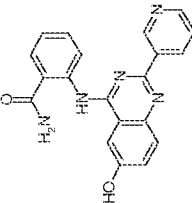
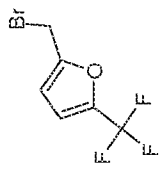
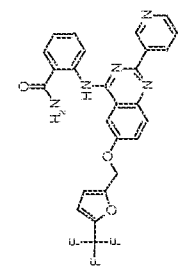
[0420] Method AM: Synthesis of 2-(6-(2-(piperidin-1-yl)ethoxy)-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (xliviv-a) To 1-(2-chloroethyl)piperidine (45 μmol) was added the solution of 2-(6-hydroxy-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (30 μmol) in NMP (200 μL). PS-BEMP (90 μmol) was added to the vials by resin dispenser. After the reaction mixture was heated at 90°C for 12 h, the residue was diluted with methanol and purified by mass triggered PREP-HPLC Condition D. The target fraction was lyophilized to afford the titled compound whose structure was finally confirmed by LCMS using LCMS **Method E**.

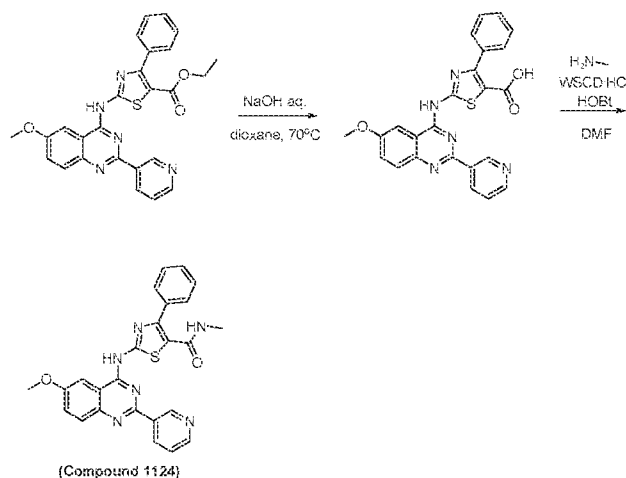
[0421] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 51**, replacing 1-(2-chloroethyl)piperidine with the appropriate alkyl halide.

Table 18:

Number	Starting Material 1	Starting Material 2	Product	Salt Type	Exact Mass	Mass Found (M+1)	Purity (%)
1116					468	469	98
1117					454	455	98
1118					456	457	98
1119					468	469	98

(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt Type	Exact Mass	Mass Found (M+1)	Purity (%)
1120					482	483	98
1121				TFA	496	497	98
1122					465	466	98
1123					505	506	98

Scheme 53: Synthesis of 2-(6-methoxy-2-(pyridin-3-yl)quinazolin-4-ylamino)-N-methyl-4-phenylthiazole-5-carboxamide dihydrochloride (Compound 1124)**[0422]**

[0423] To a suspension of ethyl 2-(6-methoxy-2-(pyridin-3-yl)quinazolin-4-ylamino)-4-phenylthiazole-5-carboxylate (1.3956 g, 2.89 mmol) in dioxane (40 mL) was added 1N NaOH (20 mL, 20.00 mmol) at room temperature to give a clear solution. The reaction mixture was stirred at room temperature for 1 h and then warmed to 50°C overnight, however starting material remained. An additional 20 mL of 1N NaOH was added and heating was continued at 50°C for 1 h and at 70 °C for 5 h 30 min. The reaction mixture was cooled to room temperature and diluted with water and ethyl acetate. The water phase was removed and adjusted to an acidic pH with 1N HCl (40 mL). A precipitate formed and was collected and washed with water. The product was *dried in vacuo* to give 1.20 g of a brown solid in a 91% yield. ¹H NMR (DMSO-d₆) ppm 12.88 (br, 2H), 9.77 (dd, J = 2.12, 0.6 Hz, 1H), 8.92-8.89 (m, 1H), 8.78 (dd, J = 4.8, 1.68 Hz, 1H), 8.32 (br, 1H), 7.96 (d, J = 9.12 Hz, 1H), 7.82-7.80 (m, 2H), 7.70-7.67 (m, 1H), 7.63 (dd, J = 9.12, 2.68 Hz, 1H), 7.50-7.44 (m, 3H), 3.98 (s, 3H).

[0424] Method AN: 2-(6-methoxy-2-(pyridin-3-yl)quinazolin-4-ylamino)-N-methyl-4-phenylthiazole-5-carboxamide To a suspension of 2-(6-methoxy-2-(pyridin-3-yl)quinazolin-4-ylamino)-4-phenylthiazole-5-carboxylic acid (291.8 mg, 0.641 mmol) in DMF (20 mL) under nitrogen atmosphere was added N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride (189 mg, 0.986 mmol) and 1-hydroxybenzotriazole hydrate (147 mg, 0.961 mmol) at room temperature. The reaction mixture was stirred at room temperature for 10 min to give a clear solution and methylamine in a methanol solution (4 mL) was added at room temperature. The reaction mixture was stirred at room temperature for 2 h 15 min. Additional methyl amine in a methanol solution (4 mL) was added at room temperature and then heated to 50°C for 1 h followed by room temperature for 2 days. The reaction mixture was partitioned between water and ethyl acetate. The water phase was collected and a solid precipitated from the water phase. The solid was filtered and dried in an oven at 60°C to give 107.9 mg (0.23 mmol) as the parent compound. The parent compound was suspended in methanol and 4N HCl in ethyl acetate (ca. 2 mL) was added to give a clear solution which precipitated over time. The solid was collected by filtration and dried in oven at 60°C for 2 days to give 82.4 mg of the HCl salt as a yellow solid in 24% yield. ¹H NMR (DMSO-d₆) δ 12.77 (brs, 1H), 9.76 (d, J = 1.72 Hz, 1H), 9.18 (d, J = 7.88 Hz, 1H), 8.92 (d J = 5.04 Hz, 1H), 8.35 (brs, 1H), 8.27 (brd, J = 4.56 Hz, 1H), 7.99-7.965 (m, 2H), 7.79 (brd, J = 7.16 Hz, 2H), 7.65 (dd, J = 9.12, 2.56 Hz, 1H), 7.50-7.41 (m, 3H), 4.00 (s, 3H), 2.76 (d, J = 4.56 Hz, 3H). The 1H of 2HCl was not observed.

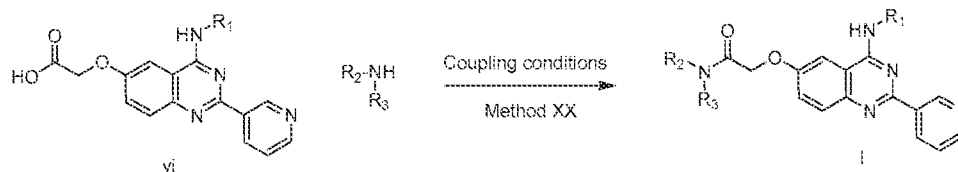
[0425] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 53**, replacing N-methyl amine with the appropriate alkyl amine.

Table 19:

Number	Starting Material 1	Starting Material 2	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1124				2 HCl	¹ H NMR (DMSO-d ₆) ppm 12.77 (brs, 1H), 9.76 (d, J = 1.72 Hz, 1H), 9.18 (d, J = 7.88 Hz, 1H), 8.92 (d, J = 5.04 Hz, 1H), 8.35 (brs, 1H), 8.27 (brd, J = 4.56 Hz, 1H), 7.99-7.965 (m, 2H), 7.79 (brd, J = 7.16 Hz, 2H), 7.65 (dd, J = 9.12, 2.56 Hz, 1H), 7.50-7.41 (m, 3H), 4.00 (s, 3H), 2.76 (d, J = 4.56 Hz, 3H). The 1H of 2HCl was not observed.	DMSO	>98	Method AN
1125				2 HCl	¹ H NMR (DMSO-d ₆) ppm 12.75 (brs, 1H), 9.77 (d, J = 1.52 Hz, 1H), 9.13 (brs, 1H), 8.88 (brd, J = 5.08 Hz, 1H), 8.34 (d, J = 2.04 Hz, 1H), 7.98 (d, J = 9.12 Hz, 1H), 7.90 (brm, 1H), 7.83-7.80 (m, 2H), 7.66-7.53 (brm, 3H), 7.51-7.42 (m, 3H), 4.00 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	Method AN
1126					¹ H NMR (DMSO-d ₆) ppm 12.66 (s, 1H), 9.76 (s, 1H), 8.88 (d, J = 7.76 Hz, 1H), 8.75 (d, J = 3.68 Hz, 1H), 8.40-8.20 (br, 2H), 7.94 (br, 1H), 7.80 (d, J = 7.16 Hz, 2H), 7.63 (m, 2H), 7.49-7.41 (m, 3H), 3.99 (s, 3H), 3.24 (q, J = 7.24 Hz, 2H), 1.07 (t, J = 7.24 Hz, 3H).	DMSO	>98	Method AN

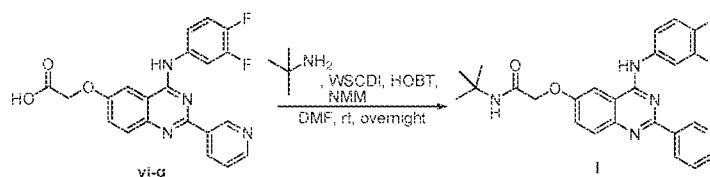
Scheme 54: General route for the synthesis of compounds with general formula I

[0426]



Scheme 55: Representative synthesis of compounds of formula I-a: (see Scheme 54)

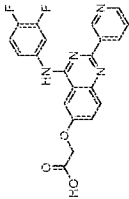
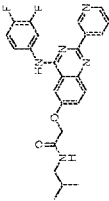
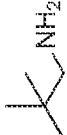
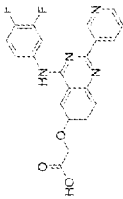
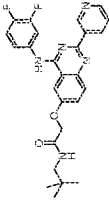
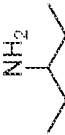
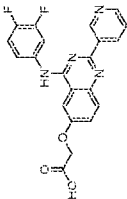
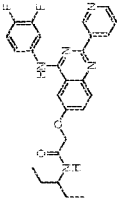
[0427]

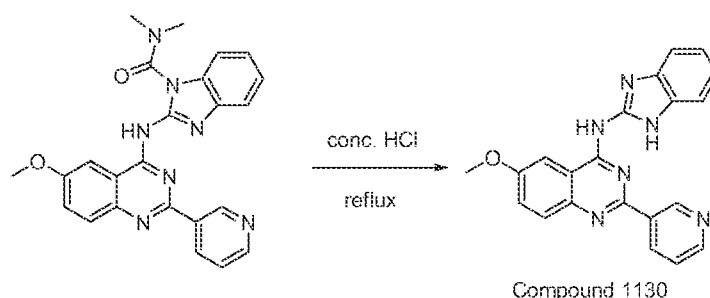
[0428] Method AO: *N*-tert-butyl-2-(4-(3,4-difluorophenyl-amino)-2-(pyridin-3-yl)quinazolin-6-yloxy)acetamide

To a solution of carboxylic acid derivative (500 mg, 1.2 mmol) in DMF was added 2-aminoisobutane (134 mg, 1.8 mmol), NMM (0.4 mL, 3.6 mmol), WSCDI (282 mg, 1.4 mmol) and HOBT (225 mg, 1.4 mmol). The reaction mixture was stirred at room temperature overnight. The resulting solution was poured into ice-water and the precipitate formed and was filtered off. The solid was washed with water and dried to give 350 mg (62% yield) of desired product. ¹H NMR (400 MHz, DMSO) δ 9.86 (s, 1H), 9.49 (d, *J* = 1.5 Hz, 1H), 8.71 - 8.57 (m, 2H), 8.11 (ddd, *J* = 13.3, 7.5, 2.5 Hz, 1H), 8.00 - 7.93 (m, 1H), 7.87 (d, *J* = 9.1 Hz, 1H), 7.67 (ddd, *J* = 11.7, 7.1, 2.1 Hz, 2H), 7.59 - 7.47 (m, 3H), 4.61 (s, 2H), 1.35 (s, 9H).

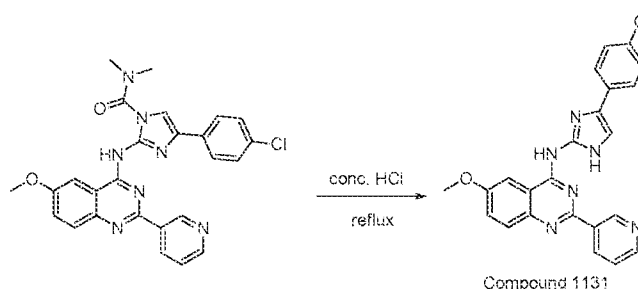
[0429] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 54**, replacing 2-aminoisobutane with the appropriate alkyl amine.

Table 20:

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1127				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.60 (s, 1H), 9.48 (d, J = 1.9 Hz, 1H), 9.14 (dt, J = 8.2, 1.6 Hz, 1H), 8.96 (dd, J = 5.5, 1.3 Hz, 1H), 8.37 (t, J = 6.0 Hz, 1H), 8.24 (d, J = 2.6 Hz, 1H), 8.11 - 7.97 (m, 3H), 7.74 (ddd, J = 11.8, 7.4, 3.3 Hz, 2H), 7.55 (dt, J = 10.5, 9.1 Hz, 1H), 4.79 (s, 2H), 3.01 (t, J = 6.5 Hz, 2H), 1.86 - 1.59 (m, 1H), 0.85 (d, J = 6.7 Hz, 6H).	DMSO	>98	Method AO
1128				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.81 (s, 1H), 9.50 (d, J = 1.9 Hz, 1H), 9.16 (d, J = 8.2 Hz, 1H), 8.98 (dd, J = 5.5, 1.3 Hz, 1H), 8.40 (d, J = 2.5 Hz, 1H), 8.29 (t, J = 6.4 Hz, 1H), 8.18 - 7.95 (m, 3H), 7.83 - 7.67 (m, 2H), 7.55 (dt, J = 10.5, 9.1 Hz, 1H), 4.88 (s, 2H), 2.98 (d, J = 6.4 Hz, 1H), 0.84 (s, 9H).	DMSO	>98	Method AO
1129				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.87 (s, 1H), 9.50 (d, J = 1.9 Hz, 1H), 9.16 (d, J = 8.2 Hz, 1H), 8.98 (dd, J = 5.5, 1.3 Hz, 1H), 8.41 (d, J = 2.5 Hz, 1H), 8.23 - 7.96 (m, 4H), 7.85 - 7.66 (m, 2H), 7.55 (dt, J = 10.5, 9.1 Hz, 1H), 4.85 (s, 2H), 3.80 - 3.43 (m, 1H), 1.65 - 1.21 (m, 4H), 0.81 (t, J = 7.4 Hz, 6H).	DMSO	>98	Method AO

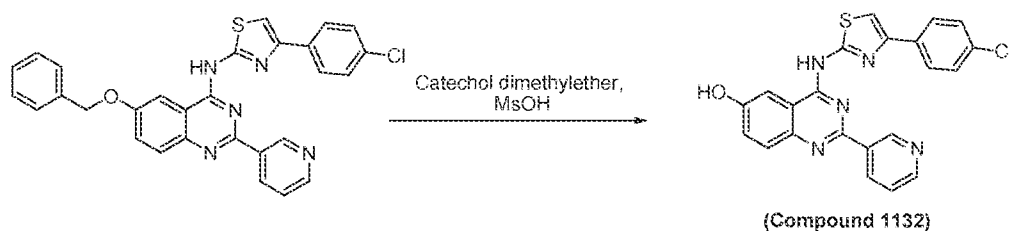
Scheme 56: Synthesis of N-(1H-benzo[d]imidazol-2-yl)-6-methoxy-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride (Compound 1130)**[0430]**

[0431] 2-(6-methoxy-2-(pyridin-3-yl)quinazolin-4-ylamino)-N,N-dimethyl-1H-benzo[d]imidazole-1-carboxamide (334.1 mg, 0.760 mmol) was dissolved in concentrated HCl (0.063 ml, 0.760 mmol). The solution was stirred at room temperature overnight and then at 50°C for 3 h 30 min followed by 100°C for 3 h. The solid was collected and dried *in vacuo* to give 116.2 mg of a yellow solid in a yield 35%. ¹H NMR (DMSO-*d*₆) δ 13.22 (br, 1H), 9.55 (s, 1H), 8.92 (dd, J = 4.96, 1.36 Hz, 1H), 8.88 (br, 1H), 8.01 (d, J = 9.12 Hz, 1H), 7.92 (br, 1H), 7.83 (br, 1H), 7.67 (dd, J = 9.12, 2.80 Hz, 1H), 7.61 (m, 2H), 7.38 (m, 2H), 3.97 (s, 3H). The 1H of 2HCl and NH - were not observed.

Scheme 57: Synthesis of N-(4-(4-chlorophenyl)-1H-imidazol-2-yl)-6-methoxy-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride (Compound 1131)**[0432]**

[0433] 4-(4-chlorophenyl)-2-(6-methoxy-2-(pyridin-3-yl)quinazolin-4-ylamino)-N,N-dimethyl-1H-imidazole-1-carboxamide and concentrated HCl_(aq) (10 mL) were added to a round bottom flask, a precipitate appeared upon refluxing the mixture for 3 h. The solid was collected (373.4 mg) and triturated with CH₂Cl₂ / methanol overnight. The product was filtered and dried to give 261.7 mg of a solid. A suspension of the product in methanol was added to a 1N solution of NaOH_(aq.) (5 mL) followed by CH₂Cl₂ and H₂O. The solid was collected and washed with methanol to give 255.7 mg (0.596 mmol) of the parent product. The free parent was suspended in CH₂Cl₂ /methanol and converted to the HCl salt by addition of a 0.3 ml solution of 4N HCl in ethyl acetate. The HCl salt was collected and dried *in vacuo* to give 248.4 mg as brown solid in a 26% yield. ¹H NMR (DMSO-*d*₆) δ 13.61 (br, 1H), 12.81 (br, 1H), 9.46 (s, 1H), 8.87 (d, J = 5.24 Hz, 1H), 8.70 (br, 1H), 7.91-7.87 (m, 5H), 7.75 (m, 1H), 7.62-7.56 (m, 3H), 3.95 (s, 3H). The 1H of 2HCl was not observed.

Scheme 58: Synthesis of 4-(4-(4-chlorophenyl)thiazol-2-ylamino)-2-(pyridin-3-yl)quinazolin-6-ol (Compound 1132)**[0434]**

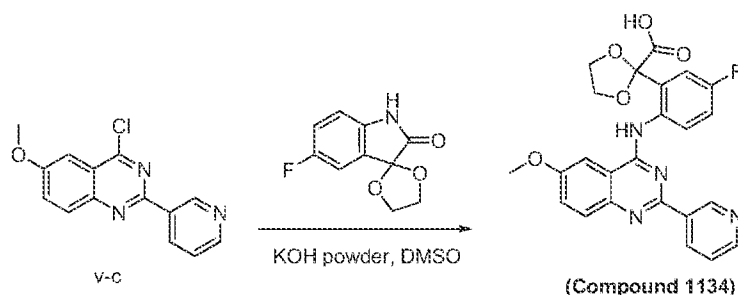


[0435] To a mixture of N-(6-(benzyloxy)-2-(pyridin-3-yl)quinazolin-4-yl)-4-(4-chlorophenyl)thiazol-2-amine and catechol, dimethyl ether (0.16 g, 1.15 mmol) was added methanesulfonic acid (4.0 mL). The mixture was stirred for 1 hr and was then poured into water. The slurry was added to a stirring sat. $\text{NaHCO}_3(\text{aq})$ solution slowly and allowed to stir for 30min. The precipitate was filtered to give a brown solid which was washed with methanol to give 4-(4-(4-chlorophenyl)thiazol-2-ylamino)-2-(pyridin-3-yl)quinazolin-6-ol (0.23g, 91.0 %).

[0436] Scheme 59: Synthesis of 4-(4-phenylthiazol-2-ylamino)-2-(pyridin-3-yl)quinazolin-6-ol (Compound 1133) Synthesis of 4-(4-phenylthiazol-2-ylamino)-2-(pyridin-3-yl)quinazolin-6-ol was performed in a similar manner to that described for 4-(4-(4-chlorophenyl)thiazol-2-ylamino)-2-(pyridin-3-yl)quinazolin-6-ol substituting for 4-(4-phenylthiazol-2-ylamino)-2-(pyridin-3-yl)quinazolin-6-ol for the N-(6-(benzyloxy)-2-(pyridin-3-yl)quinazolin-4-yl)-4-(4-chlorophenyl)thiazol-2-amine giving 4-(4-phenylthiazol-2-ylamino)-2-(pyridin-3-yl)quinazolin-6-ol. ^1H NMR (300 MHz, DMSO) δ 10.06 - 9.22 (m, 2H), 8.95 (dt, J = 7.9, 1.6 Hz, 1H), 8.62 (dd, J = 4.7, 1.4 Hz, 1H), 7.99 (d, J = 7.5 Hz, 2H), 7.87 (d, J = 2.8 Hz, 1H), 7.59 - 7.47 (m, 2H), 7.45 - 7.32 (m, 3H), 7.30 - 7.09 (m, 2H).

Scheme 60: Synthesis of 2-(5-fluoro-2-(6-methoxy-2-(pyridin-3-yl)quinazolin-4-ylamino)phenyl)-1,3-dioxolane-2-carboxylic acid (Compound 1134)

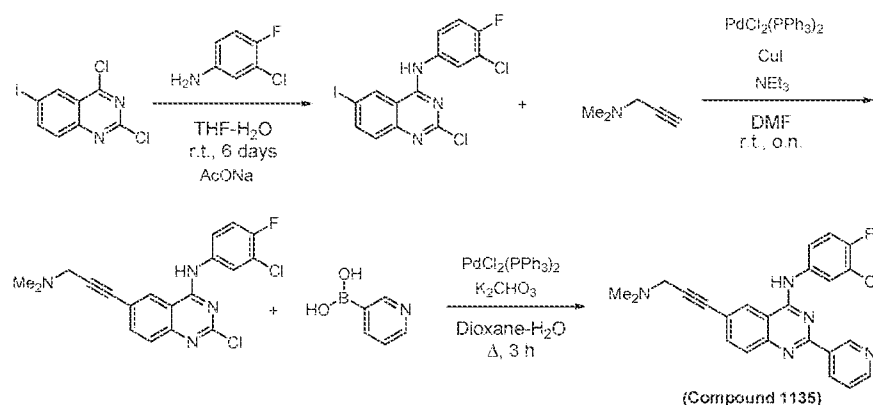
[0437]



[0438] To a mixture of 4-chloro-6-methoxy-2-(pyridin-3-yl)quinazoline (0.50 g, 1.84 mmol) and 5'-fluorospiro[[1,3]dioxolane-2,3'-indolin]-2'-one (0.42 g, 2.02 mmol) in dry DMSO (4 mL) was added KOH powder (0.11g, 2.02 mmol). The reaction mixture was stirred for 15 h at room temperature and then the reaction mixture was poured into water. The aqueous layer was washed with ethyl acetate (2x 20mL) and the resulting aqueous layer was acidified with 5N HCl to give a precipitate. The solid was filtered to give 2-(5-fluoro-2-(6-methoxy-2-(pyridin-3-yl)quinazolin-4-ylamino)phenyl)-1,3-dioxolane-2-carboxylic acid as a light yellow powder (0.27g, 0.58 mmol, 32 %). LCMS m/z = 432 ($M + 1$) (Method C) ^1H NMR (300 MHz, DMSO) δ 9.50 - 9.38 (m, 2H), 8.69 - 8.59 (m, 2H), 8.42 (dd, J = 9.7, 5.3 Hz, 1H), 7.86 (d, J = 9.1 Hz, 1H), 7.64 (d, J = 2.6 Hz, 1H), 7.61 - 7.41 (m, 4H), 4.25 - 4.11 (m, 4H), 3.98 (s, 3H).

Scheme 61: N-(3-chloro-4-fluorophenyl)-6-(3-(dimethylamino)prop-1-ynyl)-2-(pyridin-3-yl)quinazolin-4-amine (compound 1135)

[0439]



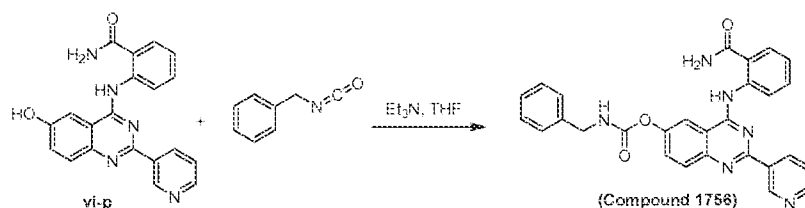
[0440] In a 50 mL round-bottomed flask was added 2,4-dichloro-6-iodoquinazoline (0.52 g, 1.6 mmol), 3-chloro-4-fluoroaniline (0.30 g, 2.1 mmol), and sodium acetate (0.20 g, 2.4 mmol) in THF (6 mL) and water (2 mL) to give a brown suspension. After being stirred at room temperature for 6 days, the reaction mixture was diluted with water (15 mL) and extracted with ethyl acetate (2 x 10 mL). The organic layers were combined and washed with brine (1 x 20 mL), dried over Na_2SO_4 , filtered and concentrated. The resulting product was washed with CH_2Cl_2 and dried to give 0.51 g of 2-chloro-N-(3-chloro-4-fluorophenyl)-6-iodoquinazolin-4-amine as a light brown solid in a 73 % yield. ^1H NMR (300 MHz, DMSO) δ 10.29 (s, 1H), 8.95 (s, 1H), 8.14 (dd, J = 1.5, 9.0 Hz, 1H), 8.06 (dd, J = 2.7, 6.6 Hz, 1H), 7.81 - 7.76 (m, 1H), 7.52 - 7.46 (m, 2H).

[0441] In a 50 mL round-bottomed flask was added 2-chloro-N-(3-chloro-4-fluorophenyl)-6-iodoquinazolin-4-amine (200 mg, 0.46 mmol), N,N-dimethylpropargylamine (99 mL, 0.92 mmol), NEt_3 (0.26 mL, 1.84 mmol), CuI (0.88 mg, 4.6 mmol) and $\text{PdCl}_2(\text{PPh}_3)_2$ (6.5 mg, 9.2 mmol) in DMF (3 mL) to give a light yellow suspension. The mixture was stirred at room temperature overnight under an argon atmosphere. The reaction mixture was diluted with water (10 mL) and ethyl acetate (10 mL) and then a precipitate formed. The resulting precipitate was removed by filtration through Celite. The filtrate was extracted with ethyl acetate (2 x 10 mL). The combined organic layer was washed with water (1 x 15 mL) and brine (1 x 15 mL) and was dried over Na_2SO_4 , filtered and concentrated. The residue was purified by column chromatography on silica gel (eluted with $\text{CH}_2\text{Cl}_2/\text{MeOH}$ 1:0 to 9:1). The desired product was washed with CH_2Cl_2 to give 61 mg of 2-chloro-N-(3-chloro-4-fluorophenyl)-6-(3-(dimethylamino)prop-1-ynyl)quinazolin-4-amine as pale yellow solid in a 34 % yield. LCMS m/z = 389 ($M+1$) (Method C) (retention time = 2.24 min). ^1H NMR (300 MHz, DMSO) δ 10.30 (s, 1H), 8.68 (s, 1H), 8.07 (dd, J = 2.7, 6.9 Hz, 1H), 7.86 (dd, J = 1.8, 8.7 Hz, 1H), 7.81 - 7.70 (m, 1H), 7.68 (d, J = 8.4 Hz, 1H), 7.49 (t, J = 9.2 Hz, 1H), 3.52 (s, 2H), 2.28 (s, 6H).

[0442] In a 50 mL round-bottomed flask was added 2-chloro-N-(3-chloro-4-fluorophenyl)-6-(3-(dimethylamino)prop-1-ynyl)quinazolin-4-amine (61 mg, 0.16 mmol), 3-pyridineboronic acid (25 mg, 0.20 mmol), K_2CO_3 (0.11 mg, 0.78 mmol) and $\text{PdCl}_2(\text{PPh}_3)_2$ (5.5 mg, 7.8 mM) in dioxane (2 mL) to give a yellow suspension. The mixture was heated at reflux for 3 h under argon. After cooling to room temperature, water (10 mL) and ethyl acetate (10 mL) were added to the mixture to form a precipitate. The resulting precipitate was filtered through Celite. The filtrate was extracted with ethyl acetate (2 x 10 mL) and the combined organic layers were washed with water (1 x 15 mL) and brine (1 x 15 mL) and then dried over Na_2SO_4 , filtered and concentrated. The residue was purified by column chromatography on silica gel (eluted with AcOEt/MeOH 1:0 to 9:1). The desired product was washed with CH_2Cl_2 to give 25 mg of N-(3-chloro-4-fluorophenyl)-6-(3-(dimethylamino)prop-1-ynyl)-2-(pyridin-3-yl)quinazolin-4-amine as a light brown solid in 37 % yield. LCMS m/z = 432 ($M+1$) (Method C) (retention time = 1.86 min). ^1H NMR (300 MHz, DMSO) δ 10.13 (s, 1H), 9.51 (s, 1H), 8.77 - 8.56 (m, 3H), 8.26 (dd, J = 6.9, 2.5 Hz, 1H), 8.02 - 7.78 (m, 3H), 7.64 - 7.47 (m, 2H), 3.54 (s, 2H), 2.30 (s, 6H).

Scheme 62: Synthesis of 4-(2-carbamoylphenylamino)-2-(pyridin-3-yl)quinazolin-6-yl benzylcarbamate (compound 1756)

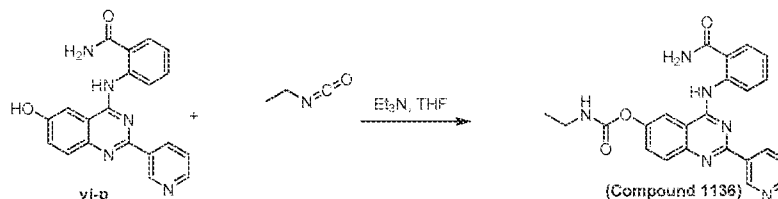
[0443]



[0444] A mixture of 2-(6-hydroxy-2-(pyridin-3-yl)quinazolin-4-ylamino)benzamide (200 mg, 0.283 mmol), (isocyanato-methyl)benzene (5 mL) and Et₃N (57 mg, 0.283 mmol) in tetrahydrofuran (THF) (10 mL) was stirred at room temperature overnight. Water (10 mL) was added to the above mixture and the mixture was concentrated *in vacuo*. The precipitate was collected by filtration and washed with water (6 mL x 2) to afford 72 mg of the desired product as a white solid in a yield 26.0%. LCMS: rt = 1.829 min, [MH]⁺ = 491.1 ¹H-NMR (400 MHz, DMSO-d₆): δ 13.05 (s, 1H), 9.63 (s, 1H), 9.07 (d, J = 8.5 Hz, 1H), 8.87 (d, J = 8.0 Hz, 1H), 8.79 (d, J = 3.7 Hz, 1H), 8.59 (t, J = 6.1 Hz, 1H), 8.49 (s, 1H), 8.02 - 7.89 (m, 4H), 7.82 - 7.68 (m, 3H), 7.46 - 7.22 (m, 6H), 4.35 (d, J = 6.1 Hz, 2H).

Scheme 63: Synthesis of 4-(2-carbamoylphenylamino)-2-(pyridin-3-yl)quinazolin-6-yl ethylcarbamate (compound 1136)

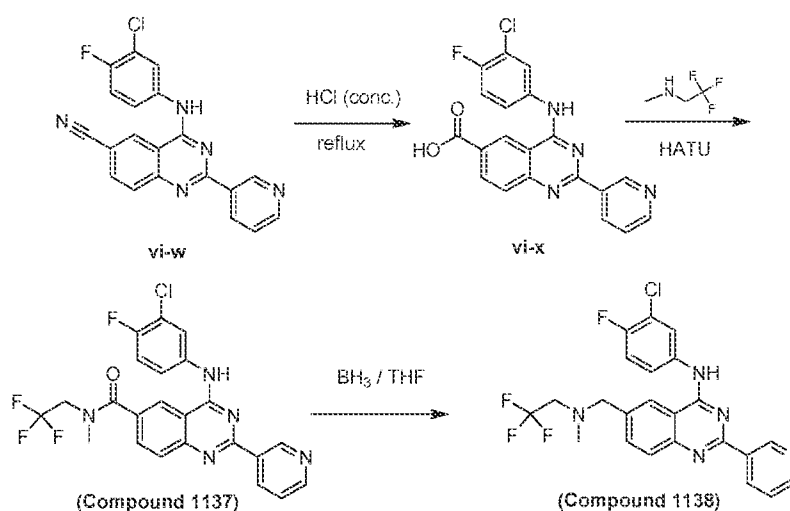
[0445]



[0446] 4-(2-carbamoylphenylamino)-2-(pyridin-3-yl)quinazolin-6-yl ethylcarbamate was synthesized in a similar manner to that described for 4-(2-carbamoylphenylamino)-2-(pyridin-3-yl)quinazolin-6-yl benzylcarbamate substituting isocyanatoethane for (isocyanatomethyl)benzene. The resulting product was analyzed by LCMS: rt = 1.11 min, [M+1]⁺ = 429.0. ¹H-NMR (400 MHz, DMSO-d₆): δ 13.06 (s, 1H), 9.66 (s, 1H), 9.10 (d, J = 8.3 Hz, 1H), 8.87 - 8.80 (m, 2H), 8.51 (s, 1H), 8.06 - 7.94 (m, 4H), 7.88 (s, 1H), 7.79 - 7.63 (m, 3H), 7.24 (t, J = 7.6 Hz, 1H), 3.22 - 3.11 (m, 2H), 1.14 (t, J = 7.2 Hz, 3H).

Scheme 64: 4-(3-chloro-4-fluorophenylamino)-N-methyl-2-(pyridin-3-yl)-N-(2,2,2-trifluoroethyl)quinazoline-6-carboxamide (compound 1137) and N-(3-chloro-4-fluorophenyl)-6-((methyl(2,2,2-trifluoroethyl)amino)methyl)-2-(pyridin-3-yl)quinazolin-4-amine (compound 1138)

[0447]



[0448] A mixture of 4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazoline-6-carbonitrile (1.5 g, 4.0 mmol) in HCl (conc, 25 mL) was heated to 100 °C and stirred overnight. After cooling and filtration, the solid was washed with water (10 mL) twice to give 1.4 g of the desired product 4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazoline-6-carboxylic acid as a yellow solid in a 89.0% yield. LCMS: r.t = 1.271 min, [MH]⁺ = 394.9

[0449] A mixture of 4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)quinazoline-6-carboxylic acid (900 mg, 2.4 mmol), DIPEA (620 mg, 4.8 mmol) and HATU (1.4 g, 3.6 mmol) in DMF (10 mL) was pre-stirred for 20 min, and 2,2,2-trifluoro-

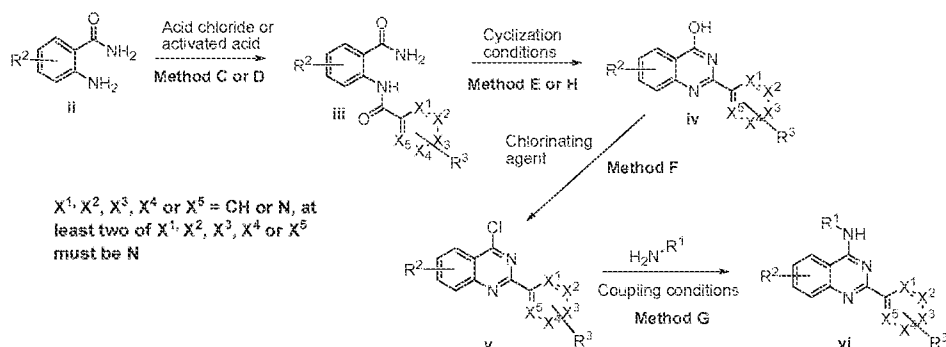
N-methylethanamine (360 mg, 2.4 mmol) was added in one portion. The resulting mixture was stirred at room temperature overnight. Water (80 mL) was added and a precipitate formed which was collected and purified by prep-HPLC to afford 490 mg of the desired product 4-(3-chloro-4-fluorophenylamino)-N-methyl-2-(pyridin-3-yl)-N-(2,2,2-trifluoroethyl)quinazoline-6-carboxamide as a white solid in 42.0% yield. LCMS: r.t = 1.970 min, [MH]⁺ = 490.0 ¹H-NMR (400 MHz, DMSO-*d*₆): δ 10.79 (brs, 1H), 9.59 (s, 1H), 9.19 (d, *J* = 7.6 Hz, 1H), 9.04 (d, *J* = 4.8 Hz, 1H), 8.90 (s, 1H), 8.22 - 8.00 (m, 5H), 7.62 (t, *J* = 9.2 Hz, 1H), 4.55 - 4.52 (m, 2H), 3.22 (s, 3H).

[0450] To a mixture of 4-(3-chloro-4-fluorophenylamino)-N-methyl-2-(pyridin-3-yl)-N-(2,2,2-trifluoroethyl)quinazoline-6-carboxamide (60 mg, 0.12 mmol) in THF (1 mL) was added BH₃-THF (2 mol/L, 1 mL). The mixture was stirred at room temperature overnight. Methanol (0.2 mL) was added to quench the reaction mixture. The mixture was purified by prep-HPLC to give the desired product N-(3-chloro-4-fluorophenyl)-6-((methyl(2,2,2-trifluoroethyl)amino)methyl)-2-(pyridin-3-yl)quinazolin-4-amine (GL001H-3309) as a white solid 15 mg in a yield of 26.0%. LCMS: r.t = 2.154 min, [MH]⁺ = 476.1. ¹H-NMR (400 MHz, DMSO-*d*₆): δ 10.09 (s, 1H), 9.53 (s, 1H), 8.70 - 8.66 (m, 2H), 8.44 (s, 1H), 8.27 - 8.25 (m, 1H), 7.93 - 7.91 (m, 3H), 7.57 - 7.55 (m, 2H), 3.93 (s, 2H), 3.39 - 3.32 (m, 2H), 2.40 (s, 3H).

[0451] Synthesis of 4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)-N-(2,2,2-trifluoroethyl)quinazoline-6-carboxamide (compound 1139) 4-(3-chloro-4-fluorophenylamino)-2-(pyridin-3-yl)-N-(2,2,2-trifluoroethyl)quinazoline-6-carboxamide was synthesized in a similar manner to that described for 4-(3-chloro-4-fluorophenylamino)-N-methyl-2-(pyridin-3-yl)-N-(2,2,2-trifluoroethyl)quinazoline-6-carboxamide substituting 2,2,2-trifluoroethylamine for 2,2,2-trifluoro-N-methylethylamine. The resulting product was analyzed. LCMS: r.t = 1.934 min, [M+1]⁺ = 476.0. ¹H-NMR (400 MHz, DMSO-*d*₆): δ 10.43 (s, 1H), 9.54 (s, 1H), 9.32 (t, *J* = 6.1 Hz, 1H), 9.14 (s, 1H), 8.72 (d, *J* = 4.6 Hz, 1H), 8.68 (d, *J* = 8.0 Hz, 1H), 8.32 (d, *J* = 8.7 Hz, 1H), 8.27 (dd, *J* = 6.7, 2.2 Hz, 1H), 7.98 (d, *J* = 8.7 Hz, 1H), 7.96 - 7.90 (m, 1H), 7.62 - 7.51 (m, 2H), 4.27 - 4.14 (m, 2H).

Scheme 65: General route for the synthesis of compounds with general formula vi

[0452]



Method E for cyclization

[0453]

E1: Sodium methoxide/Toluene

E2: NaOH/EtOH

Method G for Coupling Conditions

[0454]

G1: *i*-PrOH/85-100 °C

G2: THF/reflux

G3: *i*-AmOH-1/100-130 °C

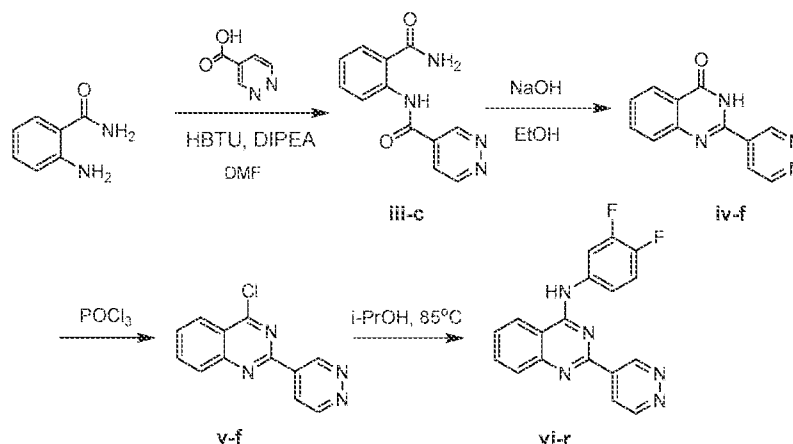
G4: MeOH/microwave/150 °C

G5: *i*-AmOH/microwave/150 °C

G6: THF/Et₃N/reflux

Scheme 66: Representative synthesis of compounds of formula vi-a with a pyridazine (see Scheme 65)

[0455]



[0456] Method D: N-(2-carbamoylphenyl) pyridazine-4-carboxamide (iii-c) A mixture of pyridazine-4-carboxylic acid (500 mg, 4.0 mmol, 1.0 eq.), 2-aminobenzamide (603 mg, 4.4 mmol, 1.1 eq.) and HBTU (3.0 g, 8.0 mmol, 2.0 eq.) was suspended in 15 mL of DMF. DIPEA (2.0 mL, 1.56 g, 12.0 mmol, 3.0 eq.) was added dropwise at room temperature and was stirred over night. After quenching with water, the resulting precipitate was collected and washed with a small amount of DCM. A white solid (388 mg) was obtained, LCMS m/z = 243.1 (M+1) (Method B) (retention time = 0.99 min), was used in the next step without further purification.

[0457] Method E1: 2-(pyridazin-4-yl) quinazolin-4(3H)-one (iv-f) A 100 mL round-bottom flask equipped with a Dean-Stark trap was charged with a mixture of N-(2-carbamoylphenyl) pyridazine-4-carboxamide (300 mg, 1.0 eq.), sodium methoxide (401 mg, 7.4 mmol, 6.0 eq.) and 10 mL of anhydrous toluene. The reaction mixture was heated to 110 °C and refluxed overnight. After cooling, the volatiles were removed *in vacuo* and the residue was quenched with a saturated aqueous solution of NH_4Cl (10 mL). The pH of the mixture was adjusted to 3 with 10 % HCl in water. The solution was extracted with DCM (50 mL x 3). The combined organic layers were washed with brine, dried over Na_2SO_4 . After filtration and evaporation, 88 mg of a yellow solid was obtained, LCMS m/z = 225.1 (M+1) (Method A) (retention time = 1.10 min) which was used in the next step without further purification.

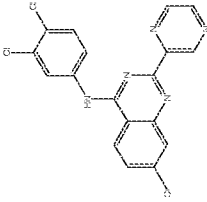
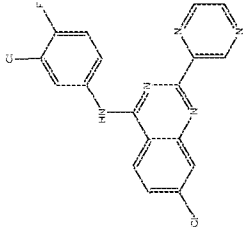
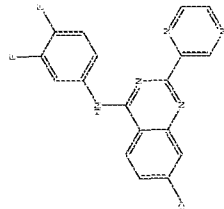
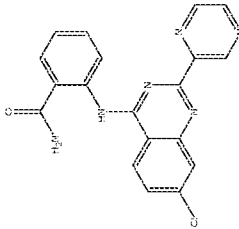
[0458] Method F5: 4-chloro-2-(pyridazin-4-yl)quinazoline (v-f) A 100-mL round-bottom flask was charged with 2-(pyridazin-4-yl) quinazolin-4(3H)-one (30 mg) which was suspended in 3 mL of POCl_3 . The reaction mixture was heated to reflux for 1 h. The reaction mixture turned to a clear brown solution. After cooling, 50 mL of ice/water was carefully added. An aqueous ammonia solution (25% by weight in water) was added dropwise to the mixture with stirring until the pH of the mixture was adjusted to 7~8. An internal temperature of 0 °C was maintained by the addition of ice. The mixture was warmed to room temperature and extracted with DCM (50 mL x 3). The combined organic layers were washed with brine, dried over Na_2SO_4 . After filtration and evaporation, 32 mg was obtained as a light brown solid. LCMS m/z = 242.9 (M+1) (Method B) (retention time = 1.65 min). The solid was used directly in the next step without further purification.

Method G1: N-(3,4-difluorophenyl)-2-(pyridazin-4-yl)quinazolin-4-amine (vi-r)

[0459] (This method is representative of method G1, G2, and G3. These three methods can be implemented in a similar way except for substitution of the appropriate solvent and temperature) The mixture of 4-chloro-2-(pyridazin-4-yl)quinazoline (10 mg, 0.041 mmol, 1 eq.) and 3,4-difluorobenzenamine (11 mg, 0.082 mmol, 2eq.) was suspended in *i*-PrOH. The mixture was heated at 85°C overnight. After cooling, the resulting precipitate was filtered and purified on HPLC (Condition C). 7.3 mg of N-(3,4-difluorophenyl)-2-(pyridazin-4-yl)quinazolin-4-amine was obtained (yield 53%). LCMS m/z = 336.0 (M+1) (Method B) (retention time = 1.731 min). $^1\text{H-NMR}$ (400 MHz, DMSO-d_6): δ 10.23 (s, 1H), 10.00 (dd, J = 2.2, 1.3 Hz, 1H), 9.44 (dd, J = 5.4, 1.2 Hz, 1H), 8.59 (d, J = 8.1 Hz, 1H), 8.39 (dd, J = 5.4, 2.4 Hz, 1H), 8.05 - 8.13 (m, 1H), 7.99 (s, 1H), 7.97 (d, J = 3.6 Hz, 1H), 7.72 - 7.79 (m, 2H), 7.51 - 7.61 (m, 1H).

[0460] The compounds in the following table were prepared in a manner analogous to that described in Scheme 1 and 64 (prepared according to method procedure A - G as designated).

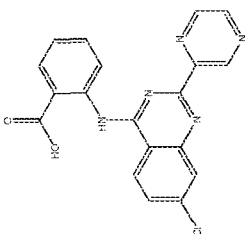
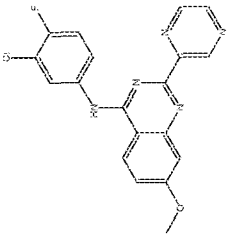
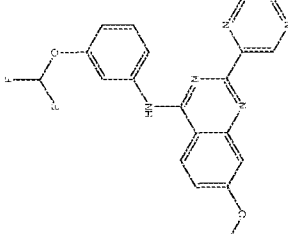
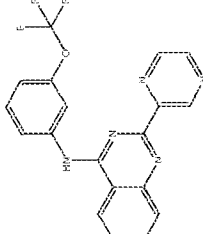
Table 21:

Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1140			402.66	¹ H-NMR (400 MHz, CD ₃ OD): δ 9.40 (s, 1H), 9.84 - 9.86 (m, 2H), 8.52 (d, J = 8.4 Hz, 1H), 8.15 (s, 1H), 8.02 (s, 1H), 7.84 (dd, J = 9.2, 2.4 Hz, 1H), 7.66 - 7.67 (m, 2H).	CD ₃ OD	402.0, 404.0, 405.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1141			386.21	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.78 (s, 1H), 9.49 (s, 1H), 8.89 (d, J = 10.8 Hz, 2H), 8.75 (d, J = 8.8 Hz, 1H), 8.44 (dd, J = 6.8, 2.4 Hz, 1H), 8.12 (d, J = 2.0 Hz, 1H), 7.94 - 7.99 (m, 1H), 7.87 (dd, J = 8.8, 2.0 Hz, 1H), 7.56 (t, J = 9.2 Hz, 1H).	DMSO	386.0, 388.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1142			369.76	¹ H-NMR (400 MHz, CD ₃ OD): δ 9.40 (s, 1H), 8.76 (d, J = 10.8 Hz, 2H), 8.40 (d, J = 8.8 Hz, 1H), 8.02 (d, J = 2.0 Hz, 1H), 7.80 - 7.86 (m, 1H), 7.68 (dd, J = 8.8, 1.6 Hz, 1H), 7.52 (d, J = 9.2 Hz, 1H), 7.33 (q, J = 9.2 Hz, 1H).	CD ₃ OD	370.0, 372.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1143			376.8	¹ H-NMR (400 MHz, CD ₃ OD): δ 9.55 (s, 1H), 8.80 (d, J = 7.2 Hz, 3H), 8.26 (d, J = 8.8 Hz, 1H), 8.04 (s, 1H), 7.85 (d, J = 6.8 Hz, 1H), 7.72 (d, J = 8.4 Hz, 1H), 7.66 (t, J = 7.2 Hz, 1H), 7.28 (t, J = 7.6 Hz, 1H).	CD ₃ OD	377.1, 379.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

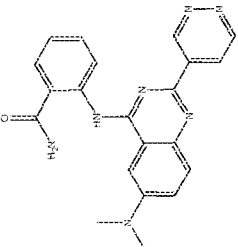
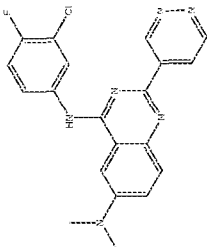
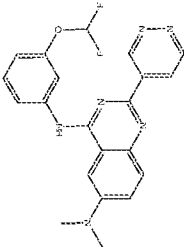
(continued)

Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1144			358.78	¹ H-NMR (400 MHz, CD ₃ OD): δ 9.39 (d, J = 1.6 Hz, 1H), 8.79 (d, J = 6.8 Hz, 2H), 8.47 (d, J = 9.2 Hz, 1H), 8.20 (s, 1H), 8.05 - 8.10 (m, 2H), 7.77 (dd, J = 8.8, 2.0 Hz, 1H), 7.63-7.65 (m, 2H).	CD ₃ OD	359.0, 361.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1145			399.78	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.63 (s, 1H), 9.52 (d, J = 1.2 Hz, 1H), 8.86 (t, J = 6.0 Hz, 2H), 8.77 (d, J = 8.4 Hz, 1H), 8.12 (d, J = 15.6 Hz, 2H), 7.85 (d, J = 8.8 Hz, 2H), 7.53 (t, J = 8.0 Hz, 1H), 7.32 (t, J = 74.0 Hz, 1H), 7.04 (dd, J = 8.0, 1.6 Hz, 1H).	DMSO	400.0, 402.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1146			402.66	¹ H-NMR (400 MHz, CD ₃ OD): δ 9.40 (s, 1H), 8.85 - 8.88 (m, 2H), 8.53 (d, J = 9.2 Hz, 1H), 8.17 (d, J = 2.0 Hz, 1H), 7.80-7.88 (m, 3H), 7.46 (s, 1H).	CD ₃ OD	401.9, 403.9, 405.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

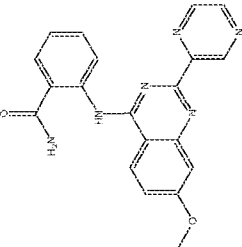
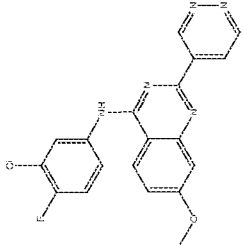
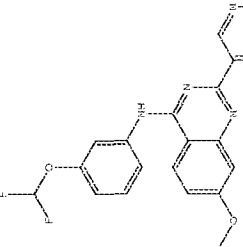
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1147			377.78	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.28 (s, 1H), 9.58 (s, 1H), 9.14 (d, J = 8.4 Hz, 1H), 8.87 (s, 1H), 8.80 (d, J = 2.0 Hz, 1H), 8.28 (d, J = 9.2 Hz, 1H), 8.07 - 8.12 (m, 2H), 7.86 (dd, J = 8.8, 2.0 Hz, 1H), 7.77 (t, J = 8.8 Hz, 1H), 7.26 (t, J = 7.6 Hz, 1H).	DMSO	378.1, 380.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1148			381.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.69 - 11.49 (m, 1H), 9.40 (d, J = 1.2 Hz, 1H), 9.00 - 8.86 (m, 3H), 8.24 (dd, J = 6.6, 2.1 Hz, 1H), 7.91 (ddd, J = 8.9, 4.2, 2.6 Hz, 1H), 7.71 (d, J = 1.4 Hz, 1H), 7.61 (t, J = 9.0 Hz, 1H), 7.49 (dd, J = 9.2, 2.3 Hz, 1H), 4.00 (s, 3H).	DMSO	382.1, 384.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1149			395.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.96 (s, 1H), 9.57 (d, J = 1.4 Hz, 1H), 8.83 - 8.74 (m, 2H), 8.57 (d, J = 9.1 Hz, 1H), 8.28 (t, J = 2.0 Hz, 1H), 7.86 (dd, J = 8.2, 1.2 Hz, 1H), 7.50 - 7.32 (m, 3H), 7.31 (t, J = 74.0 Hz, 1H), 6.94 (dd, J = 8.0, 2.2 Hz, 1H), 3.98 (s, 3H).	DMSO	396.1, 397.2 (M+2)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1150			383.33	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.27 (s, 1H), 9.57 (s, 1H), 8.80 (dd, J = 19.8, 1.7 Hz, 2H), 8.72 - 8.66 (m, 1H), 8.50 (s, 1H), 8.07 - 7.94 (m, 3H), 7.75 (t, J = 7.2 Hz, 1H), 7.57 (t, J = 8.2 Hz, 1H), 7.14 (d, J = 8.8 Hz, 1H).	DMSO	384.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

(continued)

Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1151			385.42	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.01 (s, 1H), 10.04 (s, 1H), 9.40 (d, J = 5.4 Hz, 1H), 9.20 (d, J = 8.4 Hz, 1H), 8.48 (s, 1H), 8.42 (dd, J = 5.3, 1.9 Hz, 1H), 8.04 - 7.94 (m, 2H), 7.84 (d, J = 9.2 Hz, 1H), 7.75 (t, J = 7.9 Hz, 1H), 7.58 (d, J = 9.2 Hz, 1H), 7.20 (t, J = 7.5 Hz, 1H), 7.13 (s, 1H), 3.15 (s, 6H).	DMSO	386.0 (M+1)	Method A (TFA)	95	Method D, G1
1152		HCl	394.83	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.96 (s, 1H), 9.97 (d, J = 1.1 Hz, 1H), 9.55 (d, J = 5.5 Hz, 1H), 8.61 (dd, J = 5.5, 2.2 Hz, 1H), 8.15 (dd, J = 6.8, 2.6 Hz, 1H), 8.08 (d, J = 9.3 Hz, 1H), 7.94 (ddd, J = 8.9, 4.3, 2.6 Hz, 1H), 7.74 (s, 1H), 7.62 (dd, J = 9.4, 2.5 Hz, 1H), 7.56 (t, J = 9.1 Hz, 1H), 3.16 (s, 6H).	DMSO	395.1, 397.1 (M+1)	Method A (TFA)	95	Method D, G1
1153		HCl	408.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.71 (brs, 1H), 10.01 - 9.99 (m, 1H), 9.52 (dd, J = 5.5, 1.0 Hz, 1H), 8.60 (dd, J = 5.5, 2.3 Hz, 1H), 8.07 (d, J = 9.2 Hz, 1H), 7.85 - 7.78 (m, 2H), 7.71 (s, 1H), 7.65 (dd, J = 9.3, 2.5 Hz, 1H), 7.56 (t, J = 8.2 Hz, 1H), 7.33 (t, J = 74.0 Hz, 1H), 7.09 (dd, J = 8.1, 2.1 Hz, 1H), 3.17 (s, 6H).	DMSO	409.1 (M+1)	Method A (TFA)	95	Method D, G1

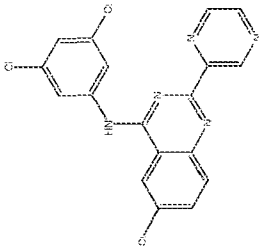
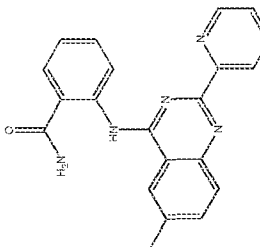
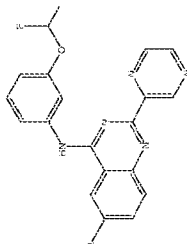
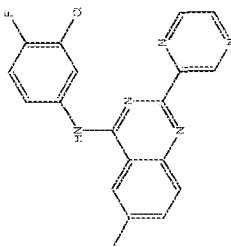
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1154			372.38	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.13 (s, 1H), 9.63 (d, J = 1.2 Hz, 1H), 9.34 (d, J = 7.6 Hz, 1H), 8.91 - 8.84 (m, 1H), 8.79 (d, J = 2.4 Hz, 1H), 8.49 (s, 1H), 8.12 (d, J = 9.6 Hz, 1H), 8.03 - 7.85 (m, 2H), 7.75 - 7.61 (m, 1H), 7.48 - 7.30 (m, 2H), 7.28 - 7.06 (m, 1H), 3.99 (s, 3H).	DMSO	373.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1155			381.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.16 (s, 1H), 9.99 - 9.95 (m, 1H), 9.42 (dd, J = 5.2, 1.2 Hz, 1H), 8.58 - 8.51 (m, 1H), 8.35 (dd, J = 5.2, 2.4 Hz, 1H), 8.21 (dd, J = 6.8, 2.4 Hz, 1H), 7.93 (ddd, J = 8.8, 4.2, 2.7 Hz, 1H), 7.53 (dd, J = 11.7, 6.4 Hz, 1H), 7.32 (dd, J = 6.6, 2.6 Hz, 2H), 3.97 (s, 3H).	DMSO	382.0, 384.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1156			395.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.07 (s, 1H), 10.01 (d, J = 1.3 Hz, 1H), 9.43 (dd, J = 5.2, 1.2 Hz, 1H), 8.55 (d, J = 10.0 Hz, 1H), 8.40 (dd, J = 5.2, 2.0 Hz, 1H), 7.91 (d, J = 2.0 Hz, 1H), 7.86 - 7.77 (m, 1H), 7.55 - 7.35 (m, 3H), 7.32 (t, J = 74.0 Hz, 1H), 7.01 (dd, J = 8.0, 2.0 Hz, 1H), 3.99 (s, 3H).	DMSO	396.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

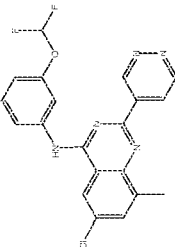
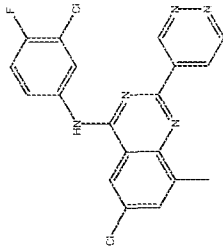
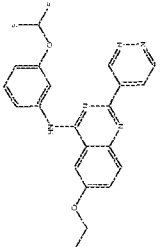
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1157			376.80	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.34 (s, 1H), 9.61 (s, 1H), 9.28 (d, J = 8.4 Hz, 1H), 8.87 (s, 1H), 8.80 (s, 1H), 8.53 (s, 1H), 8.25 (s, 1H), 7.94 - 8.06 (m, 4H), 7.70 (t, J = 8.0 Hz, 1H), 7.23 (t, J = 7.6 Hz, 1H).	DMSO	377.0, 379.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1158			358.78	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.35 (s, 1H), 9.54 (s, 1H), 8.77 - 8.84 (m, 4H), 8.32 (d, J = 8.0 Hz, 1H), 8.00 - 8.03 (m, 2H), 7.62 - 7.70 (m, 2H).	DMSO	359.0, 361.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1159			386.21	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.55 (s, 1H), 9.50 (s, 1H), 8.83 - 8.87 (m, 3H), 8.47 - 8.49 (m, 1H), 7.96 - 8.06 (m, 3H), 7.54 (t, J = 9.2 Hz, 1H).	DMSO	386.0, 388.0, 390.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1160			402.66	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.54 (s, 1H), 9.53 (s, 1H), 8.84 - 8.88 (m, 3H), 8.62 (d, J = 2.0 Hz, 1H), 8.01 - 8.07 (m, 3H), 7.73 (d, J = 8.8 Hz, 1H).	DMSO	401.9, 403.9, 406.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

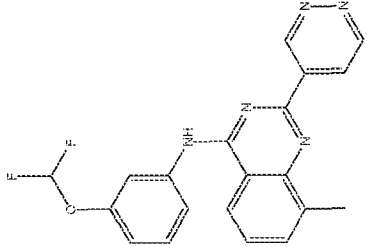
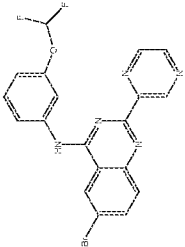
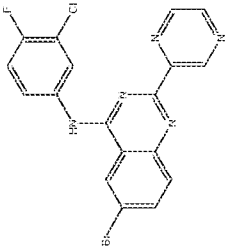
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1161			402.66	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.42 (s, 1H), 9.55 (d, J = 1.6 Hz, 1H), 8.87 - 8.88 (m, 1H), 8.82 (d, J = 2.4 Hz, 1H), 8.36 (d, J = 2.0 Hz, 2H), 8.00 - 8.07 (m, 2H), 7.40 (t, J = 1.6 Hz, 1H).	DMSO	401.9, 403.8, 405.8 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1162			468.25	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.36 (s, 1H), 9.59 (d, J = 1.2 Hz, 1H), 9.15 - 9.12 (m, 1H), 8.90 - 8.85 (m, 2H), 8.68 (d, J = 1.2 Hz, 1H), 8.52 (s, 1H), 8.28 - 8.25 (m, 1H), 8.07 (s, 1H), 7.99 - 7.96 (m, 1H), 7.83 (d, J = 8.8 Hz, 1H), 7.74 - 7.70 (m, 1H), 7.28 (t, J = 7.6 Hz, 1H).	DMSO	469.0 (M+1)	Method B (NH ₄ HCO ₃)	90	Method C, G1
1163			491.23	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.83 (s, 1H), 9.32 (s, 1H), 9.07 (s, 1H), 8.74 (s, 2H), 8.16 (dd, J = 8.0, 1.6 Hz, 1H), 7.87 (s, 1H), 7.74 - 7.64 (m, 2H), 7.39 (t, J = 8.0 Hz, 1H), 7.16 (t, J = 74.0 Hz, 1H), 6.95 - 6.92 (m, 1H).	DMSO	492.0 (M+1)	Method B (NH ₄ HCO ₃)	90	Method C, G1
1164			477.66	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.48 (s, 1H), 9.42 (s, 1H), 9.34 (s, 1H), 8.95 - 8.93 (m, 2H), 8.35 - 8.30 (m, 2H), 7.97 - 7.90 (m, 2H), 7.60 (t, J = 9.0 Hz, 1H).	DMSO	478.0, 480.0 (M+1)	Method B (NH ₄ HCO ₃)	90	Method C, G1

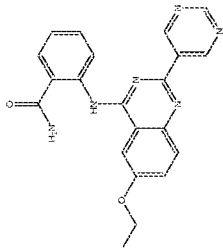
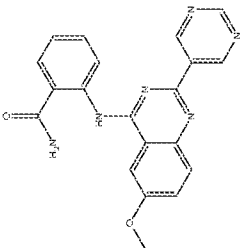
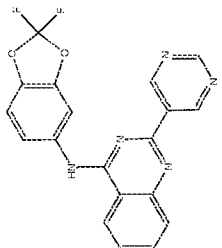
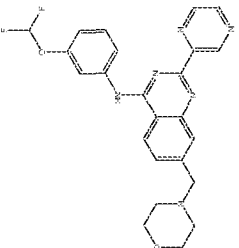
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1165			413.81	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.07 (s, 1H), 9.99 (s, 1H), 9.40 (d, J = 5.6 Hz, 1H), 8.56 (d, J = 1.6 Hz, 1H), 8.36 (dd, J = 5.2, 2.0 Hz, 1H), 7.87 (m, 1H), 7.81 (m, 1H), 7.53 (t, J = 7.8 Hz, 1H), 7.30 (t, J = 7.4 Hz, 1H), 8.36 (dd, J = 8.4, 2.0 Hz, 1H), 2.70 (s, 3H).	DMSO	414.1, 416.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1166			400.24	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.16 (s, 1H), 10.01 (s, 1H), 9.46 (d, J = 5.2 Hz, 1H), 8.56 (d, J = 1.6 Hz, 1H), 8.39 (dd, J = 5.4, 2.2 Hz, 1H), 8.20 (dd, J = 7.2, 2.6 Hz, 1H), 7.95 - 7.91 (m, 1H), 7.89 (s, 1H), 7.57 (t, J = 9.0 Hz, 1H), 2.75 (s, 3H).	DMSO	400.0, 402.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1167			409.39	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.05 - 10.3 (m, 2H), 9.45 (dd, J = 5.2, 1.2 Hz, 1H), 8.42 (dd, J = 5.2, 1.2 Hz, 1H), 8.06 (d, J = 2.8 Hz, 1H), 7.97 - 7.94 (m, 2H), 7.86 (dd, J = 8.0, 1.2 Hz, 1H), 7.66 - 7.60 (m, 2H), 7.38 (t, J = 7.4 Hz, 1H), 7.10 - 7.08 (m, 1H), 4.32 (q, J = 6.8 Hz, 2H), 1.51 (t, J = 7.0 Hz, 3H).	DMSO	410.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

(continued)

Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1168			379.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.13 (s, 1H), 10.07 (d, J = 2.0 Hz, 1H), 9.44 (dd, J = 5.2, 2.4 Hz, 1H), 8.48 - 8.44 (m, 2H), 7.93 (t, J = 2.0 Hz, 1H), 7.86 - 7.84 (m, 2H), 7.66 - 7.62 (m, 1H), 7.54 (t, J = 8.4 Hz, 1H), 7.32 (t, J = 74.0 Hz, 1H), 7.02 (dd, J = 8.0, 2.4 Hz, 1H), 2.77 (s, 3H).	DMSO	380.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1169			444.23	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.17 (s, 1H), 9.55 (s, 1H), 8.98 (s, 1H), 8.79 (d, J = 12.8 Hz, 2H), 8.26 (s, 1H), 8.08 (d, J = 9.2 Hz, 1H), 7.92 - 7.88 (m, 2H), 7.52 - 7.50 (m, 1H), 7.32 (t, J = 71.6 Hz, 1H), 6.99 (d, J = 7.6 Hz, 1H).	DMSO	444.0, 446.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1170			430.66	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.11 (s, 1H), 9.52 (s, 1H), 8.89 (d, J = 1.2 Hz, 1H), 8.83 (s, 1H), 8.77 (d, J = 2.4 Hz, 1H), 8.56 (dd, J = 6.8, 2.4 Hz, 1H), 8.05 (dd, J = 9.2, 2.0 Hz, 1H), 8.02 - 7.96 (m, 1H), 7.88 (d, J = 8.8 Hz, 1H), 7.49 (t, J = 9.2 Hz, 1H).	DMSO	430.0, 432.0, 434.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

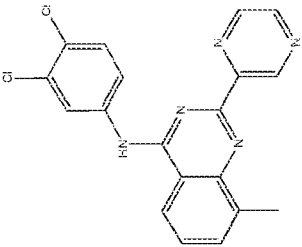
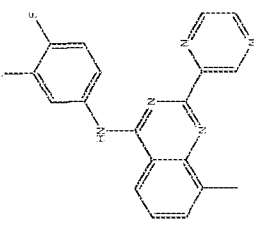
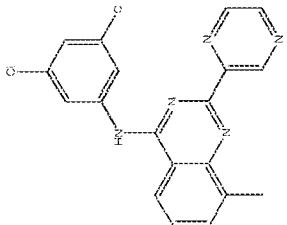
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1171			386.41	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.07 (s, 1H), 9.63 (s, 2H), 9.33 (s, 1H), 8.95 (d, J = 8.0 Hz, 1H), 8.50 (s, 1H), 7.99 - 7.94 (m, 3H), 7.76 - 7.71 (m, 1H), 7.62 - 7.59 (m, 2H), 7.24 (t, J = 7.4 Hz, 1H), 4.23 (q, J = 7.0 Hz, 2H), 1.46 (t, J = 7.0 Hz, 3H).	DMSO	387.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1172			372.38	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.11 (s, 1H), 9.61 (s, 2H), 9.34 (s, 1H), 8.91 (d, J = 8.0 Hz, 1H), 8.49 (s, 1H), 7.98 - 7.95 (m, 3H), 7.72 (t, J = 8.0 Hz, 1H), 7.62 - 7.60 (m, 2H), 7.24 (t, J = 7.6 Hz, 1H), 3.97 (s, 3H).	DMSO	373.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1173		HCl	379.32	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.55 (brs, 1H), 9.59 (s, 2H), 9.35 (s, 1H), 8.68 (d, J = 8.4 Hz, 1H), 8.09 - 8.07 (m, 1H), 8.00 - 7.96 (m, 2H), 7.76 - 7.72 (m, 1H), 7.67 (dd, J = 8.8, 2.0 Hz, 1H), 7.53 (d, J = 8.8 Hz, 1H).	DMSO	380.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1174		HCl	464.47	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.14 (s, 1H), 9.59 (s, 1H), 8.80 (m, 2H), 8.62 (d, J = 7.6 Hz, 1H), 8.30 (s, 1H), 7.90 (s, 2H), 7.70 (d, J = 7.6 Hz, 1H), 7.53 - 7.45 (m, 1H), 7.39 - 6.86 (m, 1H), 7.30 (t, J = 74.1 Hz, 1H), 3.73 (s, 2H), 3.63 - 3.58 (m, 4H), 2.48 - 2.43 (m, 4H).	DMSO	465.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

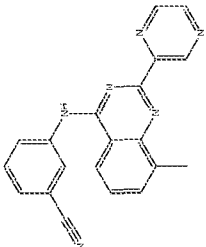
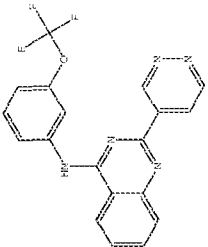
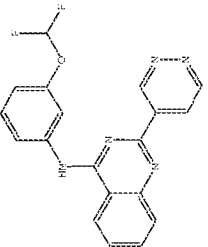
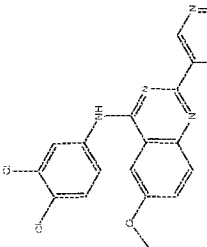
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1175			357.37	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.70 (s, 1H), 9.43 (d, J = 8.4 Hz, 1H), 8.90 (s, 1H), 8.78 (d, J = 2.0 Hz, 1H), 8.22 (d, J = 8.0 Hz, 1H), 8.11 (d, J = 8.0 Hz, 1H), 7.80 (d, J = 2.8 Hz, 1H), 7.60 - 7.65 (m, 2H), 7.09 - 7.13 (m, 3H), 2.76 (s, 3H).	DMSO	358.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1176			365.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.02 (s, 1H), 9.62 (s, 1H), 8.85 (s, 1H), 8.76 (d, J = 2.4 Hz, 1H), 8.45 (d, J = 8.0 Hz, 1H), 8.03 - 8.07 (m, 1H), 7.81 (d, J = 6.8 Hz, 1H), 7.61 (t, J = 7.6 Hz, 1H), 7.49 (t, J = 9.2 Hz, 1H), 2.75 (s, 3H).	DMSO	366.0, 368.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1177			379.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.02 (s, 1H), 9.64 (s, 1H), 8.83 (s, 1H), 8.76 (d, J = 2.0 Hz, 1H), 8.50 (d, J = 8.0 Hz, 1H), 8.38 (s, 1H), 7.92 (d, J = 8.4 Hz, 1H), 7.81 (d, J = 6.8 Hz, 1H), 7.61 (t, J = 7.8 Hz, 1H), 7.49 (t, J = 8.4 Hz, 1H), 7.32 (t, J = 74.0 Hz, 1H), 6.95 (d, J = 7.2 Hz, 1H), 2.75 (s, 3H).	DMSO	380.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

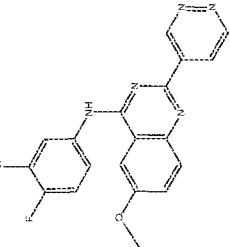
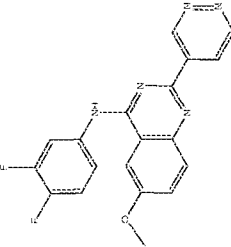
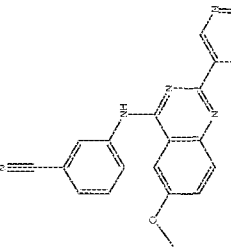
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1178			382.25	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.21 (s, 1H), 9.62 (s, 1H), 8.86 (t, J = 2.4 Hz, 1H), 8.78 (t, J = 3.2 Hz, 2H), 8.52 (d, J = 8.4 Hz, 1H), 8.10 - 8.13 (m, 1H), 7.83 (d, J = 7.2 Hz, 1H), 7.68 (d, J = 9.2 Hz, 1H), 7.63 (t, J = 7.6 Hz, 1H), 2.75 (s, 3H).	DMSO	381.9, 383.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1179			349.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.21 (s, 1H), 9.60 (s, 1H), 8.87 (s, 1H), 8.78 (d, J = 2.4 Hz, 1H), 8.50 - 8.60 (m, 2H), 7.80 - 7.85 (m, 2H), 7.61 (t, J = 7.8 Hz, 1H), 7.46 - 7.54 (m, 1H), 2.74 (s, 3H).	DMSO	350.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1180			382.25	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.20 (s, 1H), 9.62 (s, 1H), 8.85 (s, 1H), 8.77 (d, J = 2.8 Hz, 1H), 8.47 - 8.52 (m, 3H), 7.81 (d, J = 3.2 Hz, 1H), 7.60 (t, J = 7.8 Hz, 1H), 7.31 (t, J = 1.6 Hz, 1H), 2.51 (s, 3H).	DMSO	381.9, 383.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

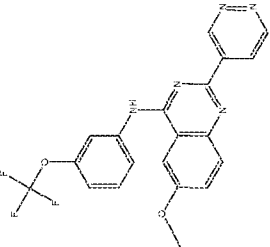
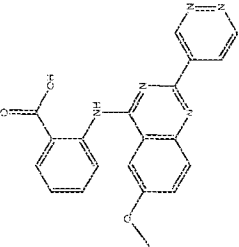
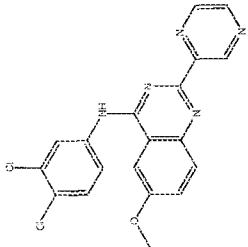
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1181			338.37	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.21 (s, 1H), 9.64 (s, 1H), 8.86 (d, J = 9.2 Hz, 2H), 8.78 (d, J = 8.0 Hz, 1H), 8.51 (d, J = 8.0 Hz, 1H), 8.38 (d, J = 8.4 Hz, 1H), 7.84 (d, J = 7.2 Hz, 1H), 7.61 - 7.65 (m, 3H), 2.76 (s, 3H).	DMSO	339.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1182			383.33	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.24 (s, 1H), 9.99 (s, 1H), 9.41 (d, J = 5.2 Hz, 1H), 8.62 (d, J = 8.4 Hz, 1H), 8.36 - 8.38 (m, 1H), 8.10 (s, 1H), 7.95 - 7.98 (m, 3H), 7.73 - 7.74 (m, 1H), 7.62 (t, J = 8.0 Hz, 1H), 7.20 (d, J = 8.0 Hz, 1H).	DMSO	384.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1183			365.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.19 (s, 1H), 10.01 (s, 1H), 9.41 - 9.43 (m, 1H), 8.63 (d, J = 8.0 Hz, 1H), 8.39 - 8.41 (m, 1H), 7.93 - 7.97 (m, 3H), 7.84 (d, J = 8.4 Hz, 1H), 7.74 (m, 1H), 7.54 - 7.56 (m, 1H), 7.32 (t, J = 7.4 Hz, 1H), 7.02 - 7.04 (m, 1H).	DMSO	366.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1184			398.25	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.27 (s, 1H), 9.96 (s, 1H), 9.44 (d, J = 4.8 Hz, 1H), 8.40 - 8.41 (m, 1H), 8.29 (d, J = 2.0 Hz, 1H), 8.06 (d, J = 2.4 Hz, 1H), 7.99 - 8.02 (m, 1H), 7.89 (d, J = 8.8 Hz, 1H), 7.71 (d, J = 8.8 Hz, 1H), 7.56 - 7.58 (m, 1H), 3.99 (s, 3H).	DMSO	398.0, 400.0, 402.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

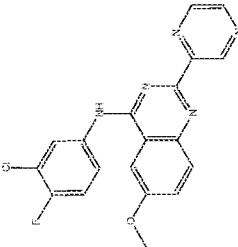
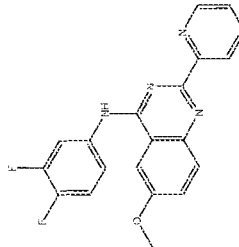
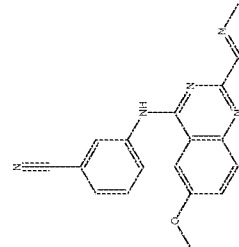
(continued)

Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1185			381.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.92 (s, 1H), 9.36 - 9.38 (m, 1H), 8.27 - 8.29 (m, 1H), 8.14 - 8.16 (m, 1H), 7.51 (s, 1H), 7.83 - 7.91 (m, 3H), 7.51 - 7.56 (m, 2H), 3.97 (s, 3H).	DMSO	382.1, 384.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1188			365.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.95 - 10.00 (m, 2H), 9.40 (d, J = 5.6 Hz, 1H), 8.32 - 8.33 (m, 1H), 8.06 (m, 1H), 7.96 (s, 1H), 7.89 (d, J = 9.2 Hz, 1H), 7.69 (m, 1H), 7.54 - 7.59 (m, 2H), 3.98 (s, 3H).	DMSO	366.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1187			354.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.00 (s, 1H), 9.90 (m, 1H), 9.35 - 9.36 (m, 1H), 8.23 - 8.30 (m, 3H), 7.84 - 7.90 (m, 2H), 7.62 - 7.71 (m, 2H), 7.54 - 7.57 (m, 1H), 3.96 (s, 3H).	DMSO	355.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

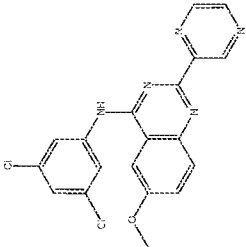
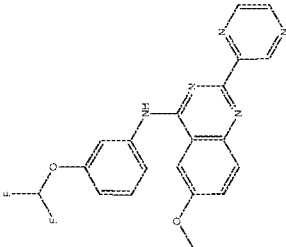
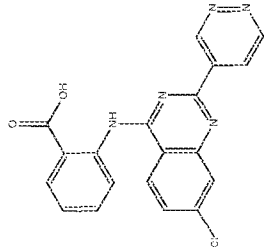
(continued)

Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1188			413.35	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.96 - 10.0 (m, 2H), 8.39 - 8.40 (m, 1H), 8.31 - 8.33 (m, 1H), 8.07 (s, 1H), 7.88 - 7.96 (m, 3H), 7.57 - 7.67 (m, 2H), 7.22 (d, J = 8.0 Hz, 1H), 4.01 (s, 3H).	DMSO	414.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1189			373.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.15 (s, 1H), 10.02 (s, 1H), 9.41 (d, J = 5.2 Hz, 1H), 8.94 (d, J = 7.6 Hz, 1H), 8.41 (m, 1H), 8.09 (d, J = 7.6 Hz, 1H), 7.93 (d, J = 8.8 Hz, 1H), 7.80 (t, J = 8.4 Hz, 1H), 7.60 - 7.62 (m, 2H), 7.25 (t, J = 7.6 Hz, 1H), 4.01 (s, 3H).	DMSO	374.1, 376.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1190			398.25	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.95 (s, 1H), 9.54 (s, 1H), 8.81 - 8.82 (m, 1H), 8.74 (d, J = 2.0 Hz, 1H), 8.66 (d, J = 2.4 Hz, 1H), 8.06 (dd, J = 8.8, 2.4 Hz, 1H), 8.00 (d, J = 2.4 Hz, 1H), 7.92 (d, J = 8.8 Hz, 1H), 7.70 (d, J = 8.8 Hz, 1H), 7.60 (dd, J = 8.8, 2.4 Hz, 1H), 3.40 (s, 3H).	DMSO	397.9, 400.0, 401.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6

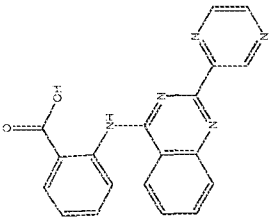
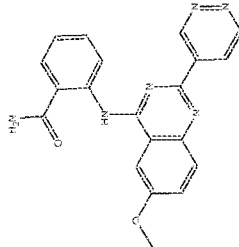
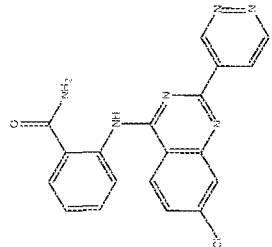
(continued)

Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1191			381.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.90 (s, 1H), 9.52 (s, 1H), 8.81 (m, 1H), 8.73 (d, J = 2.4 Hz, 1H), 8.53 (dd, J = 6.8, 2.4 Hz, 1H), 7.98 - 8.03 (m, 2H), 7.91 (d, J = 9.2 Hz, 1H), 7.59 (dd, J = 5.2, 2.4 Hz, 1H), 7.51 (t, J = 8.8 Hz, 1H), 3.40 (s, 3H).	DMSO	382.0, 384.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6
1192			365.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.89 (s, 1H), 9.52 - 9.53 (m, 1H), 8.81 - 8.82 (m, 1H), 8.73 (d, J = 2.4 Hz, 1H), 8.45 - 8.51 (m, 1H), 7.98 (m, 1H), 7.90 (d, J = 9.2 Hz, 1H), 7.75 - 7.78 (m, 1H), 7.47 - 7.59 (m, 2H), 3.40 (s, 3H).	DMSO	366.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6
1193			354.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.06 (s, 1H), 9.54 (s, 1H), 8.74 - 8.81 8.35 (d, J = 8.4 Hz, 1H), 8.04 (m, 1H), 7.92 - 7.95 (m, 1H), 7.59 - 7.69 (m, 3H), 3.40 (s, 3H).	DMSO	355.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6

(continued)

Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1194			398.25	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.91 (s, 1H), 9.55 (d, J = 1.2 Hz, 1H), 8.81 (t, J = 2.0 Hz, 1H), 8.74 (d, J = 2.4 Hz, 1H), 8.39 (d, J = 1.6 Hz, 1H), 7.97 (d, J = 2.8 Hz, 1H), 7.92 (d, J = 8.8 Hz, 1H), 7.59 (dd, J = 8.8, 2.4 Hz, 1H), 7.32 (t, J = 1.6 Hz, 1H), 3.40 (s, 3H).	DMSO	398.0, 399.9, 402.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6
1195			395.36	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.91 (s, 1H), 9.55 (m, 1H), 8.74 - 8.79 (m, 2H), 8.25 (s, 1H), 8.03 (d, J = 2.4 Hz, 1H), 7.92 (d, J = 9.2 Hz, 1H), 7.86 (d, J = 8.4 Hz, 1H), 7.60 (dd, J = 9.2, 2.4 Hz, 1H), 7.48 - 7.52 (m, 1H), 7.34 (t, J = 74.0 Hz, 1H), 6.96 - 6.98 (m, 1H), 3.40 (s, 3H).	DMSO	396.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6
1196			377.78	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.26 (s, 1H), 9.94 (s, 1H), 9.41 - 9.42 (m, 1H), 8.75 (d, J = 8.0 Hz, 1H), 8.33 - 8.34 (m, 1H), 8.19 (d, J = 8.4 Hz, 1H), 8.05 (d, J = 7.6 Hz, 1H), 7.93 (m, 1H), 7.75 - 7.77 (m, 2H), 7.05 - 7.30 (m, 2H).	DMSO	378.0, 380.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6

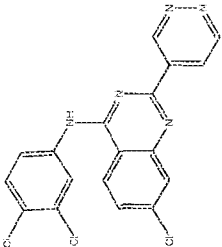
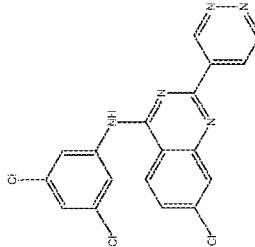
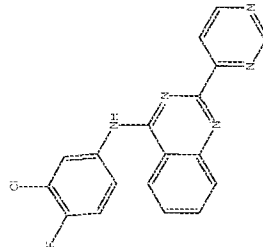
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1197			343.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.86 (s, 1H), 12.39 (s, 1H), 9.62 (d, J = 0.8 Hz, 1H), 9.29 (d, J = 8.0 Hz, 1H), 8.87 (s, 1H), 8.80 (d, J = 2.4 Hz, 1H), 8.27 (d, J = 8.0 Hz, 1H), 8.12 (dd, J = 8.0, 1.6 Hz, 1H), 7.97-8.05 (m, 3H), 7.75-7.79 (m, 2H), 7.24 (t, J = 7.6 Hz, 1H).	DMSO	344.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6
1198			372.38	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.05 (s, 1H), 9.99 (s, 1H), 9.39 (d, J = 5.2 Hz, 1H), 9.05 (d, J = 8.4 Hz, 1H), 8.47 (s, 1H), 8.36-8.38 (m, 1H), 7.85-7.98 (m, 3H), 7.71 (t, J = 7.6 Hz, 1H), 7.46-7.60 (m, 2H), 7.20 (t, J = 7.6 Hz, 1H), 3.95 (s, 3H).	DMSO	373.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1199			376.8	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.15 (s, 1H), 10.00 (s, 1H), 9.44 (d, J = 4.2 Hz, 1H), 8.95 (d, J = 8.4 Hz, 1H), 8.49 (s, 1H), 8.40-8.42 (m, 1H), 8.15 (d, J = 9.2 Hz, 1H), 7.91-7.97 (m, 3H), 7.80 (t, J = 7.6 Hz, 1H), 7.73 (t, J = 7.6 Hz, 1H), 7.25 (t, J = 7.2 Hz, 1H).	DMSO	377.0, 379.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

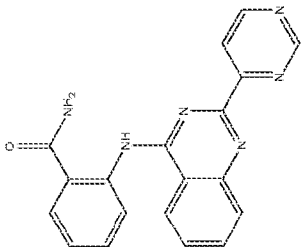
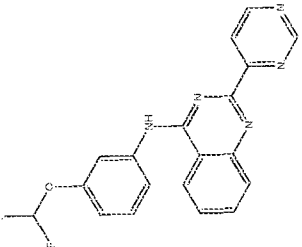
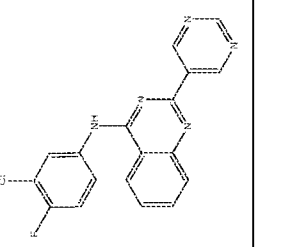
(continued)

Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1200			399.78	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.29 (s, 1H), 9.98 (s, 1H), 9.42 (d, J = 5.2 Hz, 1H), 8.66 (d, J = 9.2 Hz, 1H), 8.36 - 8.38 (m, 1H), 8.00 (d, J = 1.5 Hz, 1H), 7.89 (s, 1H), 7.79 - 7.81 (m, 2H), 7.49 - 7.57 (m, 1H), 7.31 (t, J = 74.0 Hz, 1H), 7.04 (dd, J = 8.0, 1.2 Hz, 1H).	DMSO	400.0, 401.9 (M+1)	Method A (TFA)	95	Method D, G1
1201			386.21	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.94 (s, 1H), 9.43 (d, J = 5.2 Hz, 1H), 8.72 (d, J = 8.8 Hz, 1H), 8.33 (dd, J = 5.6, 2.4 Hz, 1H), 8.21 (dd, J = 6.4, 1.6 Hz, 1H), 7.93 - 7.98 (m, 2H), 7.78 (dd, J = 8.8, 2.0 Hz, 1H), 7.54 (t, J = 9.2 Hz, 2H).	DMSO	386.0 387.8 (M+1)	Method A (TFA)	95	Method D, G1
1202			369.76	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.28 (s, 1H), 9.95 (m, 1H), 9.44 - 9.45 (m, 1H), 8.60 (d, J = 8.8 Hz, 1H), 8.32 - 8.34 (m, 1H), 8.02 - 8.08 (m, 1H), 7.98 (d, J = 6.4 Hz, 1H), 7.79 (dd, J = 9.2, 2.4 Hz, 1H), 7.70 - 7.72 (m, 1H), 7.56 (q, J = 9.2 Hz, 1H).	DMSO	369.9, 371.0 (M+1)	Method A (TFA)	95	Method D, G1
1203			358.78	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.36 (s, 1H), 9.92 (s, 1H), 9.42 (d, J = 4.8 Hz, 1H), 8.59 (d, J = 8.4 Hz, 1H), 8.23 - 8.33 (m, 3H), 7.96 (d, J = 1.6 Hz, 1H), 7.65 - 7.79 (m, 3H).	DMSO	359.0, 361.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

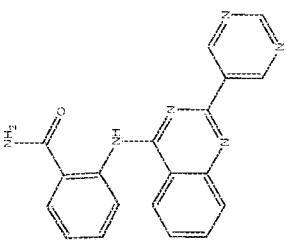
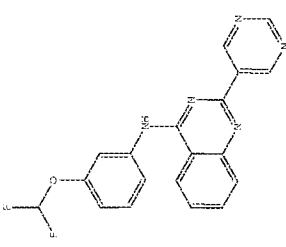
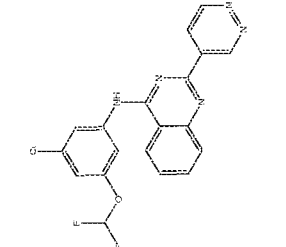
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1204			402.66	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.32 (s, 1H), 9.96 (s, 1H), 9.44 (d, J = 5.2 Hz, 1H), 8.60 (d, J = 8.4 Hz, 1H), 8.28-8.35 (m, 2H), 7.94-7.99 (m, 2H), 7.79 (d, J = 8.8 Hz, 1H), 7.73 (d, J = 8.8 Hz, 1H).	DMSO	401.9, 403.9, 405.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1205			402.66	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.90 (s, 1H), 9.42 (d, J = 4.8 Hz, 1H), 8.58 (d, J = 8.8 Hz, 1H), 8.27 (s, 1H), 8.07 (s, 2H), 7.92 (s, 1H), 7.73 (d, J = 8.0 Hz, 1H), 7.36 (s, 1H).	DMSO	401.9, 403.9, 405.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1206			351.76	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.59 (s, 1H), 9.43 (m, 1H), 9.07 (d, J = 5.2 Hz, 1H), 8.71 (d, J = 8.4 Hz, 1H), 8.50 (dd, J = 6.4, 2.0 Hz, 1H), 8.31 (dd, J = 5.2, 0.8 Hz, 1H), 8.00-8.09 (m, 3H), 7.80 (t, J = 7.2 Hz, 1H), 7.53 (t, J = 9.2 Hz, 1H).	DMSO	352.0, 354.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

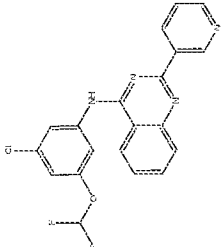
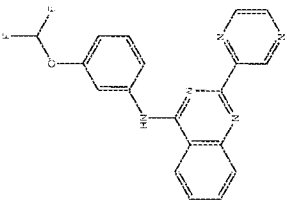
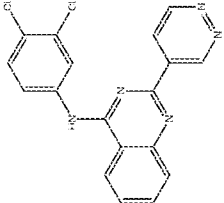
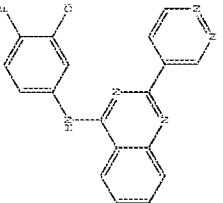
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1207			342.35	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.31 (s, 1H), 9.38 - 9.43 (m, 2H), 9.05 (d, J = 5.2 Hz, 1H), 8.52 (s, 1H), 8.47 (d, J = 5.2 Hz, 1H), 8.25 (d, J = 8.0 Hz, 1H), 7.95 - 8.04 (m, 4H), 7.82 (t, J = 7.2 Hz, 1H), 7.70 (t, J = 7.6 Hz, 1H), 7.22 (t, J = 7.6 Hz, 1H).	DMSO	343.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1208			365.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.13 (s, 1H), 9.37 (d, J = 1.2 Hz, 1H), 9.01 (d, J = 5.2 Hz, 1H), 8.68 (d, J = 8.4 Hz, 1H), 8.40 (dd, J = 5.2, 1.2 Hz, 1H), 8.28 (t, J = 1.2 Hz, 1H), 7.92 - 8.00 (m, 3H), 7.77 - 7.79 (m, 1H), 7.47 - 7.52 (m, 1H), 7.34 (t, J = 7.4 Hz, 1H), 6.97 (dd, J = 8.4, 2.4 Hz, 1 H).	DMSO	366.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1209		HCl (batch 02)	351.76	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.82 (s, 1H), 9.59 (s, 2H), 9.36 (s, 1H), 8.74 (d, J = 8.4 Hz, 1H), 8.22 (dd, J = 6.8, 2.4 Hz, 1H), 8.07 (d, J = 8.4 Hz, 1H), 8.00 (t, J = 7.6 Hz, 1H), 7.90 - 7.94 (m, 1H), 7.75 (t, J = 7.2 Hz, 1H), 7.56 (t, J = 9.2 Hz, 1 H).	DMSO	352.0, 354.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

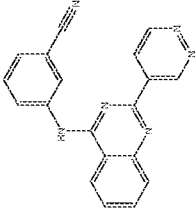
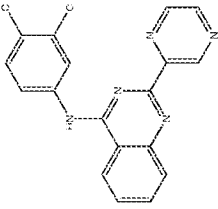
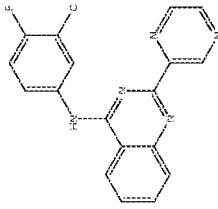
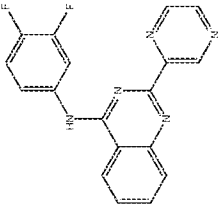
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1210			342.35	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.17 (s, 1H), 9.65 (s, 2H), 9.36 (s, 1H), 8.93 (d, J = 8.0 Hz, 1H), 8.51 (s, 1H), 8.23 (d, J = 8.4 Hz, 1H), 7.92 - 8.02 (m, 4H), 7.72 - 7.79 (m, 2H), 7.26 (t, J = 7.6 Hz, 1H).	DMSO	343.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G4
1211			365.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.15 (s, 1H), 9.62 (s, 2H), 9.32 (s, 1H), 8.63 (d, J = 8.0 Hz, 1H), 7.94 - 7.97 (m, 3H), 7.81 (dd, J = 8.4, 1.2 Hz, 1H), 7.69 - 7.74 (m, 1H), 7.50 - 7.55 (m, 1H), 7.31 (t, J = 74.0 Hz, 1H), 7.01 (dd, J = 8.4, 2.4 Hz, 1H).	DMSO	366.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6
1212			399.78	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.27 (s, 1H), 10.01 - 10.02 (m, 1H), 9.44 - 9.46 (m, 1H), 8.62 (d, J = 8.4 Hz, 1H), 8.40 (dd, J = 5.2, 2.0 Hz, 1H), 8.04 (t, J = 1.6 Hz, 1H), 8.00 (d, J = 3.2 Hz, 2H), 7.94 (m, 1H), 7.76 - 7.80 (m, 1H), 7.38 (t, J = 73.6 Hz, 1H), 7.16 (t, J = 2.0 Hz, 1H).	DMSO	400.0, 402.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6

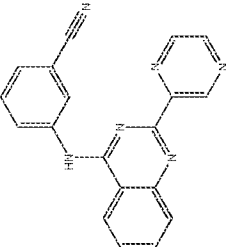
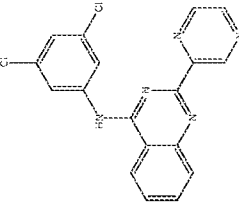
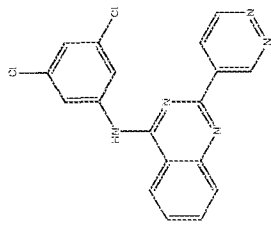
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1213			398.79	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.16 (s, 1H), 9.56 (m, 1H), 8.69 - 8.72 (m, 2H), 8.59 (d, J = 8.0 Hz, 1H), 8.08 (d, J = 1.6 Hz, 1H), 7.94 - 7.97 (m, 2H), 7.69-7.73 (m, 1H), 7.55-7.58 (m, 1H), 7.36 (t, J = 73.6 Hz, 1H), 7.13 (t, J = 2.0 Hz, 1H).	DMSO	399.0, 401.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6
1214			365.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.17 (s, 1H), 9.49 (s, 1H), 8.88 - 8.92 (m, 3H), 8.16 (d, J = 8.0 Hz, 1H), 8.07 (t, J = 7.6 Hz, 2H), 7.84 (t, J = 8.0 Hz, 2H), 7.57 (t, J = 8.0 Hz, 1H), 7.36 (t, J = 74.0 Hz, 1H), 7.10 - 7.12 (m, 1H).	DMSO	366.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6
1215			368.22	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.23 (s, 1H), 10.00 (s, 1H), 9.44 (d, J = 5.2 Hz, 1H), 8.58 (d, J = 8.0 Hz, 1H), 8.38 (dd, J = 2.4, 5.2 Hz, 1H), 8.32 (d, J = 2.0 Hz, 1H), 7.96 - 8.00 (m, 3H), 7.73 - 7.77 (m, 2H).	DMSO	367.8, 369.8, 371.8 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1216			351.76	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.21 (s, 1H), 9.99 (s, 1H), 9.43 (d, J = 5.2 Hz, 1H), 8.58 (d, J = 8.4 Hz, 1H), 8.38 (dd, J = 5.2, 2.0 Hz, 1H), 8.22 (dd, J = 7.0, 2.6 Hz, 1H), 7.89 - 7.93 (m, 3H), 7.68 - 7.72 (m, 1H), 7.55 (t, J = 8.6 Hz, 1H).	DMSO	351.9, 353.8 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

(continued)

Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1217			324.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.32 (s, 1H), 9.99 (dd, J = 2.0, 1.2 Hz, 1H), 9.44 (dd, J = 5.2, 0.8 Hz, 1H), 8.61 (d, J = 8.4 Hz, 1H), 8.38 - 8.40 (m, 2H), 8.28 - 8.31 (m, 1H), 7.99 (s, 1H), 7.97 (d, J = 3.6 Hz, 1H), 7.66 - 7.80 (m, 3H).	DMSO	325.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1
1218			368.22	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.94 (s, 1H), 9.44 (d, J = 0.8 Hz, 1H), 9.06 (d, J = 8.0 Hz, 1H), 9.00 (d, J = 2.4 Hz, 1H), 8.97 (t, J = 1.8 Hz, 1H), 8.38 (d, J = 2.4 Hz, 1H), 8.12 (t, J = 7.4 Hz, 1H), 8.00 (dd, J = 8.6, 2.2 Hz, 1H), 7.86 (t, J = 7.6 Hz, 1H), 7.82 (d, J = 8.8 Hz, 1H).	DMSO	367.9, 369.9, 371.8 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6
1219			351.76	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.62 (s, 1H), 9.40 (d, J = 0.8 Hz, 1H), 8.60 - 8.83 (m, 3H), 8.32 (dd, J = 6.6, 2.2 Hz, 1H), 8.20 (d, J = 8.8 Hz, 1H), 8.09 (t, J = 7.4 Hz, 1H), 7.95 - 8.00 (m, 1H), 7.85 (t, J = 7.8 Hz, 1H), 7.61 (t, J = 9.2 Hz, 1H).	DMSO	352.0, 354.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6
1220			335.31	¹ H-NMR (400 MHz, DMSO-d ₆): δ 11.05 (s, 1H), 9.48 (d, J = 0.8 Hz, 1H), 8.91 - 8.93 (m, 2H), 8.81 (d, J = 8.4 Hz, 1H), 8.28 - 8.33 (m, 1H), 8.14 (d, J = 8.4 Hz, 1H), 8.06 (t, J = 7.6 Hz, 1H), 7.77 - 7.85 (m, 2H), 7.56 - 7.63 (m, 1H).	DMSO	336.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6

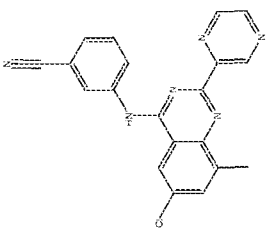
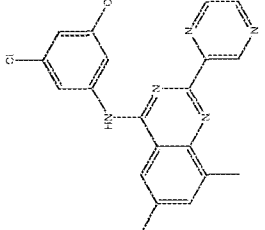
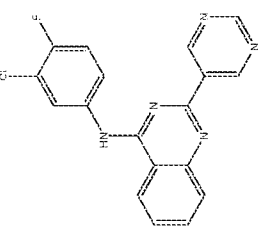
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1221			324.34	¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.09 (s, 1H), 9.40 (d, J = 1.6 Hz, 1H), 9.10 (d, J = 8.4 Hz, 1H), 9.00 (d, J = 2.0 Hz, 1H), 8.97 (t, J = 2.0 Hz, 1H), 8.49 (s, 1H), 8.26 - 8.30 (m, 2H), 8.12 - 8.06 (m, 1H), 7.87 - 7.91 (m, 1H), 7.76 - 7.84 (m, 2H)	DMSO	324.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6
1222			368.22	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.21 (s, 1H), 9.59 (d, J = 1.2 Hz, 1H), 8.85 (t, J = 1.8 Hz, 1H), 8.78 (d, J = 2.0 Hz, 1H), 8.64 (d, J = 8.4 Hz, 1H), 8.42 (d, J = 1.6 Hz, 2H), 7.96 - 8.02 (m, 2H), 7.74 - 7.78 (m, 1H), 7.35 (d, J = 1.6 Hz, 1H).	DMSO	368.0, 370.0, 372.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G6
1223			368.22	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.26 (s, 1H), 10.00 (s, 1H), 9.45 (d, J = 5.2 Hz, 1H), 8.56 (d, J = 8.4 Hz, 1H), 8.38 (dd, J = 5.0, 1.8 Hz, 1H), 8.09 (s, 2H), 7.94 (s, 1H), 7.92 (d, J = 9.2 Hz, 1H), 7.70 - 7.74 (m, 1H), 7.35 (s, 1H).	DMSO	368.1, 370.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method D, G1

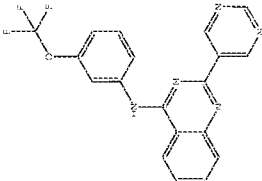
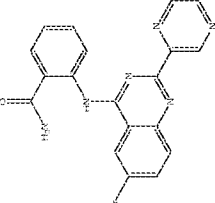
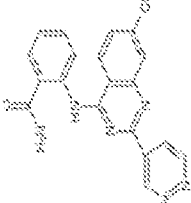
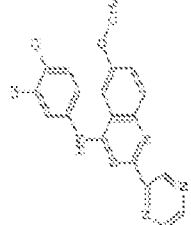
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Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1224			413.81	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.05 (s, 1H), 9.63 (s, 1H), 8.82 (s, 1H), 8.78 (s, 1H), 8.66 (s, 1H), 8.31 (s, 1H), 7.89 - 7.91 (m, 2H), 7.49 (d, J = 8.0 Hz, 1H), 7.31 (t, J = 7.3 Hz, 1H), 6.96 (d, J = 7.6 Hz, 1H), 2.74 (s, 3H).	DMSO	414.0 416.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G3
1225			400.24	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.90 (s, 1H), 9.54 (s, 1H), 8.77 - 8.80 (m, 2H), 8.60 (s, 1H), 8.50 (s, 1H), 7.99 (s, 1H), 7.76 (s, 1H), 7.44 (s, 1H), 2.67 (s, 3H).	DMSO	400.0 401.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G3
1226			390.83	¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.33 (s, 1H), 9.67 (s, 1H), 9.38 (d, J = 8.0 Hz, 1H), 8.88 (s, 1H), 8.79 (s, 1H), 8.52 (s, 1H), 8.08 (s, 1H), 8.04 (s, 1H), 7.97 (d, J = 7.2 Hz, 1H), 7.88 (s, 1H), 7.69 (d, J = 7.2 Hz, 1H), 7.21 (t, J = 7.6 Hz, 1H), 2.75 (s, 3H).	DMSO	391.0 392.0 (M+1)	Method A (TFA)	95	Method C, G3
1227			391.81	¹ H-NMR (400 MHz, DMSO-d ₆): δ 16.00 (s, 1H), 9.69 (s, 1H), 9.39 (d, J = 8.4 Hz, 1H), 8.90 (s, 1H), 8.79 (s, 1H), 8.29 (s, 1H), 8.13 (d, J = 7.6 Hz, 1H), 7.82 (s, 1H), 7.50 (t, J = 7.6 Hz, 1H), 7.06 (t, J = 7.2 Hz, 1H), 2.73 (s, 3H).	DMSO	392.0 393.1 (M+1)	Method A (TFA)	95	Method C, G3

(continued)

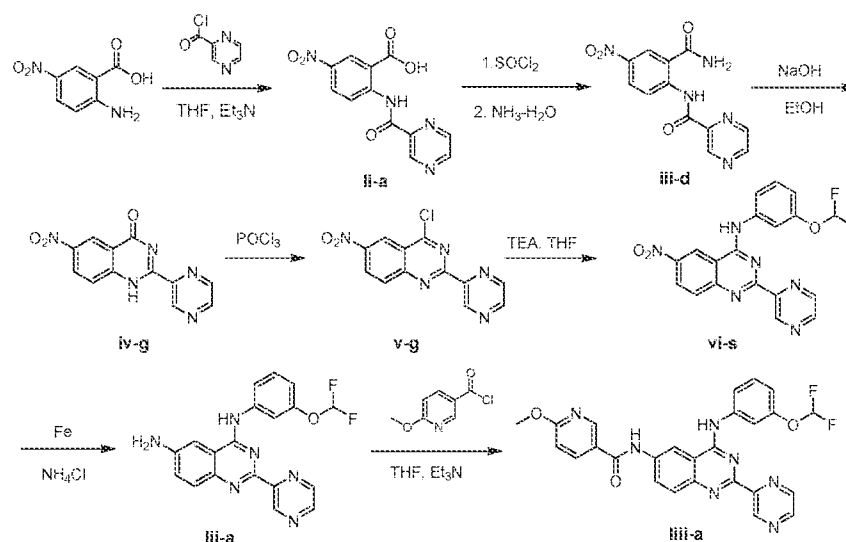
Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1228			372.81	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.07 (s, 1H), 9.57 (s, 1H), 8.82 (s, 2H), 8.76 (s, 1H), 8.57 (s, 1H), 8.32 (d, J = 7.6 Hz, 1H), 7.81 (s, 1H), 7.57 - 7.63 (m, 2H), 2.70 (s, 3H).	DMSO	373.0 374.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G3
1229			416.69	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.09 (s, 1H), 9.63 (d, J = 1.2 Hz, 1H), 8.85 (dd, J = 2.4, 1.6 Hz, 1H), 8.78 (d, J = 2.4 Hz, 2H), 8.63 (d, J = 1.6 Hz, 1H), 8.45 (s, 2H), 7.34 (t, J = 2.0 Hz, 1H), 2.74 (s, 3H).	DMSO	416.9 417.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G3
1230		HCl	351.76	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.42 (s, 1H), 9.60 (s, 2H), 9.34 (s, 1H), 8.64 (d, J = 8.4 Hz, 1H), 8.24 (dd, J = 8.4, 2.4 Hz, 1H), 7.98 - 7.92 (m, 3H), 7.76 - 7.71 (m, 1H), 7.56 (t, J = 8.8 Hz, 1H).	DMSO	352.0, 354.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

(continued)

Number	Product	Salt type	Molecular Mass	¹ H NMR	¹ H NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1231		HCl	383.33	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.58 (s, 1H), 9.61 (s, 2H), 9.34 (s, 1H), 8.71 (d, J = 8.4 Hz, 1H), 8.13 (s, 1H), 8.04 - 7.93 (m, 3H), 7.78 - 7.73 (m, 1H), 7.62 (t, J = 8.4 Hz, 1H), 7.21 (d, J = 8.0 Hz, 1H).	DMSO	384.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1232			421.25	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.34 (s, 1H), 9.58 (s, 1H), 9.27 - 9.25 (m, 1H), 8.84 - 8.77 (m, 2H), 9.52 (s, 1H), 8.35 (s, 1H), 8.07 - 7.67 (m, 5H), 7.20 (t, J = 7.5 Hz, 1H).	DMSO	421.0, 423.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G6
1233				¹ H-NMR (400 MHz, DMSO-d ₆): δ 13.15 (s, 1H), 10.00 (s, 1H), 9.44 (d, J = 5.2 Hz, 1H), 8.95 (d, J = 8.4 Hz, 1H), 8.48 (s, 1H), 8.42 - 8.40 (m, 1H), 8.16 (d, J = 9.2 Hz, 1H), 7.97 - 7.91 (m, 3H), 7.81 - 7.73 (m, 2H), 7.25 (t, J = 2.4 Hz, 1H).	DMSO	377.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method G
1234				¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.95 (s, 1H), 9.54 (s, 1H), 8.82 (s, 1H), 8.74 (d, J = 2.0 Hz, 1H), 8.66 (d, J = 2.4 Hz, 1H), 8.07 (dd, J = 8.8, 2.4 Hz, 1H), 8.00 (d, J = 2.4 Hz, 1H), 7.92 (d, J = 8.8 Hz, 1H), 7.70 (d, J = 8.8 Hz, 1H), 7.60 (dd, J = 8.8, 2.4 Hz, 1H), 4.00 (s, 3H).	DMSO	397.9, 400.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method G

Scheme 67: Representative synthesis of compounds of formula with a pyrazine liii-a (see Scheme 65)

[0461]



[0462] Method C: 5-Nitro-2-(pyrazine-2-carboxamido)benzoic acid (li-a) To a solution of pyrazine-2-carboxylic acid (1.36 g, 10.9 mmol, 1 eq.) in SOCl_2 (20 mL) was added DMF (2 drops). The mixture was stirred at 60 °C for 20 min. The volatiles were removed *in vacuo* to give crude pyrazine-2-carbonyl chloride, which was used in the next step directly. To a suspension of 2-amino-5-nitrobenzoic acid (2.00 g, 10.9 mmol, 1.0 eq.) in THF (50 mL) was added Et_3N (1.09 g) and pyrazine-2-carbonyl chloride in anhydrous THF (50 mL) dropwise. The resulting mixture was stirred at room temperature for 18 h. After the reaction was completed, the volatiles were removed. The residue was suspended in H_2O (10 mL) and the pH was adjusted to 5 by slow addition of 2N HCl in water. The resulting solid was collected and dried *in vacuo* to give 3.12 g of 5-nitro-2-(pyrazine-2-carboxamido)benzoic acid as a brown solid (99%). LCMS m/z = 289.0 (M+1) (Method B) (retention time = 1.24 min).

[0463] Method A: N-(2-carbamoyl-4-nitrophenyl)pyrazine-2-carboxamide (iii-d) A mixture of 5-nitro-2-(pyrazine-2-carboxamido)benzoic acid (3.12 g, 10.8 mmol) in SOCl_2 (20 mL) was stirred at 80 °C for 2 h. After cooling, the volatiles were removed and the residue was suspended in DCM (150 mL), and a solution of $\text{NH}_3 \cdot \text{H}_2\text{O}$ (25% by weight in water, 40 mL) was added and stirred for 4 h. The resulting precipitate was collected and dried *in vacuo* to give 2.42 g of N-(2-carbamoyl-4-nitrophenyl)pyrazine-2-carboxamide as a dark red solid (74.6%). LCMS m/z = 288.0 (M+1) (Method B) (retention time = 1.11 min).

[0464] Method E2: 6-nitro-2-(pyrazin-2-yl)quinazolin-4(1H)-one (iv-g) To a mixture of N-(2-carbamoyl-4-nitrophenyl)pyrazine-2-carboxamide (2.42 g, 8.43 mmol, 1.0 eq.) in EtOH (60 mL) was added NaOH (1.98 g, 49.5 mmol, 5.0 eq.). The resulting mixture was stirred at room temperature for 18 h. After the reaction was completed, the volatiles were removed *in vacuo*. The residue was partitioned between H_2O (50 mL) and ethyl acetate (50 mL). The aqueous layer was neutralized to pH 5 by slow addition of aqueous citric acid. The resulting precipitate was collected and dried to give 2.00 g of 6-nitro-2-(pyrazin-2-yl)quinazolin-4(3H)-one as a yellow solid (88%). LCMS m/z = 270.1 (M+1) (Method A) (retention time = 1.36 min).

[0465] Method F2: 4-Chloro-6-nitro-2-(pyrazin-2-yl)quinazoline (v-g) To a mixture of 6-nitro-2-(pyrazin-2-yl)quinazolin-4(3H)-one (1.00 g, 3.7 mmol) in POCl_3 (10 mL) was added *N,N*-dimethylbenzylamine (0.1 mL). The resulting mixture was stirred at 120 °C for 2 h. After the reaction was completed, POCl_3 was removed *in vacuo*, and the residue was co-evaporated with toluene twice to give a dark crude product, which was used for the next step without further purification.

[0466] Method G6: N-(3-(difluoromethoxy)phenyl)-6-nitro-2-(pyrazin-2-yl)quinazolin-4-amine (vi-s) A mixture of 4-chloro-6-nitro-2-(pyrazin-2-yl)quinazoline (1.00 g, crude, 3.7 mmol, 1.0 eq.), 3-(difluoromethoxy)benzenamine (600 mg, 3.7 mmol, 1.0 eq.) and Et_3N (1.00 g, 10 mmol, 3.0 eq.) in THF (80 mL) was stirred at 75 °C for 18 h. After cooling, the volatiles were removed *in vacuo* and the residue was washed with H_2O (100 mL x 2). The solid was dried *in vacuo* to afford 1.40 g of N-(3-(difluoromethoxy)phenyl)-6-nitro-2-(pyrazin-2-yl)quinazolin-4-amine as a black solid (90.2% of two steps). LCMS m/z = 411.0 (M+1) (Method A) (retention time = 1.61 min).

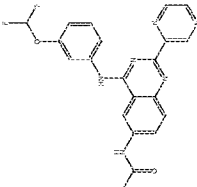
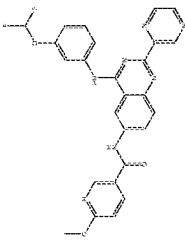
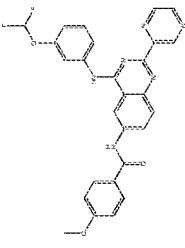
[0467] Method B: N⁴-(3-(difluoromethoxy)phenyl)-2-(pyrazin-2-yl)quinazoline-4,6-diamine (lii-a) To a mixture of N-(3-(difluoromethoxy)phenyl)-6-nitro-2-(pyrazin-2-yl)quinazolin-4-amine (1.40 g, 3.4 mmol, 1.0 eq.) in MeOH- H_2O

(v/v, 3:1, 110 mL) was added NH_4Cl (1.80 g, 34 mmol, 10.0 eq.) and Fe (1.91 g, 34 mmol, 1.0.0 eq.). The resulting mixture was stirred at 60 °C for 3 h. After the reaction was completed, the mixture was cooled to room temperature, and the iron was filtered off. The filtrate was concentrated to 15 mL and a precipitate formed and was collected and dried *in vacuo* to give 1.13 g of N^4 -(3-(difluoromethoxy)phenyl)-2-(pyrazin-2-yl)quinazoline-4,6-diamine as a pale yellow solid (87.5%). LCMS m/z = 381.1 (M+1) (Method B) (retention time = 1.60 min). $^1\text{H-NMR}$ (400 MHz, DMSO-d_6): δ 9.69 (s, 1H), 9.52 (d, J = 1.2 Hz, 1H), 8.75 (d, J = 2.4 Hz, 1H), 8.69 (d, J = 2.4 Hz, 1H), 8.30 (s, 1H), 7.87 (dd, J = 8.4, 0.8 Hz, 1H), 7.72 (d, J = 8.8 Hz, 1H), 7.49 - 7.42 (m, 2H), 7.33 - 7.32 (m, 1H), 7.31 (t, J = 74.4 Hz, 1H), 5.85 (s, 2H).

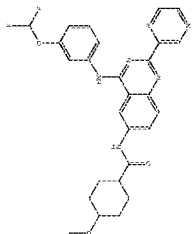
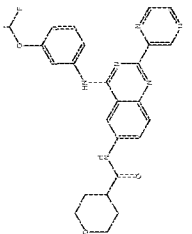
[0468] Method C: N-(4-(3-(difluoromethoxy)phenylamino)-2-(pyrazin-2-yl)quinazolin-6-yl)-6-methoxynicotinamide (liii-a) To a solution of 6-methoxynicotinic acid (100 mg, 0.65 mmol) in SOCl_2 (2 mL) was added DMF (1 drop). The mixture was stirred at 60 °C for 20 min. The volatiles were removed *in vacuo* to give 6-methoxynicotinoyl chloride, which was used for the next step directly. To a suspension of N^4 -(3-(difluoromethoxy)phenyl)-2-(pyrazin-2-yl)quinazoline-4,6-diamine (130 mg, 0.34 mmol, 0.5 eq.) in THF (5 mL) and Et_3N (101 mg, 1 mmol, 3.0 eq) was added 6-methoxynicotinoyl chloride in anhydrous THF (5 mL) dropwise. The resulting mixture was stirred at room temperature for 18 h. The volatiles were removed *in vacuo*. The residue was washed with MeOH and re-crystallized from THF/MeOH twice, and purified by reverse phase chromatography PREP-HPLC (A = $\text{NH}_4\text{HCO}_3\text{-H}_2\text{O}$, 10 mmol/L, B = MeOH) to afford 33 mg of N-(4-(3-(difluoromethoxy)phenylamino)-2-(pyrazin-2-yl)quinazolin-6-yl)-6-methoxynicotinamide as a pale yellow solid (18.8%). LCMS m/z = 516.1 (M+1), 258.6 (M/2+1) (Method A) (retention time = 1.57 min). $^1\text{H-NMR}$ (400 MHz, DMSO-d_6): δ 10.71 (s, 1H), 10.18 (s, 1H), 9.57 (d, J = 1.6 Hz, 1H), 9.01 (d, J = 1.6 Hz, 1H), 8.91 (d, J = 2.4 Hz, 1H), 8.81 (t, J = 2.0 Hz, 1H), 8.76 (d, J = 2.4 Hz, 1H), 8.34 (dd, J = 8.8, 2.8 Hz, 1H), 8.23 (s, 1H), 8.08 (dd, J = 8.8, 2.0 Hz, 1H), 8.00 (d, J = 9.2 Hz, 1H), 7.90 (d, J = 9.2 Hz, 1H), 7.46 - 7.50 (m, 1H), 7.32 (t, J = 74.4 Hz, 1H), 7.02 (d, J = 8.8 Hz, 1H), 6.95 (dd, J = 8.0, 2.0 Hz, 1H), 3.97 (s, 3H).

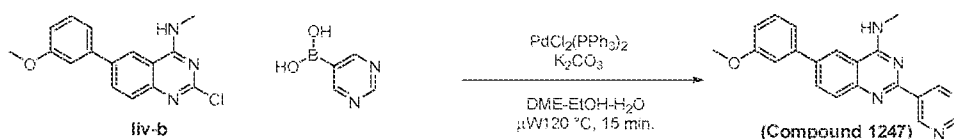
[0469] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 67** (prepared according to method **procedure A - G** as designated).

Table 22:

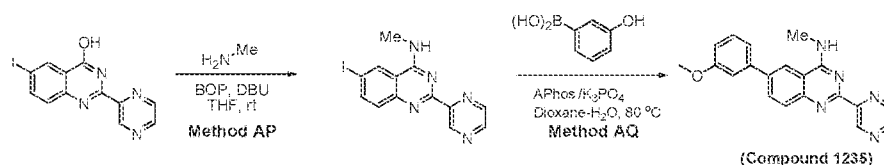
Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1242			422.39	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.10-10.45 (m, 2H), 9.68 (s, 1H), 8.45-9.10 (m, 2H), 7.68-8.37 (m, 4H), 7.36-7.52 (m, 2H), 7.02-7.08 (m, 1H), 2.21 (s, 3H).	DMSO	423.0 (M+1)	Method A (TFA)	95	Method C, G ₄ , C
1243			515.47	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.71 (s, 1H), 10.18 (s, 1H), 9.57 (d, J = 1.6 Hz, 1H), 9.01 (d, J = 1.6 Hz, 1H), 8.91 (d, J = 2.4 Hz, 1H), 8.81 (t, J = 2.0 Hz, 1H), 8.76 (d, J = 2.4 Hz, 1H), 8.34 (dd, J = 8.8, 2.8 Hz, 1H), 8.23 (s, 1H), 8.08 (dd, J = 8.8, 2.0 Hz, 1H), 8.00 (d, J = 9.2 Hz, 1H), 7.90 (d, J = 9.2 Hz, 1H), 7.46-7.50 (m, 1H), 7.32 (t, J = 74.4 Hz, 1H), 7.02 (d, J = 8.8 Hz, 1H), 6.95 (dd, J = 8.0, 2.0 Hz, 1H), 3.97 (s, 3H).	DMSO	516.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G ₄ , C
1244			514.48	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.56 (s, 1H), 10.16 (s, 1H), 9.57 (s, 1H), 9.00 (d, J = 1.6 Hz, 1H), 8.81 (d, J = 2.4 Hz, 1H), 8.76 (d, J = 2.8 Hz, 1H), 8.23 (s, 1H), 8.07-8.12 (m, 3H), 7.99 (d, J = 9.2 Hz, 1H), 7.89 (d, J = 9.2 Hz, 1H), 7.46-7.50 (m, 1H), 7.32 (t, J = 72.8 Hz, 1H), 7.11-7.13 (m, 2H), 6.95 (dd, J = 8.4, 2.4 Hz, 1H), 3.87 (s, 3H).	DMSO	515.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G ₄ , C

(continued)

Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1245			520.53	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.30 (s, 1H), 10.16 (s, 1H), 9.54 (s, 1H), 8.89 (s, 1H), 8.79 (t, J = 2.4 Hz, 1H), 8.74 (d, J = 2.8 Hz, 1H), 8.20 (s, 1H), 7.93 (s, 2H), 7.87 (d, J = 8.4 Hz, 1H), 7.48 (dd, J = 10.0, 2.0 Hz, 1H), 7.31 (t, J = 74.0 Hz, 1H), 6.94 (dd, J = 8.4, 2.0 Hz, 1H), 3.26 (s, 3H), 3.18 - 3.22 (m, 1H), 1.91 - 1.96 (m, 2H), 1.66 - 1.78 (m, 2H), 1.53 - 1.63 (m, 2H), 1.38 - 1.49 (m, 2H).	DMSO	521.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G4, C
1246			492.48	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.35 (s, 1H), 10.15 (s, 1H), 9.54 (s, 1H), 8.90 (d, J = 2.0 Hz, 1H), 8.80 (dd, J = 2.4, 1.2 Hz, 1H), 8.75 (d, J = 2.4 Hz, 1H), 8.19 (s, 1H), 7.95 (d, J = 8.8 Hz, 1H), 7.86 - 7.91 (m, 2H), 7.48 (dd, J = 10.8, 2.8 Hz, 1H), 7.31 (t, J = 74.4 Hz, 1H), 6.94 (dd, J = 7.6, 2.0 Hz, 1H), 3.94 - 3.97 (m, 2H), 3.40 - 3.43 (m, 2H), 2.70 - 2.73 (m, 1H), 1.65 - 1.78 (m, 4H).	DMSO	493.2 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G4, C

Scheme 68: Synthesis of 6-(3-methoxyphenyl)-N-methyl-2-(pyrimidin-5-yl)quinazolin-4-amine (Compound 1247)**[0470]**

[0471] Method BF : 6-(3-methoxyphenyl)-N-methyl-2-(pyrimidin-5-yl)quinazolin-4-amine (compound 1247) In a 10 mL microwave vial was added 2-chloro-6-(3-methoxyphenyl)-N-methylquinazolin-4-amine (0.080 g, 0.267 mmol), pyrimidine-5-boric acid (0.099 g, 0.801 mmol), dichlorobis(triphenylphosphine)palladium (II) ($\text{Pd}(\text{PPh}_3)_2\text{Cl}_2$) (9.37 mg, 0.013 mmol), and potassium carbonate (0.111 g, 0.801 mmol) in DME (3 mL), EtOH (1.286 mL), and water (0.857 mL) to give a yellow suspension. The vial was irradiated at 120 °C for 15 min under argon. Water (10 mL) was added to the mixture and extracted with ethyl acetate (2 x 10 mL). The organic layers were combined and washed with brine (1 x 20 mL) and then dried over MgSO_4 , filtered and concentrated. The residue was washed with $\text{MeOH-CH}_2\text{Cl}_2$ and dried to give 35 mg of 6-(3-methoxyphenyl)-N-methyl-2-(pyrimidin-5-yl)quinazolin-4-amine as a white solid (38 %). LCMS m/z = 344 ($M + 1$) (Method D) (retention time = 1.78 min). ^1H NMR (300 MHz, DMSO) δ 9.79 - 9.59 (m, 2H), 9.30 (s, 1H), 8.72 (d, J = 4.6 Hz, 1H), 8.59 (s, 1H), 8.16 (d, J = 8.6 Hz, 1H), 7.85 (d, J = 8.7 Hz, 1H), 7.53 - 7.28 (m, 3H), 7.00 (d, J = 6.8 Hz, 1H), 3.86 (s, 3H), 3.18 (d, J = 3.9 Hz, 3H).

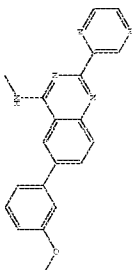
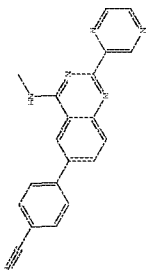
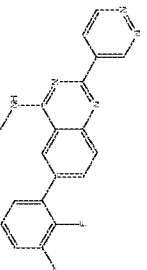
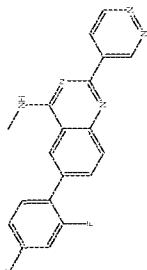
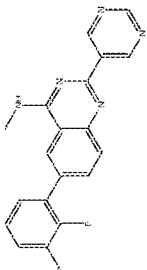
Scheme 69: Synthesis of 6-(3-methoxyphenyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine (Compound 1235)**[0472]**

[0473] In a 100 mL round-bottomed flask was added 6-iodo-2-(pyrazin-2-yl)quinazolin-4(3H)-one (0.500 g, 1.428 mmol), BOP (0.821 g, 1.857 mmol), and 1,8-diazabicyclo[5.4.0]undec-7-ene (0.426 mL, 2.86 mmol) in DMF (10 mL) to give a colorless solution. Methylamine, 2M in THF (2.142 mL, 4.28 mmol) was added and stirred at room temperature overnight. The reaction mixture was diluted with water (50 mL) and then a precipitate formed. The resulting solid was collected by filtration and dried to give 0.515 g of 6-iodo-N-methyl-2-(pyrazin-2-yl)quinazolin-4-amine as a pale brown solid in a 99 % yield. LCMS m/z = 364 ($M + 1$) (Method D) (retention time = 1.25 min). ^1H NMR (300 MHz, DMSO) δ 9.60 (s, 1H), 8.89 - 8.65 (m, 3H), 8.65 - 8.48 (m, 1H), 8.06 (d, J = 8.7 Hz, 1H), 7.59 (d, J = 8.7 Hz, 1H), 3.11 (d, J = 4.2 Hz, 3H).

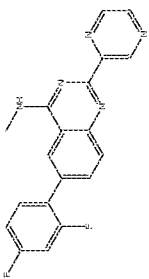
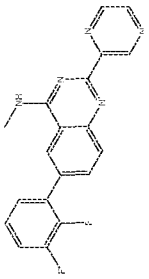
[0474] In a 50 mL round-bottomed flask was added 6-iodo-N-methyl-2-(pyrazin-2-yl)quinazolin-4-amine (0.100 g, 0.275 mmol), 3-methoxyphenylboronic acid (0.063 g, 0.413 mmol), bis(di-tert-butyl(4-dimethylaminophenyl)phosphine) dichloropalladium(II) (0.016 g, 0.022 mmol), and potassium phosphate tribasic monohydrate (0.190 g, 0.826 mmol) in dioxane (5 mL) and water (0.5 mL) to give a brown suspension. The reaction mixture was heated at 80 °C overnight under argon. After cooling to room temperature, the reaction mixture was diluted with water (10 mL) and then a precipitate formed. The resulting solid was collected by filtration and washed with ethyl acetate and dried to give 25 mg of 6-(3-methoxyphenyl)-N-methyl-2-(pyrazin-2-yl)quinazolin-4-amine as a pale yellow solid in a 26 % yield. LCMS m/z = 344 ($M + 1$) (Method D) (retention time = 1.43 min). ^1H NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.85 - 8.78 (m, 1H), 8.77 - 8.71 (m, 1H), 8.70 - 8.63 (m, 1H), 8.63 - 8.55 (m, 1H), 8.18 (d, J = 8.8 Hz, 1H), 7.89 (d, J = 8.9 Hz, 1H), 7.53 - 7.31 (m, 3H), 7.09 - 6.92 (m, 1H), 3.86 (s, 3H), 3.17 (d, J = 3.6 Hz, 3H).

[0475] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 69** (prepared according to method described for 6-(3-methoxyphenyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine).

Table 23:

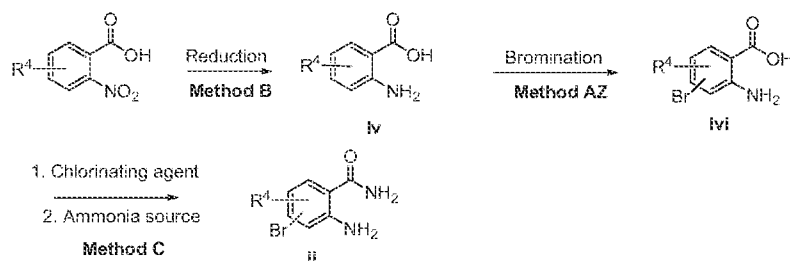
Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1235			343.38	¹ H-NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.85 - 8.78 (m, 1H), 8.77 - 8.71 (m, 1H), 8.70 - 8.63 (m, 1H), 8.63 - 8.55 (m, 1H), 8.18 (d, J = 8.8 Hz, 1H), 7.89 (d, J = 8.9 Hz, 1H), 7.53 - 7.31 (m, 3H), 7.09 - 6.92 (m, 1H), 3.86 (s, 3H), 3.17 (d, J = 3.6 Hz, 3H).	DMSO	344 (M+1)	Method D	100	Method AP/AQ
1236			338.37	¹ H-NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.87 - 8.61 (m, 4H), 8.24 (d, J = 8.8 Hz, 1H), 8.15 - 7.98 (m, 4H), 7.92 (d, J = 8.7 Hz, 1H), 3.17 (d, J = 4.0 Hz, 3H).	DMSO	339 (M+1)	Method D	100	Method C, G1
1237		2HCl		¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.17 (s, 1H), 9.54 (d, J = 5.2 Hz, 1H), 9.36 (s, 1H), 8.63 (s, 2H), 8.13 - 8.06 (m, 2H), 7.58 - 7.51 (m, 2H), 7.43 - 7.38 (m, 1H), 3.23 (d, J = 4.0 Hz, 3H).	DMSO	350.1, 351.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1238		2HCl		¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.14 (s, 1H), 9.50 (d, J = 5.2 Hz, 1H), 9.24 (s, 1H), 8.60 (dd, J = 5.4, 1.8 Hz, 1H), 8.55 (s, 1H), 8.04 (q, J = 8.8 Hz, 2H), 7.76 (dd, J = 15.4, 9.0 Hz, 1H), 7.51 - 7.45 (m, 1H), 7.32 (dt, J = 8.4, 2.0 Hz, 1H), 3.23 (d, J = 4.4 Hz, 3H).	DMSO	350.0, 351.0 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1239				¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.73 (s, 1H), 9.32 (s, 1H), 8.74 (d, J = 4.4 Hz, 1H), 8.51 (s, 1H), 8.03 (d, J = 8.7 Hz, 1H), 7.92 (d, J = 8.6 Hz, 1H), 7.54 - 7.49 (m, 2H), 7.42 - 7.37 (m, 1H), 4.12 (q, J = 5.2 Hz, 1H), 3.17 (t, J = 4.5 Hz, 3H).	DMSO	350.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

(continued)

Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for Coupling
1240		2HCl		¹ H-NMR (400 MHz, CD ₃ OD): δ 9.77 (s, 1H), 8.85 (d, J = 11.4 Hz, 2H), 8.42 (s, 1H), 8.19 - 8.03 (m, 2H), 7.60 (dd, J = 15.8, 7.8 Hz, 1H), 7.10 (t, J = 8.4 Hz, 2H), 3.38 (s, 3H).	CD ₃ OD	350.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1
1241		2HCl		¹ H-NMR (400 MHz, DMSO): δ 10.84 (d, J = 1.8 Hz, 1H), 9.87 (s, 1H), 9.06 (d, J = 13.6 Hz, 2H), 8.90 (s, 1H), 8.32 (t, J = 9.8 Hz, 2H), 7.67 - 7.35 (m, 3H), 3.39 (s, 3H).	DMSO	349.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method C, G1

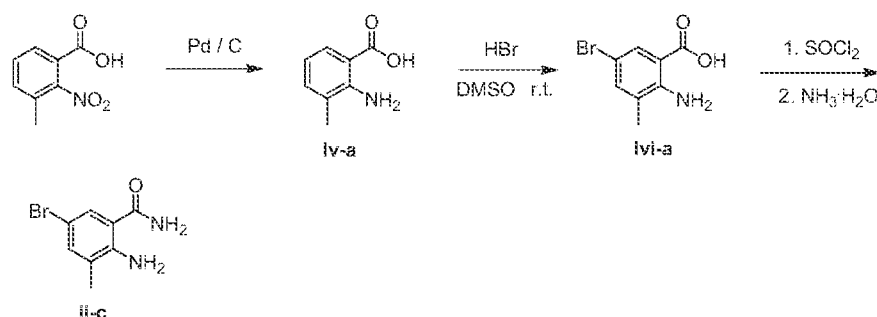
Scheme 70: General route for the synthesis of compounds with general formula ii

[0476]



Scheme 71: Representative synthesis of compounds of formula ii (see Scheme 70)

[0477]



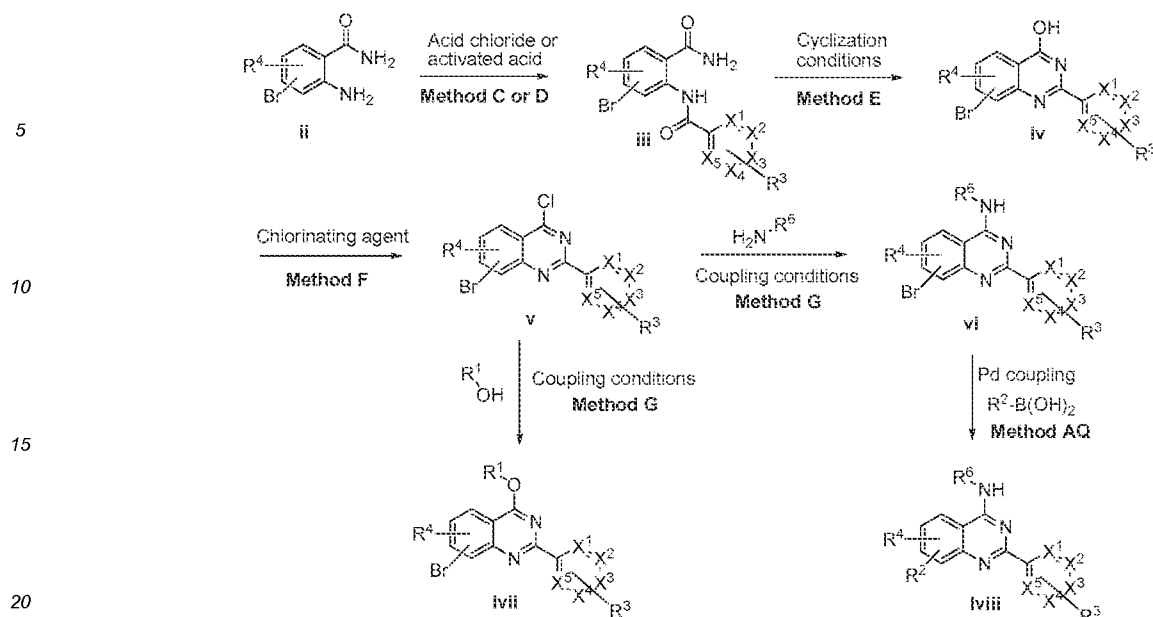
[0478] Method B2: 2-Amino-3-methylbenzoic acid (iv-a) Pd/C catalyst (150 mg) was suspended in a solution of 3-methyl-2-nitrobenzoic acid (1.50 g, 8.28 mmol, 1.0 eq) in THF (60 mL), the mixture was stirred under H₂ atmosphere at room temperature overnight. The Pd/C was removed by filtration over Celite and the THF was removed *in vacuo* to give 1.23 g of iv-a as a white solid (yield 98 %). LCMS *m/z* = 152.1 (M+1) (Method B) (retention time = 0.73 min). The product was used further without purification.

[0479] Method AZ: 2-Amino-5-bromo-3-methylbenzoic acid (ivi-a) To a solution of 2-amino-3-methylbenzoic acid (1.23 g, 8.14 mmol, 1.0 eq) in 15 mL of DMSO was added 40% HBr (6.00 mL, 44.7 mmol, 5 eq). The resulting mixture was stirred at room temperature overnight. A white precipitate formed during the course of the reaction. The reaction mixture was quenched with saturated aqueous NaHCO₃ resulting in a white solid that was filtered and dried *in vacuo* to yield 950 mg in 51% yield of ivi-a as white solid. LCMS *m/z* = 229.9 (M+1) (Method B) (retention time = 1.20 min).

[0480] Method C: 2-Amino-5-bromo-3-methylbenzamide (ii-c) A mixture of 2-amino-5-bromo-3-methylbenzoic acid (950 mg, 4.15 mmol) and SOCl₂ (20 mL) was stirred at 80°C for 2 h. After the reaction was completed, the mixture was cooled to room temperature. The SOCl₂ was removed *in vacuo* and the residue was dissolved in anhydrous THF (10 mL). The THF solution was then added dropwise to a 28% by weight solution of NH₃-H₂O (10 mL). After 1 h, the resulting precipitate was collected and dried *in vacuo* to give 820 mg of ii-c as a yellow solid (87%). LCMS *m/z* = 288.9, 230.9 (M+1) (Method B) (retention time = 1.49 min).

Scheme 72: General route for the synthesis of compounds with general formula lvii and lviii

[0481]



X^1, X^2, X^3, X^4 or $X^5 = CH$ or N , at least one X^1, X^2, X^3, X^4 or X^5 must be N

Method G for Coupling Conditions

[0482]

- G1: *i*-PrOH/85-100 °C
- G2: THF/heat
- G3: *i*-AmOH/100-130 °C
- G4: MeOH/microwave/150 °C
- G5: *i*-AmOH/ microwave/150 °C
- G6: THF/ Et_3N /reflux
- G7: THF- H_2O /NaOAc/rt-60 °C
- G8: NaH/THF
- G9: *n*-BuLi/THF
- G10: LHMDS/THF
- G11: LDA/THF
- G12: K_2CO_3 /DMF/60 °C
- G13: Cs_2CO_3 /DMA/80 °C
- G14: NaOtBu/DMF/Microwave 100 °C

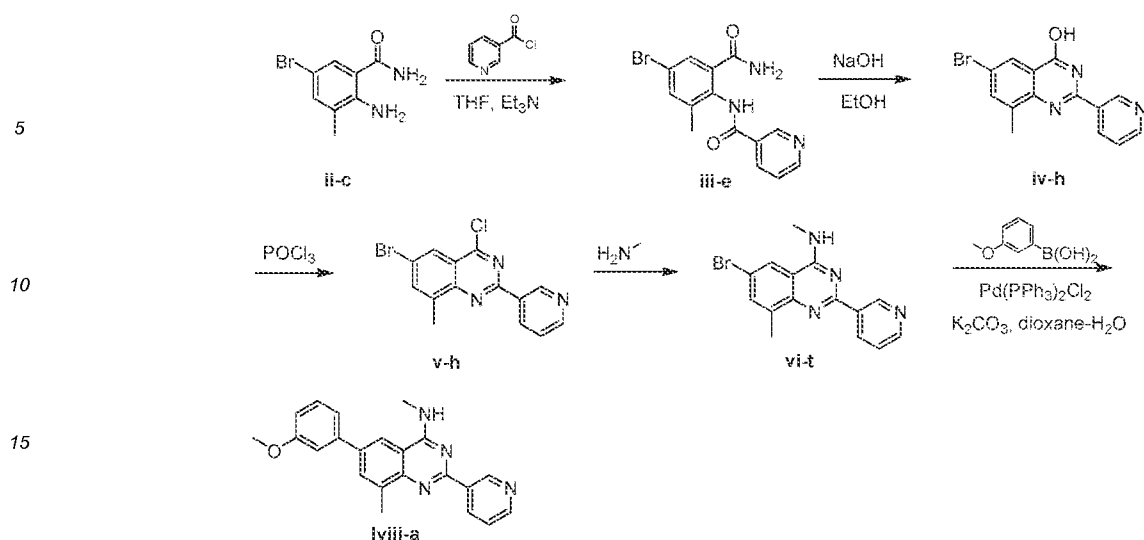
Method AQ for Coupling Conditions

[0483]

- AQ1: $Pd(PPh_3)_2Cl_2$ / K_2CO_3 / Dioxane- H_2O
- AQ2: $Pd_2(APhos)_2Cl_2$ / K_3PO_4 /Dioxane- H_2O
- AQ3: $Pd(PPh_3)_4$ / K_3PO_4 /Dioxane- H_2O
- AQ4: $Pd(dppf)Cl_2-CH_2Cl_2$ / K_3PO_4 / Dioxane- H_2O
- AQ5: $Pd(OAc)_2Cl_2$ / S-Phos/ K_3KPO_4 /Dioxane- H_2O
- AQ6: $Pd(dppf)Cl_2-CH_2Cl_2$ / Na_2CO_3 / Dioxane- H_2O

Scheme 73: Representative synthesis of compounds of formula Ivii and Iviii (see Scheme 72)

[0484]



[0485] Method C: N-(4-bromo-2-carbamoyl-6-methylphenyl)nicotinamide (iii-e) To a solution of 2-amino-5-bromo-3-methylbenzamide (820 mg, 3.55 mmol, 1.0 eq.) in THF (15 mL) and Et₃N (0.7 mL) was added nicotinoyl chloride (551 mg, 3.91 mmol, 1.1 eq.) in anhydrous THF (15 mL) dropwise. The resulting mixture was stirred at room temperature overnight. After the reaction was completed, the resultant precipitate was filtered and dried *in vacuo* to give 1.74 g of crude **iii-e** as a yellow solid. LCMS *m/z* = 333.8, 335.8 (M+1) (Method B) (retention time = 1.42 min).

[0486] Method E: 6-Bromo-8-methyl-2-(pyridin-3-yl)quinazolin-4-ol (iv-h) A mixture of N-(4-bromo-2-carbamoyl-6-methylphenyl)nicotinamide (1.74 g salt, 5.22 mmol, 1.0 eq) in EtOH (50 mL) was treated with NaOH (1.04 g, 26.1 mmol, 5.0 eq). The resulting mixture was stirred at room temperature overnight. After the reaction was completed, the volatiles were removed *in vacuo*. Water (30 mL) was added to the residue and the mixture was adjusted to pH ~ 1 or 2 by slow addition of aqueous HCl. The resultant precipitate was collected and dried to give 870 mg of **iv-h** as a yellow solid (77% yield after two steps). LCMS *m/z* = 315.7, 317.7 (M+1) (Method B) (retention time = 1.74 min).

[0487] Method F5: 6-Bromo-4-chloro-8-methyl-2-(pyridin-3-yl)quinazoline (v-h) 6-Bromo-8-methyl-2-(pyridin-3-yl)quinazolin-4-ol (870 mg, 2.76 mmol) was added to POCl₃ (10 mL). The resulting mixture was stirred at 120°C overnight. After the reaction was completed, the mixture was carefully poured into ice-water. The pH was adjusted to 7 by slow addition of NE₄OH at 0°C. The resultant solid was collected to give 1.00 g of **v-h** as a beige solid (quantitative yield). LCMS *m/z* = 333.9, 335.9 (M+1) (Method B) (retention time = 2.23 min).

[0488] Method G6: 6-Bromo-N, 8-dimethyl-2-(pyridin-3-yl)quinazolin-4-amine (vi-t) A mixture of 6-bromo-4-chloro-8-methyl-2-(pyridin-3-yl)quinazoline (200 mg, 0.60 mmol, 1.0 eq.), methylamine (81 mg, 1.20 mmol, 2.0 eq.) and Et₃N (0.2 mL) in *i*-PrOH (10 mL) was stirred at 85 °C overnight. The resultant yellow precipitate was collected to afford 125 mg of **vi-t** as a beige solid (63.4%). LCMS *m/z* = 328.8, 330.8 (M+1) (Method B) (retention time = 1.95 min).

[0489] Method G2: 7-bromo-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine (vi-u) To a suspension of 7-bromo-4-chloro-2-(pyridin-3-yl)quinazoline (5.0 g, 0.0156 mol) in THF (100 mL) was added dropwise a methylamine solution (40 wt. % in H₂O) (24 mL, 0.272 mmol) with cooling. The suspension was stirred at 60 °C for 3h, cooled, filtered, and dried to give the title compound. (3.62 g, 73.5 %)

[0490] Method AQ1: 6-(3-Methoxyphenyl)-N, 8-dimethyl-2-(pyridin-3-yl) quinazolin-4-amine (Iviii-a) (This method is representative of method AQ2 and can be implemented in a similar way except for substitution of the appropriate catalyst and base) To a mixture of 6-bromo-N,8-dimethyl-2-(pyridin-3-yl)quinazolin-4-amine (130 mg, 0.396 mmol, 1.0 eq), 3-methoxyphenylboronic acid (60 mg, 0.396 mmol, 1.0 eq), K₂CO₃ (295 mg, 2.14 mmol, 5.4 eq.) in dioxane (8 mL) and H₂O (4 mL) was added Pd(PPh₃)₂Cl₂ (15 mg, 0.021 mmol, 0.054 eq) under N₂ atmosphere. The resulting mixture was stirred at 120°C under N₂ atmosphere overnight. After the reaction was completed, the mixture was filtered, and the filtrate was concentrated *in vacuo*. The residue was purified by reverse phase HPLC column to afford 11 mg of **Iviii-a** as a white solid (yield 7.8%). LCMS *m/z* = 357.2, (M+1) (Method B) (retention time = 2.11 min). ¹H NMR (400 MHz, DMSO-*d*₆): δ 9.68 (d, *J* = 1.2 Hz, 1H), 8.75 (d, *J* = 8.0 Hz, 1H), 8.62 (d, *J* = 4.6 Hz, 1H), 8.49 (d, *J* = 4.4 Hz, 1H), 8.35 (s, 1H), 7.97 (s, 1H), 7.49 (dd, *J* = 7.8, 4.8 Hz, 1H), 7.44 - 7.29 (m, 3H), 6.93 (dd, *J* = 6.8, 2.4 Hz, 1H), 3.81 (s, 3H), 3.12 (d, *J* = 4.4 Hz, 3H), 2.68 (s, 3H).

[0491] Method AQ3: 6-(4-Fluorophenyl)-N,8-dimethyl-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride (Iviii-b) 6-Bromo-N,8-dimethyl-2-(pyridin-3-yl)quinazolin-4-amine (340mg, 1.03 mmol), 3-fluorobenzeneboronic acid (217 mg, 1.55 mmol), K₃PO₄ (658 mg, 3.10 mmol) and Pd(PPh₃)₄ (59.7 mg, 0.052 mmol) were dissolved in the mixed solvent of 1,4-dioxane (10 mL) and water (1 mL). The resulting mixture was stirred at 90°C for 6 hours under a nitrogen atmosphere.

After the reaction was completed, water was added to the mixture and stirred for 30 minutes. The resulting precipitate was collected by filtration and purified by column chromatography on NH-silica gel (eluted with THF) to give a yellow powder. The solid was suspended in ethanol and 5N HCl (1 mL) was added to the mixture. The mixture was sonicated for 10 min and the resulting precipitate was collected by filtration and dried to give 304 mg of 6-(3-fluorophenyl)-N,8-

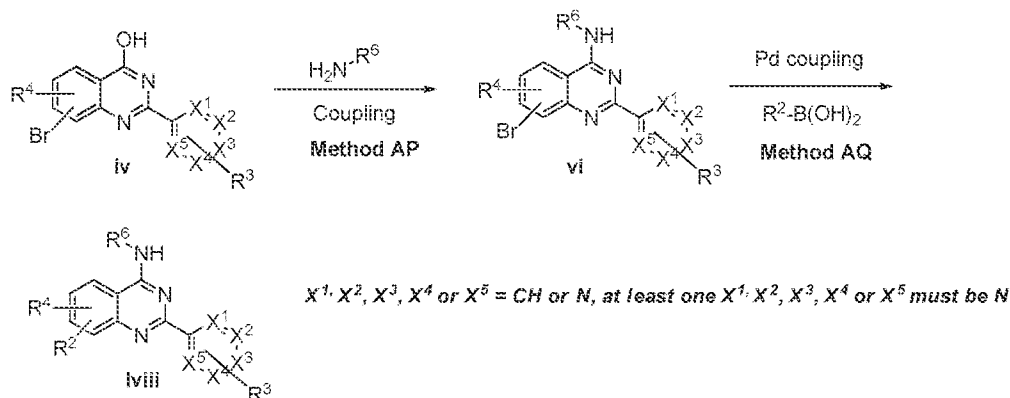
[0492] Method AQ4: 6-(2,4-Difluorophenyl)-N,5-dimethyl-2-(pyridin-3-yl)quinazolin-4-amine(Iviii-c) 6-Bromo-N,5-dimethyl-2-(pyridin-3-yl)quinazolin-4-amine (350mg, 1.06 mmol), 2,4-difluorophenylboronic acid (252 mg, 1.595 mmol), K_3PO_4 (677 mg, 3.19 mmol) and $Pd(dppf)Cl_2 \cdot CH_2Cl_2$ (87 mg, 0.106 mmol) were dissolved in the mixed solvent of 1,4-dioxane (10 mL) and water (1 mL). The resulting mixture was stirred at 90°C for 2.5 hours under N_2 . After the reaction was completed, water was added to the mixture and extracted with ethyl acetate. The combined organic layers were washed with water and brine, dried over Na_2SO_4 . After filtration and evaporation, the crude product was purified by column chromatography on NH-silica gel (eluted with isocratic 33%ethyl acetate/67%hexane) to give a white powder. The solid was suspended in ethanol and 5N HCl (1.0 mL) was added to the mixture. The mixture was sonicated for 10 min and the resultant precipitate was collected by filtration and dried to give 139 mg of 6-(2,4-difluorophenyl)-N,5-

[0493] Method AQ5: 7-(3-fluorophenyl)-N,6-dimethyl-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride (Iviii-d) A mixture of 7-chloro-N,6-dimethyl-2-(pyridin-3-yl)quinazolin-4-amine (400 mg, 1.40 mmol), 3-fluorophenylboronic acid (294 g, 2.10 mmol), $Pd(OAc)_2$ (15.8 mg, 0.070 mmol), 2-dicyclohexylphosphino-2',6'-dimethoxybiphenyl (86.7 mg, 0.211 mmol), K_3PO_4 (900 mg, 4.23 mmol) in dioxane (10 mL) and water (2 mL) was stirred under reflux for 2.5 h. Ethyl acetate (20 mL) was added to the cooled mixture and a precipitate formed and was filtered. The solid was recrystallized from DMF and water to give the title compound as free form. The solid was suspended in ethyl acetate (10 mL) and 4N HCl in ethyl acetate (1.0 mL) was added. The resulting solid was subjected to sonication for 20 min, filtered and dried to give the title compound as the bis-HCl salt (0.10 g, 18.7 %).

[0494] Method AQ6:7-(3,4-difluorophenyl)-N,8-dimethyl-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride (Iviii-e) To a suspension of 3,4-difluorophenylboronic acid (389 mg, 2.46 mmol) and 7-bromo-N,8-dimethyl-2-(pyridin-3-yl)quinazolin-4-amine (395.4 mg, 1.201 mmol) in dioxane/ H_2O (2/1) (30 mL) under a nitrogen atmosphere was added Na_2CO_3 (633.2mg, 5.97 mmol) and (1,1'-bis(diphenylphosphino)ferrocene)-dichloropalladium(II) (98 mg, 0.120 mmol) at room temperature. The mixture was stirred at 100°C for 1.5 h. Water was added to the reaction mixture and then a precipitate formed. The solid was filtered and washed with water and dried. The dried solid was then heated in a methanol/dioxane mixture to give a clear solution and filtered through Celite. The filtrate was concentrated to give the crude product. The crude product was sonicated in methanol/ CH_2Cl_2 for ca.15 min and filtered to give 305.2 mg of a pale brown solid as the parent compound. To a suspension of parent compound in methanol was added 4N HCl in ethyl acetate (ca. 4 mL) to give a clear solution. The solution was concentrated and recrystallized from ethanol to give the HCl salt. The salt was collected and dried in an oven at 60°C to give 231.3 mg in a 44% yield as pale yellow solid. 1H NMR ($DMSO-d_6$) δ 9.73 (s, 1H), 9.37 (brd, J = 8.08 Hz, 1H), 8.97 (brd, J = 5.24 Hz, 1H), 8.77 (brs, 1H), 8.20 (d, J = 8.48 Hz, 1H), 8.10-8.07 (brm, 1H), 7.63-7.55 (brm, 2H), 7.47 (d, J = 8.48 Hz, 1H), 7.32 (brm, 1H), 3.20 (d, J = 4.20 Hz, 3H), 2.65 (s, 3H). The 1H of 2HCl was not observed.

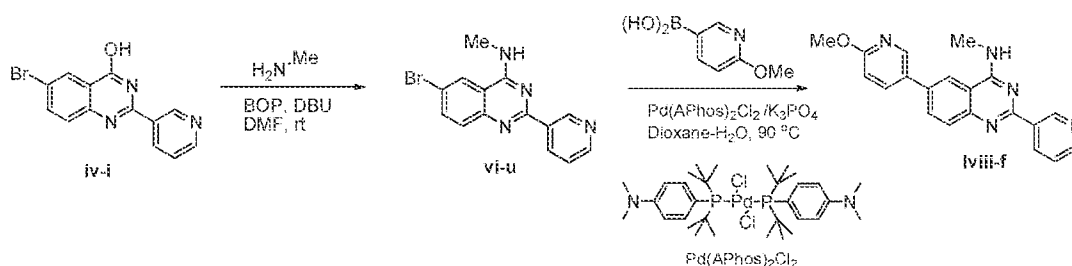
Scheme 74: General route for the synthesis of compounds with general formula Iviii

[0495]



Scheme 75: Representative synthesis of compounds of formula Iviii (see Scheme 74)

[0496]

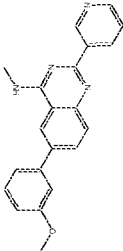
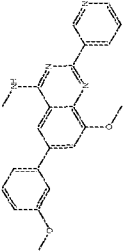


[0497] Method AP: 6-bromo-N-methyl-2-(pyridine-3-yl)quinazoline-4-amine (vi-u) To a solution of 6-bromo-2-(pyridin-3-yl)quinazolin-4(3H)-one (5.00 g, 16.6 mmol), (benzotriazol-1-yloxy)tris(dimethylamino)phosphonium hexafluorophosphate (BOP) (9.52 g, 21.5 mmol), and 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) (4.94 ml, 33.1 mmol) in DMF (50 mL) was added methylamine, 2M in THF (16.6 mL, 33.1 mmol). The mixture was stirred overnight at room temperature. Water (100 mL) was added to the mixture and stirred. The resultant precipitate was collected by filtration and dried to give 5.15 g of 6-bromo-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine as pale yellow solid (99 %). LCMS m/z = 315 ($M + 1$) (Method D) (retention time = 1.34 min). ^1H NMR (300 MHz, DMSO) δ 9.60 (dd, J = 2.1, 0.8 Hz, 1H), 8.79 - 8.70 (m, 1H), 8.67 (dd, J = 4.8, 1.7 Hz, 1H), 8.57 (s, 1H), 8.51 (d, J = 2.0 Hz, 1H), 7.90 (dd, J = 8.9, 2.1 Hz, 1H), 7.71 (d, J = 8.9 Hz, 1H), 7.52 (ddd, J = 7.9, 4.8, 0.9 Hz, 1H), 3.13 (d, J = 4.5 Hz, 3H).

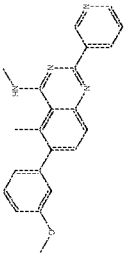
[0498] Method AQ2: 6-(6-methoxypyridin-3-yl)-N-methyl-2-(pyridine-3-yl)quinazoline-4-amine (Iviii-f) To a 1 dram reaction vial was added 6-bromo-N-methyl-2-(pyridine-3-yl)quinazoline-4-amine (35 mg, 0.111 mmol), 6-methoxypyridin-3-ylboronic acid (20.4 mg, 0.133 mmol), $\text{Pd(APhos)}_2\text{Cl}_2$ (3.2 mg, 0.004 mmol) and potassium phosphate monohydrate (77 mg, 0.33 mmol) in a mixture of dioxane-water (9:1, 2 mL). The reaction mixture was heated to 90 °C for 14 h after which it was cooled to room temperature and diluted with water (5 mL). The resultant precipitate was collected by filtration and recrystallized from methanol to give 6-(6-methoxypyridin-3-yl)-N-methyl-2-(pyridine-3-yl)quinazoline-4-amine as a pale yellow solid (19.1 mg, 51%). LCMS m/z = 344 ($M + 1$) (Method C) (retention time = 2.01 min). ^1H NMR (300 MHz, DMSO) δ 9.64 (d, J = 1.3 Hz, 1H), 8.84 - 8.74 (m, 1H), 8.68 (dd, J = 6.2, 1.7 Hz, 2H), 8.57 (d, J = 1.6 Hz, 2H), 8.16 (ddd, J = 14.4, 8.7, 2.2 Hz, 2H), 7.85 (d, J = 8.7 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 7.00 (d, J = 8.7 Hz, 1H), 3.93 (s, 3H), 3.18 (d, J = 4.3 Hz, 3H).

[0499] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 72 or 74**, replacing methylamine with the appropriate amine and 6-methoxypyridin-3-ylboronic acid with the appropriate boronic acid.

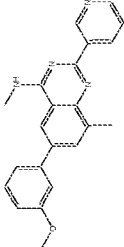
Table 24:

Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1248		HCl	342.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.66 (s, 1H), 8.50 - 8.89 (m, 3H), 8.16 (d, J = 8.2 Hz, 1H), 7.86 (d, J = 8.6 Hz, 1H), 7.30 - 7.61 (m, 4H), 7.01 (d, J = 6.4 Hz, 1H), 3.88 (s, 3H), 3.32 (brs, 1H), 3.19 (s, 3H).	DMSO	343.1 (M+1)		Method B (NH ₄ HCO ₃)	95	Method AQ1
1249			372.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.63 (s, 1H), 8.77 (d, J = 7.9 Hz, 1H), 8.67 (d, J = 3.7 Hz, 1H), 8.50 (s, 1H), 8.11 (s, 1H), 7.56 - 7.53 (m, 2H), 7.48 - 7.40 (m, 3H), 7.01 (d, J = 3.9 Hz, 1H), 4.07 (s, 3H), 3.88 (s, 3H), 3.17 (d, J = 4.0 Hz, 3H).	DMSO	372.9 (M+1)		Method B (NH ₄ HCO ₃)	95	Method AQ1

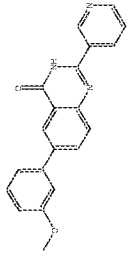
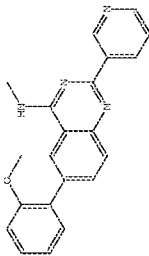
(continued)

Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1250			356.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.63 (s, 1H), 8.79 - 8.74 (m, 1H), 8.69 (d, J = 4.0 Hz, 1H), 7.67 (d, J = 8.4 Hz, 1H), 7.60 (d, J = 8.8 Hz, 1H), 7.57 - 7.51 (m, 1H), 7.42 (t, J = 8.0 Hz, 1H), 7.37-7.30 (m, 1H), 7.02-6.92 (m, 3H), 3.82 (s, 3H), 3.18 (d, J = 4.0 Hz, 3H), 2.69 (s, 3H).	DMSO	357.1 (M+1)		Method B (NH ₄ HCO ₃)	95	Method AQ1

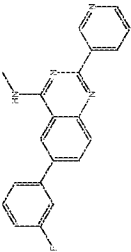
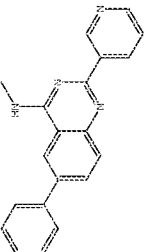
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Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1251			356.4	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.68 (d, J = 1.2 Hz, 1H), 8.75 (d, J = 8.0 Hz, 1H), 8.62 (d, J = 4.6 Hz, 1H), 8.49 (d, J = 4.4 Hz, 1H), 8.35 (s, 1H), 7.97 (s, 1H), 7.49 (dd, J = 7.8, 4.8 Hz, 1H), 7.44 (s, 1H), 7.29 (m, 3H), 6.93 (dd, J = 6.8, 2.4 Hz, 1H), 3.81 (s, 3H), 3.12 (d, J = 4.4 Hz, 3H), 2.68 (s, 3H).	DMSO	357.2 (M+1)		Method B (NH ₄ HCO ₃)	95	Method AQ1

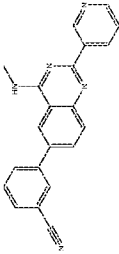
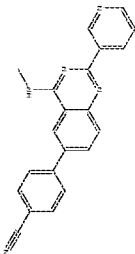
(continued)

Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1252			329.3	¹ H NMR (300 MHz, DMSO) δ 9.31 (s, 1H), 8.76 (d, J = 4.7 Hz, 1H), 8.51 (d, J = 8.0 Hz, 1H), 8.37 (d, J = 2.0 Hz, 1H), 8.18 (dd, J = 8.5, 2.2 Hz, 1H), 7.84 (d, J = 8.4 Hz, 1H), 7.59 (dd, J = 7.9, 4.8 Hz, 1H), 7.49 - 7.23 (m, 3H), 6.99 (d, J = 6.7 Hz, 1H), 3.85 (s, 3H).	DMSO	330 (M+1)	1.76	Method D	100	
1253		HCl	342.4	¹ H NMR (300 MHz, DMSO) δ 10.11 (s, 1H), 9.63 (s, 1H), 9.07 - 8.86 (m, 2H), 8.58 (s, 1H), 8.16 (s, 2H), 7.92 - 7.77 (m, 1H), 7.54 - 7.36 (m, 2H), 7.20 (d, J = 8.6 Hz, 1H), 7.16 - 7.07 (m, 1H), 3.81 (s, 3H), 3.29 (d, J = 3.9 Hz, 3H).	DMSO	343 (M+1)	1.47	Method D	100	Method AQ2

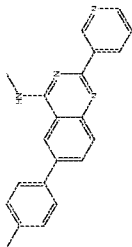
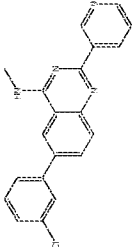
(continued)

Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1254			330.3	¹ H NMR (300 MHz, DMSO) δ 9.59 (s, 1H), 8.73 (d, J = 8.3 Hz, 1H), 8.65 (d, J = 4.7 Hz, 1H), 8.37 - 8.22 (m, 1H), 8.18 - 8.05 (m, 1H), 7.76 (d, J = 9.0 Hz, 1H), 7.71 (s, 1H), 7.56 - 7.43 (m, 2H), 4.58 (s, 2H), 3.14 (d, J = 4.4 Hz, 3H), 2.69 (d, J = 4.6 Hz, 3H).	DMSO	331 (M+1)	1.52	Method D	100	Method AQ2
1255			312.4	¹ H NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.88 - 8.47 (m, 4H), 8.25 - 8.05 (m, 1H), 7.97 - 7.72 (m, 3H), 7.67 - 7.35 (m, 4H), 3.18 (d, J = 4.4 Hz, 3H).	DMSO	313 (M+1)	1.48	Method D	100	Method AQ2

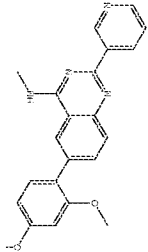
(continued)

Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1256		HCl	337.4	¹ H NMR (300 MHz, DMSO) δ 9.93 (s, 1H), 9.63 (s, 1H), 9.00 (d, J = 8.3 Hz, 1H), 8.95 - 8.82 (m, 2H), 8.46 - 8.32 (m, 2H), 8.25 (d, J = 7.0 Hz, 1H), 8.13 (d, J = 8.4 Hz, 1H), 7.92 (d, J = 7.2 Hz, 1H), 7.88 - 7.70 (m, 2H), 3.29 (d, J = 4.2 Hz, 3H).	DMSO	338 (M+1)	1.50	Method D	100	Method AQ2
1257			337.4	¹ H NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.77 (d, J = 7.9 Hz, 1H), 8.73 - 8.59 (m, 3H), 8.21 (d, J = 8.8 Hz, 1H), 8.11 - 7.94 (m, 4H), 7.87 (d, J = 8.7 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 3.18 (d, J = 4.0 Hz, 3H).	DMSO	338 (M+1)	1.50	Method D	100	Method AQ2

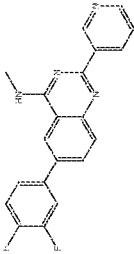
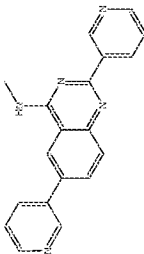
(continued)

Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1258			346.8	¹ H NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.77 (d, J = 7.9 Hz, 1H), 8.72 - 8.52 (m, 3H), 8.14 (d, J = 8.7 Hz, 1H), 7.97 - 7.74 (m, 3H), 7.68 - 7.45 (m, 3H), 3.18 (d, J = 4.0 Hz, 3H).	DMSO	347 (M+1)	1.60	1.60 Method D	100	Method AQ2
1259			346.8	¹ H NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.78 (d, J = 8.0 Hz, 1H), 8.72 - 8.54 (m, 3H), 8.17 (d, J = 8.7 Hz, 1H), 7.94 (s, 1H), 7.90 - 7.78 (m, 2H), 7.61 - 7.42 (m, 3H), 3.19 (d, J = 4.4 Hz, 3H).	DMSO	347 (M+1)	1.66	Method D	100	Method AQ2

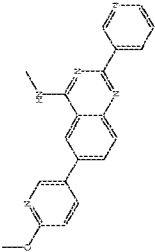
(continued)

Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1260			372.4	¹ H NMR (300 MHz, DMSO) δ 9.63 (s, 1H), 8.77 (dd, J = 7.9, 1.8 Hz, 1H), 8.67 (d, J = 4.7 Hz, 1H), 8.46 (d, J = 4.4 Hz, 1H), 8.26 (s, 1H), 7.95 - 7.84 (m, 1H), 7.75 (d, J = 8.6 Hz, 1H), 7.53 (dd, J = 7.9, 4.8 Hz, 1H), 7.35 (d, J = 8.3 Hz, 1H), 6.77 - 6.60 (m, 2H), 3.80 (ss, 6H), 3.14 (d, J = 4.3 Hz, 3H).	DMSO	373 (M+1)	1.49	Method D	100	Method AQ2 AT2

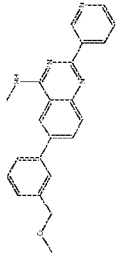
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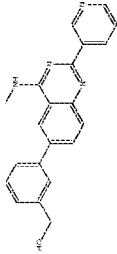
Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1261			348.3	¹ H NMR (300 MHz, DMSO) δ 9.63 (s, 1H), 8.77 (dd, J = 8.0, 1.9 Hz, 1H), 8.68 (d, J = 4.6 Hz, 1H), 8.59 (s, 2H), 8.21 - 8.09 (m, 1H), 8.02 - 7.88 (m, 1H), 7.84 (d, J = 8.7 Hz, 1H), 7.71 (s, 1H), 7.67 - 7.58 (m, 1H), 7.53 (dd, J = 7.6, 5.1 Hz, 1H), 3.18 (d, J = 4.4 Hz, 3H).	DMSO	349 (M+1)	1.61	Method D	100	Method AQ2
1262			313.4	¹ H NMR (300 MHz, DMSO) δ 9.65 (s, 1H), 9.09 (s, 1H), 8.78 (d, J = 8.0 Hz, 1H), 8.73 - 8.55 (m, 4H), 8.30 - 8.14 (m, 2H), 7.89 (d, J = 8.6 Hz, 1H), 7.63 - 7.47 (m, 2H), 3.19 (d, J = 4.3 Hz, 3H).	DMSO	314 (M+1)	1.70	Method C	96	Method AQ2

(continued)

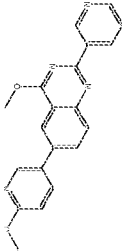
Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1263			343.4	¹ H NMR (300 MHz, DMSO) δ 9.64 (d, J = 1.3 Hz, 1 H), 8.84 – 8.74 (m, 1H), 8.68 (dd, J = 6.2, 1.7 Hz, 2H), 8.57 (d, J = 1.6 Hz, 2H), 8.16 (ddd, J = 14.4, 8.7, 2.2 Hz, 2H), 7.85 (d, J = 8.7 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 7.00 (d, J = 8.7 Hz, 1H), 3.93 (s, 3H), 3.18 (d, J 4.3 Hz, 3H).	DMSO	344.1 (M+1)	1.94	Method C	95	Method AQ2

(continued)

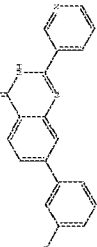
Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1264			356.4	¹ H NMR (300 MHz, DMSO) δ 10.76 (s, 1H), 9.75 (d, J = 1.6 Hz, 1H), 9.23 (d, J = 8.2 Hz, 1H), 9.05 - 8.86 (m, 2H), 8.41 - 8.24 (m, 2H), 8.24 - 8.05 (m, 2H), 7.96 (dd, J = 8.1, 5.2 Hz, 1H), 6.56 (d, J = 9.5 Hz, 1H), 4.81 (s, 2H), 3.34 (s, 3H), 3.29 (d, J = 4.3 Hz, 3H).	DMSO	357.1 (M+1)	2.13	Method C	100	Method AQ2

Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1265			342.4	¹ H NMR (300 MHz, DMSO-d ₆) 9.64 (s, 1H), 8.78 (dd, J = 8.0, 1.9 Hz, 1H), 8.67 (dd, J = 6.3, 4.7 Hz, 2H), 8.58 (s, 1H), 8.17 - 8.06 (m, 1H), 7.86 (d, J = 8.7 Hz, 1H), 7.79 (s, 1H), 7.72 (d, J = 7.6 Hz, 1H), 7.60 - 7.44 (m, 2H), 7.37 (d, J = 7.5 Hz, 1H), 5.31 (t, J = 5.4 Hz, 1H), 4.62 (d, J = 4.9 Hz, 2H), 3.18 (d, J = 4.4 Hz, 3H).	DMSO	343.1 (M+1)	1.78	Method C	99	Method AQ2

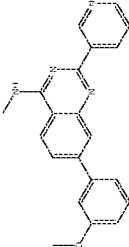
(continued)

Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1266			344.4	¹ H NMR (300 MHz, DMSO) δ 9.65 (d, J = 2.1 Hz, 1H), 8.92 - 8.59 (m, 3H), 8.59 - 8.47 (m, 1H), 8.40 - 8.11 (m, 2H), 8.04 (d, J = 8.5 Hz, 1H), 7.58 (dd, J = 7.9, 4.8 Hz, 1H), 6.95 (d, J = 8.6 Hz, 1H), 4.29 (s, 3H), 3.93 (s, 3H).	DMSO	345.1 (M+1)	2.32	Method C	100	Method AQ2

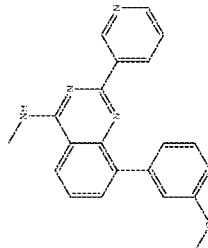
(continued)

Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Sol-vent	LCMS	Retention Time	LCMS Proto-col	Purity percent	Method for Coupling
1267			329.4	¹ H NMR (300 MHz, DMSO) δ 12.77 (s, 1H), 9.33 (d, J = 1.7 Hz, 1H), 8.77 (dd, J = 4.7, 1.5 Hz, 1H), 8.60 - 8.46 (m, 1H), 8.23 (d, J = 8.3 Hz, 1H), 8.04 (d, J = 1.5 Hz, 1H), 7.87 (dd, J = 8.3, 1.8 Hz, 1H), 7.67 - 7.53 (m, 1H), 7.52 - 7.29 (m, 3H), 7.04 (ddd, J = 7.7, 2.4, 1.5 Hz, 1H), 3.57 (s, 3H).	DMSO	330 (M+1)	1.78	Method C	100	Method AQ2

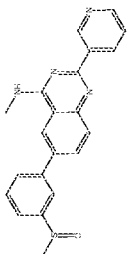
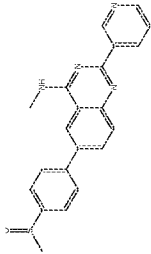
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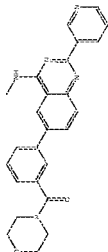
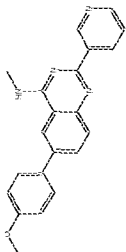
Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1268			342.4	¹ H NMR (300 MHz, DMSO) δ 9.65 (dd, J = 2.1, 0.8 Hz, 1H), 8.84 - 8.74 (m, 1H), 8.68 (dd, J = 4.7, 1.7 Hz, 1H), 8.52 (d, J = 4.4 Hz, 1H), 8.30 (d, J = 8.6 Hz, 1H), 8.03 (d, J = 1.7 Hz, 1H), 7.85 (dd, J = 8.5, 1.8 Hz, 1H), 7.54 (ddd, J = 8.0, 4.8, 0.8 Hz, 1H), 7.47 - 7.34 (m, 3H), 7.08 - 6.96 (m, 1H), 3.34 (s, 3H), 3.17 (d, J = 4.4 Hz, 3H).	DMSO	343.3 (M+1)	2.09	Method C		Method AQ2

(continued)

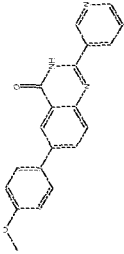
Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1269			342.4	¹ H NMR (300 MHz, DMSO) δ 9.53 (d, J = 2.0 Hz, 1H), 8.65 (dd, J = 5.9, 3.5 Hz, 2H), 8.51 (d, J = 4.2 Hz, 1H), 8.23 (d, J = 8.3 Hz, 1H), 7.84 (d, J = 7.3 Hz, 1H), 7.65 - 7.25 (m, 5H), 7.00 (dd, J = 8.1, 2.6 Hz, 1H), 3.33 (s, 3H), 3.17 (d, J = 4.2 Hz, 3H).	DMSO	343.3 (M+1)	2.16	Method C	100	Method AQ2

(continued)

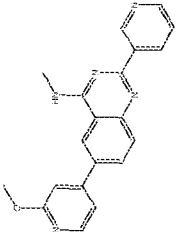
Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1270		2HCl	374.5	¹ H NMR (300 MHz, DMSO) δ 10.67 (s, 1H), 9.69 (s, 1H), 9.12 (d, J = 11.6 Hz, 2H), 8.97 (d, J = 5.0 Hz, 1H), 8.43 (d, J = 7.4 Hz, 1H), 8.31 (d, J = 8.6 Hz, 1H), 8.20 (s, 1H), 8.07 (s, 1H), 7.98 - 7.86 (m, 1H), 7.77 (s, 2H), 3.31 (d, J = 3.9 Hz, 3H), 2.90 (s, 3H).	DMSO	375.1 (M+1)	1.67	Method C	95	Method AQ2
1271			374.5	¹ H NMR (300 MHz, DMSO) δ 9.70 - 9.57 (m, 1H), 8.84 - 8.72 (m, 1H), 8.73 - 8.62 (m, 3H), 8.19 (dd, J = 8.7, 1.9 Hz, 1H), 8.06 (d, J = 8.4 Hz, 2H), 7.86 (dd, J = 10.6, 8.5 Hz, 3H), 7.60 - 7.49 (m, 1H), 3.19 (d, J = 4.4 Hz, 2H), 2.81 (s, 3H).	DMSO	375.1 (M+1)	1.64	Method C	95	Method AQ2

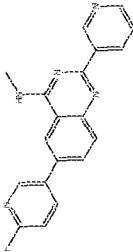
Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1272		2HCl	425.5	¹ H NMR (300 MHz, DMSO) δ 10.54 (s, 1H), 9.70 (d, J = 1.7 Hz, 1H), 9.12 (d, J = 8.1 Hz, 1H), 8.98 (dd, J = 5.5, 4.0 Hz, 2H), 8.37 (dd, J = 26.8, 7.9 Hz, 2H), 8.09 - 7.84 (m, 3H), 7.63 (t, J = 7.7 Hz, 1H), 7.49 (d, J = 7.6 Hz, 1H), 3.67 (bs, 1H), 3.31 (d, J = 4.3 Hz, 3H).	DMSO	426.2 (M+1)	1.72	Method C	95	Method AQ2
1273		HCl	342.4	¹ H NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.77 (d, J = 7.9 Hz, 1H), 8.71 - 8.65 (m, 1H), 8.60 (s, 1H), 8.53 (s, 1H), 8.10 (s, 1H), 8.10 (d, J = 8.7 Hz, 1H), 7.87 - 7.77 (m, 3H), 7.59 - 7.49 (m, 1H), 7.10 (d, J = 8.7 Hz, 2H), 3.83 (d, J = 0.9 Hz, 3H), 3.19 (s, 3H).	DMSO	343.1 (M+1)	2.06	Method C	100	Method AQ2

(continued)

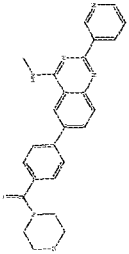
Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Sol-vent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1274			329.4	¹ H NMR (300 MHz, DMSO) δ 9.34 (d, J = 1.5 Hz, 1H), 8.74 (dd, J = 4.8, 1.6 Hz, 1H), 8.53 (d, J = 8.0 Hz, 1H), 8.32 (d, J = 2.1 Hz, 1H), 8.10 (dd, J = 8.5, 2.3 Hz, 1H), 7.79 (d, J = 8.6 Hz, 1H), 7.76 (s, 1H), 7.73 (s, 1H), 7.58 (dd, J = 8.0, 4.8 Hz, 1H), 7.07 (d, J = 8.8 Hz, 2H), 3.82 (s, 3H).	DMSO	330.0 (M+1)	1.75	Method C	100	Method AQ2

(continued)

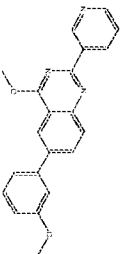
Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1275			343.4	¹ H NMR (300 MHz, DMSO) δ 9.65 (d, J = 1.5 Hz, 1H), 8.84 - 8.65 (m, 4H), 8.31 (d, J = 5.4 Hz, 1H), 8.24 (dd, J = 8.7, 1.9 Hz, 1H), 7.87 (d, J = 8.7 Hz, 1H), 7.59 - 7.46 (m, 2H), 7.33 (d, J = 0.9 Hz, 1H), 3.93 (s, 3H), 3.20 (d, J = 4.4 Hz, 3H).	DMSO	344.1 (M+1)	1.95	Method C	95	Method AQ2

Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1276			331.3	¹ H NMR (300 MHz, DMSO) δ 9.63 (d, J = 1.4 Hz, 1H), 8.81 - 8.74 (m, 1H), 8.73 (d, J 2.3 Hz, 1H), 8.68 (dd, J = 4.7, 1.7 Hz, 1H), 8.66 - 8.56 (m, 2H), 8.44 (td, J = 8.2, 2.6 Hz, 1H), 8.18 (dd, J = 8.7, 2.0 Hz, 1H), 7.87 (d, J = 8.7 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 7.38 (dd, J = 8.5, 2.9 Hz, 1H), 3.18 (d, J = 4.4 Hz, 3H).	DMSO	332.1 (M+1)	1.89	Method C	100	Method AQ2

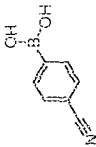
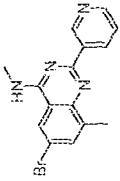
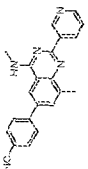
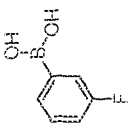
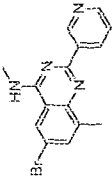
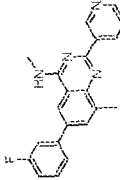
(continued)

Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1277			425.5	¹ H NMR (300 MHz, DMSO) δ 9.63 (d, J = 1.4 Hz, 1H), 8.77 (d, J = 8.0 Hz, 1H), 8.70 - 8.59 (m, 3H), 8.15 (dd, J = 8.7, 1.6 Hz, 1H), 7.92 (d, J = 8.2 Hz, 2H), 7.85 (d, J = 8.7 Hz, 1H), 7.62 - 7.48 (m, 3H), 3.62 (s, J = 54.7 Hz, 8H), 3.17 (d, J = 4.2 Hz, 3H).	DMSO	426.2 (M+1)	1.69	Method C	100	Method AQ2

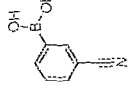
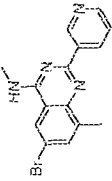
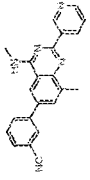
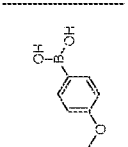
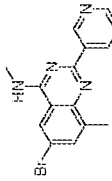
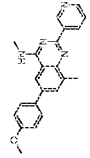
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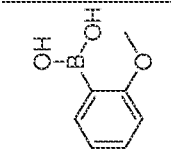
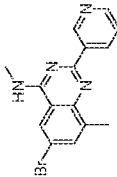
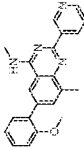
Number	Product	Salt	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1278			343.4	¹ H NMR (300 MHz, DMSO) δ 9.67 (d, J = 1.5 Hz, 1H), 8.86 - 8.77 (m, 1H), 8.74 (dd, J = 4.7, 1.6 Hz, 1H), 8.37 - 8.25 (m, 2H), 8.06 (d, J = 8.7 Hz, 1H), 7.60 (dd, J = 7.9, 4.1 Hz, 1H), 7.50 Hz, 1H), 7.30 (m, 3H), 7.02 (dd, J = 7.9, 1.4 Hz, 1H), 4.31 (s, 3H), 3.87 (s, 3H).	DMSO	344.2 (M+1)	2.47	Method C	100	Method AQ2

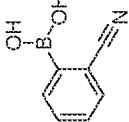
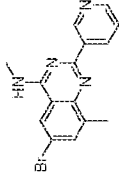
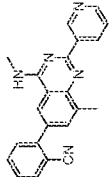
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1279				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.69 (d, J = 1.8 Hz, 1H), 9.4 - 9.33 (m, 1H), 9.03 - 8.93 (m, 2H), 8.64-8.59 (m, 1H), 8.18 - 8.14 (m, 1H), 8.13-8.05 (m, 3H), 8.04 - 7.97 (m, 2H), 3.21 (d, J = 4.4 Hz, 3H), 2.77 (s, 3H).	DMSO	>98	AQ3		
1280				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.71 - 9.65 (m, 1H), 9.44 - 9.36 (m, 1H), 9.06-8.94 (m, 2H), 8.59 - 8.54 (m, 1H), 8.19-8.10 (m, 2H), 7.79 - 7.71 (m, 2H), 7.61 - 7.54 (m, 1H), 7.29 - 7.22 (m, 1H), 3.21 (d, J = 3.9 Hz, 3H), 2.76 (s, 3H).	DMSO	>98	AQ3		

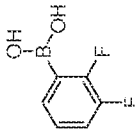
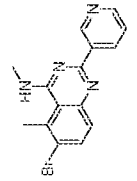
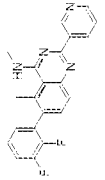
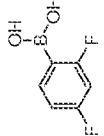
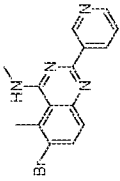
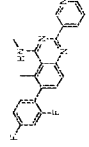
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1281				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.72 - 9.65 (m, 1H), 9.43 - 9.36 (m, 1H), 9.04 - 8.91 (m, 2H), 8.63 - 8.55 (m, 1H), 8.36 - 8.31 (m, 1H), 8.26 - 8.20 (m, 1H), 8.20 - 8.11 (m, 2H), 7.91 - 7.85 (m, 1H), 7.78 - 7.70 (m, 1H), 3.22 (d, J = 4.2 Hz, 3H), 2.76 (s, 3H).	DMSO	>98	AQ3		
1282				2HCl	¹ H NMR (400 MHz, DMSO) δ 9.70 - 9.65 (m, 1H), 9.37 (d, J = 8.2 Hz, 1H), 9.03 - 8.90 (m, 2H), 8.49 - 8.42 (m, 1H), 8.12 (dd, J = 8.1, 5.6 Hz, 1H), 8.08 - 8.03 (m, 1H), 7.88 - 7.78 (m, 2H), 7.15 - 7.04 (m, 2H), 3.84 (s, 3H), 3.21 (d, J = 3.9 Hz, 3H), 2.76 (s, 3H).	DMSO	>98	AQ3		

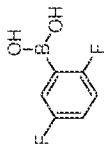
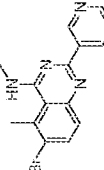
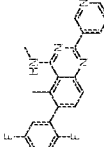
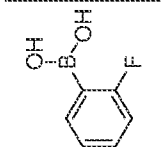
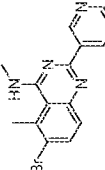
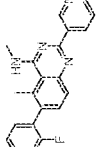
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1283				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.73 - 9.67 (m, 1H), 9.36 (d, J= 7.4 Hz, 1H), 9.02 - 8.96 (m, 1H), 8.82 (s, 1H), 8.26-8.19(m, 1H), 8.15 - 8.07 (m, 1H), 7.89-7.83(m, 1H), 7.47 - 7.38 (m, 2H), 7.21-7.15(m, 1H), 7.14 - 7.06 (m, 1H), 3.81 (s, 3H), 3.19(d, J=4.5 Hz, 3H), 2.77 (s, 3H).	DMSO	>98	AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1284				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.72 (d, J = 1.8 Hz, 1H), 9.34-9.28 (m, 1H), 8.95 (dd, J = 5.4, 1.4 Hz, 1H), 8.88 - 8.78 (m, 1H), 8.38 (d, J = 1.5 Hz, 1H), 8.07 - 8.00 (m, 2H), 7.95-7.90 (m, 1H), 7.90 - 7.83 (m, 1H), 7.76 (dd, J = 7.8, 0.7 Hz, 1H), 7.68 - 7.63 (m, 1H), 3.19 (d, J = 4.5 Hz, 3H), 2.77 (s, 3H).	DMSO	>98	AQ3		

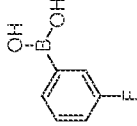
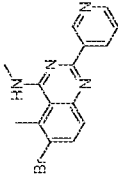
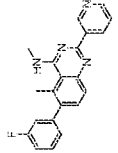
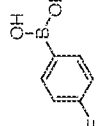
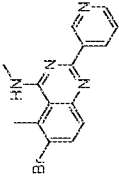
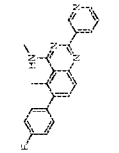
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1285					¹ H NMR (400 MHz, DMSO) δ 9.63 (dd, $J = 2.1, 0.8$ Hz, 1H), 8.80 - 8.75 (m, 1H), 8.69 (dd, $J = 3.7, 1.8$ Hz, 1H), 7.72 (d, $J = 8.5$ Hz, 1H), 7.61 (d, $J = 8.5$ Hz, 1H), 7.58 - 7.49 (m, 2H), 7.46 - 7.40 (m, 1H), 7.39 - 7.32 (m, 1H), 7.28 - 7.18 (m, 1H), 3.18 (d, $J = 4.4$ Hz, 3H), 2.64 (s, 3H).	DMSO	>98	AQ4		
1286				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.67 - 9.60 (m, 1H), 9.06 - 8.88 (m, 3H), 8.12 (d, $J = 8.8$ Hz, 1H), 7.91 - 7.79 (m, 2H), 7.53 - 7.40 (m, 2H), 7.34 - 7.24 (m, 1H), 3.34 (d, $J = 4.6$ Hz, 3H), 2.66 (s, 3H).	DMSO	>98	AQ4		

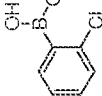
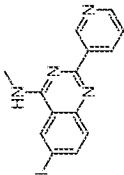
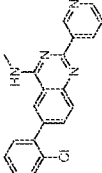
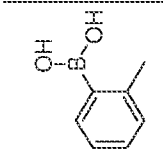
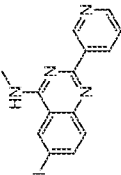
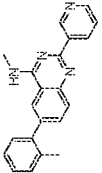
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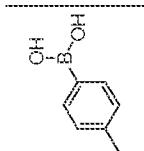
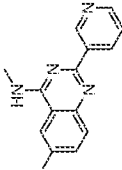
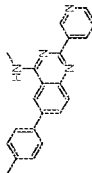
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1287				HCl	¹ H NMR (400 MHz, DMSO) δ 9.62 - 9.57 (m, 1H), 8.94 - 8.86 (m, 2H), 8.62 (s, 1H), 7.95 (d, J = 8.5 Hz, 1H), 7.86 - 7.75 (m, 2H), 7.52 - 7.36 (m, 2H), 7.35 - 7.28 (m, 1H), 3.31 (d, J = 4.5 Hz, 3H), 2.67 (s, 3H).	DMSO	>98	AQ4		
1288				HCl	¹ H NMR (400 MHz, DMSO) δ 9.63 - 9.58 (m, 1H), 8.95 - 8.86 (m, 2H), 8.73 (s, 1H), 8.01 (d, J = 8.5 Hz, 1H), 7.87 - 7.75 (m, 2H), 7.61 - 7.50 (m, 1H), 7.45 - 7.33 (m, 3H), 3.32 (d, J = 4.5 Hz, 3H), 2.66 (s, 3H).	DMSO	>98	AQ4		

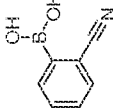
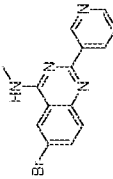
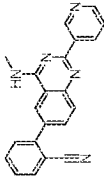
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1289				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.64 - 9.60 (m, 1H), 9.00 - 8.86 (m, 3H), 8.06 (d, J = 8.5 Hz, 1H), 7.91 - 7.79 (m, 2H), 7.65 - 7.55 (m, 1H), 7.37 - 7.29 (m, 1H), 7.29 - 7.22 (m, 2H), 3.35 (d, J = 4.6 Hz, 3H), 2.72 (s, 3H).	DMSO	>98	AQ4		
1290				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.69 - 9.64 (m, 1H), 9.14 - 8.99 (m, 2H), 8.96 (dd, J = 5.0, 1.5 Hz, 1H), 8.15 (d, J = 8.6 Hz, 1H), 7.90 - 7.81 (m, 2H), 7.50 - 7.34 (m, 4H), 3.35 (d, J = 4.6 Hz, 3H), 2.72 (s, 3H).	DMSO	>98	AQ4		

(continued)

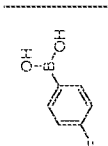
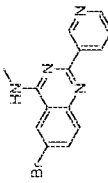
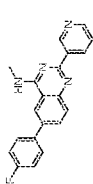
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1291				HCl	¹ H NMR (300 MHz, CDCl ₃) δ 9.63 (s, 2H), 8.96 (d, J = 8.2 Hz, 1H), 8.90 (d, J=5.0 Hz, 1H), 8.50 (s, 1H), 8.16 - 7.99 (m, 2H), 7.82 (dd, J= 8.0, 4.9 Hz, 1H), 7.71-7.62 (m, 1H), 7.61 - 7.47 (m, 3H), 3.26 (d, J=4.5 Hz, 3H).	DMSO	>98	G2/AQ3		
1292				2 HCl	¹ H NMR (300 MHz, CDCl ₃) δ 10.27 (s, 1H), 9.69 (s, 1H), 9.09 (d, J = 8.1 Hz, 1H), 8.97 (d, J=5.0 Hz, 1H), 8.56 (s, 1H), 8.32 (d, J=8.6 Hz, 1H), 8.04 (d, J=8.7 Hz, 1H), 7.89 (dd, J=8.0, 5.1 Hz, 1H), 7.45 - 7.26 (m, 4H), 3.29 (d, J = 4.2 Hz, 3H), 2.32 (s, 3H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1293				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.40 (s, 1H), 9.73 (s, 1H), 9.15 (d, J = 7.9 Hz, 1H), 8.98 (d, J = 5.1 Hz, 1H), 8.67 (s, 1H), 8.33 (d, J = 8.6 Hz, 1H), 8.18 (d, J = 8.7 Hz, 1H), 7.91 (dd, J = 7.9, 5.2 Hz, 1H), 7.45 (m, 2H), 7.20 (d, J = 8.2 Hz, 1H), 7.12 (t, J = 7.5 Hz, 1H), 3.82 (s, 3H), 3.29 (d, J = 4.1 Hz, 3H).	DMSO	>98	G2/AQ3		

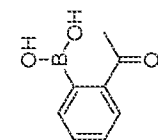
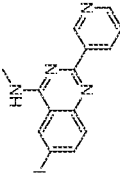
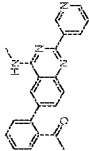
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1294					¹ H NMR (300 MHz, CDCl ₃) δ 9.66 (s, 1H), 8.80 (d, <i>J</i> = 7.9 Hz, 1H), 8.74 -8.67 (m, 1H), 8.60 (m, 1H), 8.49 (s, 1H), 8.00 (dd, <i>J</i> = 18.5, 10.4 Hz, 2H), 7.96 - 7.81 (m, 2H), 7.76 (d, <i>J</i> = 7.6 Hz, 1H), 7.65 (t, <i>J</i> = 7.6 Hz, 1H), 7.56 (dd, <i>J</i> = 7.6, 5.0 Hz, 1H), 3.18 (d, <i>J</i> = 4.2 Hz, 3H).	DMSO	>98	G2/AQ3		

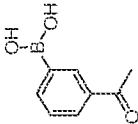
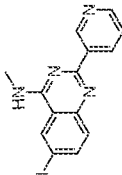
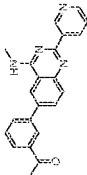
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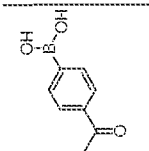
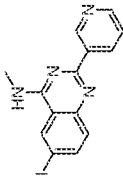
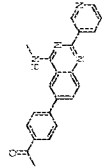
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1296				2 HCl	¹ H NMR (300 MHz, CDCl ₃) δ 9.63 (m, 2H), 8.93 (d, J = 7.7 Hz, 1H), 8.88 (d, J = 4.1 Hz, 1H), 8.76 (s, 1H), 8.29 (d, J = 8.4 Hz, 1H), 8.07 (d, J = 8.8 Hz, 1H), 7.94 (dd, J = 8.6, 5.5 Hz, 2H), 7.82 - 7.73 (m, 1H), 7.41 (t, J = 8.7 Hz, 2H), 3.28 (d, J = 4.3 Hz, 3H).	DMSO	>98	G2/AQ3		

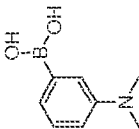
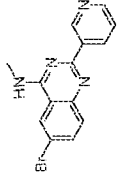
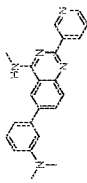
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
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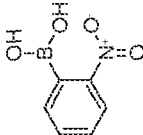
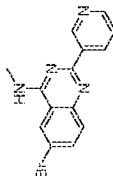
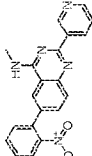
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1298				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.30 (s, 1H), 9.70 (d, 1H), 9.11 (d, 1H), 9.11 (d, 1H), 8.83 (m, 1H), 8.56 (s, 1H), 8.26 (d, $J = 8.6$ Hz, 1H), 8.01-7.80 (m, 3H), 7.78-7.64 (m, 1H), 7.60 (t, $J = 7.2$ Hz, 1H), 7.54 (d, $J = 7.2$ Hz, 1H), 3.29 (d, $J = 4.4$ Hz, 3H), 2.44 (s, 3H).	DMSO	>98	G2/AQ3		

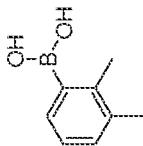
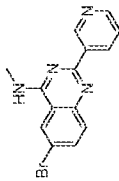
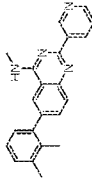
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1299					¹ H NMR (300 MHz, DMSO) δ 10.50 (s, 1H), 9.66 (s, 1H), 9.09 (d, J = 8.2 Hz, 1H), 9.01 (s, 1H), 8.94 (d, J = 5.0 Hz, 1H), 8.40 (d, J = 8.7 Hz, 1H), 8.28 (d, J = 8.8 Hz, 1H), 8.13-8.00 (m, 4H), 7.88 (dd, J = 8.0, 5.2 Hz, 1H), 3.30 (d, J = 4.1 Hz, 3H), 2.62 (s, 3H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1300				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.49 (s, 1H), 9.66 (s, 1H), 9.09 (d, J = 7.9 Hz, 1H), 9.01 (s, 1H), 8.95 (d, J = 5.0 Hz, 1H), 8.40 (d, J = 8.6 Hz, 1H), 8.29 (d, J = 8.7 Hz, 1H), 8.15-7.94 (m, 4H), 7.88 (dd, J = 7.9, 5.2 Hz, 1H), 3.31 (d, J = 4.1 Hz, 3H), 2.63 (s, 3H).	DMSO	>98	G2/AQ3		

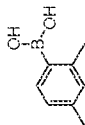
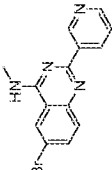
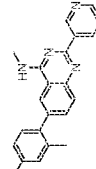
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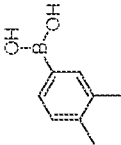
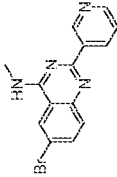
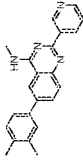
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1301				3 HCl	¹ H NMR (300 MHz, DMSO) δ 10.97 (s, 1H), 9.71 (d, J = 2.0 Hz, 1H), 9.29 (s, 1H), 9.15 (d, J = 8.0 Hz, 1H), 8.98 (dd, J = 5.0, 1.3 Hz, 1H), 8.49 - 8.38 (m, 1H), 8.34 (d, J = 8.8 Hz, 1H), 8.21 (s, 1H), 7.92 (dd, J = 8.1, 5.1 Hz, 1H), 7.84 (s, 1H), 7.69 - 7.52 (m, 2H), 3.31 (d, J = 4.3 Hz, 3H), 3.23 (s, 6H).	DMSO	>98	G2/AQ3		

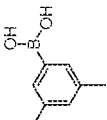
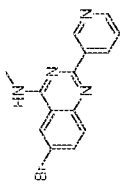
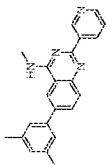
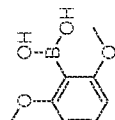
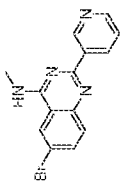
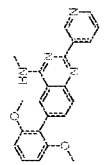
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1302				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.33 (s, 1H), 9.73 (d, J = 1.7 Hz, 1H), 9.20 (d, J = 8.0 Hz, 1H), 9.00 (d, J = 4.2 Hz, 1H), 8.68 (s, 1H), 8.32 (d, J = 8.6 Hz, 1H), 8.16 (d, J = 7.9 Hz, 1H), 8.01-7.93 (m, 2H), 7.90 (t, J = 7.2 Hz, 1H), 7.74 (dd, J = 15.4, 7.7 Hz, 2H), 3.24 (t, J = 18.2 Hz, 3H).	DMSO	>98	G2/AQ3		

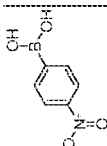
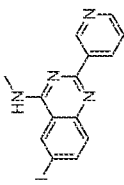
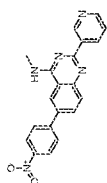
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1303				HCl	¹ H NMR (300 MHz, DMSO-d ₆) δ 10.46 (s, 1H), 9.76 (d, 1H), 9.21 (d, 1H), 9.21 (d, 1H), 9.00 (dd, 1H), 9.00 (dd, 1H), 8.58 (d, 1H), 8.41 (d, 1H), 8.41 (d, 1H), 8.08 - 7.86 (m, 2H), 7.36-7.09 (m, 3H), 3.29 (d, 3H), 2.33 (s, 3H), 2.17 (s, 3H).	DMSO	>98	G2/AQ3		

(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1304					¹ H NMR (300 MHz, DMSO) δ 9.65 (d, J = 2.1 Hz, 1H), 8.79 (dt, J = 8.0, 1.9 Hz, 1H), 8.69 (dd, J = 4.7, 1.6 Hz, 1H), 8.46 (d, J = 4.5 Hz, 1H), 8.20 (d, J = 1.7 Hz, 1H), 7.83 (d, J = 8.5 Hz, 1H), 7.75 (dd, J = 8.5, 1.6 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 7.28 Hz, 1H), 7.02 (m, 3H), 3.33 (s, 6H), 3.15 (d, J = 4.4 Hz, 3H), 2.51 (dt, J = 3.6, 1.7 Hz, 1H), 2.35 (s, 3H), 2.27 (s, 3H).	DMSO	>98	G2/AQ3		

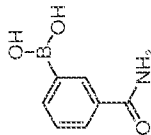
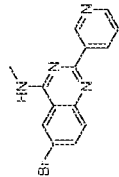
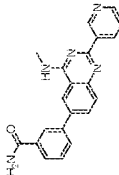
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1305				HCl	¹ H NMR (300 MHz, DMSO) δ 10.48 (s, 1H), 9.70 (s, 1H), 9.11 (d, J = 7.8 Hz, 1H), 8.97 (d, J = 4.7 Hz, 1H), 8.90 (s, 1H), 8.36 (d, J = 8.6 Hz, 1H), 8.29 (d, J = 8.7 Hz, 1H), 7.98 - 7.80 (m, 1H), 7.74 (s, 1H), 7.65 (d, J = 7.5 Hz, 1H), 7.30 (d, J = 7.8 Hz, 1H), 3.32 (d, J = 4.0 Hz, 3H), 2.33 (d, J = 10.0 Hz, 3H), 2.29 (s, 3H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1306				HCl	¹ H NMR (300 MHz, DMSO) δ 10.45 - 10.35 (m, 1H), 9.69 (s, 1H), 9.09 (s, 1H), 8.99 (s, 1H), 8.88 (s, 1H), 8.37 (d, J = 8.6 Hz, 1H), 8.28 (d, J = 8.7 Hz, 1H), 7.94 (s, 1H), 7.54 (s, 1H), 7.11 (s, 1H), 3.34 (d, J = 4.3 Hz, 3H), 2.40 (s, 3H).	DMSO	>98	G2/AQ3		
1307				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.28 (s, 1H), 9.84 - 9.66 (m, 1H), 9.17 (d, J = 8.1 Hz, 1H), 8.99 (d, J = 5.0 Hz, 1H), 8.46 (s, 1H), 8.36 (d, J = 8.6 Hz, 1H), 7.98 - 7.80 (m, 2H), 7.41 (t, J = 8.4 Hz, 1H), 6.83 (d, J = 8.4 Hz, 2H), 3.71 (s, 6H), 3.29 (d, J = 4.2 Hz, 3H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1308				HCl	¹ H NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.96-8.91 (m, 1H), 8.89-8.83 (m, 2H), 8.41 (d, J=9.1 Hz, 2H), 8.38-8.32 (m, 1H), 8.16 (d, J=9.0 Hz, 2H), 8.02 (d, J=9.1 Hz, 1H), 7.81-7.73 (m, 1H), 3.27 (d, J=4.3 Hz, 3H).	DMSO	>98	G2/AQ3		

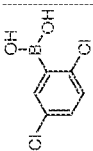
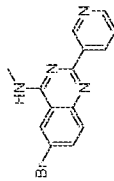
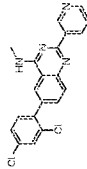
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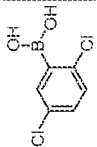
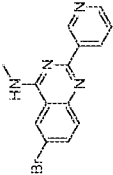
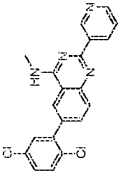
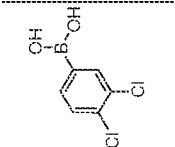
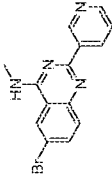
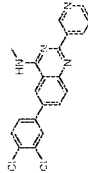
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1310				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.63 (s, 1H), 9.69 (s, 1H), 9.16 (s, 1H), 9.10 (d, 1H), 9.10 (d, 1H), 8.97 (d, 1H), 8.97 (d, 1H), 8.52 (s, 1H), 8.45 (d, 1H), 8.7 Hz, 1H), 8.29 (d, 1H), 8.07 Hz, 2H), 8.07 (d, 1H), 8.01 - 7.84 (m, 2H), 7.63 (t, 1H), 7.54 Hz, 1H), 3.31 (s, 1H), 3.31 (d, 1H), 4.3 Hz, 3H).	DMSO	>98	G2/AQ3		

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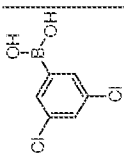
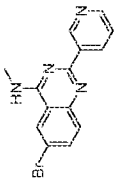
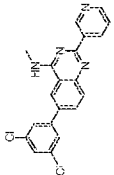
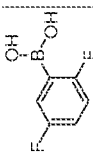
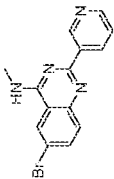
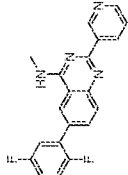
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1311				HCl	¹ H NMR (300 MHz, DMSO) δ 10.20 (s, 1H), 9.65 (s, 1H), 9.02 (s, 1H), 8.95 (s, 1H), 8.43 (d, J = 8.7 Hz, 1H), 8.26-8.14 (m, 1H), 8.04 (dd, J = 18.8, 8.1 Hz, 4H), 7.88 (s, 1H), 7.47 (s, 1H), 3.32 (s, 3H), 3.17 (d, J = 1.8 Hz, 9H).	DMSO	>98	G2/AQ3		
1312				HCl	¹ H NMR (300 MHz, DMSO) δ 9.79 (s, 1H), 9.66 (d, J = 1.7 Hz, 1H), 9.02 (d, J = 8.1 Hz, 1H), 8.96-8.86 (m, 1H), 8.57 (s, 1H), 8.18 (d, J = 8.7 Hz, 1H), 8.05 (dd, J = 8.7, 1.4 Hz, 1H), 7.82 (dd, J = 8.1, 5.0 Hz, 1H), 7.79 Hz, 7.72 (m, 1H), 7.64-7.45 (m, 2H), 3.25 (d, J = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

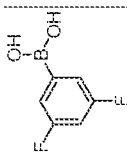
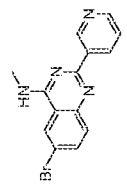
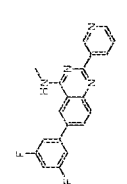
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1313				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.01 (s, 1H), 9.65 (s, 1H), 9.06 (d, <i>J</i> = 7.8 Hz, 1H), 8.95 (d, <i>J</i> = 4.8 Hz, 1H), 8.59 (s, 1H), 8.21 (d, <i>J</i> = 8.6 Hz, 1H), 8.07 (d, <i>J</i> = 9.1 Hz, 1H), 7.95-7.77 (m, 2H), 7.66-7.52 (m, 2H), 3.27 (d, <i>J</i> = 4.3 Hz, 3H), 3.17 (d, <i>J</i> = 0.7 Hz, 1H).	DMSO	>98	G2/AQ3		

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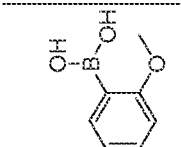
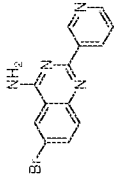
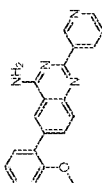
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1314				HCl	¹ H NMR (300 MHz, DMSO) δ 10.10 (s, 1H), 9.69 (s, 1H), 9.13 (d, J = 8.0 Hz, 1H), 8.98 (d, J = 5.1 Hz, 1H), 8.62 (s, 1H), 8.27 (d, J = 8.6 Hz, 1H), 8.13 (d, J = 8.3 Hz, 1H), 8.02-7.83 (m, 1H), 7.77 - 7.65 (m, 2H), 7.60 (d, J = 8.5 Hz, 1H), 3.29 (d, J = 3.9 Hz, 3H).	DMSO	>98	G2/AQ3		
1315				HCl	¹ H NMR (300 MHz, DMSO) δ 9.62 (s, 1H), 9.02-8.93 (m, 1H), 8.89 (d, J = 4.5 Hz, 1H), 8.80 (s, 1H), 8.35 (d, J = 8.7 Hz, 1H), 8.18 (s, 1H), 8.04 (d, J = 8.7 Hz, 1H), 7.96-7.75 (m, 2H), 3.29 (d, J = 4.2 Hz, 3H).	DMSO	>98	G2/AQ3		

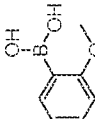
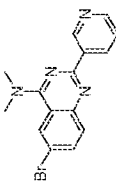
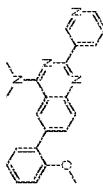
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1316				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.36 (s, 1H), 9.67 (s, 1H), 9.13 (d, J = 8.2 Hz, 1H), 9.01-8.82 (m, 2H), 8.40 (d, J = 9.1 Hz, 1H), 8.26 (d, J = 8.7 Hz, 1H), 7.97 (s, 2H), 7.90 (dd, J = 8.0, 5.2 Hz, 1H), 7.65 (s, 1H), 3.31 (d, J = 4.3 Hz, 3H).	DMSO	>98	G2/AQ3		
1317					¹ H NMR (300 MHz, DMSO) δ 9.65 (d, J = 1.9 Hz, 1H), 8.89-8.75 (m, 1H), 8.74 - 8.67 (m, 1H), 8.67-8.56 (m, 1H), 8.48 (s, 1H), 8.06 - 7.94 (m, 1H), 7.87 (d, J = 8.7 Hz, 1H), 7.68 - 7.51 (m, 2H), 7.46 (td, J = 9.6, 4.7 Hz, 1H), 7.39 - 7.21 (m, 1H), 3.18 (d, J = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

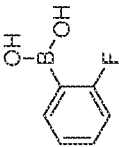
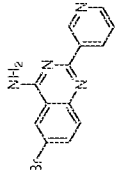
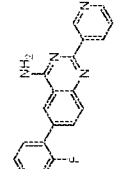
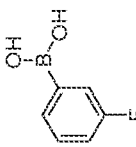
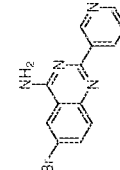
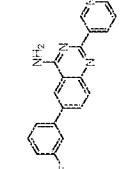
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1318				HCl	¹ H NMR (300 MHz, DMSO- <i>d</i> ₆) δ 10.57 (s, 1H), 9.69 (d, J = 2.0 Hz, 1H), 9.15 (d, J = 8.0 Hz, 1H), 9.02 (s, 1H), 8.97 (d, J = 4.4 Hz, 1H), 8.40 (d, J = 8.8 Hz, 1H), 8.29 (d, J = 8.7 Hz, 1H), 7.91 (dd, J = 8.0, 5.2 Hz, 1H), 7.77 - 7.63 (m, 2H), 7.37 - 7.18 (m, 1H), 3.29 (d, J = 4.3 Hz, 3H).	DMSO	>98	G2/AQ3		

(continued)

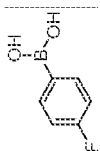
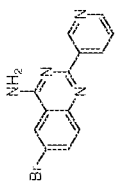
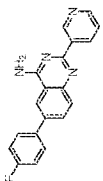
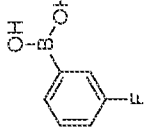
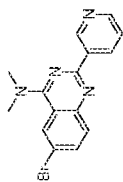
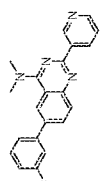
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1319					¹ H NMR (300 MHz, CDCl ₃) δ 9.58 (d, J = 2.0 Hz, 1H), 8.78-8.60 (m, 2H), 8.35 (d, J = 1.7 Hz, 1H), 8.11 - 7.84 (m, 3H), 7.78 (d, J = 8.6 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 7.48 - 7.34 (m, 2H), 7.17 (d, J = 8.1 Hz, 1H), 7.10 (t, J = 7.4 Hz, 1H), 3.81 (s, 3H).	DMSO	>98	G2/AQ3		

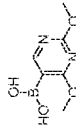
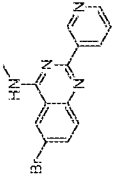
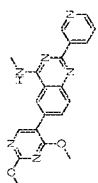
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1320					¹ H NMR (300 MHz, CDCl ₃) δ 9.62 (d, <i>J</i> = 2.1 Hz, 1H), 8.82-8.72 (m, 1H), 8.72 - 8.66 (m, 1H), 8.30 (s, 1H), 7.93 (dd, <i>J</i> = 8.7, 1.2 Hz, 1H), 7.86 (d, <i>J</i> = 8.6 Hz, 1H), 7.54 (dd, <i>J</i> = 7.9, 4.8 Hz, 1H), 7.50 - 7.33 (m, 2H), 7.18 (d, <i>J</i> = 8.3 Hz, 1H), 7.09 (t, <i>J</i> = 7.5 Hz, 1H), 3.82 (s, 3H), 3.46 (s, 6H).	DMSO	>98	G2/AQ3		

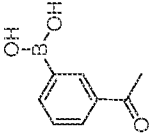
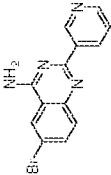
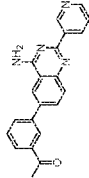
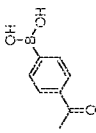
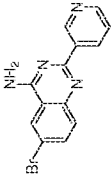
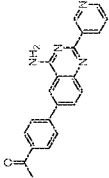
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1321					¹ H NMR (300 MHz, CDCl ₃) δ 9.59 (d, J = 2.1 Hz, 1H), 8.76-8.66 (m, 2H), 8.50 (s, 2H), 8.06 (brs, 1H), 7.99 (dt, J = 8.6, 1.7 Hz, 1H), 7.86 (d, J = 8.7 Hz, 1H), 7.75-7.63 (m, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 7.52-7.44 (m, 1H), 7.44-7.32 (m, 2H).	DMSO	>98	G2/AQ3		
1322					¹ H NMR (300 MHz, CDCl ₃) δ 9.59 (d, J = 2.1 Hz, 1H), 8.79-8.62 (m, 3H), 8.21 (dd, J = 8.8, 1.9 Hz, 1H), 8.10 (brs, 2H), 7.86 (d, J = 8.7 Hz, 1H), 7.80-7.69 (m, 2H), 7.68-7.49 (m, 2H), 7.32-7.20 (m, 1H).	DMSO	>98	G2/AQ3		

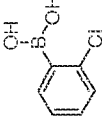
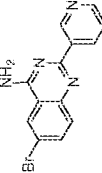
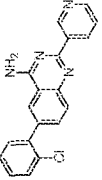
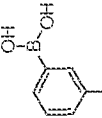
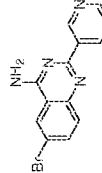
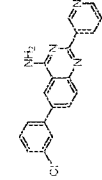
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1323					¹ H NMR (300 MHz, CDCl ₃) δ 9.58 (d, J= 2.0 Hz, 1H), 8.80-8.64 (m, 2H), 8.62 (d, J= 1.8 Hz, 1H), 8.14 (dd, J= 8.7, 1.9 Hz, 1H), 8.07 (brs, 2H), 7.98-7.87 (m, 2H), 7.85 (d, J= 8.7 Hz, 1H), 7.54 (dd, J= 7.9, 4.8 Hz, 1H), 7.38 (t, J= 8.8 Hz, 2H).	DMSO	>98	G2/AQ3		
1324					¹ H NMR (300 MHz, CDCl ₃) δ 9.65 - 9.57 (m, 1H), 8.82 - 8.70 (m, 1H), 8.70-8.63 (m, 1H), 8.36 (d, J= 1.8 Hz, 1H), 8.11 (dd, J= 8.7, 1.7 Hz, 1H), 7.89 (d, J= 8.7 Hz, 1H), 7.72 - 7.60 (m, 2H), 7.60-7.47 (m, 2H), 7.32 - 7.18 (m, 1H), 3.49 (s, 6H).	DMSO	>98	G2/AQ3		

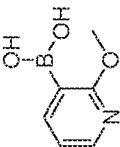
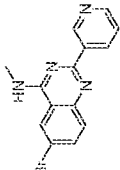
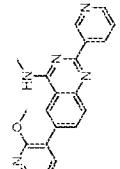
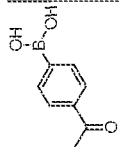
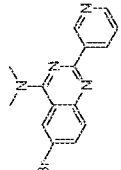
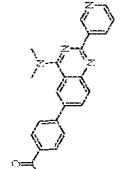
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1325					¹ H NMR (400 MHz, DMSO) δ 9.64 (d, J = 1.4 Hz, 1H), 8.78 (dt, J = 8.3, 1.8 Hz, 1H), 8.69 (dd, J = 4.7, 1.8 Hz, 1H), 8.61 - 8.46 (m, 2H), 8.40 (d, J = 1.6 Hz, 1H), 7.99 (dd, J = 9.0, 1.8 Hz, 1H), 7.83 (d, J = 9.0 Hz, 1H), 7.54 (dd, J = 8.4, 4.5 Hz, 1H), 3.99 (d, J = 8.9 Hz, 6H), 3.17 (d, J = 4.6 Hz, 3H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1326					¹ H NMR (300 MHz, DMSO) δ 9.64 - 9.55 (m, 1H), 8.81 - 8.62 (m, 3H), 8.40 (s, 1H), 8.23 (dd, J = 8.8, 1.4 Hz, 1H), 8.13 (d, J = 7.7 Hz, 1H), 8.00 (d, J = 7.7 Hz, 2H), 7.89 (d, J = 8.7 Hz, 1H), 7.69 (t, J = 7.7 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 2.71 (s, 3H).	DMSO	>98	G2/AQ3		
1327					¹ H NMR (300 MHz, DMSO) δ 9.58 (d, J = 2.1 Hz, 1H), 8.76 - 8.65 (m, 3H), 8.24 (dd, J = 8.7, 1.8 Hz, 1H), 8.19 - 7.93 (m, 6H), 7.88 (d, J = 8.8 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 2.65 (s, 3H).	DMSO	>98	G2/AQ3		

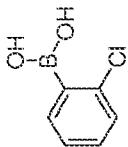
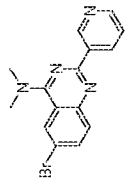
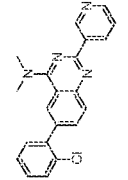
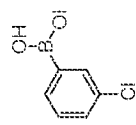
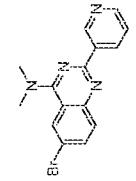
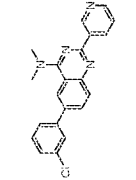
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1328					¹ H NMR (300 MHz, DMSO) δ 9.60 (d, J = 1.8 Hz, 1H), 8.73 (dt, J = 8.0, 1.8 Hz, 1H), 8.69 (dd, J = 4.8, 1.6 Hz, 1H), 8.37 (s, 1H), 7.99 (brs, 2H), 7.92-7.81 (m, 2H), 7.67-7.60 (m, 1H), 7.60-7.52 (m, 2H), 7.52-7.43 (m, 2H).	DMSO	>98	G2/AQ3		
1329					¹ H NMR (300 MHz, DMSO) δ 9.60 (d, J = 2.0 Hz, 1H), 8.77-8.65 (m, 3H), 8.19 (dd, J = 8.7, 1.8 Hz, 1H), 8.09 (brs, 2H), 7.99-7.94 (m, 1H), 7.90-7.79 (m, 2H), 7.63-7.51 (m, 2H), 7.48 (d, J = 8.1 Hz, 1H).	DMSO	>98	G2/AQ3		

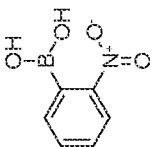
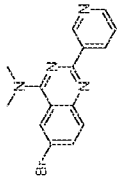
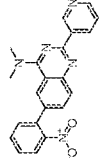
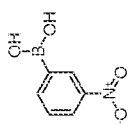
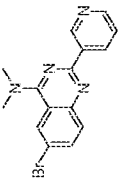
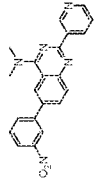
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1330					¹ H NMR (300 MHz, DMSO) δ 9.65 (s, 1H), 8.87-8.75 (m, 1H), 8.70 (s, 1H), 8.55 (s, 1H), 8.41 (s, 1H), 8.25 (s, 1H), 8.09-7.76 (m, 3H), 7.62-7.44 (m, 1H), 7.18 (s, 1H), 3.93 (s, 3H), 3.17 (s, 3H).	DMSO	>98	G2/AQ3		
1331					¹ H NMR (300 MHz, DMSO) δ 9.61 (s, 1H), 8.75 (d, J = 7.6 Hz, 1H), 8.68 (s, 1H), 8.45 (s, 1H), 8.18 (d, J = 9.2 Hz, 1H), 8.07 (d, J = 8.2 Hz, 2H), 8.01-7.87 (m, 3H), 7.55 (s, 1H), 3.51 (s, 6H), 2.63 (s, 3H).	DMSO	>98	G2/AQ3		

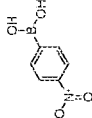
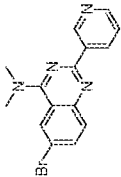
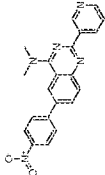
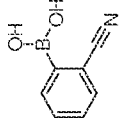
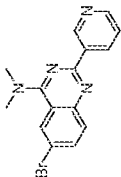
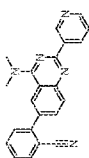
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1332					¹ H NMR (300 MHz, DMSO) δ 9.63 (s, 1H), 8.83-8.72 (m, 1H), 8.70 (d, J = 3.8 Hz, 1H), 8.21 (s, 1H), 7.89 (q, J = 8.6 Hz, 2H), 7.74-7.41 (m, 5H), 3.47 (s, 6H).	DMSO	>98	G2/AQ3		
1333					¹ H NMR (300 MHz, DMSO) δ 9.62 (d, J = 2.0 Hz, 1H), 8.76 (dt, J = 7.9, 1.8 Hz, 1H), 8.70 (dd, J = 4.8, 1.6 Hz, 1H), 8.38 (d, J = 1.9 Hz, 1H), 8.13 (dd, J = 8.7, 1.9 Hz, 1H), 7.98 - 7.84 (m, 2H), 7.78 (d, J = 7.6 Hz, 1H), 7.66-7.41 (m, 3H), 3.51 (s, 6H).	DMSO	>98	G2/AQ3		

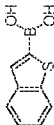
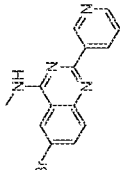
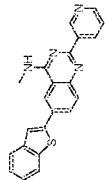
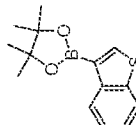
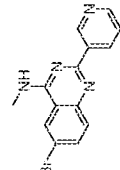
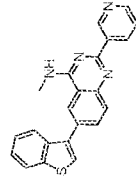
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1334				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.81 - 9.63 (m, 1H), 9.18 (d, J=8.4 Hz, 1H), 9.07 - 8.89 (m, 1H), 8.39 (d, J=8.7 Hz, 1H), 8.31 (d, J=1.7 Hz, 1H), 8.22 - 8.07 (m, 1H), 8.02 (dd, J=8.7, 1.7 Hz, 1H), 7.98 - 7.91 (m, 1H), 7.91 - 7.83 (m, 1H), 7.80 - 7.68 (m, 2H), 3.65 (s, 6H).	DMSO	>98	G2/AQ3		
1335					¹ H NMR (300 MHz, DMSO) δ 9.65 - 9.62 (m, 1H), 8.82 - 8.75 (m, 1H), 8.73 - 8.68 (m, 1H), 8.61 - 8.56 (m, 1H), 8.50 (d, J=2.0 Hz, 1H), 8.34 - 8.20 (m, 3H), 7.97 (d, J=8.7 Hz, 1H), 7.82 (t, J=8.0 Hz, 1H), 7.57 (dd, J=8.3, 5.2 Hz, 1H), 3.54 (s, 6H).	DMSO	>98	G2/AQ3		

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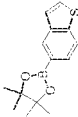
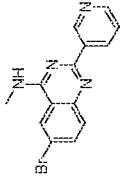
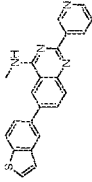
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1336					¹ H NMR (300 MHz, DMSO) δ 9.61 (d, J = 1.8 Hz, 1H), 8.80-8.72 (m, 1H), 8.72 - 8.67 (m, 1H), 8.47 (d, J = 1.6 Hz, 1H), 8.39-8.29 (m, 2H), 8.19 (dd, J = 8.7, 1.8 Hz, 1H), 8.14 - 8.07 (m, 2H), 7.94 (d, J = 8.7 Hz, 1H), 7.55 (dd, J = 7.9, 4.8 Hz, 1H), 3.52 (s, 6H).	DMSO	>98	G2/AQ3		
1337				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.67 - 9.61 (m, 1H), 8.82 - 8.73 (m, 1H), 8.73-8.65 (m, 1H), 8.39 (s, 1H), 8.03 - 7.96 (m, 3H), 7.90-7.72 (m, 2H), 7.72 - 7.60 (m, 1H), 7.56 (dd, J = 7.6, 5.1 Hz, 1H), 3.53 (s, 6H).	DMSO	>98	G2/AQ3		

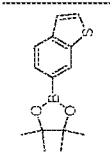
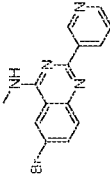
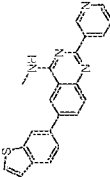
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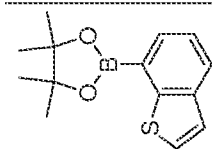
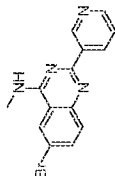
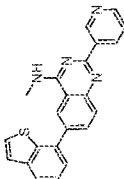
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1338					¹ H NMR (300 MHz, DMSO) δ 9.65 (d, J = 2.1 Hz, 1H), 8.78 (dt, J = 8.1, 1.8 Hz, 1H), 8.71 - 8.59 (m, 3H), 8.19 (dd, J = 8.6, 1.7 Hz, 1H), 8.07 - 7.95 (m, 2H), 7.93-7.81 (m, 2H), 7.54 (dd, J = 7.9, 4.7 Hz, 1H), 7.47 - 7.34 (m, 2H), 3.23-3.20 (m, 3H).	DMSO	>98	G2/AQ3		
1339					¹ H NMR (300 MHz, DMSO) δ 9.68 - 9.64 (m, 1H), 8.83 - 8.75 (m, 1H), 8.69 (d, J = 4.7 Hz, 1H), 8.56-8.45 (m, 2H), 8.17 - 8.05 (m, 1H), 8.05-7.87 (m, 4H), 7.58 - 7.40 (m, 3H), 3.18 (d, J = 4.1 Hz, 3H).	DMSO	>98	G2/AQ3		

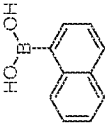
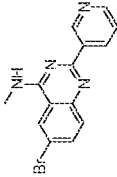
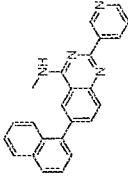
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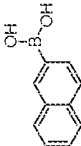
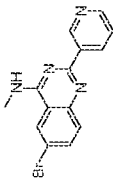
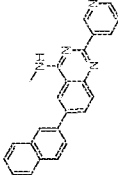
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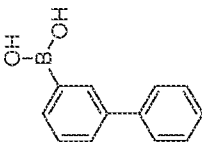
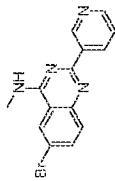
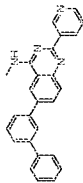
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1341					¹ H NMR (300 MHz, DMSO) δ 9.66 (d, <i>J</i> = 1.5 Hz, 1H), 8.79 (dt, <i>J</i> = 7.9, 1.9 Hz, 1H), 8.68 (dd, <i>J</i> = 4.8, 1.7 Hz, 1H), 8.65 (d, <i>J</i> = 1.9 Hz, 1H), 8.61-8.53 (m, 1H), 8.35 (d, <i>J</i> = 1.6 Hz, 1H), 8.21 (dd, <i>J</i> = 8.7, 2.0 Hz, 1H), 8.16 (d, <i>J</i> = 8.4 Hz, 1H), 7.92-7.80 (m, 3H), 7.59 - 7.48 (m, 2H), 3.21 (d, <i>J</i> = 4.5 Hz, 3H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1342					¹ H NMR (300 MHz, DMSO) δ 9.66 (d, J = 1.6 Hz, 1H), 8.79 (dt, J = 8.0, 1.9 Hz, 1H), 8.73 - 8.61 (m, 3H), 8.50 (s, 1H), 8.24 (dd, J = 8.7, 1.9 Hz, 1H), 8.04 (d, J = 8.4 Hz, 1H), 7.96 - 7.82 (m, 3H), 7.60 - 7.50 (m, 2H), 3.21 (d, J = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

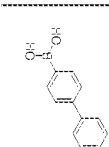
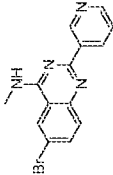
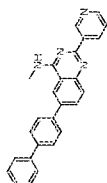
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1343				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.41 (brs, 1H), 9.75 (d, J= 1.7 Hz, 1H), 9.24 - 9.19 (m, 1H), 9.01 (dd, J = 5.2, 1.5 Hz, 1H), 8.90 (d, J = 1.5 Hz, 1H), 8.46 (d, J = 8.7 Hz, 1H), 8.39 (dd, J= 8.7, 1.7 Hz, 1H), 8.06 - 7.91 (m, 2H), 7.86 (d, J=5.5 Hz, 1H), 7.66 - 7.52 (m, 3H), 3.29 (d, J=4.5 Hz, 3H).	DMSO	>98	G2/AQ3		

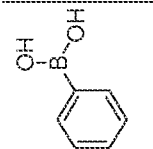
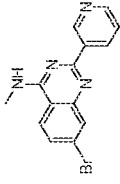
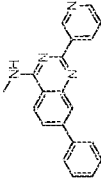
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1344				HCl	¹ H NMR (300 MHz, DMSO) δ 10.33 (brs, 1H), 9.78 (d, J = 2.0 Hz, 1H), 9.23 (d, J = 8.1 Hz, 1H), 9.01 (d, J = 5.0 Hz, 1H), 8.73 (s, 1H), 8.48 (d, J = 8.6 Hz, 1H), 8.14 (d, J = 8.6 Hz, 1H), 8.05 (d, J = 8.0 Hz, 2H), 7.95 (dd, J = 8.1, 5.2 Hz, 1H), 7.79 (d, J = 8.1 Hz, 1H), 7.72-7.50 (m, 4H), 3.27 (d, J = 4.3 Hz, 3H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1345					¹ H NMR (300 MHz, DMSO) δ 9.68 (d, <i>J</i> = 1.4 Hz, 1H), 8.85-8.77 (m, 1H), 8.77 - 8.62 (m, 3H), 8.40 (s, 1H), 8.30 (dd, <i>J</i> = 8.7, 1.6 Hz, 1H), 8.14 - 7.95 (m, 4H), 7.91 (d, <i>J</i> = 8.7 Hz, 1H), 7.63-7.50 (m, 3H), 3.22 (d, <i>J</i> = 4.3 Hz, 3H).	DMSO	>98	G2/AQ3		

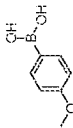
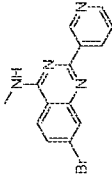
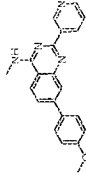
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1346					¹ H NMR (300 MHz, DMSO) δ 9.67 (d, <i>J</i> = 2.0 Hz, 1H), 8.80 (dt, <i>J</i> = 8.0, 1.9 Hz, 1H), 8.69 (dd, <i>J</i> = 4.7, 1.7 Hz, 1H), 8.67 Hz, 1H), 8.67 -8.55 (m, 2H), 8.25 (dd, <i>J</i> = 8.7, 1.8 Hz, 1H), 8.13 - 8.09 (m, 1H), 7.92-7.77 (m, 4H), 7.71 (d, <i>J</i> = 7.8 Hz, 1H), 7.67 - 7.47 (m, 4H), 7.47-7.37 (m, 1H), 3.20 (d, <i>J</i> = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

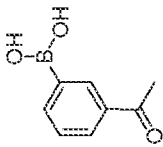
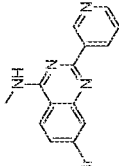
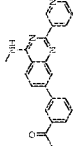
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1347					¹ H NMR (300 MHz, DMSO) δ 9.68 - 9.65 (m, 1H), 8.79 (dt, J = 8.0, 1.9 Hz, 1H), 8.69 (dd, J = 4.8, 1.7 Hz, 1H), 8.68 - 8.60 (m, 2H), 8.20 (dd, J = 8.7, 1.9 Hz, 1H), 7.99 - 7.95 (m, 2H), 7.92 - 7.81 (m, 3H), 7.80 - 7.73 (m, 2H), 7.61 - 7.46 (m, 3H), 7.46 - 7.36 (m, 1H), 3.20 (d, J = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

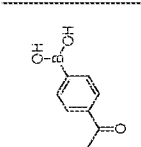
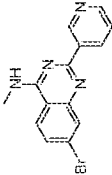
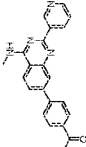
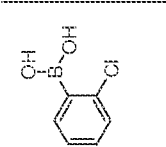
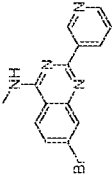
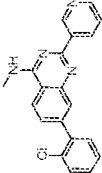
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1348					¹ H NMR (300 MHz, DMSO) δ 9.66 (d, J = 1.5 Hz, 1H), 8.80 (dt, J = 7.9, 1.9 Hz, 1H), 8.70 (dd, J = 4.7, 1.7 Hz, 1H), 8.55 (d, J = 4.6 Hz, 1H), 8.32 (d, J = 8.6 Hz, 1H), 8.04 (d, J = 1.8 Hz, 1H), 7.92 - 7.84 (m, 3H), 7.60 - 7.50 (m, 3H), 7.50 - 7.42 (m, 1H), 3.18 (d, J = 4.5 Hz, 3H).	DMSO	>98	G2/AQ3		

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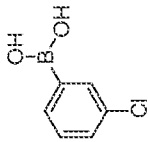
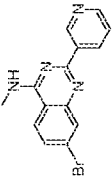
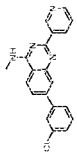
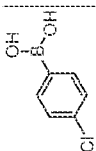
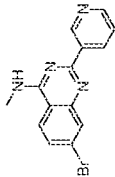
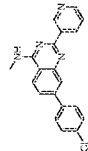
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1349				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.23 (brs, 1H), 9.67 (d, J = 2.1 Hz, 1H), 9.04 (d, J = 7.7 Hz, 1H), 8.98-8.91 (m, 1H), 8.57 (d, J = 8.7 Hz, 1H), 8.48 (s, 1H), 8.08 (d, J = 8.3 Hz, 1H), 7.91-7.76 (m, 3H), 7.22 - 7.10 (m, 2H), 3.85 (s, 3H), 3.31 (d, J = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

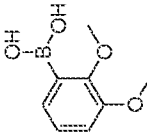
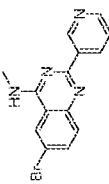
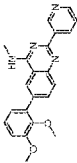
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1350				2 HCl	¹ H NMR (300 MHz, DMSO- <i>d</i> ₆) δ 10.48 (brs, 1H), 9.74 (d, <i>J</i> = 1.9 Hz, 1H), 9.19 (d, <i>J</i> = 8.3 Hz, 1H), 9.07 - 1H), 9.07 - 8.97 (m, 1H), 8.71 (d, <i>J</i> = 8.8 Hz, 1H), 8.64 (s, 1H), 8.36 (s, 1H), 8.19 (d, <i>J</i> = 7.6 Hz, 1H), 8.16-8.06 (m, 2H), 7.95 (dd, <i>J</i> = 8.1, 5.0 Hz, 1H), 7.75 (t, <i>J</i> = 7.7 Hz, 1H), 3.31 (d, <i>J</i> = 4.4 Hz, 3H), 2.71 (s, 3H).	DMSO	>98	G2/AQ3		

(continued)

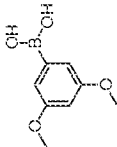
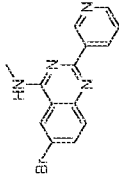
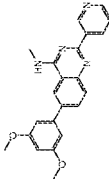
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1351				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.47 (brs, 1H), 9.73 (d, J = 1.3 Hz, 1H), 9.19 (d, J = 8.3 Hz, 1H), 9.03 -8.94 (m, 1H), 8.69-8.62 (m, 2H), 8.15 - 8.07 (m, 3H), 8.02-7.77 (m, 3H), 3.29 (d, J = 4.3 Hz, 3H), 2.65 (s, 3H).	DMSO	>98	G2/AQ3		
1352				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.44 (brs, 1H), 9.74 (s, 1H), 9.19 (d, J = 7.3 Hz, 1H), 8.99 (d, J = 5.2 Hz, 1H), 8.68 (d, J = 8.4 Hz, 1H), 8.37 (s, 1H), 8.03 - 7.90 (m, 1H), 7.83 (d, J = 8.2 Hz, 1H), 7.72-7.62 (m, 1H), 7.53 (d, J = 9.6 Hz, 3H), 3.32 (d, J = 4.0 Hz, 3H).	DMSO	>98	G2/AQ3		

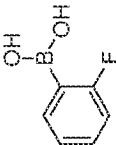
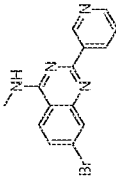
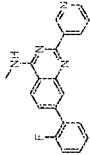
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1353				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.48 (brs, 1H), 9.77 (s, 1H), 9.29 - 9.23 (m, 1H), 9.05-8.97 (m, 1H), 8.74 - 8.59 (m, 2H), 8.20-8.08 (m, 1H), 8.02 - 7.95 (m, 1H), 7.90 (s, 1H), 7.84-7.79 (m, 1H), 7.68 - 7.57 (m, 2H), 3.35-3.29 (m, 3H).	DMSO	>98	G2/AQ3		
1354				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.35 (brs, 1H), 9.72 (d, J = 2.0 Hz, 1H), 9.21-9.11 (m, 1H), 8.98 (dd, J = 5.1, 1.4 Hz, 1H), 8.65 (d, J = 8.7 Hz, 1H), 8.59 (s, 1H), 8.09 (dd, J = 8.8, 1.7 Hz, 1H), 7.95 - 7.82 (m, 3H), 7.70-7.60 (m, 2H), 3.30 (d, J = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

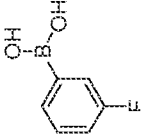
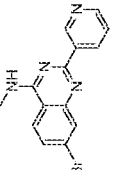
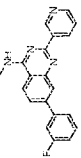
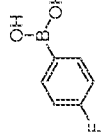
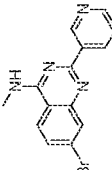
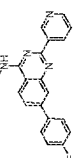
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1355				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.51 (brs, 1H), 9.77 (d, <i>J</i> = 1.9 Hz, 1H), 9.22 (d, <i>J</i> = 8.2 Hz, 1H), 9.01 (dd, <i>J</i> = 5.1, 1.4 Hz, 1H), 8.69 (s, 1H), 8.40 (d, <i>J</i> = 8.7 Hz, 1H), 8.17 (dd, <i>J</i> = 8.7, 1.6 Hz, 1H), 7.96 (dd, <i>J</i> = 8.1, 5.2 Hz, 1H), 7.30 - 7.15 (m, 2H), 7.10 (dd, <i>J</i> = 7.3, 1.9 Hz, 1H), 3.89 (s, 3H), 3.59 (s, 3H), 3.30 (d, <i>J</i> = 4.5 Hz, 3H).	DMSO	>98	G2/AQ3		

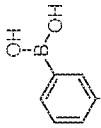
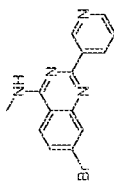
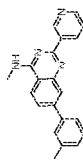
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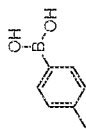
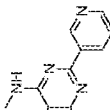
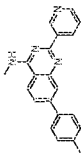
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1357				HCl	¹ H NMR (300 MHz, DMSO) δ 10.21 (brs, 1H), 9.63 (s, 1H), 8.99 (d, J = 8.0 Hz, 1H), 8.93 (d, J = 3.7 Hz, 1H), 8.87 (s, 1H), 8.37 (d, J = 8.9 Hz, 1H), 8.15 (d, J = 8.2 Hz, 1H), 7.91-7.78 (m, 1H), 7.05 (d, J = 2.1 Hz, 2H), 6.66-6.58 (m, 1H), 3.87 (s, 6H), 3.31 (d, J = 4.2 Hz, 3H).	DMSO	>98	G2/AQ3		

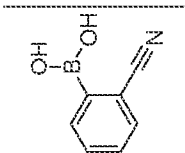
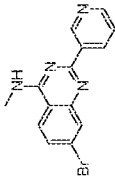
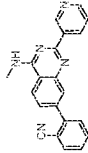
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Meth-od
1358				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.43 (brs, 1H), 9.73 (d, <i>J</i> = 1.8 Hz, 1H), 9.17 (d, <i>J</i> = 8.2 Hz, 1H), 8.99 (dd, <i>J</i> = 5.0, 1.3 Hz, 1H), 8.69 (d, <i>J</i> = 8.6 Hz, 1H), 8.49 (s, 1H), 8.05-7.88 (m, 2H), 7.71 (t, <i>J</i> = 7.9 Hz, 1H), 7.64-7.50 (m, 1H), 7.50 - 7.33 (m, 2H), 3.32 (d, <i>J</i> = 4.3 Hz, 3H).	DMSO	>98	G2/AQ3		

(continued)

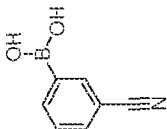
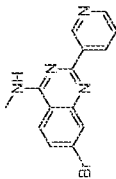
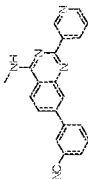
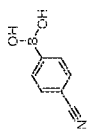
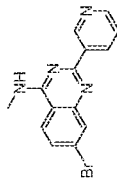
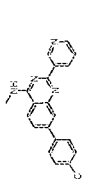
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1359				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.29 (s, 1H), 9.70 (d, J = 1.7 Hz, 1H), 9.14 (d, J = 8.3 Hz, 1H), 8.97 (dd, J = 5.1, 1.3 Hz, 1H), 8.63 (d, J = 8.7 Hz, 1H), 8.58 (s, 1H), 8.15-8.04 (m, 1H), 7.90 (dd, J = 8.0, 5.1 Hz, 1H), 7.75-7.57 (m, 3H), 7.43-7.30 (m, 1H), 3.30 (d, J = 4.3 Hz, 3H).	DMSO	>98	G2/AQ3		
1360				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.41 (brs, 1H), 9.73 (d, J = 1.9 Hz, 1H), 9.16 (d, J = 8.3 Hz, 1H), 8.98 (dd, J = 5.1, 1.4 Hz, 1H), 8.65 (d, J = 8.8 Hz, 1H), 8.60 (s, 1H), 8.12-8.02 (m, 1H), 8.00-7.84 (m, 3H), 7.49-7.33 (m, 2H), 3.30 (d, J = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

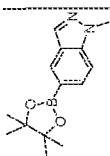
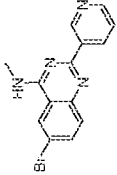
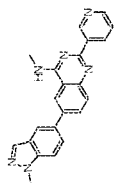
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1361				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.39 (s, 1H), 9.71 (d, J = 1.8 Hz, 1H), 9.17 - 9.08 (m, 1H), 8.97 (dd, J = 5.1, 1.5 Hz, 1H), 8.64 (d, J = 8.7 Hz, 1H), 8.58 (s, 1H), 8.07 (dd, J = 8.6, 1.7 Hz, 1H), 7.88 (dd, J = 8.1, 5.2 Hz, 1H), 7.70 Hz, 1H), 7.46 (t, J = 7.6 Hz, 1H), 7.33 (d, J = 7.4 Hz, 1H), 3.30 (d, J = 4.4 Hz, 3H), 2.43 (s, 3H).	DMSO	>98	G2/AQ3		

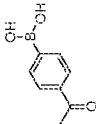
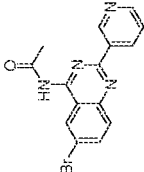
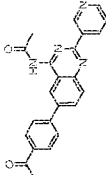
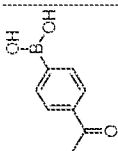
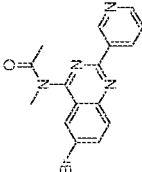
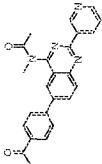
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1362				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.37 (brs, 1H), 9.70 (d, J = 1.7 Hz, 1H), 9.10 (d, J = 8.1 Hz, 1H), 8.96 (dd, J = 5.0, 1.4 Hz, 1H), 8.62 (d, J = 8.7 Hz, 1H), 8.58 (s, 1H), 8.08 (dd, J = 8.6, 1.6 Hz, 1H), 7.87 (dd, J = 7.9, 5.0 Hz, 1H), 7.74 (d, J = 8.2 Hz, 2H), 7.39 (d, J = 8.0 Hz, 2H), 3.30 (d, J = 4.4 Hz, 3H), 2.40 (s, 3H).	DMSO	>98	G2/AQ3		

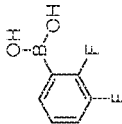
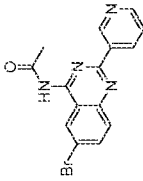
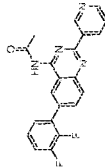
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1363				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.82 (brs, 1H), 9.65 (s, 1H), 9.14 - 9.02 (m, 1H), 9.02-8.90 (m, 1H), 8.64 - 8.51 (m, 1H), 8.20 (s, 1H), 8.07 (d, J=7.8 Hz, 1H), 8.00 - 7.85 (m, 3H), 7.85-7.66 (m, 2H), 3.30 (s, 3H).	DMSO	>98	G2/AQ3		

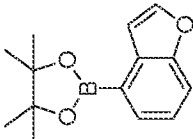
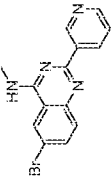
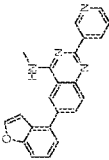
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1364				HCl	¹ H NMR (300 MHz, DMSO) δ 9.83 (brs, 1H), 9.68-9.63 (m, 1H), 9.12 - 9.02 (m, 1H), 9.01-8.91 (m, 1H), 8.56 (d, J=9.2 Hz, 1H), 8.42 (s, 1H), 8.35 (s, 1H), 8.19 (d, J=8.7 Hz, 1H), 8.13 (d, J=9.3 Hz, 1H), 7.97 (d, J=7.8 Hz, 1H), 7.93-7.84 (m, 1H), 7.79 (t, J=8.2 Hz, 1H), 3.28 (d, J=4.5 Hz, 3H).	DMSO	>98	G2/AQ3		
1365				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.27 (brs, 1H), 9.72 (s, 1H), 9.23 - 9.10 (m, 1H), 8.98 (s, 1H), 8.77-8.55 (m, 2H), 8.18 - 7.86 (m, 6H), 3.34-3.27 (m, 3H).	DMSO	>98	G2/AQ3		

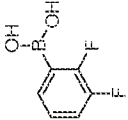
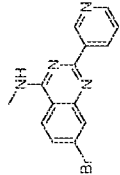
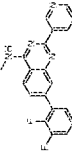
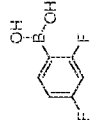
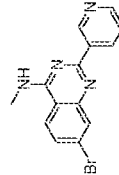
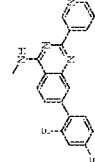
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1366					¹ H NMR (300 MHz, DMSO) δ 9.66 (d, J = 2.0 Hz, 1H), 8.79 (dt, J = 7.9, 1.9 Hz, 1H), 8.69 (dd, J = 4.7, 1.6 Hz, 1H), 8.65 - 8.53 (m, 2H), 8.22-8.12 (m, 3H), 7.92 (dd, J = 8.8, 1.5 Hz, 1H), 7.86 (d, J = 8.7 Hz, 1H), 7.80 (d, J = 8.8 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 4.11 (s, 3H), 3.20 (d, J = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1367					¹ H NMR (300 MHz, DMSO) δ 11.10 (s, 1H), 9.63 (d, 1H), 8.85 (d, 1H), 8.81-8.70 (m, 2H), 8.39 (dd, 1H), 8.15 (m, 1H), 8.02 (m, 5H), 7.60 (dd, 1H), 8.0, 4.8 Hz, 1H), 2.65 (s, 3H), 2.61 (s, 3H).	DMSO	>98	G2/AQ3		
1368				HCl	¹ H NMR (300 MHz, DMSO) δ 9.72 (d, 1H), 9.27-9.21 (m, 1H), 8.97 (dd, 1H), 8.50 (dd, 1H), 8.50 (dd, 1H), 8.34 (d, 1H), 8.30 (d, 1H), 8.30 (d, 1H), 8.15 (m, 1H), 8.09 (m, 2H), 8.09-7.98 (m, 3H), 3.55 (s, 3H), 2.65 (s, 3H), 2.22 (s, 3H).	DMSO	>98	G2/AQ3		

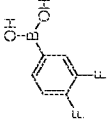
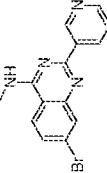
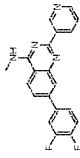
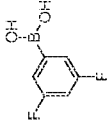
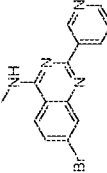
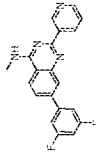
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1369				HCl	¹ H NMR (300 MHz, DMSO- <i>d</i> ₆) δ 11.16 (s, 1H), 9.67 (s, 1H), 9.20 (d, 1H), 8.2 Hz, <i>J</i> = 8.2 Hz, 1H), 8.97 (d, <i>J</i> = 4.3 Hz, 1H), 8.75 (s, 1H), 8.23 (d, <i>J</i> = 8.7 Hz, 1H), 8.14 (d, <i>J</i> = 8.8 Hz, 1H), 8.02 (dd, <i>J</i> = 8.0, 5.5 Hz, 1H), 7.66 - 7.47 (m, 2H), 7.47 - 7.34 (m, 1H), 2.56 (s, 3H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1370				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.17 (brs, 1H), 9.68 (d, J = 2.0 Hz, 1H), 9.06 (d, J = 8.0 Hz, 1H), 8.95 (dd, J = 5.0, 1.4 Hz, 1H), 8.77 (s, 1H), 8.31 (s, 2H), 8.16 (d, J = 2.1 Hz, 1H), 7.87 (dd, J = 7.8, 5.2 Hz, 1H), 7.72 (d, J = 7.8 Hz, 1H), 7.61 - 7.47 (m, 2H), 7.19 (d, J = 1.4 Hz, 1H), 3.31 (d, J = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

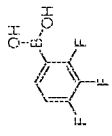
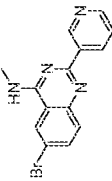
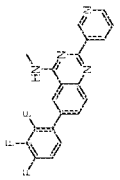
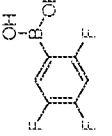
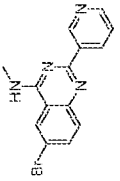
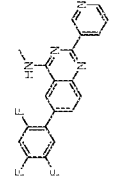
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1371				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.28 (brs, 1H), 9.71 (s, 1H), 9.16 (d, J = 7.9 Hz, 1H), 8.98 (d, J = 4.2 Hz, 1H), 8.67 (d, J = 8.9 Hz, 1H), 8.45 (s, 1H), 8.01-7.83 (m, 2H), 7.68-7.50 (m, 2H), 7.50-7.35 (m, 1H), 3.31 (d, J = 4.1 Hz, 3H).	DMSO	>98	G2/AQ3		
1372				HCl	¹ H NMR (300 MHz, DMSO) δ 9.72-9.45 (m, 2H), 8.95 (d, J = 7.6 Hz, 1H), 8.88 (d, J = 3.8 Hz, 1H), 8.49 (d, J = 8.5 Hz, 1H), 8.13 (s, 1H), 7.91-7.75 (m, 3H), 7.53-7.43 (m, 1H), 7.36-7.25 (m, 1H), 3.27 (d, J = 4.3 Hz, 3H).	DMSO	>98	G2/AQ3		

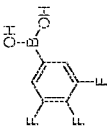
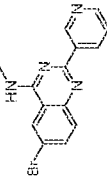
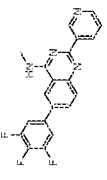
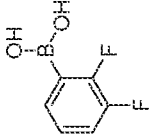
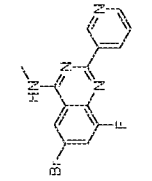
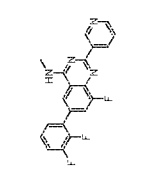
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1373				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.04 (brs, 1H), 9.70 (s, 1H), 9.22 - 9.07 (m, 1H), 8.93 (s, 1H), 8.62 (d, J=7.6 Hz, 1H), 8.55 (s, 1H), 8.06 (d, J=8.9 Hz, 1H), 7.99 - 7.79 (m, 2H), 7.79 - 7.55 (m, 2H), 3.33 - 3.27 (m, 3H).	DMSO	>98	G2/AQ3		
1374				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 9.34 (brs, 1H), 9.06 - 8.92 (m, 1H), 8.86 (s, 1H), 8.50 - 8.37 (m, 1H), 8.25 (s, 1H), 8.11 - 7.95 (m, 1H), 7.85 - 7.74 (m, 1H), 7.74 - 7.57 (m, 2H), 7.43 - 7.29 (m, 1H), 3.28 - 3.21 (m, 3H).	DMSO	>98	G2/AQ3		

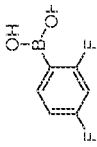
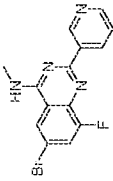
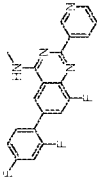
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1375				HCl	¹ H NMR (300 MHz, DMSO) δ 9.63 (d, J = 1.6 Hz, 1H), 9.38 (brs, 1H), 9.01 - 8.91 (m, 1H), 8.84 (dd, J = 5.0, 1.6 Hz, 1H), 8.60 (s, 1H), 8.08 (s, 2H), 7.76 (dd, J = 7.9, 5.1 Hz, 1H), 7.64 - 7.38 (m, 2H), 3.26 (d, J = 3.7 Hz, 3H).	DMSO	>98	G2/AQ3		
1376				HCl	¹ H NMR (300 MHz, DMSO) δ 9.88 (brs, 1H), 9.65 (s, 1H), 9.18 - 8.99 (m, 1H), 8.90 (s, 1H), 8.70 (s, 1H), 8.31 - 8.03 (m, 2H), 7.94 - 7.77 (m, 2H), 7.77 - 7.57 (m, 1H), 3.28 (s, 3H).	DMSO	>98	G2/AQ3		

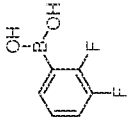
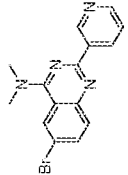
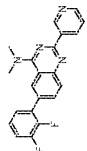
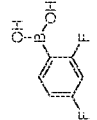
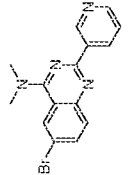
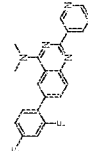
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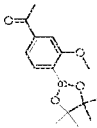
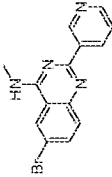
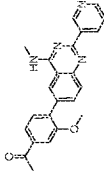
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1377				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.33 (s, 1H), 9.67 (s, 1H), 9.10 (d, 1H), 9.10 (d, 1H), 9.02 (s, 1H), 9.02 (s, 1H), 8.92 (d, 1H), 8.92 (d, 1H), 8.35 (d, 1H), 8.35 (d, 1H), 8.8 Hz, 1H), 8.25 (d, 1H), 8.25 (d, 1H), 8.03 - 7.76 (m, 3H), 3.31 (s, 3H).	DMSO	>98	G2/AQ3		
1378				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.66 (d, J = 1.7 Hz, 1H), 9.56-9.34 (m, 1H), 9.26 - 9.10 (m, 1H), 9.05 (d, J=4.7 Hz, 1H), 8.44 (s, 1H), 8.19 (dd, J = 8.1, 5.7 Hz, 1H), 7.95 (d, J= 11.5 Hz, 1H), 7.63-7.45 (m, 2H), 7.45 - 7.26 (m, 1H), 3.19 (d, J=4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

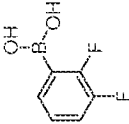
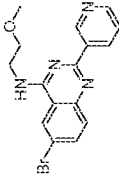
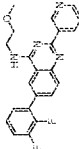
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1379				HCl	¹ H NMR (300 MHz, DMSO) δ 9.67 - 9.59 (m, 1H), 9.38 (d, J = 8.3 Hz, 1H), 9.22 - 9.10 (m, 1H), 9.03 (d, J = 4.9 Hz, 1H), 8.37 (s, 1H), 8.16 (dd, J = 8.1, 5.6 Hz, 1H), 7.89 (d, J = 11.6 Hz, 1H), 7.78 (td, J = 8.9, 6.6 Hz, 1H), 7.54 - 7.36 (m, 1H), 7.30 (td, J = 8.3, 2.0 Hz, 1H), 3.18 (d, J = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

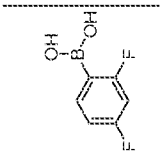
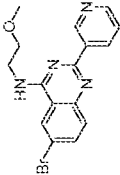
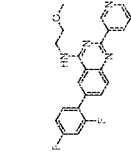
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1380				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.72 (d, J = 1.6 Hz, 1H), 9.25-9.14 (m, 1H), 8.99 (dd, J = 5.2, 1.5 Hz, 1H), 8.51 (s, 1H), 8.44 (d, J = 8.7 Hz, 1H), 8.26-8.11 (m, 1H), 7.94 (dd, J = 8.0, 5.2 Hz, 1H), 7.65-7.49 (m, 2H), 7.49-7.29 (m, 1H), 3.69 (s, 6H).	DMSO	>98	G2/AQ3		
1381				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.71 (s, 1H), 9.18 (d, J = 8.0 Hz, 1H), 8.98 (d, J = 5.1 Hz, 1H), 8.49-8.36 (m, 2H), 8.16 (d, J = 8.8 Hz, 1H), 7.99-7.86 (m, 1H), 7.86-7.72 (m, 1H), 7.46 (t, J = 10.2 Hz, 1H), 7.29 (t, J = 8.5 Hz, 1H), 3.69 (s, 6H).	DMSO	>98	G2/AQ3		

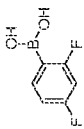
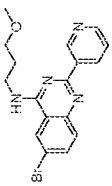
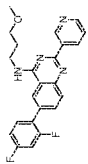
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1382				HCl	¹ H NMR (300 MHz, DMSO) δ 10.25 (brs, 1H), 9.66 (d, J = 1.6 Hz, 1H), 9.04 (d, J = 8.2 Hz, 1H), 8.94 (dd, J = 5.0, 1.6 Hz, 1H), 8.70 (s, 1H), 8.23 (d, J = 8.8 Hz, 1H), 8.16 (dd, J = 8.7, 1.5 Hz, 1H), 7.84 (dd, J = 8.0, 5.1 Hz, 1H), 7.71 (dd, J = 8.0, 1.4 Hz, 1H), 7.65-7.55 (m, 2H), 3.90 (s, 3H), 3.29 (d, J = 4.5 Hz, 3H), 2.64 (s, 3H).	DMSO	>98	G2/AQ3		

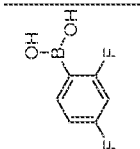
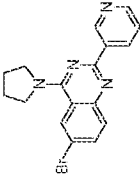
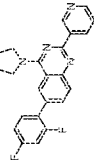
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Meth-od
1383				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.18 (brs, 1H), 9.68 (d, <i>J</i> = 1.8 Hz, 1H), 9.17 (d, <i>J</i> = 8.0 Hz, 1H), 8.99 (dd, <i>J</i> = 5.2, 1.4 Hz, 1H), 8.86 (s, 1H), 8.31 (d, <i>J</i> = 8.7 Hz, 1H), 8.22 (d, <i>J</i> = 8.8 Hz, 1H), 7.97 (dd, <i>J</i> = 8.0, 5.3 Hz, 1H), 7.64 - 7.49 (m, 2H), 7.49 - 7.34 (m, 1H), 4.06 - 3.92 (m, 2H), 3.73 (t, <i>J</i> = 5.5 Hz, 2H), 3.33 (s, 3H).	DMSO	>98	G2/AQ3		

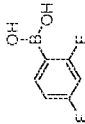
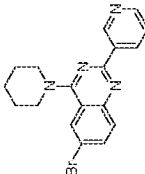
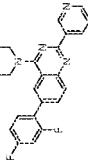
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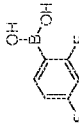
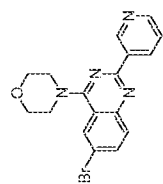
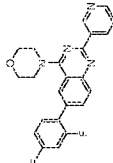
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Meth-od
1384				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.18 (brs, 1H), 9.67 (d, J = 1.6 Hz, 1H), 9.14 (d, J=8.1 Hz, 1H), 8.99 Hz, 1H), 8.99 (dd, J= 5.2, 1.5 Hz, 1H), 8.80 (s, 1H), 8.29 (d, J=8.7 Hz, 1H), 8.18 (d, J = 8.7 Hz, 1H), 7.95 (dd, J = 8.2, 5.2 Hz, 1H), 7.79 (td, J = 8.9, 6.6 Hz, 1H), 7.50 (ddd, J = 11.6, 9.3, 2.6 Hz, 1H), 7.33 (td, J = 8.3, 2.3 Hz, 1H), 4.08-3.94 (m, 3H), 3.73 (t, J = 5.5 Hz, 3H), 3.32 (s, 3H).	DMSO	>98	G2/AQ3		

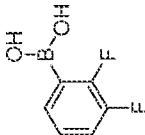
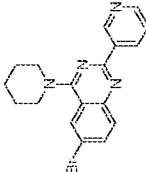
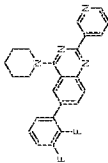
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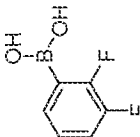
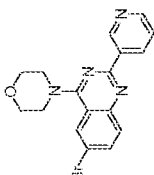
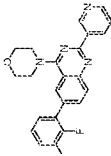
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1385				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.33 (brs, 1H), 9.70 (d, J = 1.7 Hz, 1H), 9.15 (d, J = 8.2 Hz, 1H), 8.99 (dd, J = 5.2, 1.5 Hz, 1H), 8.81 (s, 1H), 8.33 (d, J = 8.7 Hz, 1H), 8.18 (d, J = 8.7 Hz, 1H), 7.95 (dd, J = 8.1, 5.2 Hz, 1H), 7.81 (d, J = 8.9, 6.6 Hz, 1H), 7.49 (ddd, J = 11.5, 9.3, 2.5 Hz, 1H), 7.32 (td, J = 8.5, 2.2 Hz, 1H), 3.96 (-3.78 (m, 2H), 3.48 (t, J = 6.1 Hz, 2H), 3.27 (s, 3H), 2.14 (-1.93 (m, 2H).	DMSO	>98	G2/AQ3		

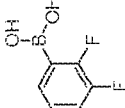
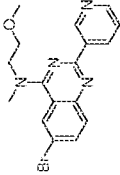
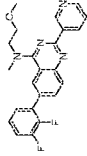
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1386				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.73 (s, 1H), 9.17 (d, J = 8.4 Hz, 1H), 8.99 (s, 1H), 8.52 (s, 1H), 8.43 (d, J = 8.8 Hz, 1H), 8.21 (d, J = 9.0 Hz, 1H), 8.02-7.90 (m, 1H), 7.87 - 7.78 (m, 1H), 7.57-7.38 (m, 1H), 7.38 - 7.24 (m, 1H), 4.31-4.18 (m, 4H), 2.08 (s, 4H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1387				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.68 (d, J = 1.7 Hz, 1H), 9.17 (d, J = 8.3 Hz, 1H), 8.99 (dd, J = 5.2, 1.5 Hz, 1H), 8.34 (d, J = 8.6 Hz, 1H), 8.24 - 8.12 (m, 2H), 7.97 (dd, J = 8.0, 5.2 Hz, 1H), 7.81 (td, J = 8.9, 6.6 Hz, 1H), 7.49 (ddd, J = 11.6, 9.3, 2.5 Hz, 1H), 7.30 (td, J = 8.4, 2.1 Hz, 1H), 4.16 (brs, 4H), 1.80 (brs, 6H).	DMSO	>98	G2/AQ3		

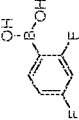
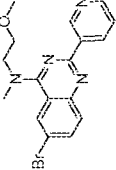
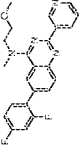
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1388				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.70 (d, J = 1.7 Hz, 1H), 9.26 (d, J = 8.3 Hz, 1H), 9.01 (dd, J = 5.3, 1.4 Hz, 1H), 8.37 - 8.19 (m, 2H), 8.15 (d, J = 8.7 Hz, 1H), 8.02 (dd, J = 8.1, 5.3 Hz, 1H), 7.81 (td, J = 8.9, 6.5 Hz, 1H), 7.48 (ddd, J = 11.6, 9.3, 2.6 Hz, 1H), 7.30 (td, J = 8.4, 2.3 Hz, 1H), 4.23 - 4.12 (m, 4H), 3.90 - 3.80 (m, 4H).	DMSO	>98	G2/AQ3		

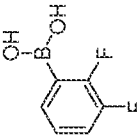
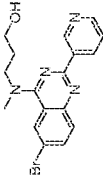
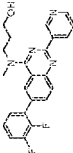
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1389				2HCl	¹ H NMR (300 MHz, DMSO) δ 9.68 (d, J = 1.7 Hz, 1H), 9.17 (d, J = 8.2 Hz, 1H), 8.99 (dd, J = 5.2, 1.5 Hz, 1H), 8.34 (d, J = 8.7 Hz, 1H), 8.27-8.13 (m, 2H), 7.97 (dd, J = 8.1, 5.4 Hz, 1H), 7.67-7.48 (m, 2H), 7.41 (ddd, J = 14.7, 9.9, 4.9 Hz, 1H), 4.16 (brs, 4H), 1.80 (brs, 6H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1390				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.70 (d, <i>J</i> = 1.8 Hz, 1H), 9.26 (d, <i>J</i> = 8.3 Hz, 1H), 9.01 (dd, <i>J</i> = 5.3, 1.4 Hz, 1H), 8.36 - 8.23 (m, 2H), 8.18 (d, <i>J</i> = 8.7 Hz, 1H), 8.03 (dd, <i>J</i> = 8.0, 5.4 Hz, 1H), 7.64 - 7.46 (m, 2H), 7.46 - 7.29 (m, 1H), 4.23 - 4.16 (m, 4H), 3.85 (d, <i>J</i> = 4.7 Hz, 4H).	DMSO	>98	G2/AQ3		

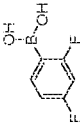
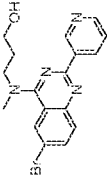
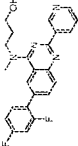
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1391				2 HCl	¹ H NMR (300 MHz, DMSO- <i>d</i> ₆) δ 9.68 (d, <i>J</i> = 1.7 Hz, 1H), 9.18 (d, <i>J</i> = 8.2 Hz, 1H), 9.00 (dd, <i>J</i> = 5.2, 1.2 Hz, 1H), 8.62 (s, 1H), 8.39 (d, <i>J</i> = 8.7 Hz, 1H), 8.21 (d, <i>J</i> = 8.8 Hz, 1H), 7.98 (dd, <i>J</i> = 8.0, 5.2 Hz, 1H), 7.64 - 7.47 (m, 2H), 7.46 - 7.27 (m, 1H), 4.26 (t, <i>J</i> = 5.2 Hz, 2H), 3.85 (t, <i>J</i> = 5.3 Hz, 2H), 3.72 (s, 3H), 3.33 (s, 3H)	DMSO	>98	G2/AQ3		

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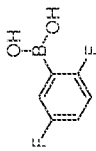
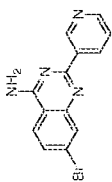
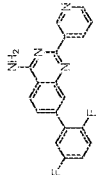
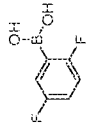
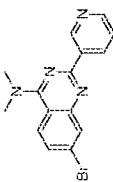
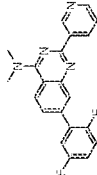
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1392				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.67 (d, J = 1.7 Hz, 1H), 9.15 (d, J = 8.1 Hz, 1H), 8.99 (d, J = 4.1 Hz, 1H), 8.56 (s, 1H), 8.36 (d, J = 8.7 Hz, 1H), 8.17 (d, J = 8.7 Hz, 1H), 7.96 (dd, J = 8.1, 5.2 Hz, 1H), 7.86 - 7.70 (m, 1H), 7.55-7.40 (m, 1H), 7.30 (td, J = 8.5, 2.2 Hz, 1H), 4.36 -4.18 (m, 2H), 3.88-3.77 (m, 2H), 3.72 (s, 3H), 3.33 (s, 3H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1393				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.70 (d, J =1.7 Hz, 1H), 9.24-9.06 (m, 1H), 8.99 (dd, J =5.1, 1.5 Hz, 1H), 8.52 (s, 1H), 8.40 (d, J =8.7 Hz, 1H), 8.30-8.14 (m, 1H), 7.93 (dd, J =8.0, 5.2 Hz, 1H), 7.63-7.46 (m, 2H), 7.46-7.25 (m, 1H), 4.22-4.11 (m, 2H), 3.73 (s, 3H), 3.59 (t, J =5.9 Hz, 2H), 2.10-1.92 (m, 2H).	DMSO	>98	G2/AQ3		

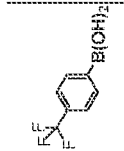
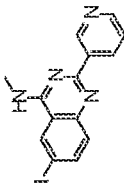
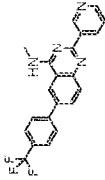
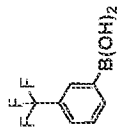
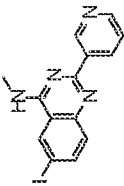
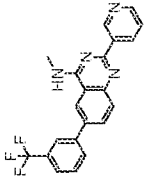
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Meth-od
1394				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.68 (d, J = 1.7 Hz, 1H), 9.11 (d, J = 8.3 Hz, 1H), 8.97 (dd, J = 5.1, 1.5 Hz, 1H), 8.46 (s, 1H), 8.36 (d, J = 8.7 Hz, 1H), 8.17 (d, J = 8.8 Hz, 1H), 7.90 (dd, J = 8.1, 5.2 Hz, 1H), 7.80 (td, J = 8.9, 6.6 Hz, 1H), 7.47 (ddd, J = 11.6, 9.3, 2.6 Hz, 1H), 7.29 (td, J = 8.4, 2.1 Hz, 1H), 4.22 -4.08 (m, 2H), 3.72 (s, 3H), 3.59 (t, J = 5.9 Hz, 2H), 2.10 -1.94 (m, 2H).	DMSO	>98	G2/AQ3		

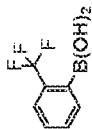
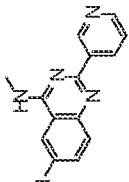
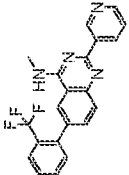
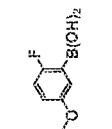
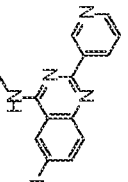
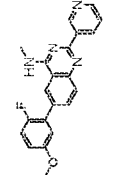
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1395				HCl	¹ H NMR (300 MHz, DMSO) δ 9.97 - 9.66 (m, 2H), 9.66 - 9.58 (m, 1H), 9.17 - 9.05 (m, 1H), 8.99 (dd, J = 5.2, 1.5 Hz, 1H), 8.66 (d, J = 8.6 Hz, 1H), 8.50 (s, 1H), 8.01 - 7.88 (m, 2H), 7.62 (ddd, J = 9.1, 6.1, 3.1 Hz, 1H), 7.58 - 7.36 (m, 2H).	DMSO	>98	G2/AQ3		
1396				HCl	¹ H NMR (300 MHz, DMSO) δ 9.70 (d, J = 1.8 Hz, 1H), 9.16 (d, J = 8.3 Hz, 1H), 8.98 (d, J = 3.9 Hz, 1H), 8.58 - 8.42 (m, 2H), 8.00 - 7.82 (m, 2H), 7.69 - 7.38 (m, 3H), 3.69 (s, 6H).	DMSO	>98	G2/AQ3		

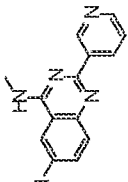
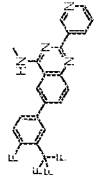
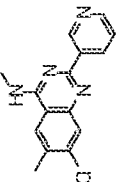
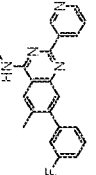
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1397				HCl	¹ H NMR (400 MHz, DMSO) δ 10.47 (s, 1H), 9.68 (d, J = 1.8 Hz, 1H), 9.11 (d, J = 8.0 Hz, 1H), 9.04 (s, 1H), 8.96 (dd, J = 5.1, 1.4 Hz, 1H), 8.41 (dd, J = 8.8, 1.6 Hz, 1H), 8.30 (d, J = 8.7 Hz, 1H), 8.14 (d, J = 8.2 Hz, 2H), 7.90 (d, J = 8.4 Hz, 3H), 3.34 (t, J = 17.1 Hz, 3H).	DMSO	>98	AQ4		
1398				HCl	¹ H NMR (400 MHz, DMSO) δ 10.15 (s, 1H), 9.65 (d, J = 1.7 Hz, 1H), 9.06 (d, J = 7.9 Hz, 1H), 8.99 - 8.90 (m, 2H), 8.44 (d, J = 7.2 Hz, 1H), 8.28-8.17 (m, 3H), 7.94 - 7.76 (m, 3H), 3.32 (d, J = 4.5 Hz, 3H).	DMSO	>98	AQ4		

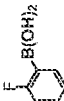
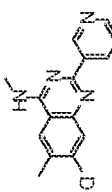
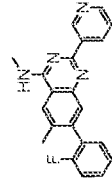
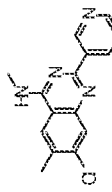
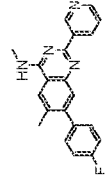
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1399				HCl	¹ H NMR (400 MHz, DMSO) δ 10.05 (s, 1H), 9.65 (d, J =2.1 Hz, 1H), 9.10 - 9.00 (m, 1H), 8.96 (dd, J =5.1, 1.5 Hz, 1H), 8.65 (s, 1H), 8.26 - 8.17 (m, 1H), 8.16-8.07 (m, 1H), 7.90 (dd, J =7.8, 5.3 Hz, 1H), 7.77 - 7.69 (m, 1H), 7.68-7.55 (m, 3H), 3.29 (d, J =4.6 Hz, 3H).	DMSO	>98	AQ4		
1400				HCl	¹ H NMR (400 MHz, DMSO) δ 9.77 (s, 1H), 9.63 (d, J =1.7 Hz, 1H), 9.04 - 8.85 (m, 2H), 8.63 (s, 1H), 8.12 (q, J =8.7 Hz, 2H), 7.83 (dd, J =7.8, 4.9 Hz, 1H), 7.38-7.25 (m, 2H), 7.25 - 7.17 (m, 1H), 3.92 (s, 3H).	DMSO	>98	AQ4		

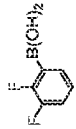
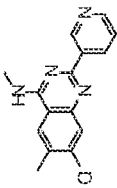
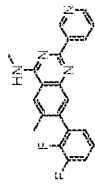
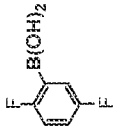
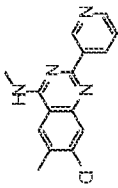
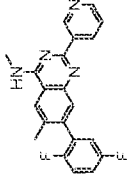
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1401				HCl	¹ H NMR (400 MHz, DMSO) δ 10.05 (s, 1H), 9.64 (d, J = 1.7 Hz, 1H), 9.04 (d, J = 7.6 Hz, 1H), 8.99-8.92 (m, 1H), 8.89 (s, 1H), 8.41 (d, J = 8.8 Hz, 1H), 8.33 - 8.20 (m, 2H), 8.16 (d, J = 8.6 Hz, 1H), 7.94 - 7.84 (m, 1H), 7.81 - 7.68 (m, 1H), 3.32 (d, J = 4.5 Hz, 3H).	DMSO	>98	AQ4		
1402				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.30 (s, 1H), 9.68 (d, J = 1.8 Hz, 1H), 9.15-9.05 (m, 1H), 8.97 (dd, J = 5.1, 1.4 Hz, 1H), 8.52 (s, 1H), 8.10 (s, 1H), 7.96 - 7.82 (m, 1H), 7.67-7.55 (m, 1H), 7.42 - 7.26 (m, 3H), 3.31 (d, J = 4.5 Hz, 3H), 2.40 (s, 3H).	DMSO	>98	AQ5		

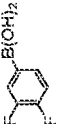
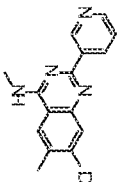
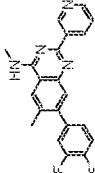
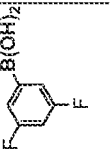
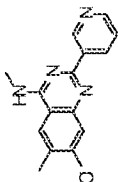
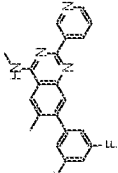
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1403				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.87 (s, 1H), 9.59 (d, J =1.5 Hz, 1H), 9.00 (m, 2H), 8.89 (m, 2H), 8.42 (s, 1H), 7.93-7.80 (m, 2H), 7.62-7.54 (m, 1H), 7.49-7.36 (m, 3H), 3.30 (d, J =4.6 Hz, 3H), 2.32 (s, 3H).	DMSO	>98	AQ5		
1404				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.19 (s, 1H), 9.68-9.61 (m, 1H), 9.05-8.98 (m, 1H), 8.94 (dd, J =5.0, 1.6 Hz, 1H), 8.48 (d, J =5.7 Hz, 1H), 8.02 (s, 1H), 7.91-7.82 (m, 1H), 7.59-7.50 (m, 2H), 7.43-7.36 (m, 2H), 3.31 (d, J =4.6 Hz, 3H), 2.40 (s, 3H).	DMSO	>98	AQ5		

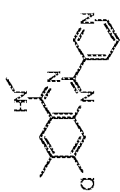
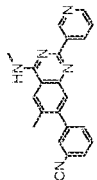
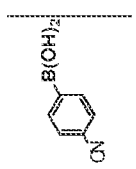
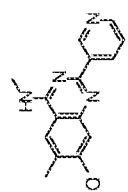
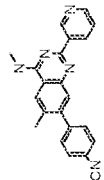
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1405				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.11 (s, 1H), 9.69 (m, 1H), 9.61 (m, 1H), 9.11-9.02 (m, 1H), 8.95 (dd, 1H), 8.52 (s, 1H), 8.05 (s, 1H), 7.89 (dd, 1H), 7.81 (dd, 1H), 7.48 (m, 1H), 7.38 (m, 1H), 7.33-7.25 (m, 1H), 3.31 (d, 1H), 4.6 Hz, 3H), 2.33 (s, 3H).	DMSO	>98	AQ5		
1406				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.03 (s, 1H), 9.63 (dd, 1H), 9.63 (dd, 1H), 9.09 (m, 1H), 8.95 (dd, 1H), 8.95 (dd, 1H), 8.95 (dd, 1H), 8.52 (s, 1H), 8.05 (s, 1H), 7.89 (dd, 1H), 7.81 (dd, 1H), 7.48 (m, 1H), 7.38 (m, 1H), 7.33-7.25 (m, 1H), 3.31 (d, 1H), 4.6 Hz, 3H), 2.33 (s, 3H).	DMSO	>98	AQ5		

(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1407				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.09 (s, 1H), 9.63 (d, J = 1.6 Hz, 1H), 9.08-8.98 (m, 1H), 8.95 (dd, J = 5.1, 1.5 Hz, 1H), 8.46 (s, 1H), 7.98 (s, 1H), 7.87 (dd, J = 8.1, 5.1 Hz, 1H), 7.64 (ddt, J = 16.8, 14.3, 5.3 Hz, 2H), 7.40-7.30 (m, 1H), 3.30 (d, J = 4.6 Hz, 3H), 2.41 (s, 3H).	DMSO	>98	AQ5		
1408				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.92 (s, 1H), 9.61 (d, J = 1.6 Hz, 1H), 9.03-8.97 (m, 1H), 8.94 (dd, J = 5.1, 1.5 Hz, 1H), 8.42 (s, 1H), 7.93 (s, 1H), 7.87 (dd, J = 8.1, 5.0 Hz, 1H), 7.44-7.36 (m, 1H), 7.33-7.23 (m, 2H), 3.29 (d, J = 4.6 Hz, 3H), 2.41 (s, 3H).	DMSO	>98	AQ5		

(continued)

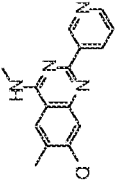
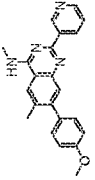
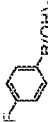
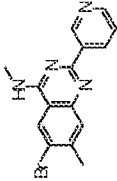
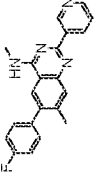
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1409				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.11 (s, 1H), 9.64 (d, J = 2.2 Hz, 1H), 9.10-9.03 (m, 1H), 8.96 (dd, J = 5.1, 1.5 Hz, 1H), 8.49 (s, 1H), 8.05-7.94 (m, 3H), 7.94-7.88 (m, 1H), 7.88-7.82 (m, 1H), 7.77 (t, J = 7.7 Hz, 1H), 3.31 (d, J = 4.5 Hz, 3H), 2.40 (s, 3H).	DMSO	>98	AQ5		
1410				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.22 (s, 1H), 9.67 (d, J = 1.6 Hz, 1H), 9.14-9.07 (m, 1H), 8.97 (dd, J = 5.1, 1.5 Hz, 1H), 8.52 (s, 1H), 8.10-8.00 (m, 3H), 7.91 (dd, J = 7.9, 5.3 Hz, 1H), 7.75-7.67 (m, 2H), 3.31 (d, J = 4.5 Hz, 3H), 2.39 (s, 3H).	DMSO	>98	AQ5		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1411				2 HCl	¹ H NMR (400 MHz, DMSO- <i>d</i> ₆) δ 10.20-9.85 (m, 1H), 9.58 (s, 1H), 8.94-8.85 (m, 2H), 8.38 (s, 1H), 7.88-7.76 (m, 2H), 7.54-7.44 (m, 1H), 7.24-7.16 (m, 2H), 7.12 (td, 2H), 7.4, 0.9 Hz, 1H), 3.76 (s, 3H), 3.31 (d, J=4.5 Hz, 3H), 2.24 (s, 3H).	DMSO	>98%	AQ5		

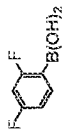
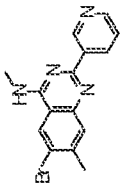
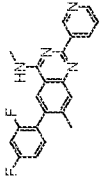
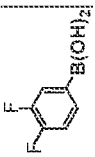
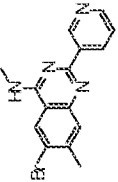
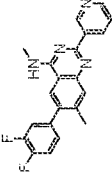
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1412				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.20 (s, 1H), 9.69 - 9.61 (m, 1H), 9.01 (d, J = 7.8 Hz, 1H), 8.93 (dd, J = 5.0, 1.6 Hz, 1H), 8.49 (s, 1H), 8.06 (s, 1H), 7.83 (dd, J = 8.0, 5.1 Hz, 1H), 7.51-7.41 (m, 1H), 7.13 - 7.05 (m, 1H), 7.05-6.95 (m, 2H), 3.81 (d, J = 10.8 Hz, 3H), 3.31 (d, J = 4.6 Hz, 3H), 2.41 (s, 3H).	DMSO	>98%	AQ5		

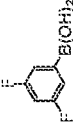
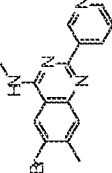
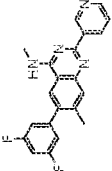
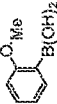
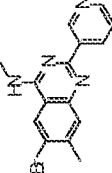
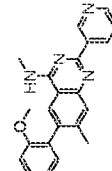
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1413				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.09 (s, 1H), 9.60 (s, 1H), 9.00 - 8.88 (m, 2H), 8.42 (s, 1H), 7.94 (s, 1H), 7.86 - 7.78 (m, 1H), 7.48 - 7.39 (m, 2H), 7.16 - 7.06 (m, 2H), 3.85 (s, 3H), 3.31 (d, J = 4.6 Hz, 3H), 2.43 (s, 3H).	DMSO	>98%	AQ5		
1414				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.02 (s, 1H), 9.63 (s, 1H), 9.05 - 8.91 (m, 2H), 8.36 (s, 1H), 8.10 - 7.99 (m, 1H), 7.91 - 7.80 (m, 1H), 7.59 - 7.50 (m, 2H), 7.44 - 7.32 (m, 2H), 3.27 (d, J = 4.5 Hz, 3H), 2.45 (s, 3H).	DMSO	>98%	AQ5		

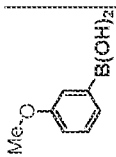
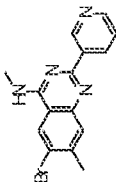
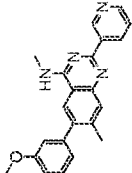
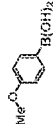
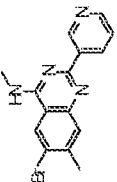
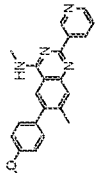
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1415				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.01 (s, 1H), 9.65 (s, 1H), 9.10 - 8.99 (m, 1H), 8.97 (d, J = 5.0 Hz, 1H), 8.41 (s, 1H), 8.09 (s, 1H), 7.97 - 7.83 (m, 1H), 7.61 - 7.41 (m, 2H), 7.31 (td, J = 8.4, 2.2 Hz, 1H), 3.26 (d, J = 4.5 Hz, 3H), 2.35 (s, 3H).	DMSO	>98	AQ5		
1416				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.01 - 9.69 (m, 1H), 9.61 (s, 1H), 8.94 (m, J = 5.0 Hz, 2H), 8.36 (s, 1H), 7.99 (s, 1H), 7.92 - 7.81 (m, 1H), 7.71 - 7.57 (m, 2H), 7.41 - 7.31 (m, 1H), 3.26 (d, J = 4.5 Hz, 3H), 2.47 (s, 3H).	DMSO	>98	AQ5		

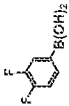
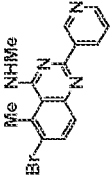
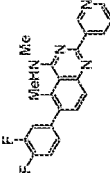
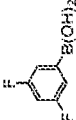
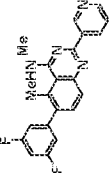
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1417				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.90 (s, 1H), 9.63 (d, J = 1.8 Hz, 1H), 9.08-8.89 (m, 2H), 8.40 (s, 1H), 8.02 (s, 1H), 7.92-7.81 (m, 1H), 7.44-7.25 (m, 3H), 3.27 (d, J = 4.5 Hz, 3H), 2.49 (s, 3H).	DMSO	>98	AQ5		
1418				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.99 (s, 1H), 9.62 (s, 1H), 8.95 (d, J = 5.0 Hz, 2H), 8.27 (s, 1H), 8.11-7.99 (m, 1H), 7.89-7.80 (m, 1H), 7.53-7.45 (m, 1H), 7.24 (dd, J = 7.4, 1.7 Hz, 1H), 7.19 (d, J = 8.0 Hz, 1H), 7.12 (td, J = 7.4, 0.9 Hz, 1H), 3.74 (d, J = 8.9 Hz, 3H), 3.26 (d, J = 4.6 Hz, 3H), 2.28 (s, 3H).	DMSO	>98	AQ5		

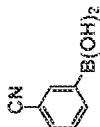
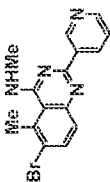
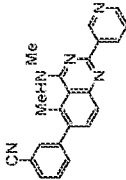
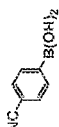
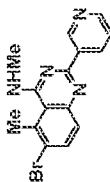
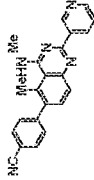
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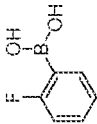
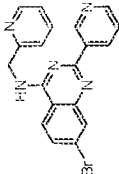
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1419				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.30 (s, 1H), 9.72 (d, J = 1.6 Hz, 1H), 9.20-9.10 (m, 1H), 8.98 (dd, J = 5.1, 1.5 Hz, 1H), 8.45 (s, 1H), 8.21 (s, 1H), 7.91 (dd, J = 8.0, 5.1 Hz, 1H), 7.48 - 7.40 (m, 1H), 7.08-7.01 (m, 3H), 3.84 (s, 3H), 3.26 (d, J = 4.5 Hz, 3H), 2.45 (s, 3H).	DMSO	>98	AQ5		
1420				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.08 - 9.71 (m, 1H), 9.58 (d, J = 2.0 Hz, 1H), 8.96 - 8.83 (m, 2H), 8.31 (s, 1H), 7.94 (s, 1H), 7.87 - 7.79 (m, 1H), 7.47 - 7.39 (m, 2H), 7.10 (d, J = 8.8 Hz, 2H), 3.84 (s, 3H), 3.27 (d, J = 4.5 Hz, 3H), 2.47 (s, 3H).	DMSO	>98	AQ5		

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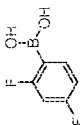
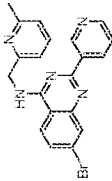
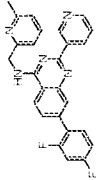
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1421				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.69 - 9.63 (m, 1H), 9.10 - 8.93 (m, 3H), 8.12 (d, J = 8.4 Hz, 1H), 7.90 - 7.82 (m, 2H), 7.68 - 7.58 (m, 1H), 7.56 - 7.48 (m, 1H), 7.30 - 7.22 (m, 1H), 3.35 (d, J = 4.6 Hz, 3H), 2.73 (s, 3H).	DMSO	>98	AQ4		
1422				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.62 (s, 1H), 8.95 (d, J = 5.0 Hz, 3H), 8.03 (s, 1H), 7.85 (t, J = 9.9 Hz, 2H), 7.44 - 7.34 (m, 1H), 7.17 (dd, J = 8.2, 2.2 Hz, 2H), 3.34 (d, J = 4.5 Hz, 3H), 2.73 (s, 3H).	DMSO	>98	AQ4		

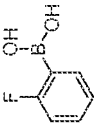
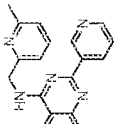
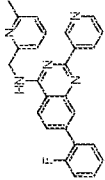
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1423				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.62 (d, J = 2.1 Hz, 1H), 9.00-8.75 (m, 3H), 8.08 - 7.99 (m, 1H), 7.99-7.93 (m, 1H), 7.93 - 7.79 (m, 3H), 7.79-7.73 (m, 2H), 3.34 (d, J = 4.5 Hz, 3H), 2.70 (s, 3H).	DMSO	>98	AQ4		
1424				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.61 (s, 1H), 8.99-8.63 (m, 3H), 8.07 - 7.98 (m, 3H), 7.89-7.78 (m, 2H), 7.65 - 7.59 (m, 2H), 3.33 (d, J = 4.5 Hz, 3H), 2.70 (s, 3H).	DMSO	>98	AQ4		

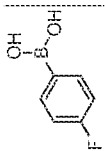
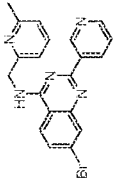
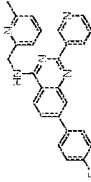
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1425				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.49 (s, 1H), 9.56 (s, 1H), 9.19 (s, 1H), 8.95 (d, J = 5.3 Hz, 1H), 8.83 (d, J = 5.5 Hz, 1H), 8.73 (d, J = 6.4 Hz, 1H), 8.41 (d, J = 5.6 Hz, 1H), 8.26 (s, 1H), 8.09 (s, 1H), 8.02-7.91 (m, 2H), 7.85 (s, 1H), 7.78-7.68 (m, 1H), 7.61-7.51 (m, 1H), 7.48-7.36 (m, 2H), 5.36 (s, 2H).	DMSO	>98	Method AQ3		

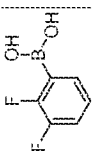
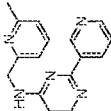
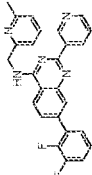
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1426				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.53 (s, 1H), 9.67 (d, J = 1.8 Hz, 1H), 9.32 (dd, J = 5.8, 4.1 Hz, 1H), 8.97 (dd, J = 5.4, 1.4 Hz, 1H), 8.75 (d, J = 8.7 Hz, 1H), 8.35 (t, J = 7.9 Hz, 1H), 8.24 (d, J = 13.5 Hz, 1H), 8.03 (dd, J = 8.1, 5.4 Hz, 1H), 7.92 (dd, J = 13.5, 4.8 Hz, 2H), 7.85 - 7.73 (m, 2H), 7.53 - 7.43 (m, 1H), 7.36 - 7.25 (m, 1H), 5.40 (d, J = 5.6 Hz, 2H), 2.85 (s, 3H).	DMSO	>98	Method AQ3		

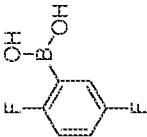
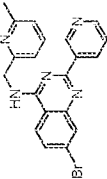
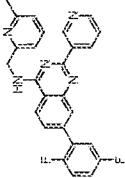
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1427				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.42 (s, 1H), 9.64 (d, J = 1.7 Hz, 1H), 9.26 (d, J = 8.1 Hz, 1H), 8.95 (dd, J = 5.3, 1.4 Hz, 1H), 8.71 (d, J = 8.7 Hz, 1H), 8.30 (d, J = 18.0 Hz, 2H), 8.04 - 7.84 (m, 3H), 7.74 (dd, J = 7.8, 6.0 Hz, 2H), 7.62 - 7.33 (m, 3H), 5.38 (d, J = 5.5 Hz, 2H), 2.84 (s, 3H).	DMSO	>98	Method AQ3		

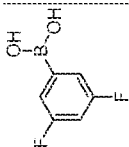
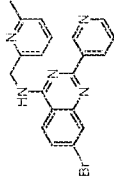
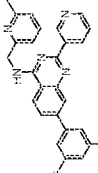
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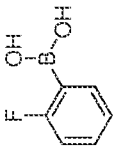
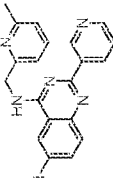
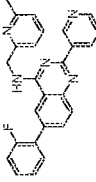
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1428				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.40 (s, 1H), 9.63 (d, J = 1.6 Hz, 1H), 9.24 (d, J = 8.1 Hz, 1H), 8.95 (dd, J = 5.3, 1.4 Hz, 1H), 8.69 (d, J = 8.7 Hz, 1H), 8.41-8.22 (m, 2H), 8.09 (dd, J = 8.7, 1.8 Hz, 1H), 8.01-7.84 (m, 4H), 7.74 (d, J = 7.8 Hz, 1H), 7.46-7.33 (m, 2H), 5.36 (d, J = 5.5 Hz, 2H), 2.83 (s, 3H).	DMSO	>98	Method AQ3		

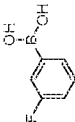
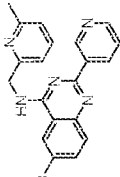
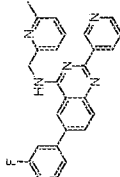
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1429				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.21 (s, 1H), 9.62 (s, 1H), 9.19 (d, <i>J</i> = 7.8 Hz, 1H), 8.93 (d, <i>J</i> = 4.6 Hz, 1H), 8.68 (d, <i>J</i> = 8.6 Hz, 1H), 8.26 (dd, <i>J</i> = 28.0, 20.1 Hz, 2H), 8.04 - 7.80 (m, 3H), 7.72 (d, <i>J</i> = 7.8 Hz, 1H), 7.64 - 7.50 (m, 2H), 7.46 - 7.33 (m, 1H), 5.33 (d, <i>J</i> = 5.3 Hz, 2H), 2.81 (s, 3H).	DMSO	>98	Method AQ3		

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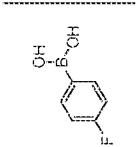
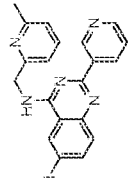
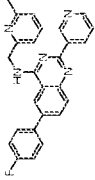
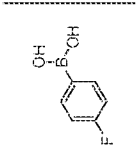
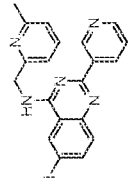
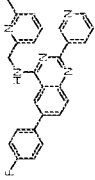
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Meth-od
1430				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.40 (s, 1H), 9.66 (d, J= 1.8 Hz, 1H), 9.31 (d, J= 8.2 Hz, 1H), 8.96 (dd, J= 5.4, 1.4 Hz, 1H), 8.73 (d, J= 8.7 Hz, 1H), 8.33 (t, J= 7.9 Hz, 1H), 8.23 (s, 1H), 8.03 (dd, J= 8.1, 5.4 Hz, 1H), 7.93 (dd, J= 11.7, 8.3 Hz, 2H), 7.76 (d, J= 7.9 Hz, 1H), 7.70 - 7.60 (m, 1H), 7.54-7.35 (m, 2H), 5.38 (d, J= 5.5 Hz, 2H), 2.85 (s, 3H).	DMSO	>98	Method AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1431				3 HCl	¹ H NMR (400 MHz, DMSO- <i>d</i> ₆) δ 10.15 (s, 1H), 9.61 (s, 1H), 9.22 (t, 1H), 9.22 (t, 1H), 8.98 - 8.90 (m, 1H), 8.65 (dd, <i>J</i> = 8.6, 3.0 Hz, 1H), 8.36 - 8.26 (m, 2H), 8.13 (dd, <i>J</i> = 8.7, 1.9 Hz, 1H), 8.05 - 7.95 (m, 1H), 7.88 (d, <i>J</i> = 7.9 Hz, 1H), 7.76 - 7.64 (m, 3H), 7.43 - 7.33 (m, 1H), 5.33 (s, 2H), 2.82 (s, 3H).	DMSO	>98	Method AQ3		

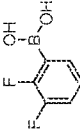
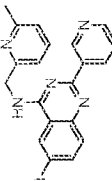
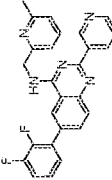
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1432				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.26 (s, 1H), 9.62 (d, J = 1.8 Hz, 1H), 9.23 (d, J = 8.2 Hz, 1H), 8.94 (dd, J = 5.4, 1.5 Hz, 1H), 8.78 (s, 1H), 8.30 (t, J = 7.9 Hz, 1H), 8.20-8.15 (m, 1H), 8.10 (d, J = 8.7 Hz, 1H), 8.00 (dd, J = 7.9, 5.3 Hz, 1H), 7.88 (d, J = 7.9 Hz, 1H), 7.80-7.71 (m, 2H), 7.59-7.50 (m, 1H), 7.47-7.35 (m, 2H), 5.35 (d, J = 5.5 Hz, 2H), 2.82 (s, 3H).	DMSO	>98	Method AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Meth-od
1433				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.39 (s, 1H), 9.61 (d, J = 1.7 Hz, 1H), 9.19 (d, J = 8.2 Hz, 1H), 9.01 (d, J = 1.8 Hz, 1H), 8.93 (dd, J = 5.3, 1.5 Hz, 1H), 8.38 (dd, J = 8.8, 1.9 Hz, 1H), 8.29 (t, J = 7.9 Hz, 1H), 8.09 (d, J = 8.7 Hz, 1H), 7.97 (dd, J = 8.1, 5.3 Hz, 1H), 7.92 - 7.80 (m, 3H), 7.72 (d, J = 7.8 Hz, 1H), 7.66 - 7.57 (m, 1H), 7.35 - 7.26 (m, 1H), 5.36 (d, J = 5.5 Hz, 2H), 2.81 (s, 3H).	DMSO	>98	Method AQ3		

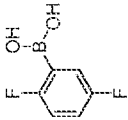
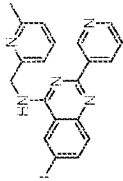
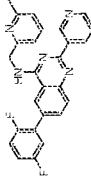
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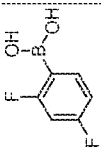
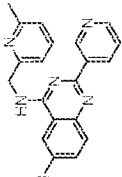
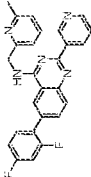
5	1434				Salt type	1H NMR 1H NMR (400 MHz, DMSO) δ 10.45 (s, 1H), 9.60 (d, J= 1.9 Hz, 1H), 9.19 (d, J= 8.2 Hz, 1H), 9.01-8.89 (m, 2H), 8.36 - 8.25 (m, 2H), 8.10 (d, J = 8.7 Hz, 1H), 8.05-7.94 (m, 3H), 7.88 (dd, J = 12.8, 5.4 Hz, 1H), 7.72 (d, J= 7.8 Hz, 1H), 7.46 - 7.35 (m, 2H), 5.36 (d, J=5.5 Hz, 2H), 2.82 (s, 3H).	1H NMR Solvent DMSO	Purity per-cent >98	Method of Coupling Method AQ3	LCMS	LCMS Method
10											
15											
20											
25											
30											
35	1434				Salt type 3 HCl	1H NMR 1H NMR (400 MHz, DMSO) δ 10.45 (s, 1H), 9.60 (d, J= 1.9 Hz, 1H), 9.19 (d, J= 8.2 Hz, 1H), 9.01-8.89 (m, 2H), 8.36 - 8.25 (m, 2H), 8.10 (d, J = 8.7 Hz, 1H), 8.05-7.94 (m, 3H), 7.88 (dd, J = 12.8, 5.4 Hz, 1H), 7.72 (d, J= 7.8 Hz, 1H), 7.46 - 7.35 (m, 2H), 5.36 (d, J=5.5 Hz, 2H), 2.82 (s, 3H).	1H NMR Solvent DMSO	Purity per-cent >98	Method of Coupling Method AQ3	LCMS	LCMS Method
40											
45											
50											
55											

(continued)

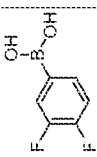
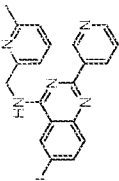
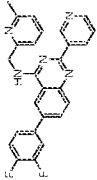
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1435				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.28 (s, 1H), 9.63 (d, J = 1.8 Hz, 1H), 9.24 (d, J = 8.2 Hz, 1H), 8.95 (d, 1H), 8.83 (s, 1H), 8.30 (t, J = 7.9 Hz, 1H), 8.19 (dd, J = 8.7, 1.8 Hz, 1H), 8.11 (d, J = 8.7 Hz, 1H), 8.00 (dd, J = 8.1, 5.4 Hz, 1H), 7.88 (d, J = 8.0 Hz, 1H), 7.73 (d, J = 7.8 Hz, 1H), 7.64-7.51 (m, 2H), 7.46-7.37 (m, 1H), 5.36 (d, J = 5.5 Hz, 2H), 2.82 (s, 3H).	DMSO	>98	Method AQ3		

(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1436				3 HCl	¹ H NMR (400 MHz, DMSO) δ 10.02 (s, 1H), 9.56 (d, J = 1.6 Hz, 1H), 9.06 (d, J = 8.0 Hz, 1H), 8.88 (dd, J = 5.2, 1.5 Hz, 1H), 8.75 (s, 1H), 8.25 - 8.13 (m, 2H), 8.04 (d, J = 8.7 Hz, 1H), 7.89 (dd, J = 7.8, 5.3 Hz, 1H), 7.79 (d, J = 8.0 Hz, 1H), 7.70 - 7.60 (m, 2H), 7.53 - 7.44 (m, 1H), 7.38 (ddd, J = 12.2, 8.4, 3.5 Hz, 1H), 5.27 (d, J = 5.3 Hz, 2H), 2.77 (s, 3H).	DMSO	>98	Method AQ3		

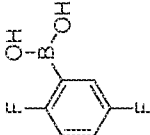
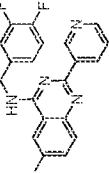
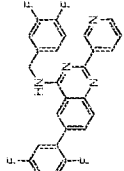
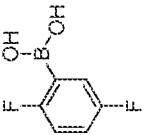
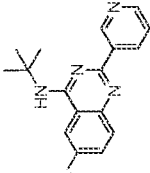
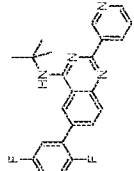
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1437				3 HCl	¹ H NMR (400 MHz, DMSO- <i>d</i> ₆) δ 10.20 (s, 1H), 9.60 (d, <i>J</i> = 1.7 Hz, 1H), 9.19 (d, <i>J</i> = 8.1 Hz, 1H), 8.93 (dd, <i>J</i> = 5.4, 1.5 Hz, 1H), 8.75 (s, 1H), 8.29 (t, <i>J</i> = 7.9 Hz, 1H), 8.17 - 8.05 (m, 2H), 7.98 (dd, <i>J</i> = 8.0, 5.4 Hz, 1H), 7.83 (ddd, <i>J</i> = 15.5, 8.4, 6.0 Hz, 2H), 7.72 (d, <i>J</i> = 7.8 Hz, 1H), 7.54 - 7.44 (m, 1H), 7.38 - 7.28 (m, 1H), 5.33 (d, <i>J</i> = 5.5 Hz, 2H), 2.81 (s, 3H).	DMSO	>98	Method AQ3		

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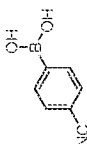
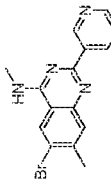
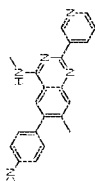
5	1438				4 HCl	¹ H NMR (400 MHz, DMSO) δ 10.45 (s, 1H), 9.62 (d, J= 1.7 Hz, 1H), 9.21 (d, J= 8.1 Hz, 1H), 9.03 (d, J= 1.7 Hz, 1H), 8.93 (dd, J = 5.3, 1.4 Hz, 1H), 8.39 -8.26(m, 2H), 8.15-8.04 (m, 2H), 8.00 - 7.81 (m, 3H), 7.76- 7.59(m, 2H), 5.36(d, J =5.5 Hz, 2H), 2.82 (s, 3H).	DMSO	>98	Method AQ3	LCMS	LCMS Method
10											
15											
20											
25											
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40											
45											
50											
55											

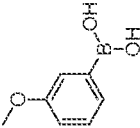
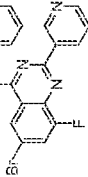
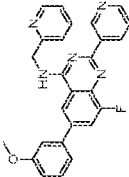
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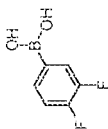
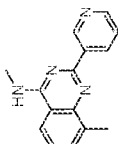
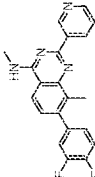
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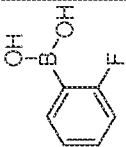
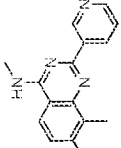
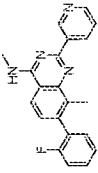
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1440				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.41 (s, 1H), 9.63 (d, J = 1.7 Hz, 1H), 9.15 (d, J = 8.1 Hz, 1H), 8.97 (dd, J = 5.2, 1.4 Hz, 1H), 8.83 (s, 1H), 8.20 (s, 1H), 7.97 (dd, J = 8.0, 5.3 Hz, 1H), 7.73 - 7.56 (m, 2H), 7.54 - 7.28 (m, 4H), 5.03 (d, J = 5.7 Hz, 2H).	DMSO	>98	Method AQ3		
1441				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.59 (d, J = 1.6 Hz, 1H), 9.07 - 8.94 (m, 2H), 8.84 (s, 1H), 8.76 (s, 1H), 8.22 - 8.13 (m, 2H), 7.96 (dd, J = 8.0, 5.3 Hz, 1H), 7.66 (ddd, J = 9.2, 6.1, 3.2 Hz, 1H), 7.53 - 7.32 (m, 2H), 1.69 (s, 9H).	DMSO	>98	Method AQ3		

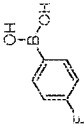
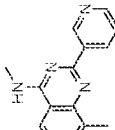
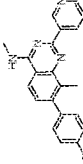
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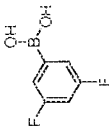
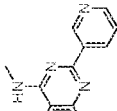
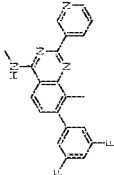
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
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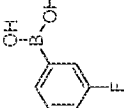
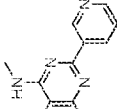
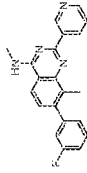
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1444				2 HCl	¹ H NMR (400 MHz, DMSO) δ 9.76 - 9.69 (m, 1H), 9.37 (t, J = 5.4 Hz, 1H), 8.99 (dd, J = 11.7, 7.5 Hz, 1H), 8.81 (dt, J = 7.9, 4.0 Hz, 1H), 8.70 (t, J = 7.3 Hz, 1H), 8.52 (d, J = 9.2 Hz, 1H), 8.35-8.27 (m, 1H), 8.10 (dd, J = 12.8, 7.3 Hz, 1H), 8.01 Hz, 1H), 8.01 - 7.90 (m, 2H), 7.77 - 7.71 (m, 1H), 7.50 - 7.37 (m, 3H), 7.06 - 7.01 (m, 1H), 5.19 (t, J = 17.9 Hz, 2H), 3.88 (s, 3H).	DMSO	>98	Method AQ3, F, G2 (reflux)		

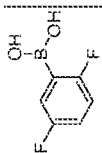
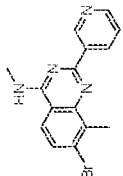
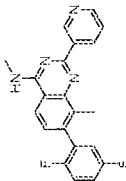
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1445				2 HCl	¹ H NMR (DM-SO-d6) ppm 9.73 (s, 1H), 9.37 (brd, J = 8.08 Hz, 1H), 8.97 (brd, J = 5.24 Hz, 1H), 8.77 (brs, 1H), 8.20 (d, J = 8.48 Hz, 1H), 8.10-8.07 (brm, 1H), 7.63-7.55 (brm, 2H), 7.47 (d, J = 8.48 Hz, 1H), 7.32 (brm, 1H), 3.20 (d, J = 4.20 Hz, 3H), 2.65 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	AQ6		

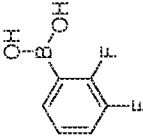
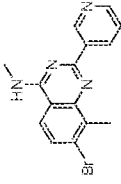
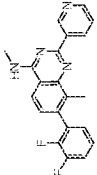
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1446				3 HCl	¹ H NMR (DMF-d6) ppm 9.73 (d, J = 1.76 Hz, 1H), 9.39 (brd, J = 8.16 Hz, 1H), 8.98 (dd, J = 5.44, 1.20 Hz, 1H), 8.80 (brs, 1H), 8.22 (d, J = 8.48 Hz, 1H), 8.12-8.08 (brm, 1H), 7.57-7.51 (m, 1H), 7.46-7.35 (m, 4H), 3.21 (d, J = 4.32 Hz, 3H), 2.55 (s, 3H). The 1H of 3HCl was not observed.	DMSO	>98	AQ6		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1447				3 HCl	¹ H NMR (DM-SO-d6) ppm 9.72 (d, J = 1.68 Hz, 1H), 9.26 (brd, J = 8.48 Hz, 1H), 8.91 (dd, J = 5.28, 1.40 Hz, 1H), 8.69 (brs, 1H), 8.18 (d, J = 8.56 Hz, 1H), 8.00-7.96 (brm, 1H), 7.53-7.49 (m, 2H), 7.44 (d, J = 8.56 Hz, 1H), 7.38-7.33 (m, 2H), 3.20 (d, J = 4.40 Hz, 3H), 2.63 (s, 3H). The 1H of 3HCl was not observed.	DMSO	>98	AO6		

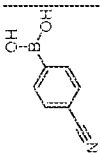
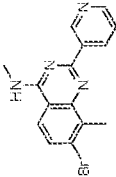
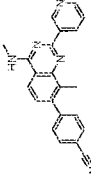
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Meth-od
1448				2 HCl	¹ H NMR (DM-SO-d6) ppm 9.73 (d, J = 1.44 Hz, 1H), 9.36 (brd, J = 7.88 Hz, 1H), 8.96 (d, J = 5.04 Hz, 1H), 8.77 (brs, 1H), 8.21 (d, J = 8.56 Hz, 1H), 8.07 (brn, 1H), 7.48 (d, J = 8.56 Hz, 1H), 7.38-7.32 (m, 1H), 7.27-7.22 (m, 2H), 3.20 (d, J = 4.4 Hz, 3H), 2.65 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	AQ6		

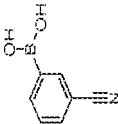
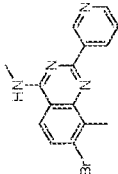
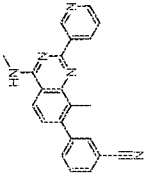
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1449				3 HCl	¹ H NMR (DMF-d6) ppm 9.73 (d, J = 1.72 Hz, 1H), 9.32(brd, J = 7.56 Hz, 1H), 8.94 (brd, J = 5.48 Hz, 1H), 8.74 (brs, 1H), 8.20 (d, J = 8.56 Hz, 1H), 8.08-8.02 (m, 1H), 7.60-7.54 (m, 1H), 7.47(d,J = 7.35-7.27 (m, 3H), 3.20 (d, J = 4.4 Hz, 3H), 2.65 (s, 3H). The 1H of 3HCl was not observed.	DMSO	>98	AQ6		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1450				2 HCl	¹ H NMR (DM-SO-d6) ppm 9.74 (d, J = 1.76 Hz, 1H), 9.37 (brd, J = 8.16 Hz, 1H), 8.97 (brdd, J = 5.44, 1.24 Hz, 1H), 8.79 (brs, 1H), 8.22 (d, J = 8.52 Hz, 1H), 8.10-8.06 (brm, 1H), 7.48-7.35 (m, 4H), 3.21 (d, J = 4.36 Hz, 3H), 2.56 (brs, 3H). The 1H of 2HCl was not observed.	DMSO	>98	AO6		

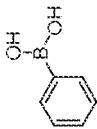
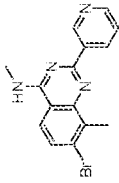
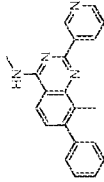
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1451				3 HCl	¹ H NMR (DM-SO-d6) ppm 9.74 (d, J = 1.52 Hz, 1H), 9.37 (d, J = 8.08 Hz, 1H), 8.97 (dd, J = 5.54, 1.16 Hz, 1H), 8.80 (brs, 1H), 8.23 (d, J = 8.52 Hz, 1H), 8.10-8.07 (m, 1H), 7.60-7.53 (m, 1H), 7.48 (d, J = 8.52 Hz, 1H), 7.41-7.36 (m, 1H), 7.30-7.26 (m, 1H), 3.21 (d, J = 4.36 Hz, 3H), 2.56 (s, 3H). The 1H of 3HCl was not observed.	DMSO	>98	AQ6		

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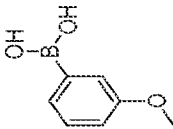
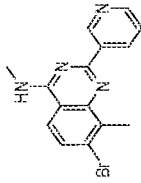
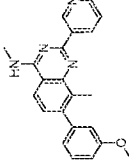
5	1452	Starting Material 1		Starting Material 2		Product		Salt type	3 HCl	¹H NMR 1H NMR (DM- SO-d6) ppm 9.73 (d, J = 1.68 Hz, 1H), 9.26 (brd, J = 7.8 Hz, 1H), 8.91 (brdd, J = 5.28, 1.36 Hz, 1H), 8.71 (brs, 1H), 8.21 (d, J = 8.52 Hz, 1H), 8.00 (d, J = 8.4 Hz, 2H), 7.99-7.95 (m, 1H), 7.69 (d, 8.4 Hz, 2H), 7.46 (d, J = 8.52 Hz, 1H), 3.20 (d, J = 4.4 Hz, 3H), 2.63 (s, 3H). The 1H of 3HCl was not observed.	¹H NMR Solvent DMSO	Purity per- cent >98	Method of Coupling AQ6	LCMS	LCMS Meth- od
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1453				2 HCl	¹ H NMR (DMF-d ₆) ppm 9.73 (d, J = 1.72 Hz, 1H), 9.33 (brd, J = 7.64 Hz, 1H), 8.95 (brd, J = 5.28 Hz, 1H), 8.76 (brs, 1H), 8.22 (d, J = 8.48 Hz, 1H), 8.05 (brm, J = 7.64, 5.28 Hz, 1H), 7.98-7.93 (m, 2 H), 7.84-7.81 (m, 1H), 7.74 (dd, J = 7.84, 7.76 Hz, 1H), 7.49 (d, J = 8.84 Hz, 1H), 3.21 (d, J = 4.4 Hz, 3H), 2.63 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	AQ6		

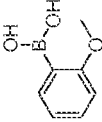
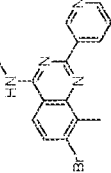
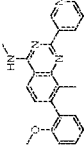
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1454				2 HCl	¹ H NMR (DM-SO-d ₆) ppm 9.72 (d, J = 1.76 Hz, 1H), 9.42 (brd, J = 8.00 Hz, 1H), 9.00 (d, J = 5.52 Hz, 1H), 8.83 (bsr, 1H), 8.22 (d, J = 8.56 Hz, 1H), 8.13 (brm, 1H), 7.55-7.51 (m, 2H), 7.48-7.43 (m, 4H), 3.21 (brd, J = 4.16 Hz, 3H), 2.64 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	AQ6		

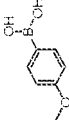
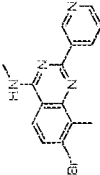
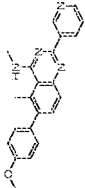
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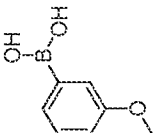
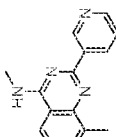
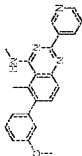
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1456				2 HCl	¹ H NMR (DM-SO-d ₆) ppm 9.72 (d, J = 1.64 Hz, 1H), 9.38 (d, J = 8.04 Hz, 1H), 8.98 (dd, J = 5.48; 1.24 Hz, 1H), 8.79 (brs, 1H), 8.20 (d, J = 8.52 Hz, 1H), 8.12-8.09 (m, 1H), 7.48-7.42 (m, 2H), 7.04-6.97 (m, 3H), 3.82 (s, 3H), 3.21 (d, J = 4.28 Hz, 3H), 2.65 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	AQ6		

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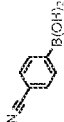
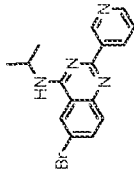
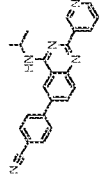
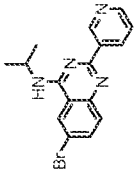
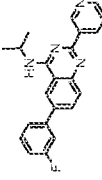
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1457				2 HCl	¹ H NMR (DM-SO-d ₆) ppm 9.71 (d, J = 1.32 Hz, 1H), 9.40 (d, J = 7.64 Hz, 1H), 8.99 (dd, J = 5.52, 1.16 Hz, 1H), 8.82 (brs, 1H), 8.17 (d, J = 8.48 Hz, 1H), 8.13 (m, 1H), 7.47-7.43 (m, 1H), 7.37 (d, J = 8.48 Hz, 1H), 7.22-7.16 (m, 2H), 7.11-7.07 (m, 1H), 3.73 (s, 3H), 3.21 (d, J = 4.12 Hz, 3H), 2.47 (s, 3H). The 1H of 2HCl was not observed.	DMSO	>98	AQ6		

(continued)

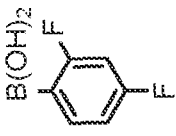
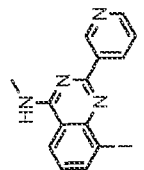
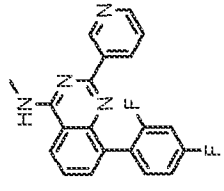
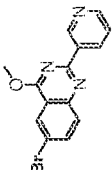
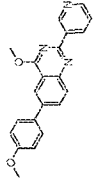
5	1458	Starting Material 1		Starting Material 2		Product		Salt type	2 HCl	¹ H NMR	¹ H NMR (DM-SO-d ₆) ppm 9.57 (d, J = 1.88 Hz, 1H), 8.93 (dd, J = 4.88, 1.56 Hz, 1H), 8.87 (brd, J = 6.52 Hz, 1H), 7.95 (d, J = 8.36 Hz, 1H), 7.84 (d, J = 8.36 Hz, 1H), 7.81 (m, 1H), 7.34 (d, J = 8.72 Hz, 2H), 7.11 (d, J = 8.72 Hz, 2H), 3.84 (s, 3H), 3.34 (d, J = 4.56 Hz, 3H), 2.72 (s, 3H). The 1H of 2HCl and NH- were not observed.	¹ H NMR Solvent	DMSO	Purity per-cent	>98	Method of Coupling	AQ6	LCMS	LCMS Meth-od
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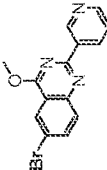
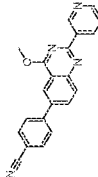
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1459				2 HCl	¹ H NMR (DM-SO-d ₆) ppm 9.59 (s, 1H), 8.94 (dd, J = 4.92, 1.48 Hz, 1H), 8.89 (br, 1H), 7.98 (d, J = 8.48 Hz, 1H), 7.86 (d, J = 8.48 Hz, 1H), 7.82 (m, 1H), 7.46 (t, J = 7.96 Hz, 1H), 7.06-7.03 (m, 1H), 6.98-6.94 (m, 2H), 3.83 (s, 3H), 3.34 (d, J = 4.56 Hz, 3H), 2.72 (s, 3H). The 1H of 2HCl and NH- were not observed.	DMSO	>98	AQ6		

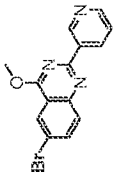
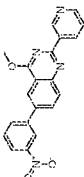
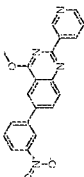
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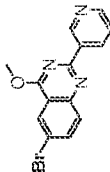
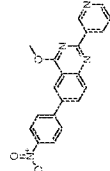
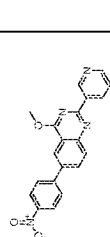
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1460					¹ H NMR (300 MHz, DMSO) δ 9.60 (s, 1H), 8.84-8.57 (m, 3H), 8.30 (d, J = 7.0 Hz, 1H), 8.18 (d, J = 8.8 Hz, 1H), 8.12-7.93 (m, 4H), 7.85 (dd, J = 8.7, 1.5 Hz, 1H), 7.60-7.46 (m, 1H), 4.85-4.58 (m, 1H), 1.37 (d, J = 6.5 Hz, 6H).	DMSO	100	AQ1	366 (M+1)	Method A (Formic add)
1461				HCl	¹ H NMR (300 MHz, DMSO) δ 9.81 (s, 1H), 9.60 (s, 1H), 8.95 (s, 3H), 8.39 (d, J = 8.9 Hz, 1H), 8.19 (s, 1H), 7.96-7.73 (m, 3H), 7.68-7.55 (m, 1H), 7.39-7.25 (m, 1H), 4.99-4.81 (m, 1H), 1.42 (d, J = 6.5 Hz, 6H).	DMSO	100	AQ1	359 (M+1)	Method A (Formic add)

(continued)

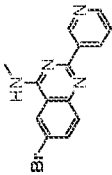
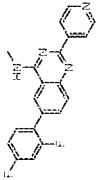
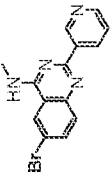
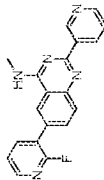
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1462					¹ H NMR (300 MHz, DMSO) δ 9.43 (s, 1H), 8.75-8.47 (m, 3H), 8.29 (d, J = 7.2 Hz, 1H), 7.79 (d, J = 7.1 Hz, 1H), 7.72-7.54 (m, 2H), 7.48 (dd, J = 7.8, 4.9 Hz, 1H), 7.44-7.32 (m, 1H), 7.30-7.12 (m, 1H), 3.32 (s, 3H), 3.16 (d, J = 4.3 Hz, 3H).	DMSO	100	AQ1	349 (M+1)	Method A (Formic add)
1463					¹ H NMR (300 MHz, DMSO) δ 9.68 (dd, J = 2.1, 0.8 Hz, 1H), 8.86-8.78 (m, 1H), 8.74 (dd, J = 4.8, 1.7 Hz, 1H), 8.34-8.25 (m, 2H), 8.09-8.02 (m, 1H), 7.80 (d, J = 8.9 Hz, 2H), 7.60 (ddd, J = 8.0, 4.8, 0.8 Hz, 1H), 7.09 (d, J = 8.9 Hz, 2H), 4.31 (s, 3H), 3.83 (s, 3H).	DMSO	99	Method AQ2	344.0 (M+1)	Method C

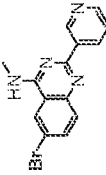
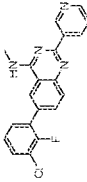
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1464					¹ H NMR (300 MHz, DMSO) δ 9.67 (d, J = 2.1 Hz, 1H), 8.82 (d, J = 7.9 Hz, 1H), 8.74 (dd, J = 4.8, 1.7 Hz, 1H), 8.46 (d, J = 1.7 Hz, 1H), 8.36 (dd, J = 8.8, 2.2 Hz, 1H), 8.15 - 7.94 (m, 5H), 7.60 (ddd, J = 8.0, 4.8, 0.8 Hz, 1H), 4.31 (s, 3H).	DMSO	99	Method AQ2	338.9 (M+1)	Method C

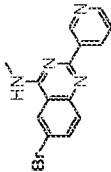
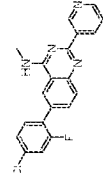
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Meth-od
1465				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.71 (d, J = 1.8 Hz, 1H), 9.15 (d, J = 8.0 Hz, 1H), 8.90 (dd, J = 5.2, 1.4 Hz, 1H), 8.58 (s, J = 1.8 Hz, 1H), 8.48 (d, J = 1.6 Hz, 1H), 8.43 (dd, J = 8.7, 2.2 Hz, 1H), 8.32 (dd, J = 7.8, 0.9 Hz, 1H), 8.28 (dd, J = 8.2, 2.2 Hz, 1H), 8.14 (d, J = 8.7 Hz, 1H), 7.92 (dd, J = 8.0, 5.2 Hz, 1H), 7.81 (t, J = 8.0 Hz, 1H), 4.34 (s, 3H).	DMSO	99	Method AQ2	359.0 (M+1)	Method C

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1466				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.71 (d, J = 2.0 Hz, 1H), 9.12 (dd, J = 8.0, 1.4 Hz, 1H), 8.89 (dd, J = 5.2, 1.5 Hz, 1H), 8.49 (d, J = 1.6 Hz, 1H), 8.42 (dd, J = 8.8, 2.2 Hz, 1H), 8.34 (d, J = 9.0 Hz, 2H), 8.14 (dd, J = 9.2, 2.4 Hz, 3H), 7.88 (dd, J = 7.8, 5.5 Hz, 1H), 4.34 (s, 3H).	DMSO	99	Method AQ2	359.0 (M+1)	Method C

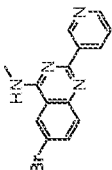
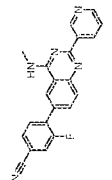
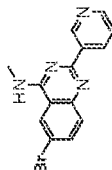
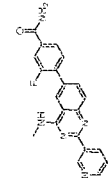
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1467				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.36 (s, 1H), 9.00 (d, J = 6.5 Hz, 2H), 8.78 (d, J = 4.5 Hz, 2H), 8.58 (s, 1H), 8.06 (s, 2H), 7.75 (dd, J = 15.6, 8.9 Hz, 1H), 7.47 (ddd, J = 11.7, 9.4, 2.6 Hz, 1H), 7.31 (td, J = 8.4, 2.7 Hz, 1H), 3.22 (d, J = 4.2 Hz, 3H).	DMSO	99	Method AQ2	349.6 (M+1)	Method C
1468				3 HCl	¹ H NMR (300 MHz, DMSO) δ 9.97 (s, 1H), 9.63 (d, J = 1.6 Hz, 1H), 9.04 (d, J = 7.0 Hz, 1H), 8.94 (dd, J = 5.1, 1.5 Hz, 1H), 8.76 (d, J = 0.7 Hz, 1H), 8.41-8.05 (m, 4H), 7.89 (dd, J = 7.1, 4.8 Hz, 1H), 7.59 (ddd, J = 7.1, 4.9, 1.8 Hz, 1H), 3.27 (d, J = 4.3 Hz, 3H).	DMSO	99	Method AQ2	332.4 (M+1)	Method C

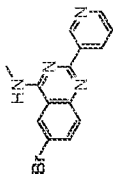
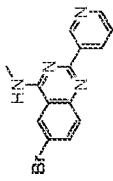
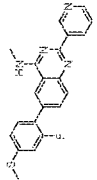
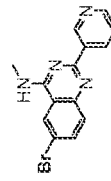
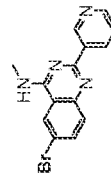
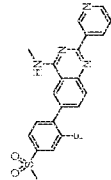
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1469				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.04 (s, 1H), 9.65 (d, J = 1.5 Hz, 1H), 9.06 (d, J = 7.3 Hz, 1H), 8.95 (dd, J = 5.1, 1.5 Hz, 1H), 8.71 (s, 1H), 8.19 (s, 2H), 7.89 (dd, J = 7.6, 5.1 Hz, 1H), 7.76 - 7.61 (m, 2H), 7.42 (td, J = 7.9, 0.9 Hz, 1H), 3.27 (d, J = 4.4 Hz, 3H).	DMSO	95	Method AQ2	365.3 (M+1)	Method C

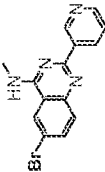
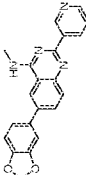
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1470				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.03 (s, 1H), 9.63 (d, J = 1.5 Hz, 1H), 9.04 (d, J = 7.9 Hz, 1H), 8.94 (dd, J = 5.1, 1.5 Hz, 1H), 8.68 (s, 1H), 8.16 (s, 2H), 7.88 (dd, J = 7.6, 5.1 Hz, 1H), 7.74 (t, J = 8.5 Hz, 1H), 7.67 (dd, J = 10.8, 2.0 Hz, 1H), 7.51 (dd, J = 8.3, 1.6 Hz, 1H), 3.27 (d, J = 4.4 Hz, 3H).	DMSO	94	Method AQ2	365.3 (M+1)	Method C

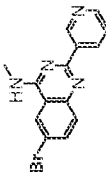
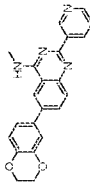
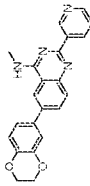
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1471				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.95 (s, 1H), 9.64 (d, J = 1.6 Hz, 1H), 9.07 (d, J = 7.9 Hz, 1H), 8.94 (dd, J = 5.1, 1.5 Hz, 1H), 8.75 (s, 1H), 8.18 (s, 2H), 8.09 (d, J = 11.1 Hz, 1H), 7.92 (d, J = 4.0 Hz, 3H), 3.27 (d, J = 4.4 Hz, 3H).	DMSO	96	Method AQ2	356.4 (M+1)	Method C
1472				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.01 (s, 1H), 9.63 (d, J = 1.5 Hz, 1H), 9.03 (d, J = 7.8 Hz, 1H), 8.94 (dd, J = 5.1, 1.5 Hz, 1H), 8.73 (s, 1H), 8.25 - 8.12 (m, 3H), 7.93 - 7.76 (m, 4H), 7.63 (d, J = 0.8 Hz, 1H), 3.28 (d, J = 4.4 Hz, 3H).	DMSO	95	Method AQ2	374.4 (M+1)	Method C

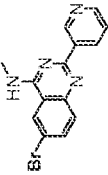
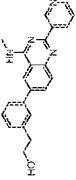
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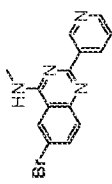
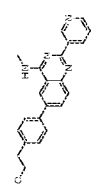
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1473				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.97 (s, 1H), 9.61 (d, J = 2.1 Hz, 1H), 8.98 (d, J = 4.6 Hz, 1H), 8.93 (dd, J = 5.1, 1.5 Hz, 1H), 8.68 - 8.60 (m, 1H), 8.13 (d, J = 6.1 Hz, 2H), 7.85 (dd, J = 7.7, 5.2 Hz, 1H), 7.63 (t, J = 8.3 Hz, 1H), 7.37 - 7.23 (m, 2H), 3.27 (d, J = 4.5 Hz, 3H), 2.56 (s, J = 5.2 Hz, 3H).	DMSO	95	Method AQ2	377.5 (M+1)	Method C
1474				2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.98 (s, 1H), 9.64 (d, J = 1.6 Hz, 1H), 9.06 (d, J = 8.8 Hz, 1H), 8.94 (dd, J = 5.1, 1.5 Hz, 1H), 8.75 (s, 1H), 8.26 - 8.11 (m, 2H), 8.04 - 7.85 (m, 5H), 3.35 (s, 3H), 3.27 (d, J = 4.1 Hz, 3H).	DMSO	96	Method AQ2	409.5 (M+1)	Method C

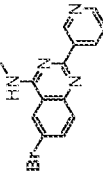
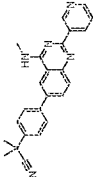
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1475				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.25 (s, 1H), 9.63 (d, J = 1.6 Hz, 1H), 9.00 (d, J = 7.1 Hz, 1H), 8.93 (dd, J = 5.0, 1.5 Hz, 1H), 8.79 (s, 1H), 8.31 (dd, J = 8.8, 1.5 Hz, 1H), 8.17 (d, J = 8.5 Hz, 1H), 7.84 (dd, J = 7.5, 4.8 Hz, 1H), 7.51 (d, J = 1.8 Hz, 1H), 7.40 (dd, J = 8.2, 1.9 Hz, 1H), 7.09 (d, J = 8.1 Hz, 1H), 6.10 (s, 2H), 3.29 (d, J = 4.3 Hz, 3H).	DMSO	99	Method AQ2	357.5 (M+1)	Method C

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1476				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.26 (s, 1H), 9.63 (d, J = 1.6 Hz, 1H), 9.01 (d, J = 8.6 Hz, 1H), 8.93 (dd, J = 5.0, 1.5 Hz, 1H), 8.77 (s, 1H), 8.31 (dd, J = 8.8, 1.4 Hz, 1H), 8.18 (d, J = 8.6 Hz, 1H), 7.84 (dd, J = 6.9, 5.8 Hz, 1H), 7.44 (d, J = 2.2 Hz, 1H), 7.39 (dd, J = 8.4, 2.3 Hz, 1H), 7.01 (d, J = 8.4 Hz, 1H), 4.30 (s, 4H), 3.29 (d, J = 4.4 Hz, 3H).	DMSO	99	Method AQ2	371.5 (M+1)	Method C

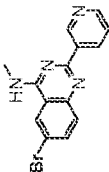
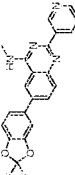
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1477				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.19 (s, 1H), 9.61 (d, J = 1.6 Hz, 1H), 8.98 (d, J = 8.8 Hz, 1H), 8.93 (dd, J = 5.0, 1.5 Hz, 1H), 8.82 (s, 1H), 8.35 (dd, J = 8.8, 1.3 Hz, 1H), 8.16 (d, J = 8.5 Hz, 1H), 7.85 (dd, J = 8.1, 5.2 Hz, 1H), 7.76 (m, 2H), 7.66 (m, 2H), 7.46 (t, J = 7.6 Hz, 1H), 7.31 (d, J = 7.7 Hz, 1H), 3.69 (t, J = 7.0 Hz, 2H), 3.31 (d, J = 4.4 Hz, 3H), 2.84 (t, J = 7.0 Hz, 2H).	DMSO	99	Method AQ2	357.5 (M+1)	Method C

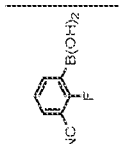
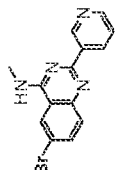
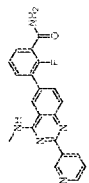
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1478				2 HCl	¹ H NMR (300 MHz, DMSO-d ₆) δ 10.11 (s, 1H), 9.60 (d, J = 1.9 Hz, 1H), 8.93 (dd, J = 10.3, 5.4 Hz, 2H), 8.80 (d, J = 0.6 Hz, 1H), 8.34 (dd, J = 8.0, 0.5 Hz, 1H), 8.10 (d, J = 8.3 Hz, 1H), 7.88–7.75 (m, 3H), 7.40 (d, J = 8.3 Hz, 2H), 3.65 (t, J = 6.9 Hz, 2H), 3.30 (d, J = 4.4 Hz, 3H), 2.79 (t, J = 6.9 Hz, 2H).	DMSO	99	Method AQ2	357.5 (M+1)	Method C

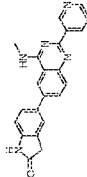
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1479				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.27 (s, 1H), 9.62 (d, J = 1.5 Hz, 1H), 8.99 (s, 1H), 8.96-8.88 (m, 2H), 8.37 (d, J = 8.9 Hz, 1H), 8.18 (d, J = 8.6 Hz, 1H), 7.96 (d, J = 8.5 Hz, 2H), 7.85 (s, 1H), 7.70 (d, J = 8.5 Hz, 2H), 3.30 (d, J = 4.3 Hz, 3H), 1.74 (s, 6H).	DMSO	99	Method AQ2	380.5 (M+1)	Method C

(continued)

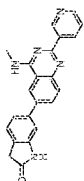
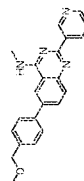
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Meth-od
1480				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.23 (s, 1H), 9.62 (d, J = 1.8 Hz, 1H), 9.02 (d, J = 6.6 Hz, 1H), 8.93 (dd, J = 5.0, 1.5 Hz, 1H), 8.88 (d, J = 0.5 Hz, 1H), 8.33 (d, J = 8.5 Hz, 1H), 8.16 (d, J = 8.5 Hz, 1H), 7.99 (d, J = 1.7 Hz, 1H), 7.86 (dd, J = 7.4, 5.1 Hz, 1H), 7.75 (dd, J = 8.5, 1.8 Hz, 1H), 7.60 (d, J = 8.5 Hz, 1H), 3.29 (d, J = 4.3 Hz, 3H).	DMSO	99	Method AQ2	393.4 (M+1)	Method C

(continued)

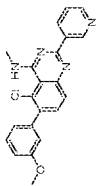
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1481					¹ H NMR (300 MHz, DMSO) δ 9.63 (d, J = 1.4 Hz, 1H), 8.81-8.73 (m, 1H), 8.68 (dd, J = 4.7, 1.7 Hz, 1H), 8.61 (d, J = 4.3 Hz, 1H), 8.45 (s, 1H), 7.97 (d, J = 8.6 Hz, 1H), 7.86 (d, J = 8.7 Hz, 2H), 7.77-7.61 (m, 3H), 7.53 (dd, J = 7.9, 4.8 Hz, 1H), 7.40 (t, J = 7.6 Hz, 1H), 3.15 (d, J = 4.3 Hz, 3H).	DMSO	99	Method AQ2, followed by hydrolysis with 30% H ₂ O ₂ and NaOH in ethanol	374.5 (M+1)	Method C

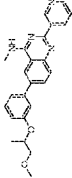
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1482				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.60 (s, 1H), 10.29 (s, 1H), 9.62 (d, J = 1.5 Hz, 1H), 9.00 (d, J = 7.1 Hz 1H), 8.92 (dd, J = 4.9, 1.4 Hz, 1H), 8.76 (s, 1H), 8.29 (d, J = 8.9 Hz, 1H), 8.20 (d, J = 8.4 Hz, 1H), 7.84 (dd, J = 7.8, 4.8 Hz, 1H), 7.79 - 7.68 (m, 2H), 6.95 (d, J = 8.0 Hz, 1H), 3.57 (s, 2H), 3.29 (d, J = 4.3 Hz, 3H).	DMSO	94	Method AQ1, except that dioxane water was replaced with DME/water /EtOH in the microwave at 120°C for 10 min	368.5 (M+1)	Method C

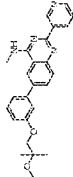
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1483				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.67 (s, 1H), 9.60 (d, J = 2.4 Hz, 1H), 8.99-8.88 (m, 2H), 8.73 (d, J = 1.0 Hz, 1H), 8.27 (d, J = 9.2 Hz, 1H), 8.10 (d, J = 7.5 Hz, 1H), 7.83 (dd, J = 6.9, 4.3 Hz, 1H), 7.40 (q, J = 8.3 Hz, 2H), 7.20 (s, 1H), 3.56 (s, 2H), 3.29 (d, J = 4.1 Hz, 3H).	DMSO	99	Method AQ1, except that di-oxane water was replaced with DME/water /EtOH in the microwave at 120°C for 10 min	368.5 (M+1)	Method C
1484				2 HCl	¹ H NMR (300 MHz, DMSO) δ 10.12 (s, 1H), 9.61 (d, J = 1.9 Hz, 1H), 9.04-8.89 (m, 2H), 8.84 (d, J = 1.2 Hz, 1H), 8.36 (dd, J = 8.7, 1.3 Hz, 1H), 8.14 (d, J = 8.1 Hz, 1H), 7.94-7.78 (m, 3H), 7.50 (d, J = 8.3 Hz, 2H), 4.49 (s, 2H), 3.39-3.23 (m, 6H).	DMSO	99	Method AQ1, except that di-oxane water was replaced with DME/water /EtOH in the microwave at 120°C for 10 min	357.5 (M+1)	Method C

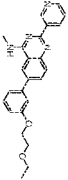
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1485					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.63 (s, 1H), 8.76 (td, J = 8.0, 1.6 Hz, 1H), 8.71 (dd, J = 4.8, 1.6 Hz, 1H), 8.37 (s, 1H), 7.81 (d, J = 8.8 Hz, 1H), 7.72 (d, J = 8.8 Hz, 1H), 7.59-7.54 (m, 1H), 7.43 (t, J = 8.0 Hz, 1H), 7.06-6.98 (m, 3H), 3.82 (s, 3H).	DMSO	95	Method G1,AQ1	377.0 379.0 (M+1)	Method B (NH ₄ HCO ₃)

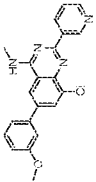
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1486				2HCl	¹ H-NMR(400 MHz, DMSO-d ₆): δ 9.81 (s, 1H), 9.38 (d, J = 8.4 Hz, 1H), 9.15 (d, J = 5.2 Hz, 1H), 8.69 (d, J = 1.6 Hz, 1H), 8.41 (dd, J = 8.8, 2.0 Hz, 1H), 8.29 - 8.26 (m, 1H), 8.14 (d, J = 8.8 Hz, 1H), 7.50 - 7.41 (m, 3H), 7.11 - 7.08 (m, 1H), 4.80 - 4.76 (m, 1H), 3.66 - 3.57 (m, 2H), 3.50 (s, 3H), 3.44 (s, 3H), 1.35 (d, J = 6.4 Hz, 3H).	DMSO	95	Method G1,AQ1	401.1 (M+1)	Method B (NH ₄ HCO ₃)

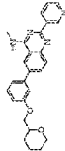
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1487				2HCl	¹ H-NMR(400 MHz, CD3OD): δ 9.55 (s, 1H), 8.92 (d, J = 4.9 Hz, 1H), 8.89 (dt, J = 8.2, 1.8 Hz, 1H), 8.62 (s, 1H), 8.37 (d, J = 8.8 Hz, 1H), 8.03 (d, J = 8.7 Hz, 1H), 7.83 (dd, J = 8.0, 5.0 Hz, 1H), 7.49 (t, J = 8.1 Hz, 1H), 7.43 (s, 2H), 7.10 (d, J = 7.6 Hz, 1H), 4.01 (s, 2H), 3.46 (s, 3H), 3.35 (s, 3H), 1.37 (s, 6H).	MeOD	95	Method G1,AQ1	415.1 (M+1)	Method B (NH ₄ HCO ₃)

(continued)

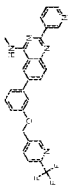
5	1488 	Starting Material 1	Starting Material 2	Product	Salt type 2HCl	¹ H NMR (400 MHz, DMSO-d6): δ 10.34 (s, 1H), 9.65 (s, 1H), 9.04 (s, 1H), 9.04 (d, J = 6.9 Hz, 1H), 8.99 - 8.87 (m, 2H), 8.39 (d, J = 9.2 Hz, 1H), 8.24 (d, J = 8.2 Hz, 1H), 7.86 (s, 1H), 7.47-7.43 (m, 3H), 7.05 (d, J = 6.8 Hz, 1H), 4.26-4.22 (m, 2H), 3.77 - 3.73 (m, 2H), 3.56-3.50 (m, 2H), 3.32 (d, J = 4.3 Hz, 3H), 1.15 (t, J = 7.0 Hz, 3H).	¹ H NMR Solvent DMSO	Purity per-cent 95	Method of Coupling Method G1,AQ1	LCMS 400.9 (M+1)	LCMS Method Method B (NH ₄ HCO ₃)
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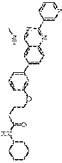
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1489					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.68 (s, 1H), 8.81 - 8.78 (m, 2H), 8.71 (d, J = 3.6 Hz, 1H), 8.54 (s, 1H), 8.32 (d, J = 1.2 Hz, 1H), 7.57 (dd, J = 7.6, 4.8 Hz, 1H), 7.46 - 7.42 (m, 3H), 7.04 - 7.01 (m, 1H), 3.88 (s, 3H), 3.19 (d, J = 4.0 Hz, 3H).	DMSO	95	Method G1, AQ1	377.1, 379.1 (M+1)	Method B (NH ₄ HCO ₃)

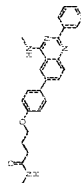
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1490				2HCl	¹ H-NMR(400 MHz, DMSO-d ₆): δ 10.50 (bs, 1H), 9.68 (d, J = 2.0 Hz, 1H), 9.09 (d, J = 8.0 Hz, 1H), 8.96 (dd, J = 5.2, 1.2 Hz, 1H), 8.94 (s, 1H), 8.40 (d, J = 9.2 Hz, 1H), 8.29 (d, J = 8.8 Hz, 1H), 7.89 (dd, J = 8.0, 1.6 Hz, 1H), 7.48 - 7.43 (m, 3H), 7.03 (d, J = 7.2 Hz, 1H), 4.06-4.05 (m, 2H), 3.92 (d, J = 11.6 Hz, 1H), 3.69 - 3.66 (m, 1H), 3.42 - 3.39 (m, 1H), 3.32 (d, J = 4.4 Hz, 3H), 1.86 - 1.83 (m, 1H), 1.69 (d, J = 12.4 Hz, 1H), 1.52 - 1.51 (m, 3H), 1.38 - 1.35 (m, 1H).	DMSO	95	Method G1,AQ1	427.2 (M+1)	Method B (NH ₄ HCO ₃)

(continued)

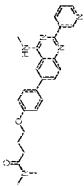
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1492				3HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.80 (s, 1H), 9.70 (s, 1H), 9.13 (s, 1H), 8.0 Hz, 1H), 9.07 (s, 1H), 8.97 (d, J = 4.4 Hz, 1H), 8.94 (s, 1H), 8.39 (d, J = 8.8 Hz, 1H), 8.32 (d, J = 8.8 Hz, 1H), 8.23 (d, J = 8.0 Hz, 1H), 7.98 (t, J = 8.0 Hz, 1H), 7.90 (t, J = 6.2 Hz, 1H), 7.68 (s, 1H), 7.53 (d, J = 8.0 Hz, 1H), 7.46 (d, J = 7.8 Hz, 1H), 7.12 (d, J = 8.0 Hz, 1H), 5.50 (s, 2H), 3.30 (d, J = 2.4 Hz, 3H).	DMSO	95	Method G1,AQ1	488.0 (M+1)	Method B (NH ₄ HCO ₃)

(continued)

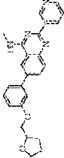
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1493				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.58 (s, 1H), 9.71 (s, 1H), 9.14 (d, J = 7.6 Hz, 1H), 8.98 (d, J = 8.0 Hz, 2H), 8.40 (d, J = 8.8 Hz, 1H), 8.31 (d, J = 8.8 Hz, 1H), 7.92 (t, J = 6.2 Hz, 1H), 7.81 (t, J = 7.6 Hz, 1H), 7.47 - 7.43 (m, 3H), 7.02 (d, J = 7.2 Hz, 1H), 4.10 (t, J = 6.0 Hz, 2H), 3.53 (d, J = 6.4 Hz, 1H), 3.32 (d, J = 2.4 Hz, 3H), 2.26 (t, J = 7.0 Hz, 2H), 2.00 - 1.93 (m, 2H), 1.73 - 1.63 (m, 4H), 1.55 - 1.51 (m, 1H), 1.28 - 1.02 (m, 5H).	DMSO	95	Method G1,AQ1	496.2 (M+1)	Method B (NH ₄ HCO ₃)

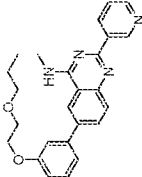
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1494				2HCl	¹ H-NMR(400 MHz, DMSO-d6): δ 10.07 (s, 1H), 9.60 (s, 1H), 8.92 (s, 1H), 8.92 (d, J = 4.4 Hz, 2H), 8.76 (s, 1H), 8.32 (d, J = 8.4 Hz, 1H), 8.10 (d, J = 8.0 Hz, 1H), 7.85-7.83 (m, 4H), 7.11 (d, J = 8.8 Hz, 2H), 4.04 (t, J = 6.4 Hz, 2H), 3.30 (d, J = 4.0 Hz, 3H), 2.58 (d, J = 4.0 Hz, 3H), 2.26 (t, J = 7.4 Hz, 2H), 2.00 - 1.93 (m, 2H).	DMSO	95	Method G1,AQ1	428.1, 429.1, (M+1)	Method B (NH ₄ HCO ₃)

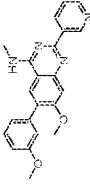
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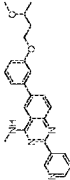
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1495				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.63 (s, 1H), 9.73 (s, 1H), 9.17 (s, 1H), 9.17 (d, J = 7.6 Hz, 1H), 8.99 (s, 1H), 8.91 (s, 1H), 8.34 (s, 1H), 7.93 (t, J = 6.0 Hz, 1H), 7.88 (d, J = 7.2 Hz, 2H), 7.09 (d, J = 7.6 Hz, 2H), 4.08-4.06 (m, 2H), 3.32 (s, 3H), 2.98 (s, 3H), 2.84 (s, 3H), 2.47 - 2.46 (m, 2H), 1.98-1.94 (m, 2H).	DMSO	95	Method G1, AQ1	442.1, 443.1, (M+1)	Method B (NH ₄ HCO ₃)

(continued)

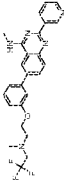
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1496				2HCl	¹ H - NMR (400 MHz, DMSO-d ₆): δ 10.26 (s, 1H), 9.64 (s, 1H), 9.01 (d, J = 7.0 Hz, 1H), 8.94 (s, 1H), 8.89 (s, 1H), 8.39 (d, J = 8.2 Hz, 1H), 8.19 (d, J = 8.1 Hz, 1H), 7.85 (s, 1H), 7.47 (d, J = 5.8 Hz, 3H), 7.05 (d, J = 3.4 Hz, 1H), 4.21 (dt, J = 10.8, 5.5 Hz, 1H), 4.08 (qd, J = 10.1, 5.2 Hz, 2H), 3.82 (dd, J = 14.4, 7.0 Hz, 1H), 3.71 (dd, J = 14.1, 7.4 Hz, 1H), 3.31 (d, J = 4.2 Hz, 3H), 2.08-2.00 (m, 1H), 1.98-1.79 (m, 2H), 1.72-1.68 (m, 1H).	DMSO	95	Method G1, AQ1	413.1, 414.1 (M+1)	Method B (NH ₄ HCO ₃)

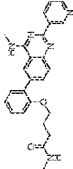
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1497				2HCl	1H-NMR(400 MHz, DMSO-d6): δ 10.53 (s, 1H), 9.68 (s, 1H), 9.09 (s, 1H), 9.09 (d, J = 7.7 Hz, 1H), 8.96(d, J = 5.9 Hz, 2H), 8.39 (d, J = 8.7 Hz, 1H), 8.30 (d, J = 8.3 Hz, 1H), 7.94-7.79(m, 1H), 7.48 - 7.45 (m, 3H), 7.04 (d, J = 7.5 Hz, 1H), 4.30-4.18(m, 2H), 3.80-3.70 (m, 2H), 3.44 (t, J = 6.6 Hz, 2H), 3.31 (d, J = 4.2 Hz, 3H), 1.56 - 1.54 (m, 2H), 0.89 (t, J = 7.4 Hz, 3H).	DMSO-d ₆	95	Method G1,AQ1	415.1 (M+1)	Method B (NH ₄ HCO ₃)

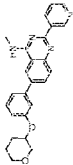
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1498				2HCl	¹ H-NMR(400 MHz, DMSO-d6): δ 10.07 (s, 1H), 9.67 (s, 1H), 9.03 (s, 1H), 9.03 (d, J = 8.0 Hz, 1H), 8.95(d, = 4.5 Hz, 1H), 8.51 (s, 1H), 7.85 (s, 2H), 7.41 (t, J = 7.9 Hz, 1H), 7.18 (dd, J = 7.8, 5.1Hz, 2H), 7.02 (dd, J = 8.2, 2.3 Hz, 1H), 3.97 (s, 3H), 3.84 (s, 3H).	DMSO	95	Method G1,AQ1	373.1 (M+1)	Method B (NH ₄ HCO ₃)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1499				2HCl	¹ H-NMR(400 MHz, DMSO-d ₆): δ 10.06 (s, 1H), 9.66 (s, 1H), 9.02 (s, 1H), 9.02 (d, J = 7.5 Hz, 1H), 8.95 (s, 1H), 8.89 (s, 1H), 8.39(d, J = 8.5 Hz, 1H), 8.22 (d, J = 8.6 Hz, 1H), 7.85 (dd, J = 7.0, 5.4 Hz, 1H), 7.52 - 7.43 (m, 3H), 7.09-7.01 (m, 1H), 4.17 - 4.15 (m, 2H), 3.32-3.30 (m, 4H), 3.25 (s, 3H), 1.98 - 1.82 (m, 2H), 1.17 (d, J = 6.1 Hz, 3H).	DMSO	95	Method G1,AQ1	415.1 (M+1)	Method B (NH ₄ HCO ₃)

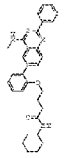
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1500				2HCl	1H-NMR(400 MHz, CD3OD): δ 9.80 (s, 1H), 9.37 (d, J = 8.2 Hz, 1H), 9.13 (d, J = 5.3 Hz, 1H), 8.90 (s, 1H), 8.45 (d, J = 8.4 Hz, 1H), 8.25 (dd, J = 7.9, 5.6 Hz, 1H), 8.17 (d, J = 8.7 Hz, 1H), 7.64 (s, 1H), 7.54 (d, J = 5.1 Hz, 2H), 7.17 (dd, J = 7.5, 3.6 Hz, 1H), 4.70 - 4.62 (m, 2H), 4.42 (q, J = 9.0 Hz, 2H), 3.93-3.86(m, 2H), 3.51 (s, 3H), 3.24 (s, 3H).	CD ₃ OD	95	Method G1,AQ1	468.1 (M+1)	Method B (NH ₄ HCO ₃)

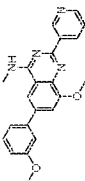
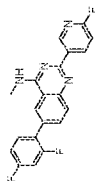
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1501			 <chem>CN(C)C(=O)CCOC1=CC=C(C=C1)Oc2ccc(cc2)OC</chem>	2HCl	¹ H-NMR(400 MHz,DMSO): δ 10.45 (brs, 1H), 9.72 (s, 1H), 9.12 (d, J = 7.6 Hz, 1H), 8.97 (d, J = 4.0 Hz, 1H), 8.70 (s, 1H), 8.35 (d, J = 8.8 Hz, 1H), 8.23 (d, J = 8.4 Hz, 1H), 7.90 - 7.87 (m, 2H), 7.49 (d, J = 7.2 Hz, 1H), 7.43 (t, J = 7.2 Hz, 1H), 7.18 (d, J = 8.4 Hz, 1H), 7.11 (t, J = 7.2 Hz, 1H), 4.03 (t, J = 6.4 Hz, 2H), 3.30 (d, J = 4.4 Hz, 3H), 2.54(d, J = 4.0 Hz, 3H), 2.19 (t, J = 7.4 Hz, 2H), 1.93 - 1.86 (m, 2H).	DMSO	95	Method G1,AQ1	428.1 (M+1)	Method B (NH4HCO ₃)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1502					1H-NMR(400 MHz, DMSO-d ₆): δ 9.65(s, 1H), 8.88(s, 1H), 8.81(d, J = 8 Hz, 1H), 8.73(s, 1H), 8.62(s, 1H), 8.19(d, J = 8 Hz, 1H), 7.88(d, J = 9.2 Hz, 1H), 7.61 - 7.58(m, 1H), 7.47-7.44(m, 3H), 7.06 - 7.04(m, 1H), 4.56-4.54(m, 1H), 3.87(dd, J = 11.2, 2.0 Hz, 1H), 3.68 - 3.64(m, 1H), 3.56-3.53(m, 2H), 3.21(d, J = 4.4 Hz, 3H), 2.09-2.05(m, 1H), 1.83 - 1.72(m, 2H), 1.61-1.55(m, 1H).	DMSO	95	Method G1,AQ1	413.1 (M+1)	Method B (NH ₄ HCO ₃)

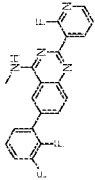
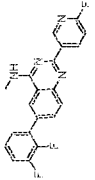
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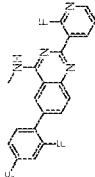
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1503				2HCl	¹ H-NMR(400 MHz, CD3OD): δ 9.77 (s, 1H), 9.43 (d, J = 8.1 Hz, 1H), 9.08 (d, J = 5.6 Hz, 1H), 8.46 (s, 1H), 8.34-8.25(m, 1H), 8.20(d, J = 8.7 Hz, 1H), 8.02 (d, J = 8.7 Hz, 1H), 7.37 (d, J = 7.5 Hz, 1H), 7.32 (t, J = 7.4 Hz, 1H), 7.05 (d, J = 8.3 Hz, 1H), 7.00 (t, J = 7.5 Hz, 1H), 3.98 (t, J = 6.1 Hz, 2H), 3.44 (t, J = 11.1 Hz, 1H), 3.35 (s, 3H), 2.17 (t, J = 7.3 Hz, 2H), 1.93 --1.89 (m, 2H), 1.70 - 1.53(m, 4H), 1.48 (d, J = 12.5 Hz, 1H), 1.22-1.12(m, 2H), 1.08 - 1.02 (m, 3H).	CD ₃ OD	95	Method G1,AQ1	496.2 (M+1)	Method B (NH ₄ HCO ₃)

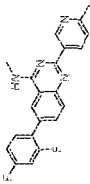
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1504					1H-NMR (400 MHz, DMSO-d ₆): δ 9.63 (s, 1H), 8.76 (d, J = 8.0 Hz, 1H), 8.67 (d, J = 3.6 Hz, 1H), 8.50 (d, J = 4.0 Hz, 1H), 8.10 (s, 1H), 7.56 - 7.53 (m, 2H), 7.46 - 7.42 (m, 3H), 7.03 - 7.01 (m, 1H), 4.07 (s, 3H), 3.88 (s, 3H), 3.17 (d, J = 4.0 Hz, 1H).	DMSO	95	Method G1, AQ1	372.9 (M+1)	Method B (NH ₄ HCO ₃)
1505					1H-NMR (400 MHz, DMSO): δ 9.28 (s, 1H), 8.94 (t, J = 7.6 Hz, 1H), 8.62 (d, J = 4.2 Hz, 1H), 8.42 (s, 1H), 7.94 (d, J = 8.8 Hz, 1H), 7.84 (d, J = 8.8 Hz, 1H), 7.73 - 7.71 (m, 1H), 7.46 (t, J = 10.0 Hz, 1H), 7.33 - 7.27 (m, 2H), 3.16 (d, J = 4.4 Hz, 3H).	DMSO	95	Method G1, AQ1	367.0 (M+1)	Method B (NH ₄ HCO ₃)

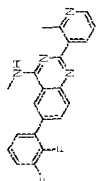
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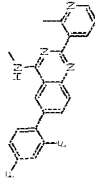
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1506					1H-NMR (400 MHz, DMSO): δ 8.77 (s, 1H), 8.66 (t, J = 8.6 Hz, 1H), 8.52 (s, 1H), 8.44 - 8.31 (m, 1H), 8.03 (d, J = 8.6 Hz, 1H), 7.88 (d, J = 8.6 Hz, 1H), 7.62-7.49 (m, 3H), 7.42 - 7.37 (m, 1H), 3.12 (d, J = 3.0 Hz, 3H).	DMSO	95	Method G1,AQ1	366.9 (M+1)	Method B (NH ₄ HCO ₃)
1507					1H-NMR (400 MHz, DMSO): δ 9.29 (d, J = 2.0 Hz, 1H), 8.95 (dt, J = 8.4, 2.4 Hz, 1H), 8.65 (d, J = 4.4 Hz, 1H), 8.49 (s, 1H), 7.94 - 7.93 (m, 2H), 7.60-7.26 (m, 4H), 3.16 (d, J = 4.4 Hz, 3H).	DMSO	95	Method G1,AQ1	367.1 (M+1)	Method B (NH ₄ HCO ₃)

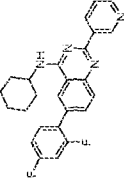
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1508				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.46 (s, 1H), 8.77 (s, 1H), 8.65 (s, 1H), 8.65 (t, J = 8.5 Hz, 1H), 8.55 (s, 1H), 8.19 (d, J = 8.7 Hz, 1H), 8.07 (d, J = 8.6 Hz, 1H), 7.79 (dd, J = 15.8, 7.8 Hz, 1H), 7.71 Hz, 1H), -7.66 (m, 1H), 7.51 (t, J = 10.2 Hz, 1H), 7.34 (t, J = 8.5 Hz, 1H), 3.23 (d, J = 3.3 Hz, 3H).	DMSO	95	Method G1, AQ1	367.0 (M+1)	Method B (NH ₄ HCO ₃)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1509				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.98 (s, 1H), 9.55 (d, <i>J</i> = 1.8 Hz, 1H), 9.07 (d, <i>J</i> = 7.2 Hz, 1H), 8.67 (s, 1H), 8.16 (dd, <i>J</i> = 18.0, 8.4 Hz, 2H), 7.84 (d, <i>J</i> = 8.3 Hz, 1H), 7.78 (td, <i>J</i> = 8.9, 6.6 Hz, 1H), 7.55 - 7.45 (m, 1H), 7.33 (td, <i>J</i> = 8.4, 2.0 Hz, 1H), 3.28 (d, <i>J</i> = 4.4 Hz, 3H), 2.74 (s, 3H).	DMSO	95	Method G1, AQ1	363.1 (M+1)	Method B (NH ₄ HCO ₃)

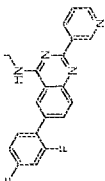
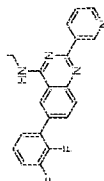
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1510				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.17 (s, 1H), 8.82 (dd, J = 7.9, 6.7 Hz, 2H), 8.65 (s, 1H), 8.21 (d, J = 8.5 Hz, 1H), 8.07 (d, J = 8.6 Hz, 1H), 7.80 (s, 1H), 7.65-7.50 (m, 2H), 7.42 (dd, J = 12.7, 7.6 Hz, 1H), 3.20 (d, J = 4.4 Hz, 3H), 2.94 (s, 3H).	DMSO	95	Method G1,AQ1	363.0 (M+1)	Method B (NH ₄ HCO ₃)

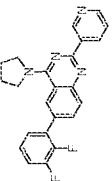
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1511				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.24 (s, 1H), 8.84 (d, J=4.0 Hz, 1H), 8.77 (s, 1H), 8.64 (s, 1H), 8.18 (d, J=8.6 Hz, 1H), 8.08 (d, J=8.6 Hz, 1H), 7.79 (td, J=8.9, 6.6 Hz, 2H), 7.57 - 7.46 (m, 1H), 7.34 (td, J=8.5, 2.2 Hz, 1H), 3.20 (d, J=4.5 Hz, 3H), 2.93 (s, 3H).	DNLSO	95	Method G1,AQ1	362.9 (M+1)	Method B (NH ₄ HCO ₃)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1512				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.57 (s, 1H), 8.83 (dd, J = 8.0, 1.6 Hz, 1H), 8.63 (dd, J = 4.8, 2.0 Hz, 1H), 8.38 (s, 1H), 7.95 - 7.86 (m, 2H), 7.67 - 7.57 (m, 2H), 7.15 - 7.09 (m, 2H), 4.44 (brs, 1H), 2.17 (t, J = 7.6 Hz, 1H), 1.91 - 1.89 (m, 1H), 1.78 (d, J = 12.0 Hz, 1H), 1.56 - 1.48 (m, 4H), 1.32 - 1.28 (m, 1H).	DMSO	95	Method G1,AQ1	417.1 (M+1)	Method B (NH ₄ HO ₃)

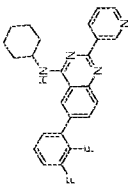
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1513				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.55 (s, 1H), 8.94 - 8.89 (m, 2H), 8.58 (s, 1H), 8.23 (d, J = 8.8 Hz, 1H), 8.04 (d, J = 8.8 Hz, 1H), 7.85 (t, J = 6.8 Hz, 1H), 7.73 (dd, J = 14.4, 8.0 Hz, 1H), 7.25 - 7.19 (m, 2H), 4.01 (q, J = 7.2 Hz, 2H), 1.48 (t, J = 7.2 Hz, 3H).	DMSO	95	Method G1, AQ1	363.1 (M+1)	Method B (NH ₄ HCO ₃)
1514				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.71 (s, 1H), 9.21 (d, J = 8.4 Hz, 1H), 9.08 (d, J = 4.4 Hz, 1H), 8.68 (s, 1H), 8.30 (d, J = 8.8 Hz, 1H), 8.15 - 8.12 (m, 2H), 7.50 - 7.38 (m, 3H), 4.04 (q, J = 7.2 Hz, 2H), 1.49 (t, J = 7.2 Hz, 3H).	DMSO	95	Method G1, AQ1	363.1 (M+1)	Method B (NH ₄ HCO ₃)

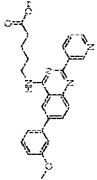
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1515				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.67 (s, 1H), 9.25 (d, J = 8.0 Hz, 1H), 9.03 (d, J = 5.2 Hz, 1H), 8.60 (s, 1H), 8.21-8.14 (m, 2H), 8.06 (d, J = 8.4 Hz, 1H), 7.40-7.26 (m, 3H), 4.26 (brs, 4H), 2.14 (brs, 4H).	DMSO-d ₆	95	Method G1,AQ1	389.1 (M+1)	Method B (NH ₄ HCO ₃)

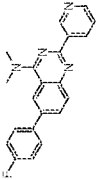
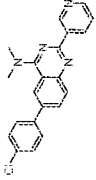
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1516				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.60 (s, 1H), 9.14 (d, J = 8.0 Hz, 1H), 9.10 (d, J = 5.2 Hz, 1H), 8.69 (s, 1H), 8.19 (d, J = 8.4 Hz, 1H), 8.09 (dd, J = 8.0, 5.2 Hz, 1H), 8.03 (d, J = 8.8 Hz, 1H), 7.39-7.27 (m, 3H), 4.60 - 4.50 (m, 1H), 2.08 (d, J = 11.6 Hz, 2H), 1.84 (d, J = 12.0 Hz, 2H), 1.69 (d, J = 8.4 Hz, 1H), 1.57-1.45 (m, 4H), 1.18 - 1.10 (m, 1H).	DMSO	95	Method G1,AQ1	417.1 (M+1)	Method B (NH ₄ HCO ₃)

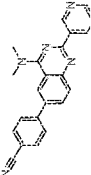
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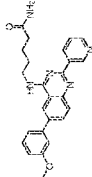
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1517					¹ H-NMR (400 MHz, DMSO-d ₆): δ 12.04 (brs, 1H), 9.58 (s, 1H), 9.38 (brs, 1H), 8.84 (d, J = 5.6 Hz, 2H), 8.72 (s, 1H), 8.28 (d, J = 9.2 Hz, 1H), 7.75 (t, J = 6.8 Hz, 1H), 7.51 - 7.41 (m, 3H), 7.07 - 7.03 (m, 1H), 3.88 (s, 3H), 3.80 (q, J = 5.6 Hz, 2H), 2.32 (t, J = 7.2 Hz, 2H), 1.86 - 1.74 (m, 2H), 1.73 - 1.64 (m, 2H).	DMSO	95	Method G1, AQ1	428.9 (M+1)	Method B (NH ₄ HCO ₃)

(continued)

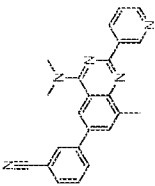
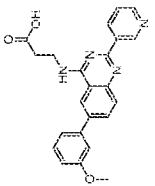
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1518					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.61 (s, 1H), 8.74 (td, J = 8.0, 1.6 Hz, 1H), 8.71 Hz, 1H), 8.71 - 8.66 (m, 1H), 8.31 (d, J = 1.6 Hz, 1H), 8.08 (dd, J = 8.8, 2.0 Hz, 1H), 7.90 (d, J = 8.8 Hz, 1H), 7.87-7.81 (m, 2H), 7.58 - 7.49 (m, 1H), 7.39-7.30 (m, 2H), 3.49 (s, 6H).	DMSO	95	Method G1, AQ1	345.1 (M+1)	Method B (NH ₄ HCO ₃)
1519					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.61 (s, 1H), 8.74 (d, J = 7.2 Hz, 1H), 8.69 (d, J = 3.6 Hz, 1H), 8.32 (s, 1H), 8.08 (d, J = 8.4 Hz, 1H), 7.89 (d, J = 8.4 Hz, 1H), 7.82 (d, J = 7.6 Hz, 2H), 7.62-7.48 (m, 3H), 3.48 (s, 6H).	DMSO	95	Method G1, AQ1	361.1 363.1	Method B (NH ₄ HCO ₃)

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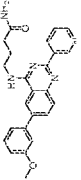
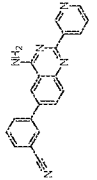
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1520					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.61 (s, 1H), 8.74 (td, J = 8.0, 1.6 Hz, 1H), 8.69 (dd, J = 4.8, 1.6 Hz, 1H), 8.42 (d, J = 2.0 Hz, 1H), 8.16 (dd, J = 8.8, 2.0 Hz, 1H), 8.04 - 7.89 (m, 5H), 7.54 (q, J = 5.2 Hz, 1H), 3.51 (s, 6H).	DMSO	95	Method G1,AQ1	352.1 (M+1)	Method B (NH ₄ HCO ₃)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1521					¹ H-NMR(400 MHz, DMSO-d ₆): δ 9.61 (s, 1H), 8.76 (d, J = 8.0 Hz, 1H), 8.68-8.60 (m, 2H), 8.15 (d, J = 8.8 Hz, 1H), 7.85 (d, J = 8.8 Hz, 1H), 7.56 (q, J = 4.8 Hz, 1H), 7.49-7.40 (m, 3H), 7.30 (s, 1H), 7.06 - 6.97 (m, 1H), 6.73 (s, 1H), 3.87 (s, 5H), 2.14 (t, J = 7.2 Hz, 2H), 1.80 - 1.71 (m, 2H), 1.71-1.62 (m, 2H).	DMSO	95	Method G1,AQ1	428.1 (M+1)	Method B (NH ₄ HCO ₃)

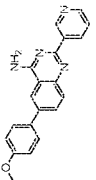
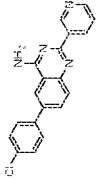
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1522				2 HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.67 (s, 1H), 9.39 (d, J = 8.2 Hz, 1H), 9.02 (d, J = 5.3 Hz, 1H), 8.31 (d, J = 9.3 Hz, 2H), 8.24 - 8.10 (m, 3H), 7.88 (d, J = 7.7 Hz, 1H), 7.72 (t, J = 7.8 Hz, 1H), 3.54 (s, 6H), 2.78 (s, 3H).	DMSO	95	Method G1, AQ1	366.1 (M+1)	Method B (NH ₄ HCO ₃)
1523					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.63 (d, J = 1.7 Hz, 1H), 8.88 - 8.32 (m, 4H), 8.16 (dd, J = 8.7, 1.8 Hz, 1H), 7.86 (t, J = 9.2 Hz, 1H), 7.55 (dd, J = 7.9, 4.8 Hz, 1H), 7.50 - 7.34 (m, 3H), 7.01 (dt, J = 7.3, 2.1 Hz, 1H), 3.96 - 3.84 (m, 5H), 2.78 (t, J = 6.9 Hz, 2H).	DMSO	95	Method G1, AQ1	400.9 (M+1)	Method B (NH ₄ HCO ₃)

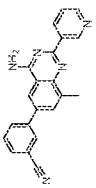
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1524					¹ H-NMR (400 MHz, DMSO-d ₆): 9.63 (s, 1H), 8.77 (d, J = 8 Hz, 1H), 8.66-8.63 (m, 3H), 8.15 (d, J = 8.8 Hz, 1H), 7.85 (d, J = 8.8 Hz, 1H), 7.45-7.41 (m, 3H), 7.33 (s, 1H), 7.01 (d, J = 7.0 Hz, 1H), 6.79 (s, 1H), 3.88 (s, 3H), 3.72 (d, J=6.0 Hz, 2H), 2.24 (t, J = 7.2 Hz, 2H), 2.04 - 1.94 (m, 2H).	DMSO	95	Method G1,AQ1	414.1 (M+1)	Method B (NH ₄ HCO ₃)
1525					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.58 (d, J = 1.2 Hz, 1H), 8.73 - 8.68 (m, 3H), 8.37 (s, 1H), 8.27-8.22 (m, 4H), 7.89 - 7.87 (m, 2H), 7.75 (t, J=8.0 Hz, 1H), 7.55 (dd, J = 8.0, 5.2 Hz, 1H).	DMSO	95	Method G1,AQ1	324.1 (M+1)	Method B (NH ₄ HCO ₃)

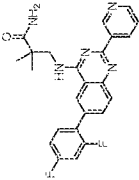
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1526					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.57 (d, J = 1.6 Hz, 1H), 8.72 - 8.66 (m, 2H), 8.57 (d, J = 1.6 Hz, 1H), 8.19 - 7.91 (m, 3H), 7.83 (d, J = 8.0 Hz, 3H), 7.54 (dd, J = 7.6, 4.8 Hz, 1H), 7.10 (d, J = 8.4 Hz, 2H), 3.83 (s, 3H).	DMSO	95	Method G1,AQ1	329.1 (M+1)	Method B (NH ₄ HCO ₃)
1527					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.58 (s, 1H), 8.72 - 8.66 (m, 3H), 8.16 (d, J = 8.4 Hz, 3H), 7.91 (d, J = 8.4 Hz, 2H), 7.85 (d, J = 8.8 Hz, 1H), 7.60 (d, J = 8.4 Hz, 2H), 7.54 (dd, J = 7.6, 4.8 Hz, 1H).	DMSO	95	Method G1,AQ1	333.1, 335.0 (M+1)	Method B (NH ₄ HCO ₃)

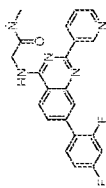
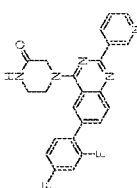
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1528				2HCl	¹ H-NMR (400 MHz, DMSO- d ₆): δ 9.59 (s, 1H), 9.28 (d, J = 8.0 Hz, 1H), 9.01 (d, J = 5.6 Hz, 1H), 8.65 (s, 2H), 8.37 (s, 2H), 8.25 (d, J = 8.0 Hz, 1H), 8.22 (s, 1H), 8.13 (t, J = 6.8 Hz, 1H), 7.88 (d, J = 7.6 Hz, 1H), 7.74 (t, J = 8.0 Hz, 1H), 2.77 (s, 3H).	DMSO	95	Method G1, AQ1	338.1 (M+1)	Method B (NH ₄ HCO ₃)

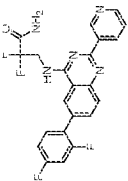
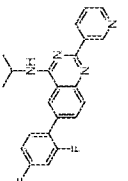
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1529				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.78 (s, 1H), 9.72 (d, J = 1.8 Hz, 1H), 9.18 (d, J = 8.1 Hz, 1H), 8.98 (dd, J = 5.1, 1.2 Hz, 1H), 8.78 (s, 1H), 8.29 (d, J = 8.6 Hz, 1H), 8.17 (d, J = 8.6 Hz, 1H), 7.95 (dd, J = 7.9, 5.3 Hz, 1H), 7.80 (td, J = 8.9, 6.6 Hz, 1H), 7.55 - 7.45 (m, 1H), 7.34 (td, J = 8.4, 2.3 Hz, 1H), 7.26 (s, 1H), 6.96 (s, 1H), 3.98 (d, J = 6.1 Hz, 2H), 1.24 (s, 6H).	DMSO	95	Method G1,AQ1	434.1 (M+1)	Method B (NH ₄ HCO ₃)

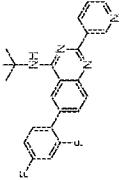
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1530				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.58 (s, 1H), 9.04 - 8.89 (m, 2H), 8.71 - 8.66 (m, 1H), 8.15 - 8.11 (m, 2H), 7.82 - 7.76 (m, 2H), 7.51 (t, J = 9.2 Hz, 1H), 7.34 (t, J = 8.8 Hz, 1H), 4.64 (s, 2H), 3.19 (s, 3H), 2.91 (s, 3H).	DMSO	95	Method G1,AQ1	420.1 (M+1) Method B (NH ₄ HCO ₃)	
1531					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.63 (s, 1H), 8.78 (d, J = 7.9 Hz, 1H), 8.72 (d, J = 3.2 Hz, 1H), 8.19 (s, 2H), 8.08 - 7.97 (m, 2H), 7.80 (dd, J = 15.5, 8.9 Hz, 1H), 7.58 (dd, J = 7.6, 4.9 Hz, 1H), 7.52 - 7.41 (m, 1H), 7.29 (td, J = 8.3, 1.9 Hz, 1H), 4.48 (s, 2H), 4.15 (s, 2H), 3.48 (s, 2H).	DMSO	95	Method G1,AQ1	418.1 (M+1) Method B (NH ₄ HCO ₃)	

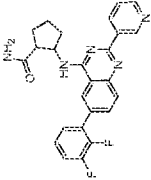
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1532					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.65 (s, 1H), 8.87 (s, 1H), 8.79 (d, J = 7.7 Hz, 1H), 8.70 (s, 1H), 8.59 (s, 1H), 8.23 (s, 1H), 8.05-7.89 (m, 3H), 7.75 (dd, J = 15.1, 7.6 Hz, 1H), 7.56 (s, 1H), 7.48 (t, J = 9.9 Hz, 1H), 7.32 (t, J = 8.1 Hz, 1H), 4.46 (t, J = 13.4 Hz, 2H).	DMSO	95	Method G1, AQ1	442.0 (M+1)	Method B (NH ₄ HCO ₃)
1533				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.63 (s, 1H), 8.95 - 9.03 (m, 2H), 8.78 (s, 1H), 8.14-8.24 (m, 2H), 7.88 (t, J = 6.0 Hz, 1H), 7.79 (d, J = 8.8 Hz, 1H), 7.54-7.48 (m, 1H), 7.35 - 7.32 (m, 1H), 4.89 (m, 1H), 1.39 (d, J = 1.2 Hz, 6H).	DMSO	95	Method G1, AQ1	377.1 (M+1)	Method B (NH ₄ HCO ₃)

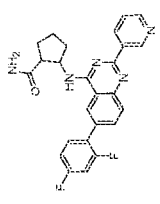
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Meth-od
1534				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.61 (s, 1H), 8.69 - 8.74 (m, 2H), 8.59 (s, 1H), 7.92 (d, J = 8.0 Hz, 1H), 7.84 (d, J = 8.4 Hz, 1H), 7.71 - 7.77 (m, 1H), 7.66 (s, 1H), 7.56 - 7.60 (m, 1H), 7.42 - 7.48 (m, 1H), 7.27 - 7.32 (m, 1H), 1.66 (s, 9H).	DMSO	95	Method G1,AQ1	391.1 (M+1)	Method B (NH ₄ HCO ₃)

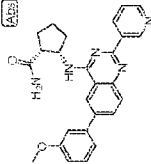
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1535				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.74 (s, 1H), 9.66 (s, 1H), 9.22 (d, J = 7.6 Hz, 1H), 8.99 (d, J = 4.8 Hz, 1H), 8.83 (s, 1H), 8.33 (d, J = 8.4 Hz, 1H), 8.20 (d, J = 8.8 Hz, 1H), 7.95 (t, J = 6.2 Hz, 1H), 7.58 (q, J = 8.6 Hz, 1H), 7.51 (t, J = 7.0 Hz, 1H), 7.46-7.41 (m, 1H), 7.25 (s, 1H), 6.72 (s, 1H), 5.13 - 5.06 (m, 1H), 3.21 (q, J = 7.3 Hz, 1H), 2.22-1.92 (m, 5H), 1.68 - 1.57 (m, 1H).	DMSO	95	Method G1,AQ1	446.0 (M+1)	Method B (NH ₄ HCO ₃)

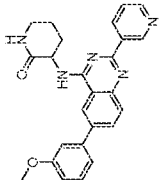
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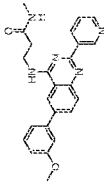
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1536				2HCl	¹ H-NMR (400 MHz, DMSO- d ₆): δ 9.81 (s, 2H), 9.32 (d, J = 8.0 Hz, 1H), 9.02 (d, J = 5.2 Hz, 1H), 8.81 (s, 1H), 8.42 (d, J = 8.4 Hz, 1H), 8.18 (d, J = 8.4 Hz, 1H), 8.00 (dd, J = 7.8, 5.8 Hz, 1H), 7.76 (dd, J = 15.4, 8.6 Hz, 1H), 7.51 (t, J = 10.0 Hz, 1H), 7.35 (t, J = 8.2 Hz, 1H), 7.29 (s, 1H), 6.73 (s, 1H), 5.15-5.07 (m, 1H), 3.21 (q, J = 7.3 Hz, 1H), 2.23-1.87 (m, 5H), 1.68 - 1.57 (m, 1H).	DMSO	95	Method G1,AQ1	446.1 (M+1)	Method B (NH ₄ HCO ₃)

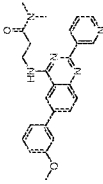
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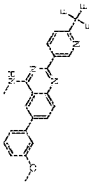
5	Number	Starting Material 1	Starting Material 2	Product	Salt type	1H NMR	1H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
10											
15	1537					¹ H-NMR (400 MHz, DMSO- _d ₆): δ 9.66 (s, 1H), 8.79 (d, J = 7.6 Hz, 1H), 8.69 (dd, J = 4.4, 1.2 Hz, 1H), 8.61 (d, J = 1.6 Hz, 1H), 8.40 (d, J = 7.2 Hz, 1H), 8.13 (dd, J = 8.6, 1.4 Hz, 1H), 7.85 (d, J = 8.4 Hz, 1H), 7.55 (dd, J = 7.6, 4.8 Hz, 1H), 7.49 - 7.40 (m, 3H), 7.11 (s, 1H), 7.02 (d, J = 7.2 Hz, 1H), 6.69 (s, 1H), 5.03-4.96 (m, 1H), 3.87 (s, 3H), 3.20 (dd, J = 15.0, 7.4 Hz, 1H), 2.22 - 1.88 (m, 5H), 1.70-1.56 (m, 1H).	DMSO	95	Method G1,AQ1		Method B (NH ₄ HCO ₃)
20											
25											
30											
35											
40											
45											
50											
55											

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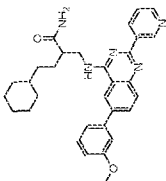
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1538					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.60 (s, 1H), 8.86 (d, J = 7.7 Hz, 1H), 8.74 (d, J = 8.0 Hz, 1H), 8.69 (d, J = 3.3 Hz, 1H), 8.65 (s, 1H), 8.19 (d, J = 7.4 Hz, 1H), 7.87 (d, J = 8.8 Hz, 2H), 7.54 (dd, J = 7.8, 4.8 Hz, 1H), 7.50 - 7.40 (m, 3H), 7.01 (dd, J = 7.3, 3.9 Hz, 1H), 4.85 (dd, J = 16.9, 8.7 Hz, 1H), 3.88 (s, 3H), 2.16 (s, 2H), 1.96 (s, 2H).	DMSO	95	Method G1,AQ1	426.1, 427.1 (M+1)	Method B (NH ₄ HCO ₃)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1539				2HCl	¹ H-NMR (400 MHz, DMSO- d ₆): δ 10.34 (s, 1H), 9.67 (s, 1H), 9.07 (s, 1H), 8.97 (s, 1H), 8.97 (d, J = 5.0 Hz, 1H), 8.92 (s, 1H), 8.39 (d, J = 8.8 Hz, 1H), 8.25 (d, J = 7.4 Hz, 1H), 7.99 (s, 1H), 7.90 (s, 1H), 7.48 (d, J = 5.8 Hz, 3H), 7.09-6.99 (m, 1H), 4.03 (d, J = 6.1 Hz, 2H), 3.89 (s, 3H), 2.65 (t, J = 6.8 Hz, 2H), 2.56 (d, J = 4.4 Hz, 3H).	DMSO	95	Method G1,AQ1	414.0, 415.0 (M+1)	Method B (NH ₄ HCO ₃)

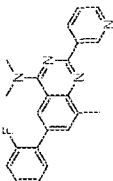
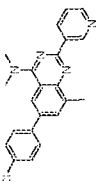
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1540				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.48 (s, 1H), 9.64 (s, 1H), 9.05 (s, 1H), 9.05 (d, J = 7.9 Hz, 1H), 9.01 - 8.92 (m, 2H), 8.38 (d, J = 8.8 Hz, 1H), 8.28 (d, J = 8.6 Hz, 1H), 7.93-7.83 (m, 1H), 7.55 - 7.43 (m, 3H), 7.08-6.98 (m, 1H), 4.04 (dd, J = 12.4, 6.6 Hz, 2H), 3.89 (s, 3H), 2.98 (s, 3H), 2.89 (t, J = 7.0 Hz, 2H), 2.84 (s, 3H).	DMSO	95	Method G1,AQ1	428.1, 429.1 (M+1)	Method B (NH ₄ HCO ₃)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1541				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.02 (brs, 1H), 9.75 (d, J = 1.6 Hz, 1H), 9.07 (dd, J = 8.0, 1.6 Hz, 1H), 8.83 (s, 1H), 8.35 (d, J = 9.2 Hz, 1H), 8.19 (d, J = 8.0 Hz, 1H), 8.14 (d, J = 8.8 Hz, 1H), 7.48-7.46 (m, 3H), 7.06 - 7.03 (m, 1H), 3.89 (s, 3H), 3.30 (d, J = 4.0 Hz, 3H).	DMSO	95	Method G1, AQ1	411.1 (M+1)	Method B (NH ₄ HCO ₃)

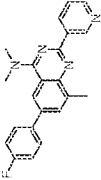
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1542				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.22 (brs, 1H), 9.64 (s, 1H), 9.01 (d, J = 6.4 Hz, 1H), 8.95 (d, J = 4.0 Hz, 1H), 8.89 (s, 1H), 8.39 (d, J = 8.4 Hz, 1H), 8.23 (d, J = 8.0 Hz, 1H), 7.85 (t, J = 5.2 Hz, 1H), 7.52 - 7.42 (m, 4H), 7.08-7.05 (m, 1H), 6.89 (s, 1H), 3.89 (s, 4H), 3.83 - 3.76 (m, 2H), 2.83-2.79 (m, 1H), 1.65 - 1.56 (m, 7H), 1.24-1.07 (m, 6H), 0.89 - 0.81 (m, 2H).	DMSO	95	Method G1,AQ1	510.2 (M+1)	Method B (NH ₄ HCO ₃)

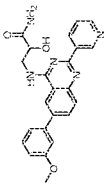
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1543					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.67 (d, J = 1.6 Hz, 1H), 8.80 (dt, J = 8.0, 2.0 Hz, 1H), 8.70 (dd, J = 4.4, 2.0 Hz, 1H), 8.15 (s, 1H), 7.87 (s, 1H), 7.71-7.67 (m, 1H), 7.58 - 7.55 (m, 1H), 7.47 (d, J = 2.4 Hz, 1H), 7.34 (m, 2H), 3.45 (s, 6H), 2.76 (s, 3H).	DMSO	95	Method G1,AQ1	359.1 (M+1)	Method B (NH ₄ HCO ₃)
1544					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.66 (s, 1H), 8.82 - 8.79 (m, 1H), 8.70 (d, J = 4.0 Hz, 1H), 8.20 (d, J = 1.6 Hz, 1H), 8.02 (s, 1H), 7.86-7.84 (m, 2H), 7.58 - 7.55 (m, 3H), 3.48 (s, 6H), 2.77 (s, 3H).	DMSO	95	Method G1,AQ1	375.1 (M+1)	Method B (NH ₄ HCO ₃)

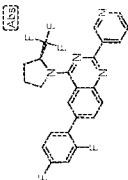
(continued)

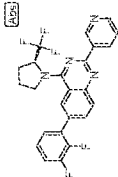
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1545					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.68 (s, 1H), 9.01 (d, J = 8.0 Hz, 1H), 8.82 (d, J = 4.4 Hz, 1H), 8.18 (d, J = 1.6 Hz, 1H), 8.03 (s, 1H), 7.88-7.84 (m, 2H), 7.80 - 7.77 (m, 1H), 7.37-7.33 (m, 2H), 3.50 (s, 6H), 2.78 (s, 3H).	DMSO	95	Method G1,AQ1	359.1 (M+1)	Method B (NH ₄ HCO ₃)

(continued)

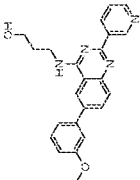
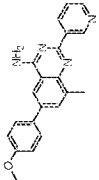
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1546					¹ H-NMR (400 MHz, CD ₃ OD): δ 9.51 (s, 1H), 8.81 (d, J = 7.8 Hz, 1H), 8.54 (s, 1H), 8.35 (s, 1H), 8.02 (d, J = 8.8 Hz, 1H), 7.83 (d, J = 8.8 Hz, 1H), 7.50 (d, J = 7.2 Hz, 1H), 7.29 (dt, J = 11.2, 4.8 Hz, 3H), 6.88 (d, J = 7.5 Hz, 1H), 4.43 (dd, J = 7.1, 4.0 Hz, 1H), 4.16 (dd, J = 13.8, 4.0 Hz, 1H), 3.86 (dd, J = 13.7, 7.4 Hz, 1H), 3.80 (s, 3H).	CD ₃ OD	95	Method G1,AQ1	416.1 (M+1)	Method B (NH ₄ HCO ₃)

(continued)

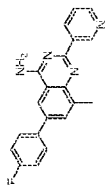
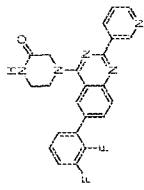
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1547				2HCl	¹ H-NMR (400 MHz, CD ₃ OD): δ 9.74 (s, 1H), 9.44 (d, J = 8.2 Hz, 1H), 9.02 (d, J = 5.5 Hz, 1H), 8.50 (s, 1H), 8.24 (dd, J = 8.0, 5.9 Hz, 1H), 8.16 (q, J = 8.7 Hz, 2H), 7.70-7.58 (m, 1H), 7.12-7.00 (m, 2H), 6.13 (s, 1H), 4.44-4.41 (m, 1H), 4.29-4.27 (m, 1H), 2.37-2.16 (m, 3H), 2.09-2.07 (m, 1H).	CD ₃ OD	95	Method G1, AQ1	457.0 (M+1)	Method B (NH ₄ HCO ₃)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1548				2HCl	¹ H-NMR (400 MHz, CD ₃ OD): δ 9.85 (s, 1H), 9.57 (d, J = 8.0 Hz, 1H), 9.10 (d, J = 4.8 Hz, 1H), 8.64 (s, 1H), 8.34-8.22 (m, 3H), 7.51 (t, J = 6.9 Hz, 1H), 7.47-7.33 (m, 2H), 6.23 (s, 1H), 4.53 - 4.51 (m, 1H), 4.38-4.36 (m, 1H), 2.51 - 2.30 (m, 3H), 2.19-2.18 (m, 1H).	CD ₃ OD	95	Method G1,AQ1	457.1 (M+1)	Method B (NH ₄ HCO ₃)

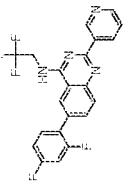
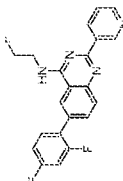
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1549				2HCl	¹ H-NMR (400 MHz, DMSO): δ 10.38 (brs, 1H), 9.63 (d, J = 1.6 Hz, 1H), 9.00 (d, J = 8.0 Hz, 1H), 8.95-8.94 (m, 2H), 8.39 (d, J = 9.2 Hz, 1H), 8.23 (d, J = 8.4 Hz, 1H), 7.88-7.85 (m, 1H), 7.49 - 7.48 (m, 3H), 7.07-7.04 (m, 1H), 3.94 - 3.89 (m, 5H), 3.60-3.57 (m, 2H), 1.96 - 1.93 (m, 2H).	DMSO	95	Method G1,AQ1	387.1 (M+1)	Method B (NH ₄ HCO ₃)
1550					¹ H-NMR (400 MHz, DMSO): δ 9.62 (s, 1H), 8.72-8.70 (m, 2H), 8.40 (s, 1H), 8.11 - 7.94 (m, 2H), 7.91-7.74 (m, 3H), 7.54 (dd, J = 7.6, 4.8 Hz, 1H), 7.09 (d, J = 8.6 Hz, 2H), 3.83 (s, 3H), 2.74 (s, 3H).	DMSO	95	Method G1,AQ1	343.1 (M+1)	Method B (NH ₄ HCO ₃)

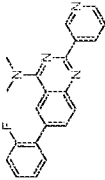
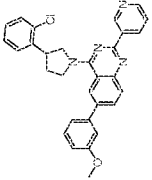
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1551					¹ H-NMR (400 MHz, DMSO): δ 9.64 (s, 1H), 8.76 (d, J = 8.0 Hz, 1H), 8.70 (d, J = 4.4 Hz, 1H), 8.46 (s, 1H), 8.12-7.83 (m, 5H), 7.57 (dd, J = 7.8, 4.8 Hz, 1H), 7.39 (t, J = 8.7 Hz, 2H), 2.84 (s, 3H).	DMSO	95	Method G1,AQ1	331.1 (M+1)	Method B (NH ₄ HCO ₃)
1552					¹ H-NMR (400 MHz, DMSO): δ 9.62 (s, 1H), 8.77 (d, J = 8.0 Hz, 1H), 8.71 (d, J = 4.4 Hz, 1H), 8.23 (s, 1H), 8.20 (s, 1H), 8.07-8.00 (m, 2H), 7.60 - 7.50 (m, 3H), 7.40-7.37 (m, 1H), 4.48 (s, 2H), 4.12 - 4.10 (m, 2H), 3.48-3.46 (m, 2H).	DMSO	95	Method G1,AQ1	418.1 (M+1)	Method B (NH ₄ HCO ₃)

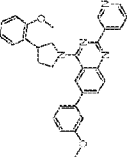
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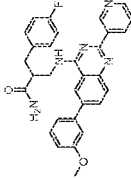
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1553				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.74 (s, 1H), 9.45 (s, 1H), 9.27 (d, J = 8.8 Hz, 1H), 8.97 (s, 1H), 8.69 (s, 1H), 8.12-8.00 (m, 3H), 7.83 - 7.76 (m, 1H), 7.51-7.32 (m, 2H), 4.80 - 4.64 (m, 2H).	DMSO	95	Method G1, AQ1	417.1 (M+1)	Method B (NH ₄ HCO ₃)
1554				2HCl	¹ H-NMR (400 MHz, CD ₃ OD): δ 9.86 (s, 1H), 9.48 (d, J = 8.8 Hz, 1H), 9.18 (d, J = 5.2 Hz, 1H), 8.72 (s1H), 8.92 (s, 1H), 8.37-8.31 (m, 2H), 8.22 (d, J = 8.8 Hz, 1H), 7.80-7.55 (m, 1H), 7.26 - 7.20 (m, 2H), 4.92 (d, J=4.8 Hz, 1H), 4.81 (t, J = 4.8 Hz, 1H), 4.39 - 4.30 (m, 2H).	CD ₃ OD	95	Method G1, AQ1	380.9 (M+1)	Method B (NH ₄ HCO ₃)

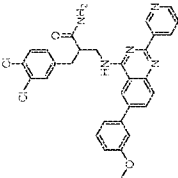
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1555					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.62 (s, 1H), 8.7-8.68 (m, 2H), 8.32 (s, 1H), 7.97-7.93 (m, 2H), 7.91-7.36 (m, 5H), 3.47 (s, 6H).	DMSO	95	Method G1,AQ1	345.1 (M+1)	Method B (NH ₄ HCO ₃)
1556					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.61 (s, 1H), 8.75 (d, J = 8.0 Hz, 1H), 8.67 (d, J = 3.6 Hz, 1H), 8.48 (s, 1H), 8.13 (d, J = 8.8 Hz, 1H), 7.91 (d, J = 8.8 Hz, 1H), 7.59-7.49 (m, 3H), 7.45-7.30 (m, 5H), 7.02-6.92 (m, 1H), 4.50 (q, J = 3.2 Hz, 1H), 4.39-4.20 (m, 2H), 4.14-4.09 (m, 1H), 3.98-3.83 (m, 1H), 3.38 (s, 3H), 2.49-2.40 (m, 1H), 2.35-2.18 (m, 1H).	DMSO	95	Method G1,AQ1	493.0 495.0 (M+1)	Method B (NH ₄ HCO ₃)

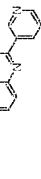
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1557					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.60 (s, 1H), 8.73 (dd, J = 7.6, 1.2 Hz, 1H), 8.67 (dd, J = 8.4, 1.2 Hz, 1H), 8.47 (s, 1H), 8.11 (dd, J = 8.4, 1.6 Hz, 1H), 7.89 (d, J = 8.8 Hz, 1H), 7.54-7.50 (m, 1H), 7.42 (t, J = 8.0 Hz, 1H), 7.38-7.32 (m, 2H), 7.32-7.23 (m, 2H), 7.05 (d, J = 8.4 Hz, 1H), 7.00-6.92 (m, 2H), 4.45 (t, J = 8.4 Hz, 1H), 4.24 (s, 2H), 4.08-3.95 (m, 1H), 3.88-3.80 (m, 7H), 2.44-2.33 (m, 1H), 2.29-2.15 (m, 1H).	DMSO	95	Method G1,AQ1	489.1 (M+1)	Method B (NH ₄ HCO ₃)

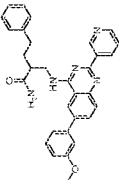
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1558					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.56 (s, 1H), 9.02 (s, 1H), 8.19 (d, J = 7.5 Hz, 1H), 7.86 (d, J = 8.7 Hz, 1H), 7.58 (dd, J = 7.7, 4.9 Hz, 1H), 7.52 - 7.37 (m, 3H), 7.37 - 7.24 (m, 3H), 7.12 (t, J = 8.8 Hz, 2H), 7.02 (d, J = 7.5 Hz, 1H), 6.84 (s, 1H), 3.89-3.78 (m, 4H), 3.72 - 3.66 (m, 1H), 3.13 (s, 1H), 2.97-2.90 (m, 1H), 2.86 - 2.78 (m, 1H).	DMSO	95	Method G1,AQ1	506.1 (M+1)	Method B (NH ₄ HCO ₃)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1559				2HCl	¹ H-NMR(400 MHz, DMSD-d ₆): δ 10.48 - 10.00 (m, 1H), 9.70 (s, 1H), 9.12 - 8.77 (m, 3H), 8.42-8.32 (m, 1H), 8.30 - 8.17 (m, 1H), 7.95-7.79 (m, 1H), 7.55 - 7.44 (m, 6H), 7.26 (d, J = 7.2 Hz, 1H), 7.05 (s, 1H), 6.94 (s, 1H), 3.89 (s, 5H), 3.22 (s, 1H), 2.97-2.81 (m, 2H).	DMSO	95	Method G1,AQ1	558.1 560.0 562.0 (M+1)	Method B (NH ₄ HCO ₃)

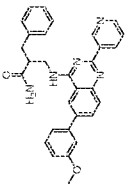
(continued)

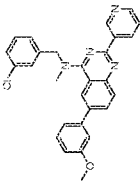
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1560				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.25 (s, 1H), 9.61 (d, J = 1.5 Hz, 1H), 8.96 (d, J = 4.0 Hz, 1H), 8.87 (s, 1H), 8.39 (d, J = 8.5 Hz, 1H), 8.25 (d, J = 7.7 Hz, 1H), 7.87 (s, 1H), 7.52–7.44 (m, 4H), 7.36 (t, J = 7.6 Hz, 1H), 7.28 (dd, J = 13.5, 6.0 Hz, 1H), 7.22–7.10 (m, 2H), 7.06 (d, J = 7.5 Hz, 1H), 6.89 (s, 1H), 3.95–3.81 (m, 5H), 3.21 (d, J = 9.0 Hz, 1H), 2.94 (d, J = 7.3 Hz, 2H).	DMSO	95	Method G1,AQ1	508.2 (M+1)	Method B (NH ₄ HCO ₃)

(continued)

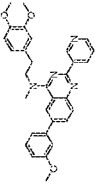
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1561					¹ H-NMR (400 MHz, DMSO- d ₆): δ 9.64 (s, 1H), 8.74 (m, 2H), 8.69 (d, J = 3.5 Hz, 1H), 8.64 (s, 1H), 8.15 (d, J = 8.4 Hz, 1H), 7.85 (d, J = 8.7 Hz, 1H), 7.58-7.49 (m, 1H), 7.43 (m, 4H), 7.28 - 7.18 (m, 4H), 7.15 (m, 1H), 7.01 (d, J = 7.2 Hz, 1H), 6.94 (s, 1H), 3.82-3.73 (m, 5H), 2.90 - 2.87 (m, 1H), 2.68-2.65 (m, 2H), 1.89 - 1.86 (m, 2H).	DMSO	95	Method G1,AQ1	503.9 (M+1)	Method B (NH ₄ HCO ₃)

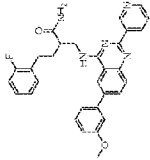
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1562					¹ H-NMR (400 MHz, DMSO-d ₆): 9.85 (s, 1H), 8.76 (s, 1H), 8.65 (d, J = 3.6 Hz, 1H), 8.64 (s, 1H), 8.53 (d, J = 8.0 Hz, 1H), 8.14 (d, J = 8.6 Hz, 1H), 7.83 (d, J = 8.7 Hz, 1H), 7.51 (dd, J = 8.0, 4.7 Hz, 1H), 7.45 (d, J = 6.7 Hz, 2H), 7.40 (s, 1H), 7.31-7.27 (m, 5H), 7.24 (d, J = 6.6 Hz, 1H), 7.00 (d, J = 6.9 Hz, 1H), 6.80 (s, 1H), 3.77-3.70 (m, 5H), 3.17 - 3.15 (m, 1H), 2.98-2.81 (m, 2H).	DMSO	95	Method G1,AQ1	489.9 (M+1)	Method B (NH ₄ HCO ₃)

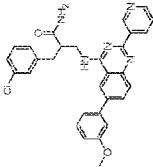
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1563					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.57 (s, 1H), 8.74 - 8.68 (m, 2H), 8.21 (s, 1H), 8.15 (dd, J = 8.7, 1.6 Hz, 1H), 7.95 (d, 3H), 7.52 (m, 2H), 7.46 - 7.31 (m, 4H), 7.12 (d, J = 6.5 Hz, 2H), 6.95-6.93 (m, 1H), 5.16 (s, 2H), 3.78 (s, 3H), 3.53 (s, 3H).	DMSO	95	Method G1, AQ1	467.1, 469.1 (M+1)	Method B (NH ₄ HCO ₃)

(continued)

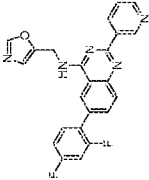
5	1564									1H-NMR (400 MHz, DMSO-d6): δ 9.64 (d, J = 1.4 Hz, 1H), 8.75 (d, J = 7.9 Hz, 1H), 8.70 (d, J = 4.6 Hz, 1H), 8.32 (s, 1H), 8.12 (d, J = 8.7 Hz, 1H), 7.91 (d, J = 8.7 Hz, 1H), 7.55 (dd, J = 7.9, 4.8 Hz, 1H), 7.43 (t, J = 7.9 Hz, 1H), 7.36-7.29 (m, 2H), 7.01 - 6.98 (m, 1H), 6.88 (s, 1H), 6.83 (s, 2H), 4.12-4.03 (m, 2H), 3.85 (s, 3H), 3.69 (s, 3H), 3.62 (s, 3H), 3.56 (d, J = 8.0 Hz, 3H), 3.16-3.02 (m, 2H).	DMSO	95	Method G1,AQ1	507.2 (M+1)	Method B (NH4HCO3)
10															
15															
20															
25															
30															
35															
40															
45															
50															
55															

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1565					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.64 (d, J = 2.0 Hz, 1H), 8.77-8.69 (m, 2H), 8.68 (d, J = 1.6 Hz, 1H), 8.64 (d, J = 1.6 Hz, 1H), 8.16 (dd, J = 8.8, 1.6 Hz, 1H), 7.85 (d, J = 8.8 Hz, 1H), 7.55-7.52 (m, 1H), 7.46 - 7.41 (m, 4H), 7.29 (t, J = 7.2 Hz, 1H), 7.23 - 7.19 (m, 1H), 7.14-7.06 (m, 2H), 7.01 (dt, J = 9.2, 2.0 Hz, 1H), 6.96 (s, 1H), 3.87 - 3.76 (m, 5H), 2.94-2.93 (m, 1H), 2.68 (t, J = 7.8 Hz, 2H), 1.89-1.83 (m, 2H).	DMSO	95	Method G1,AQ1	522.3 (M+1)	Method B (NH ₄ HCO ₃)

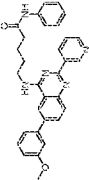
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1566					¹ H-NMR (400 MHz, DMSO- d ₆): δ 9.59 (s, 1H), 8.76 (t, J = 5.2 Hz, 1H), 8.69 (dd, J = 4.4, 1.2 Hz, 1H), 8.63 (s, 1H), 8.56 (d, J = 8.4 Hz, 1H), 8.15 (dd, J = 8.8, 1.4 Hz, 1H), 7.84 (d, J = 8.4 Hz, 1H), 7.53-7.50 (m, 1H), 7.48 - 7.40 (m, 3H), 7.36-7.32 (m, 3H), 7.30 - 7.24 (m, 2H), 7.02-7.00 (m, 1H), 6.85 (s, 1H), 3.92 - 3.83 (m, 4H), 3.72-3.69 (m, 1H), 3.18 - 3.14 (m, 1H), 2.96-2.84 (m, 2H).	DMSO	95	Method G1,AQ1	524.2, 526.2 (M+1)	Method B (NH ₄ HCO ₃)

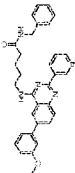
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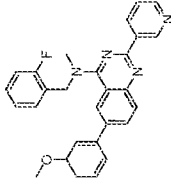
5	1567 	Number		Starting Material 1	Starting Material 2	Product	Salt type	1H NMR	1H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
10								1H-NMR (400 MHz, DMSO-d ₆): δ 9.64 (s, 1H), 9.08 (s, 1H), 8.79 - 8.69 (m, 2H), 8.51 (s, 1H), 8.31 (s, 1H), 8.00 - 7.89 (m, 2H), 7.85 - 7.69 (m, 1H), 7.67 - 7.42 (m, 2H), 7.30 - 7.21 (m, 2H), 5.00 (d, J = 5.1 Hz, 2H).	DMSO	95	Method G1,AQ1	416.0 (M+1)	Method B (NH ₄ HCO ₃)
15													
20													
25													
30													
35													
40													
45													
50													
55													

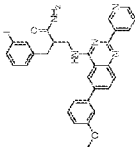
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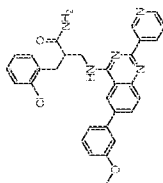
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1568				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.58 (s, 1H), 10.09 (d, J=4.4 Hz, 1H), 9.65 (s, 1H), 9.05 - 9.02 (m, 2H), 8.95 (d, J = 4.4 Hz, 1H), 8.37 (d, J = 8.4 Hz, 1H), 8.30-8.27 (m, 1H), 7.85 (t, J = 6.2 Hz, 1H), 7.59 (d, J = 8.0 Hz, 2H), 7.49-7.43 (m, 3H), 7.24 (t, J = 7.8 Hz, 2H), 7.03-6.97 (m, 2H), 3.89 (s, 3H), 3.86 - 3.84 (m, 2H), 2.44 (t, J=7.0 Hz, 2H), 1.87 - 1.77 (m, 4H).	DMSO	95	Method G1,AQ1	504.1 (M+1)	Method B (NH ₄ HCO ₃)

(continued)

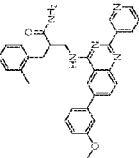
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1569				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.62 (s, 1H), 9.67 (s, 1H), 9.08 (s, 1H), 9.08 (d, J=7.2 Hz, 1H), 9.03 (s, 1H), 8.97 (d, J=5.2 Hz, 1H), 8.44 (t, J=5.6 Hz, 1H), 8.40 (dd, J=8.8, 0.8 Hz, 1H), 8.32 (dd, J=9.2, 2.8 Hz, 1H), 7.89 (t, J=6.4 Hz, 1H), 7.50-7.44 (m, 3H), 7.25-7.14 (m, 5H), 7.03 (d, J=7.2 Hz, 1H), 4.24 (d, J=6.0 Hz, 2H), 3.89 (s, 3H), 3.87-3.82 (m, 2H), 2.26 (t, J=7.0 Hz, 2H), 1.84-1.69 (m, 4H).	DMSO	95	Method G1,AQ1	518.1 (M+1)	Method B (NH ₄ HCO ₃)

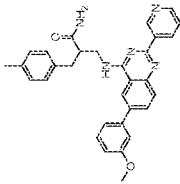
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1570					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.55 (s, 1H), 8.69 (dd, J = 11.3, 6.2 Hz, 2H), 8.23 (s, 1H), 8.15 (d, J = 8.7 Hz, 1H), 7.95 (d, J = 8.7 Hz, 1H), 7.59-7.50 (m, 2H), 7.44-7.26 (m, 3H), 7.23 (t, J = 7.5 Hz, 1H), 7.15 (s, 1H), 7.12 (d, J = 7.7 Hz, 1H), 6.95 (d, J = 8.1 Hz, 1H), 5.18 (s, 2H), 3.79 (s, 3H), 3.54 (s, 3H).	DMSO	95	Method G1, AQ1	451.2, 452.2, 453.2 (M+1)	Method B (NH ₄ HCO ₃)

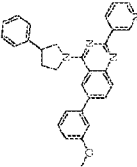
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1571				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.62 (s, 1H), 9.60 (s, 1H), 8.88 (d, J = 4.3 Hz, 1H), 8.82 (s, 1H), 8.78 (d, J = 8.1 Hz, 1H), 8.31 (d, J = 8.6 Hz, 1H), 8.12 (d, J = 7.9 Hz, 1H), 7.75 (dd, J = 7.8, 5.1 Hz, 1H), 7.53 - 7.39 (m, 4H), 7.33 (dd, J = 14.3, 7.9 Hz, 1H), 7.12 (t, J = 7.8 Hz, 2H), 7.03 (dd, J = 11.0, 6.3 Hz, 2H), 6.90 (s, 1H), 3.97 - 3.90 (m, 4H), 3.82 (dd, J = 18.0, 9.5 Hz, 1H), 3.19 (dd, J = 13.4, 7.5 Hz, 1H), 2.98 (dd, J = 13.6, 7.9 Hz, 1H), 2.87 (dd, J = 13.6, 6.7 Hz, 1H).	DMSO	95	Method G1,AQ1	508.1, 509.1, 510.1 (M+1)	Method B (NH ₄ HCO ₃)

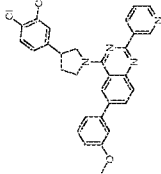
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1572				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.20 (s, 1H), 9.61 (s, 1H), 8.94 (s, 1H), 8.94 (d, J = 4.8 Hz, 1H), 8.86 (s, 2H), 8.37 (d, J = 8.4 Hz, 1H), 8.22 (s, 1H), 7.83 (s, 1H), 7.56-7.32 (m, 6H), 7.31 - 7.19 (m, 2H), 7.06 (d, J = 7.2 Hz, 1H), 6.89 (s, 1H), 4.03-3.93 (m, 1H), 3.88 (s, 3H), 3.82 (dd, J = 14.4, 5.5 Hz, 1H), 3.22 (s, 1H), 3.02 (d, J = 7.2 Hz, 2H).	DMSO	95	Method G1,AQ1	524.1, 525.1, 526.0 (M+1)	Method B (NH ₄ HCO ₃)

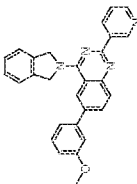
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1573				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.31 (s, 1H), 9.63 (s, 1H), 8.96 (s, 1H), 8.96 (d, J = 4.4 Hz, 1H), 8.90 (s, 1H), 8.84 (d, J = 7.7 Hz, 1H), 8.39 (d, J = 8.7 Hz, 1H), 8.27 (d, J = 8.0 Hz, 1H), 8.0 Hz, 1H), 7.91-7.80 (m, 1H), 7.48 (dd, J = 15.6, 7.7 Hz, 3H), 7.36 (s, 1H), 7.24 - 7.03 (m, 5H), 6.85 (s, 1H), 4.03-3.93 (m, 1H), 3.88 (s, 3H), 3.83 (dd, J = 14.7, 7.6 Hz, 1H), 3.15 (s, 1H), 2.94 (dd, J = 13.8, 8.0 Hz, 1H), 2.84 (dd, J = 13.8, 6.6 Hz, 1H), 2.29 (s, 3H).	DMSO	95	Method G1, AQ1	504.1, 505.2, 506.1 (M+1)	Method B (NH ₄ HCO ₃)

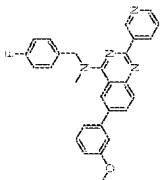
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1574				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.43 (s, 1H), 9.71 (s, 1H), 9.05 (s, 1H), 9.01 (d, J = 7.6 Hz, 1H), 8.96 (s, 1H), 8.45 (d, J = 8.6 Hz, 1H), 8.38 (d, J = 8.4 Hz, 1H), 7.99-7.89 (m, 1H), 7.61 - 7.41 (m, 4H), 7.23 (d, J = 7.8 Hz, 2H), 7.13 (d, J = 7.6 Hz, 3H), 6.93 (s, 1H), 4.01-3.84 (m, 5H), 3.31 - 3.21 (m, 1H), 3.02-2.97 (m, 7.5 Hz, 1H), 2.87-2.81 (m, 1H), 2.29 (s, 3H).	DMSO	95	Method G1,AQ1	504.1 (M+1)	Method B (NH ₄ HCO ₃)

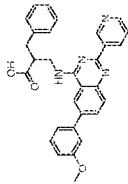
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1575					¹ H-NMR(400 MHz, CD ₃ OD): δ 9.39 (s, 1H), 8.69 (d, <i>J</i> = 8.0 Hz, 1H), 8.65 (s, 1H), 8.38 (s, 1H), 8.07 (d, <i>J</i> =7.9 Hz, 1H), 7.85 (d, <i>J</i> =8.6 Hz, 1H), 7.60 - 7.53 (m, 1H), 7.35-7.25(m, 5H), 7.18(dd, <i>J</i> = 13.6, 7.2 Hz, 2H), 7.12 (s, 1H), 6.88 (dd, <i>J</i> = 7.9, 2.1 Hz, 1H), 4.49 (dd, <i>J</i> = 11.3, 7.9 Hz, 1H), 4.36 - 4.17 (m, 2H), 4.04 (t, <i>J</i> = 10.4 Hz, 1H), 3.74 (s, 3H), 3.60-3.49(m, 1H), 2.50 - 2.39 (m, 1H), 2.27-2.14 (m, 1H).	CD3OD	95	Method G1,AQ1	459.0 (M+1)	Method B (NH ₄ HCO ₃)

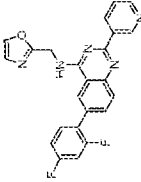
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1576					¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.67 (d, J = 1.7 Hz, 1H), 9.05 (s, 1H), 8.94 (d, J = 3.7 Hz, 1H), 8.59 (s, 1H), 8.40-8.22 (m, 2H), 7.85 (s, 1H), 7.79 (d, J = 1.9 Hz, 1H), 7.66 (d, J = 8.3 Hz, 1H), 7.52-7.36 (m, 4H), 7.04 (d, J = 7.6 Hz, 1H), 4.74-4.72 (m, 2H), 4.48 - 4.43 (m, 3H), 3.85 (s, 3H), 3.74-3.60 (m, 1H), 2.25 - 2.23 (m, 1H).	DMSO	95	Method G1,AQ1	527.0, 529.0 (M+1)	Method B (NH ₄ HCO ₃)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1577				2HCl	¹ H-NMR (400 MHz, DMSO): δ 9.69 (d, <i>J</i> = 1.6 Hz, 1H), 8.89 (d, <i>J</i> = 8.0 Hz, 1H), 8.79 (d, <i>J</i> = 1.6 Hz, 1H), 8.73 (s, 1H), 8.21 (d, <i>J</i> = 9.2 Hz, 1H), 8.01 (d, <i>J</i> = 8.8 Hz, 1H), 7.67 (dd, <i>J</i> = 7.6, 4.8 Hz, 1H), 7.56 - 7.38 (m, 7H), 7.05 - 7.02 (m, 1H), 5.59 (s, 4H), 3.89 (s, 3H).	DMSO	95	Method G1, AQ1	431.1 (M+1)	Method B (NH ₄ HCO ₃)

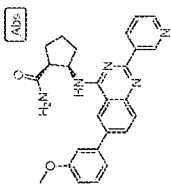
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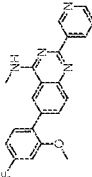
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1578					¹ H-NMR (400 MHz, DMSO): δ 9.59 (d, J = 1.6 Hz, 1H), 8.81-8.64 (m, 2H), 8.28 - 8.10 (m, 2H), 7.96 (d, J=8.6 Hz, 1H), 7.55 (td, J = 8.5, 5.0 Hz, 3H), 7.40-7.06 (m, 5H), 6.94 (dd, J = 8.2, 2.0 Hz, 1H), 5.15 (s, 2H), 3.78 (s, 3H), 3.51 (s, 3H).	DMSO	95	Method G1, AQ1	451.2 (M+1)	Method B (NH ₄ HCO ₃)

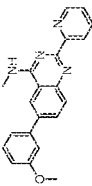
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1579				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 10.61 (s, 1H), 9.66 (s, 1H), 9.00 (s, 1H), 9.00 (d, J = 13.5 Hz, 3H), 8.43 - 8.31 (m, 2H), 7.89 (s, 1H), 7.48 (dd, J = 14.6, 7.2 Hz, 3H), 7.33 - 7.22 (m, 4H), 7.21 - 7.14 (m, 1H), 7.05 (d, J = 7.2 Hz, 1H), 4.10 - 3.85 (m, 2H), 3.90 (s, 3H), 3.37 - 3.27 (m, 1H), 3.07 - 2.91 (m, 2H).	DMSO	95	Method G1,AQ1	491.2 (M+1)	Method B (NH ₄ HCO ₃)

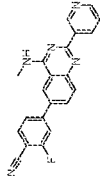
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1580				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.91 (s, 1H), 9.60 (s, 1H), 9.11 (s, 1H), 8.95 (s, 1H), 8.66 (s, 1H), 8.20 - 8.02 (m, 3H), 7.97 (s, 1H), 7.78 (dd, J = 13.3, 6.5 Hz, 1H), 7.50 (t, J = 11.3 Hz, 1H), 7.33 (t, J = 8.1 Hz, 1H), 7.21 (s, 1H), 5.08 (d, J = 3.8 Hz, 2H).	DMSO	95	Method G1,AQ1	416.0 (M+1)	Method B (NH ₄ HCO ₃)

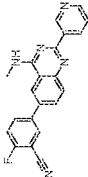
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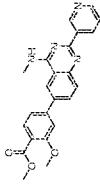
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1581				2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.84 (s, 1H), 9.71 (s, 1H), 9.12 (d, J = 6.2 Hz, 1H), 8.96 (d, J = 4.8 Hz, 1H), 8.91 (s, 1H), 8.37 (d, J = 8.4 Hz, 1H), 8.30 (s, 1H), 7.88 (s, 1H), 7.57-7.38 (m, 3H), 7.24 (s, 1H), 7.08 (d, J = 7.9 Hz, 1H), 6.73 (s, 1H), 5.18-5.05 (m, 1H), 3.88 (s, 3H), 3.23 (dd, J = 14.8, 7.4 Hz, 1H), 2.31-2.17 (m, 1H), 2.15-1.89 (m, 4H), 1.71-1.57 (m, 1H).	DMSO	95	Method G1,AQ1	440.2 (M+1)	Method B (NH ₄ HCO ₃)

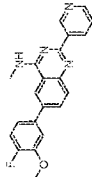
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1583				2HCl	¹ H-NMR(300 MHz,DMSO): δ 10.39 (s, 1H), 9.71 (s, 1H), 9.14 (d, J = 8.1 Hz, 1H), 8.98 (d, J = 4.0 Hz, 1H), 8.71 (s, 1H), 8.31 (d, J = 8.8 Hz, 1H), 8.25-8.19(m, 1H), 7.96 - 7.89 (m, 1H), 7.38 (dd, J = 9.2, 3.0 Hz, 1H), 7.34 - 7.14 (m, 2H), 3.81 (s, 4H), 3.30 (d, J = 4.4 Hz, 4H).	DMSO	95	Method AQ2, AP	361.7 (M+1)	Method C

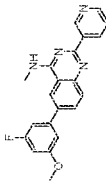
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1584				2HCl	¹ H-NMR(300 MHz,DMSO): δ 10.84 (s, 1H), 9.01 (s, 1H), 8.95 (d, J = 3.9 Hz, 1H), 8.72 (d, J = 7.8 Hz, 1H), 8.50-8.40 (m, 1H), 8.33 (d, J = 8.8 Hz, 1H), 8.24 (td, J = 7.8, 1.7 Hz, 1H), 7.85 (dd, J = 6.5, 4.8 Hz, 1H), 7.57-7.43 (m, 3H), 7.09-7.02 (m, 1H), 3.90 (s, 3H), 3.38 (s, 3H)	DMSO	95	Method AQ2, AP	343.5 (M+1)	Method C

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1585				2HCl	¹ H-NMR(300 MHz, DMSO): δ 10.44 - 10.13 (m, 1H), 9.66 (s, 1H), 9.08 (d, J = 10.1 Hz, 2H), 8.96 (d, J = 3.7 Hz, 1H), 8.44 (d, J = 8.9 Hz, 1H), 8.22 (d, J = 8.4 Hz, 1H), 8.13 (dd, J = 13.3, 5.3 Hz, 2H), 8.00 (d, J = 8.2 Hz, 1H), 7.91 (s, 1H), 3.30 (d, J = 4.4 Hz, 3H).	DMSO	95	Method AQ2, AP	356.4 (M+1)	Method C

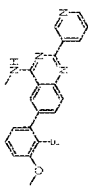
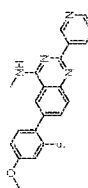
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1586					¹ H-NMR (300 MHz, DMSO): δ 9.64 (s, 1H), 8.78 (d, J = 8.0 Hz, 1H), 8.73-8.69 (m, 1H), 8.63 (s, 2H), 8.47 - 8.37 (m, 1H), 8.32-8.25 (m, 1H), 8.23 - 8.12 (m, 1H), 7.87 (d, J = 8.6 Hz, 1H), 7.73 (t, J = 9.0 Hz, 1H), 7.55 (dd, J = 7.9, 4.7 Hz, 1H), 3.20 (d, J = 4.3 Hz, 3H).	DMSO	95	Method AQ2, AP	356.4 (M+1)	Method C

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1587					¹ H-NMR(300 MHz, DMSO): δ 9.65 (s, 1H), 8.79 (d, J = 8.0 Hz, 1H), 8.72-8.57 (m, 2H), 8.24 (d, J = 8.7 Hz, 1H), 7.88 (d, J = 8.7 Hz, 1H), 7.82 (d, J = 8.0 Hz, 1H), 7.76 (d, J=8.0 Hz, 1H), 7.53 (t, J=10.9 Hz, 2H), 7.45 - 7.36 (m, 1H), 3.99 (s, 3H), 3.94 (s, 3H), 3.21 (d, J = 4.1 Hz, 3H).	DMSO	95	Method AQ2, AP	401.5 (M+1)	Method C

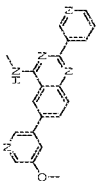
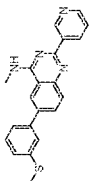
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1588				2HCl	¹ H-NMR(300 MHz,DMSO): δ 10.77 (s, 1H), 9.66 (s, 1H), 9.06 (s, 1H), 8.95(d, J = 3.8 Hz, 1H), 8.38 (d, J = 8.8 Hz, 1H), 8.26 (d, J = 8.8 Hz, 1H), 7.87 (s, 1H), 7.71 (d, J = 8.1 Hz, 1H), 7.38 (dd, J = 22.8, 11.6 Hz, 2H), 4.03 (s, 3H), 3.31(d, J = 4.1 Hz, 3H).	DMSO	95	Method AQ2, AP	361.5 (M+1)	Method C

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1589				2HCl	1H-NMR (300 MHz, DMSO): δ 10.23 - 10.05 (m, 1H), 9.64 (s, 1H), 9.00 (s, 1H), 8.94 (s, 1H), 8.89 (s, 1H), 8.38 (d, J = 8.6 Hz, 1H), 8.14 (d, J = 8.5 Hz, 1H), 7.85 (s, 1H), 7.36 (d, J = 8.7 Hz, 2H), 6.95 (d, J = 10.9 Hz, 1H), 3.90 (s, 1H), 3.30 (d, J = 4.3 Hz, 3H).	DMSO	95	Method AQ2, AP	361.5 (M+1)	Method C

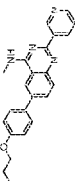
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1590				2HCl	¹ H-NMR (300 MHz, DMSO): δ 10.34 (s, 1H), 9.70 (s, 1H), 9.14 (d, J = 7.4 Hz, 1H), 8.98 (d, J = 3.9 Hz, 1H), 8.75 (s, 1H), 8.32 (d, J = 8.8 Hz, 1H), 8.17 (d, J = 8.8 Hz, 1H), 7.93 (d, J = 4.8 Hz, 1H), 7.27 (dd, J = 19.5, 6.3 Hz, 3H), 3.88 (s, 3H), 3.28 (d, J = 4.1 Hz, 3H).	DMSO	95	Method AQ2, AP	361.3 (M+1)	Method C
1591				2HCl	¹ H-NMR (300 MHz, DMSO): δ 10.21 (s, 1H), 9.66 (d, J = 1.6 Hz, 1H), 9.04 (d, J = 7.3 Hz, 1H), 8.96 (d, J = 3.5 Hz, 1H), 8.67 (s, 1H), 8.28-8.12 (m, 2H), 7.88 (s, 1H), 7.66 (t, J = 8.8 Hz, 1H), 7.11-6.97 (m, 2H), 3.86 (s, 3H), 3.30 (d, J = 4.4 Hz, 3H).	DMSO	95	Method AQ2, AP	361.3 (M+1)	Method C

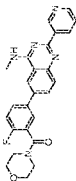
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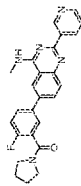
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1592				3HCl	¹ H-NMR (300 MHz, DMSO): δ 10.77 (s, 1H), 9.69 (s, 1H), 9.39 (s, 1H), 9.16 (d, J = 7.8 Hz, 1H), 9.04 (s, 1H), 8.97 (d, J = 3.6 Hz, 1H), 8.63 (d, J = 2.5 Hz, 1H), 8.53 (s, 2H), 8.31 (d, J = 8.5 Hz, 1H), 7.91 (d, J = 5.4 Hz, 1H), 4.11 (s, 3H), 3.30 (d, J = 4.4 Hz, 3H).	DMSO	95	Method AQ2, AP	343.1 (M+1)	Method C
1593				2HCl		DMSO	95	Method AQ2, AP	359.1 (M+1)	Method C

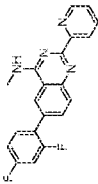
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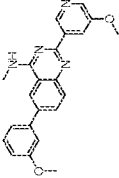
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1594					¹ H-NMR (300 MHz, DMSO): δ 9.64 (d, J = 1.3 Hz, 1H), 8.82-8.75 (m, 1H), 8.68 (dd, J = 4.8, 1.7 Hz, 1H), 8.60 (s, 1H), 8.53 (d, J = 1.7 Hz, 1H), 8.10 (dd, J = 8.7, 1.9 Hz, 1H), 7.86 - 7.77 (m, 4H), 7.58-7.48 (m, 2H), 7.09 (d, J = 8.8 Hz, 2H), 6.97 (d, J = 8.8 Hz, 1H), 3.99 (d, J = 6.0 Hz, 2H), 3.18 (d, J = 4.4 Hz, 3H), 1.82-1.69 (m, 2H), 1.07 (d, J = 6.7 Hz, 3H).	DMSO	95	Method AQ2, AP	371.1 (M+1)	Method C

(continued)

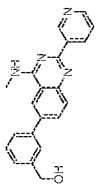
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1595				2HCl	¹ H-NMR (300 MHz, DMSO): δ 10.24 - 10.10 (bs, 1H), 9.65 (s, 1H), 9.02 (s, 1H), 8.96 (d, J = 3.8 Hz, 1H), 8.90 (s, 1H), 8.39 (s, 1H), 8.20 (s, 1H), 8.07 (s, 1H), 7.98 (s, 1H), 7.89 (s, 1H), 7.54 (t, J = 9.1 Hz, 1H), 3.70 (bs, 4H), 3.59 (bs, 4H), 3.32 (d, J = 4.4 Hz, 3H).	DMSO	95	Method AQ2, AP	443.5 (M+1)	Method C

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1596				2HCl	¹ H-NMR(300 MHz,DMSO): δ 10.37 (s, 1H), 9.69(d, <i>J</i> = 1.7 Hz, 1H), 9.10 (d, <i>J</i> = 7.0 Hz, 1H), 9.03-8.93(m, 2H), 8.42(d, <i>J</i> = 8.6 Hz, 1H), 8.28 (d, <i>J</i> = 9.3Hz, 1H), 8.05 (d, <i>J</i> = 3.5 Hz, 1H), 7.99 (dd, <i>J</i> = 6.2, 2.4 Hz, 2H), 7.52 (t, <i>J</i> = 9.1 Hz, 1H), 3.54-3.43(m, 4H), 3.30 (t, <i>J</i> = 10.4 Hz, 3H), 1.98 - 1.81 (m, 4H).	DMSO	95	Method AQ2, AP	428.5 (M+1)	Method C

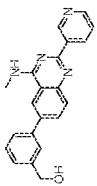
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1597				2HCl	¹ H-NMR(300 MHz,DMSO): δ 10.71 (s, 1H), 8.96 (d, J = 3.9 Hz, 1H), 8.81 (s, 1H), 8.74 (d, J = 7.9 Hz, 1H), 8.36 (d, J = 8.7 Hz, 1H), 8.29-8.19(m, 2H), 7.87 (dd, J = 6.5, 4.7 Hz, 1H), 7.80 (dd, J = 15.5, 8.9 Hz, 1H), 7.61-7.50(m, 1H), 7.36 (d, J = 8.1 Hz, 1H), 3.37 (d, J = 4.5 Hz, 3H).	DMSO	95	Method AQ2, AP	349.4 (M+1)	Method C

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1598					¹ H-NMR(300 MHz,DMSO): δ 9.27 (d, J = 1.5 Hz, 1H), 8.61 (dd, J = 12.6, 3.0 Hz, 2H), 8.41 (d, J = 2.9 Hz, 1H), 8.34-8.27 (m, 1H), 8.16 (dd, J = 8.7, 1.9 Hz, 1H), 7.86 (d, J = 8.7 Hz, 1H), 7.45 (dd, J = 16.0, 9.2 Hz, 3H), 7.05-6.97 (m, 1H), 3.96 (s, 3H), 3.87 (s, 3H), 3.19 (d, J = 4.3 Hz, 3H).	DMSO	95	Method AQ2, AP	373.4 (M+1)	Method C

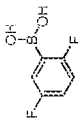
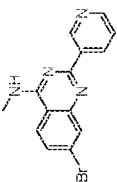
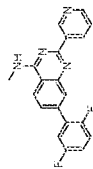
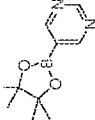
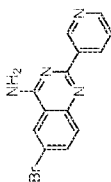
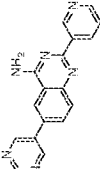
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1599				2HCl	¹ H-NMR (300 MHz, DMSO): δ 10.44 - 10.13 (m, 1H), 9.65 (s, 1H), 9.02 (s, 1H), 8.96 (d, 1H), 8.87 (s, 1H), 8.37 (d, J = 9.1 Hz, 1H), 8.24 (s, 1H), 7.86 (s, 2H), 7.78 (d, J = 7.9 Hz, 1H), 7.54 (d, J = 7.6 Hz, 1H), 7.43 (d, J = 7.5 Hz, 1H), 4.64 (s, 2H), 3.33 (d, J = 4.2 Hz, 3H), 3.17 (s, 1H).	DMSO	95	Method AQ2, AP	343.5 (M+1)	Method C

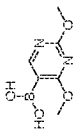
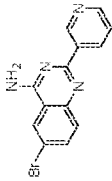
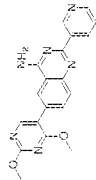
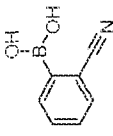
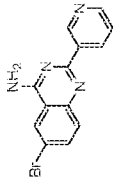
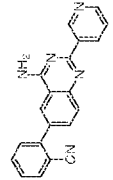
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1600				MSA	¹ H-NMR (300 MHz, DMSO): δ 9.90 (s, 1H), 9.57 (d, J = 1.6 Hz, 1H), 8.93 (dd, J = 5.0, 1.5 Hz, 2H), 8.73 (s, 1H), 8.32 (d, J = 8.7 Hz, 1H), 8.01 (d, J = 8.8 Hz, 1H), 7.85 (dd, J = 10.1, 7.0 Hz, 2H), 7.73 (d, J = 7.5 Hz, 1H), 7.52 (t, J = 7.6 Hz, 1H), 7.42 (d, J = 7.5 Hz, 1H), 4.63 (s, 2H), 3.30 (d, J = 4.3 Hz, 3H), 2.32 (s, 3H).	DMSO	95	Method AQ2, AP	343.5 (M+1)	Method C

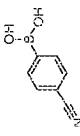
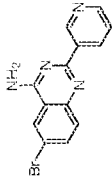
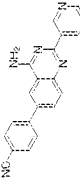
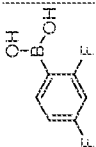
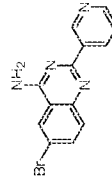
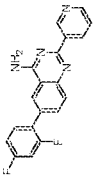
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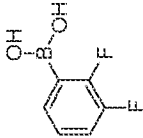
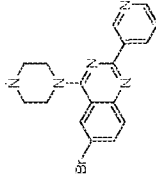
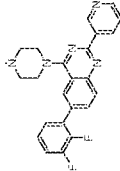
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1601				MsOH	¹ H NMR (300 MHz, DMSO) δ 9.72 (brs, 1H), 9.58 (d, J = 1.6 Hz, 1H), 9.03-8.89 (m, 2H), 8.45 (d, J = 8.7 Hz, 1H), 8.11 (s, 1H), 7.99-7.84 (m, 2H), 7.64 (ddd, J = 9.2, 6.1, 3.1 Hz, 1H), 7.58 - 7.31 (m, 2H), 3.28 (d, J = 4.5 Hz, 3H), 2.39 (s, 3H).	DMSO	>98	G2/AQ3		
1602					¹ H NMR (300 MHz, DMSO) δ 9.59 (d, J = 2.0 Hz, 1H), 9.33 (s, 2H), 9.24 (s, 1H), 8.79 (d, J = 1.5 Hz, 1H), 8.76-8.64 (m, 2H), 8.30 (dd, J = 8.7, 1.5 Hz, 1H), 8.12 (brs, 2H), 7.91 (d, J = 8.7 Hz, 1H), 7.55 (dd, J = 7.9, 4.8 Hz, 1H).	DMSO	>98	G2/AQ3		

(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1603					¹ H NMR (300 MHz, DMSO) δ 9.64 - 9.53 (m, 1H), 8.77 - 8.58 (m, 2H), 8.53 (s, 1H), 8.43 (d, J = 1.5 Hz, 1H), 8.11 - 7.90 (m, 3H), 7.82 (d, J = 8.7 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 4.00 (s, 3H), 3.98 (s, 3H).	DMSO	>98	G2/AQ3		
1604					¹ H NMR (300 MHz, DMSO) δ 9.59 (d, J = 2.0 Hz, 1H), 8.83 - 8.66 (m, 2H), 8.55 (s, 1H), 8.27 - 7.99 (m, 4H), 7.99 - 7.82 (m, 2H), 7.77 (d, J = 7.7 Hz, 1H), 7.66 (t, J = 7.6 Hz, 1H), 7.56 (dd, J = 7.9, 4.8 Hz, 1H).	DMSO	>98	G2/AQ3		

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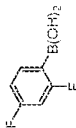
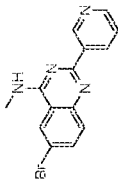
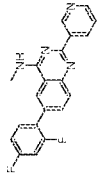
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1605					¹ H NMR (300 MHz, DMSO) δ 9.58 (d, J = 2.1 Hz, 1H), 8.80-8.64 (m, 2H), 8.25 (dd, J = 8.7, 1.6 Hz, 1H), 8.10 (d, J = 8.3 Hz, 2H), 8.02 (d, J = 8.3 Hz, 2H), 7.88 (d, J = 8.7 Hz, 1H), 7.55 (dd, J = 7.9, 4.8 Hz, 1H).	DMSO	>98	G2/AQ3		
1606				2HCl	¹ H NMR (300 MHz, DMSO) δ 9.95 - 9.43 (m, 3H), 9.05 (d, J = 8.2 Hz, 1H), 8.99 (dd, J = 5.1, 1.5 Hz, 1H), 8.74 (s, 1H), 8.33 (d, J = 8.7 Hz, 1H), 8.21 (d, J = 8.7 Hz, 1H), 7.93 (dd, J = 8.0, 5.1 Hz, 1H), 7.79 (td, J = 8.9, 6.6 Hz, 1H), 7.51 (ddd, J = 11.5, 9.3, 2.6 Hz, 1H), 7.33 (td, J = 8.3, 1.9 Hz, 1H).	DMSO	>98	G2/AQ3		

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1607				3HCl	¹ H NMR (300 MHz, DMSO) δ 11.88 (s, 1H), 9.77 (d, 1H), 9.56 - 9.28 (m, 1H), 9.19 - 9.01 (m, 1H), 8.31 - 8.11 (m, 4H), 7.75 - 7.47 (m, 2H), 7.47 - 7.23 (m, 1H), 4.83 - 4.67 (m, 2H), 3.96 (t, J = 12.2 Hz, 2H), 3.57 (d, J = 11.9 Hz, 2H), 3.46 - 3.19 (m, 2H), 2.83 (d, J = 3.1 Hz, 3H).	DMSO	>98	G2/AQ03		

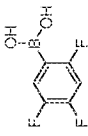
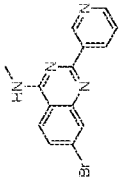
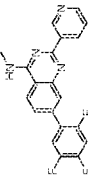
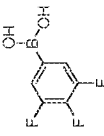
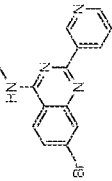
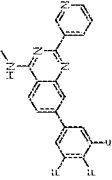
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Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1608				HCl	¹ H NMR (300 MHz, DMSO) δ 14.45 (s, 1H), 10.36 (s, 1H), 9.31 (d, J = 2.2 Hz, 1H), 8.82 (s, 1H), 8.73 (ddd, J = 8.8, 2.5, 0.9 Hz, 1H), 8.36 (d, J = 9.0 Hz, 1H), 8.16 (d, J = 8.4 Hz, 1H), 7.56-7.36 (m, 3H), 7.11 (d, J = 8.4 Hz, 1H), 7.03 (d, J = 7.1 Hz, 1H), 3.99 (s, 3H), 3.87 (s, 3H), 3.30 (d, J = 3.9 Hz, 3H).	DMSO	100	Method AQ2/BF		
1609				HCl	¹ H NMR (300 MHz, DMSO) δ 10.97 (s, 1H), 9.56 - 9.35 (m, 2H), 9.04-8.81 (m, 2H), 8.66 (t, J = 7.7 Hz, 1H), 8.61-8.45 (m, 2H), 8.15 (d, J = 8.6 Hz, 1H), 8.10-7.96 (m, 1H), 7.77 - 7.58 (m, 1H), 3.25 (d, J = 4.1 Hz, 3H).	DMSO	100	Method AQ2/BF		

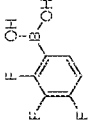
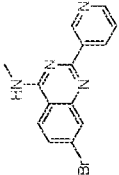
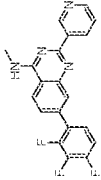
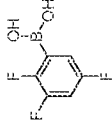
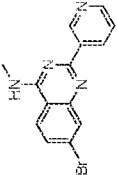
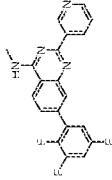
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1610				2 MsOH	¹ H NMR (300 MHz, CDCl ₃) δ 9.93 (brs, 1H), 9.61 (d, J= 1.8 Hz, 1H), 9.10 - 8.96 (m, 2H), 8.64 (s, 1H), 8.21 (d, J= 8.7 Hz, 1H), 8.06 (d, J= 8.7 Hz, 1H), 7.99 (dd, J= 8.0, 5.3 Hz, 1H), 7.69 - 7.47 (m, 2H), 7.47 - 7.34 (m, 1H), 3.30 (d, J= 4.5 Hz, 3H), 2.42 (s, 6H).	DMSO	>98	G2/AQ3	Metho d 3	

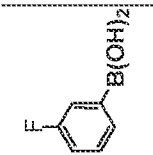
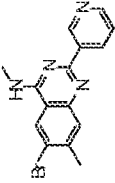
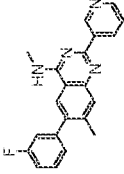
Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1611				2HCl	¹ H NMR (300 MHz, CDCl ₃) δ 10.42 (brs, 1H), 9.71 (d, <i>J</i> = 1.9 Hz, 1H), 9.17 (d, <i>J</i> = 8.1 Hz, 1H), 8.99 (dd, <i>J</i> = 5.1, 1.3 Hz, 1H), 8.78 (s, 1H), 8.33 (d, <i>J</i> = 8.7 Hz, 1H), 8.17 (d, <i>J</i> = 8.7 Hz, 1H), 7.94 (dd, <i>J</i> = 8.1, 5.2 Hz, 1H), 7.79 (td, <i>J</i> = 8.9, 6.7 Hz, 1H), 7.55 - 7.37 (m, 1H), 7.31 (td, <i>J</i> = 8.4, 2.2 Hz, 1H), 3.29 (d, <i>J</i> = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1612				HCl	¹ H NMR (300 MHz, DMSO) δ 9.95 (brs, 1H), 9.65 (s, 1H), 9.18 - 9.02 (m, 1H), 8.95 (d, J = 4.7 Hz, 1H), 8.58 (d, J = 8.7 Hz, 1H), 8.28 (s, 1H), 8.09 - 7.70 (m, 4H), 3.29 (d, J = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		
1613					¹ H NMR (300 MHz, DMSO) δ 9.64 (d, J = 1.5 Hz, 1H), 8.77 (dt, J = 7.9, 1.9 Hz, 1H), 8.69 (dd, J = 4.8, 1.7 Hz, 1H), 8.61 Hz, 1H), 8.61 - 8.48 (m, 1H), 8.31 (d, J = 8.7 Hz, 1H), 8.11 (d, J = 1.8 Hz, 1H), 8.00 - 7.83 (m, 3H), 7.55 (dd, J = 7.9, 4.8 Hz, 1H), 3.17 (d, J = 4.4 Hz, 3H).	DMSO	>98	G2/AQ3		

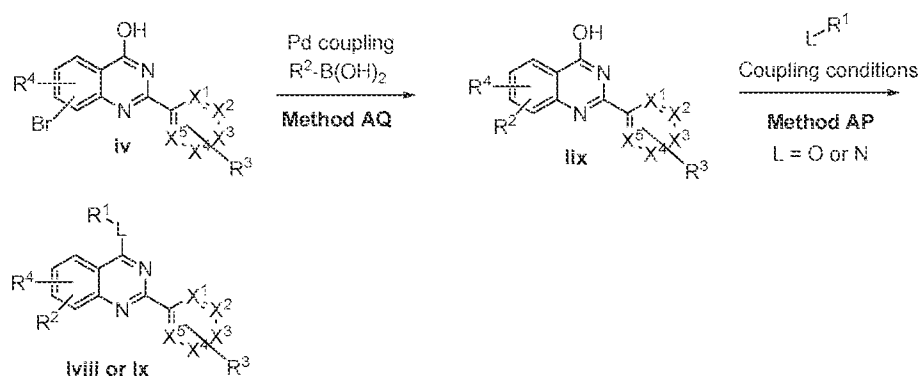
(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per- cent	Method of Coupling	LCMS	LCMS Meth- od
1614					¹ H NMR (300 MHz, DMSO) δ 9.65 (dd, J = 2.1, 0.8 Hz, 1H), 8.78 (dt, J = 8.0, 1.9 Hz, 1H), 8.69 (dd, J = 4.8, 1.7 Hz, 1H), 8.65-8.50 (m, 1H), 8.33 (d, J = 8.6 Hz, 1H), 8.04-7.85 (m, 1H), 7.69 (dt, J = 8.5, 1.8 Hz, 1H), 7.65-7.37 (m, 3H), 3.18 (d, J = 4.5 Hz, 3H).	DMSO	>98	G2/AQ3	Method 3	
1615					¹ H NMR (300 MHz, DMSO) δ 9.65 (d, J = 1.7 Hz, 1H), 8.84-8.74 (m, 1H), 8.69 (dd, J = 4.7, 1.5 Hz, 1H), 8.65-8.52 (m, 1H), 8.34 (d, J = 8.6 Hz, 1H), 7.98 (s, 1H), 7.72 (d, J = 8.5 Hz, 1H), 7.69-7.43 (m, 3H), 3.18 (d, J = 4.3 Hz, 3H).	DMSO	>98	G2/AQ3	Method 3	

Number	Starting Material 1	Starting Material 2	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity per-cent	Method of Coupling	LCMS	LCMS Method
1760				2 HCl	¹ H NMR (400 MHz, DMSO) δ 10.04 (s, 1H), 9.65 (s, 1H), 9.11 - 1H), 9.11 - 8.99 (m, 1H), 8.96 (dd, J= 5.0, 1.4 Hz, 1H), 8.41 (s, 1H), 8.13 - 8.05 (m, 1H), 7.92 - 7.82 (m, 1H), 7.64 - 7.55 (m, 1H), 7.44 - 7.26 (m, 3H), 3.27 (d, J= 4.6 Hz, 3H), 2.47 (s, 3H).	DMSO	>98	AQ5	ND	4

Scheme 76: General route for the synthesis of compounds with general formula Iviii and Ix

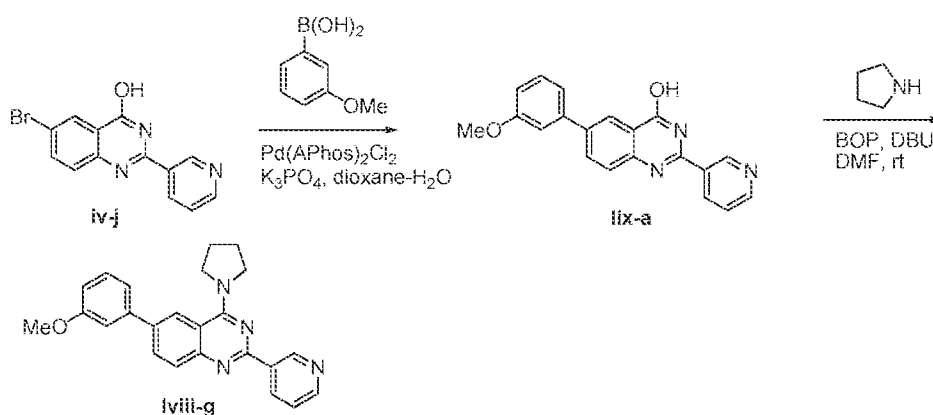
[0500]



X^1, X^2, X^3, X^4 or $X^5 = CH$ or N , at least one X^1, X^2, X^3, X^4 or X^5 must be N

Scheme 77: Representative synthesis of compounds of formula Iviii (see Scheme 76)

[0501]

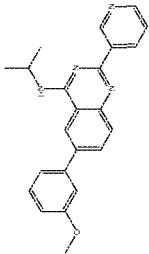
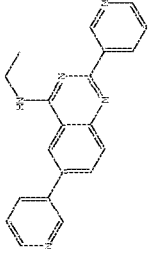


[0502] 6-(3-methoxyphenyl)-2-(pyridine-3-yl)quinazoline-4-ol (Iix-a) 6-(3-methoxyphenyl)-2-(pyridine-3-yl)quinazoline-4-ol was prepared from 6-bromo-2-(pyridine-3-yl)quinazolin-4-ol (synthesized following Scheme 70 substituting 2-amino-5-bromobenzamide for 2-amino-5-bromo-3-methylbenzamide) and 3-methoxyphenylboronic acid as described in Scheme 72 using method AQ2. The resultant product, 6-(6-methoxypyridin-3-yl)-N-methyl-2-(pyridine-3-yl)quinazoline-4-amine, was a pale yellow solid (19.1 mg, 51%). LCMS $m/z = 344$ ($M+1$) (Method C) (retention time = 2.01 min). ¹H NMR (300 MHz, DMSO) δ 9.64 (d, $J = 1.3$ Hz, 1H), 8.84 - 8.74 (m, 1H), 8.68 (dd, $J = 6.2, 1.7$ Hz, 2H), 8.57 (d, $J = 1.6$ Hz, 2H), 8.16 (ddd, $J = 14.4, 8.7, 2.2$ Hz, 2H), 7.85 (d, $J = 8.7$ Hz, 1H), 7.54 (dd, $J = 7.9, 4.8$ Hz, 1H), 7.00 (d, $J = 8.7$ Hz, 1H), 3.93 (s, 3H), 3.18 (d, $J = 4.3$ Hz, 3H).

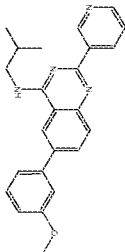
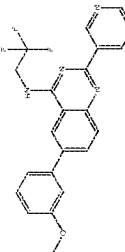
[0503] Method AP: 6-(3-methoxyphenyl)-2-(pyridine-3-yl)-4-(pyrrolidin-1-yl)quinazoline (Iviii-g) 6-(3-methoxyphenyl)-2-(pyridine-3-yl)-4-(pyrrolidin-1-yl)quinazoline was prepared from 6-(3-methoxyphenyl)-2-(pyridine-3-yl)quinazoline-4-ol and pyrrolidine in a manner analogous to that described for 6-bromo-N-methyl-2-(pyridine-3-yl)quinazoline-4-amine using Method AP in Scheme 72. 6-(3-methoxyphenyl)-2-(pyridine-3-yl)-4-(pyrrolidin-1-yl)quinazoline was a pale yellow solid (43 mg, 31%). LCMS $m/z = 383$ ($M+1$) (Method C) (retention time = 2.49 min). ¹H NMR (300 MHz, DMSO) δ 9.62 (s, 1H), 8.94 (d, $J = 5.0$ Hz, 2H), 8.56 (s, 1H), 8.32 (dd, $J = 19.9, 8.5$ Hz, 2H), 7.83 (s, 1H), 7.56 - 7.30 (m, 3H), 7.04 (d, $J = 6.8$ Hz, 1H), 4.27 (s, 4H), 3.86 (s, 3H), 2.08 (s, 4H).

The compounds in the following table were prepared in a manner analogous to that described in Scheme 76, replacing pyrrolidine with the appropriate amine and 3-methoxyphenylboronic acid with the appropriate boronic acid.

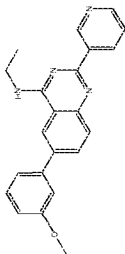
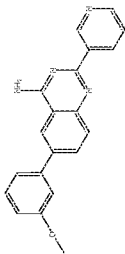
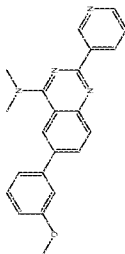
Table 25:

Number	Product	Salt	¹ H-NMR	¹ H NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity Percent	Method for Coupling
1616		HCl	¹ H NMR (300 MHz, DMSO) δ 9.82 (s, 1H), 9.59 (s, 1H), 9.08 - 8.81 (m, 3H), 8.36 (d, J = 8.8 Hz, 1H), 8.17 (d, J = 8.6 Hz, 1H), 7.95 - 7.77 (m, 1H), 7.58 - 7.37 (m, 3H), 7.05 (d, J = 6.5 Hz, 1H), 5.01 - 4.79 (m, 1H), 3.87 (s, 3H), 1.42 (d, J = 6.5 Hz, 6H).	DMSO	371 (M+1)	2.36	Method C	100	Method AP
1617		HCl	¹ H NMR (300 MHz, DMSO) δ 10.38 (s, 1H), 9.64 (s, 1H), 9.43 (s, 1H), 9.28 (s, 1H), 9.07 (d, J = 7.9 Hz, 1H), 9.00 - 8.76 (m, 3H), 8.48 (d, J = 7.5 Hz, 1H), 8.27 (d, J = 8.4 Hz, 1H), 8.08 - 7.95 (m, 1H), 7.94 - 7.81 (m, 1H), 3.97 - 3.74 (m, 2H), 1.38 (t, J = 7.2 Hz, 3H).	DMSO	328 (M+1)	1.82	Method C	100	Method AP

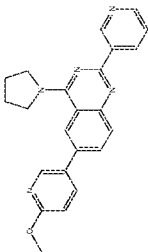
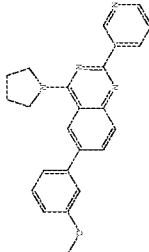
(continued)

Number	Product	Salt	¹ H-NMR	¹ H NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity Percent	Method for Coupling
1818		HCl	¹ H NMR (300 MHz, DMSO) δ 10.11 (s, 1H), 9.58 (s, 1H), 9.03 - 8.79 (m, 3H), 8.35 (d, J = 8.7 Hz, 1H), 8.23 - 8.09 (m, 1H), 7.90 - 7.75 (m, 1H), 7.58 - 7.36 (m, 1H), 7.04 (d, J = 3.3 Hz, 1H), 3.88 (s, 3H), 3.65 (t, J = 6.0 Hz, 2H), 2.29 - 2.09 (m, 1H), 1.02 (d, J = 6.7 Hz, 6H).	DMSO	385 (M+1)	1.76	Method D	100	Method AP
1619		HCl	¹ H NMR (300 MHz, DMSO) δ 9.83 - 9.52 (m, 2H), 9.27 (d, J = 7.1 Hz, 1H), 8.95 (d, J = 5.1 Hz, 1H), 8.87 (s, 1H), 8.32 (d, J = 8.9 Hz, 1H), 8.15 - 7.92 (m, 2H), 7.59 - 7.33 (m, 3H), 7.04 (d, J = 3.4 Hz, 1H), 4.86 - 4.57 (m, 2H), 3.88 (s, 3H).	DMSO	411 (M+1)	1.99	Method D	100	Method AP

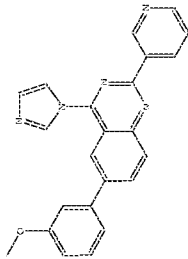
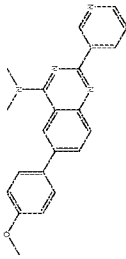
(continued)

Number	Product	Salt	¹ H-NMR	¹ H NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity Percent	Method for Coupling
1620		HCl	¹ H NMR (300 MHz, DMSO) δ 10.50 (s, 1H), 9.64 (d, J = 1.4 Hz, 1H), 9.12 - 8.85 (m, 3H), 8.46 - 8.33 (m, 1H), 8.27 (d, J = 8.9 Hz, 1H), 7.88 (dd, J = 7.9, 5.1 Hz, 1H), 7.57 - 7.37 (m, 3H), 7.11 - 6.94 (m, 1H), 4.00 - 3.76 (m, 5H), 1.37 (t, J = 7.2 Hz, 3H).	DMSO	MS not work	1.56	Method D	100	Method AP
1621			¹ H NMR (300 MHz, DMSO) δ 9.57 (s, 1H), 8.78 - 8.50 (m, 3H), 8.33-7.89 (m, 3H), 7.83 (d, J = 8.7 Hz, 1H), 7.53 (dd, J = 7.9, 4.8 Hz, 1 H), 7.47 - 7.32 (m, 3H), 7.04 - 6.91 (m, 1 H), 3.86 (s, 3H).	DMSO	329 (M+1)	1.47	Method D	100	Method AP
1622		HCl	¹ H NMR (300 MHz, DMSO) δ 9.65 (s, 1H), 9.14 - 8.87 (m, 2H), 8.50 (s, 1H), 8.33 (s, 2H), 7.94 - 7.77 (m, 1H), 7.52 - 7.25 (m, 3H), 7.02 (d, J = 7.9 Hz, 1H), 3.84 (s, 3H), 3.71 (s, 6H).	DMSO	357 (M+1)	1.55	Method D	100	Method AP

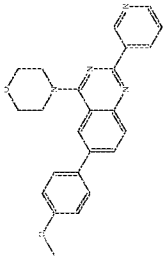
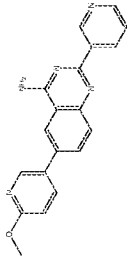
(continued)

Number	Product	Salt	¹ H-NMR	¹ H NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity Percent	Method for Coupling
1623			¹ H NMR (300 MHz, DMSO) δ 9.60 (d, J = 1.4 Hz, 1H), 8.81 - 8.55 (m, 3H), 8.42 (d, J = 1.8 Hz, 1H), 8.11 (ddd, J = 18.8, 8.7, 2.2 Hz, 2H), 7.86 (d, J = 8.7 Hz, 1H), 7.52 (dd, J = 7.9, 4.8 Hz, 1H), 6.95 (d, J = 8.6 Hz, 1H), 4.27 (s, 4H), 3.92 (s, 3H), 2.08 (s, 4H).	DMSO	384.1 (M+1)	2.33	Method C	100	Method AP
1624		2HCl	¹ H NMR (300 MHz, DMSO) δ 9.62 (s, 1H), 8.94 (d, J = 5.0 Hz, 2H), 8.56 (s, 1H), 8.32 (dd, J = 19.9, 8.5 Hz, 2H), 7.83 (s, 1H), 7.56 - 7.30 (m, 3H), 7.04 (d, J = 6.8 Hz, 1H), 4.27 (s, 4H), 3.86 (s, 3H), 2.08 (s, 4H).	DMSO	383.2 (M+1)	2.47	Method C	95	Method AP

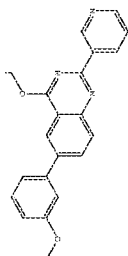
(continued)

Number	Product	Salt	¹ H-NMR	¹ H NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity Percent	Method for Coupling
1625			¹ H NMR (300 MHz, DMSO) δ 9.70 (d, J = 1.4 Hz, 1H), 8.85 (d, J = 7.9 Hz, 1H), 8.80 - 8.72 (m, 2H), 8.45 (d, J = 8.7 Hz, 1H), 8.27 (dd, J = 14.9, 5.9 Hz, 3H), 7.63 (dd, J = 7.3, 4.8 Hz, 1H), 7.52 - 7.33 (m, 4H), 7.04 (d, J = 7.1 Hz, 1H), 3.88 (s, 3H).	DMSO	380.1 (M+1)	2.01	Method C	100	Method AP
1626		HCl	¹ H NMR (300 MHz, DMSO) δ 9.61 (s, 1H), 8.75 (d, J = 7.7 Hz, 1H), 8.68 (d, J = 4.6 Hz, 1H), 8.30 (s, 1H), 8.08 (d, J = 8.6 Hz, 1H), 7.89 (d, J = 8.8 Hz, 1H), 7.75 (d, J = 8.7 Hz, 2H), 7.54 (dd, J = 7.8, 5.0 Hz, 1H), 7.07 (d, J = 8.7 Hz, 2H), 3.82 (s, 4H), 3.49 (s, 7H).	DMSO	358.0 (M+1)	2.25	Method C	100	Method AP

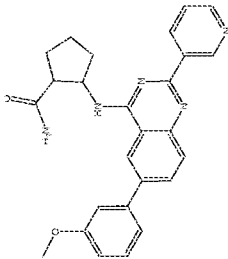
(continued)

Number	Product	Salt	¹ H-NMR	¹ H NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity Percent	Method for Coupling
1627		HCl	¹ H NMR (300 MHz, DMSO) δ 9.62 (s, 1H), 8.76 (d, J = 7.9 Hz, 1H), 8.73 - 8.66 (m, 1H), 8.19 - 8.09 (m, 2H), 7.97 (d, J = 8.6 Hz, 1H), 7.82 - 7.73 (m, 2H), 7.61 - 7.52 (m, 1H), 7.14-7.05 (m, 2H), 3.94 (s, 5H), 3.89 - 3.80 (m, 9H).	DMSO	399.2 (M+1)	2.19	Method C	100	Method AP
1628			¹ H NMR (300 MHz, DMSO) δ 9.55 (dd, J = 2.1, 0.8 Hz, 1H), 8.76 - 8.62 (m, 3H), 8.60 (d, J = 1.7 Hz, 1H), 8.24 - 8.10 (m, 2H), 8.02 (s, 2H), 7.83 (d, J = 8.7 Hz, 1H), 7.56 - 7.46 (m, 1H), 6.98 (d, J = 9.2 Hz, 1H), 3.91 (s, 3H).	DMSO	330.1 (M+1)	1.72	Method C	100	Method AP

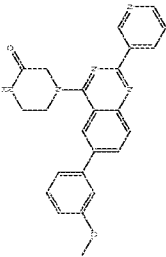
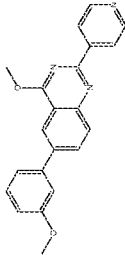
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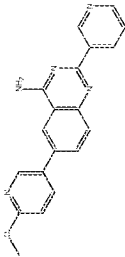
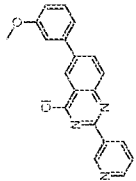
(continued)

Number	Product	Salt	¹ H-NMR	¹ H NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity Percent	Method for Coupling
1630		2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.68 (d, J = 1.8 Hz, 1H), 9.07 (d, J = 5.5 Hz, 1H), 8.96 (d, J = 3.5 Hz, 1H), 8.90 (s, 1H), 8.37 (d, J = 7.5 Hz, 1H), 8.22 (d, J = 8.3 Hz, 1H), 7.94 - 7.83 (m, 1H), 7.57 - 7.39 (m, 3H), 7.22 (d, J = 1.0 Hz, 1H), 7.08 (d, J = 7.3 Hz, 1H), 6.72 (s, 1H), 5.18 - 5.04 (m, 1H), 3.88 (s, J = 2.7 Hz, 3H), 3.23 (dd, J = 13.7, 6.1 Hz, 1H), 2.31 - 2.17 (m, 1H), 2.17 - 1.88 (m, 4H), 1.74 - 1.55 (m, 1H).	DMSO	440.6 (M+1)		Method C	99	Method AP

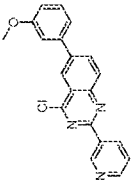
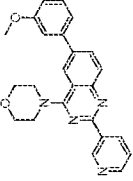
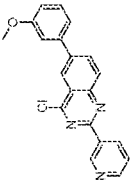
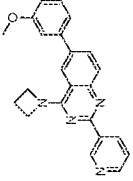
(continued)

Number	Product	Salt	¹ H-NMR	¹ H NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity Percent	Method for Coupling
1631		2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.70 (d, J = 1.6 Hz, 1H), 9.24 (d, J = 7.4 Hz, 1H), 8.99 (dd, J = 5.3, 1.4 Hz, 1H), 8.34 (dd, J = 15.7, 6.1 Hz, 3H), 8.21 (d, J = 8.4 Hz, 1H), 8.01 (dd, J = 7.7, 5.1 Hz, 1H), 7.51 - 7.33 (m, 3H), 7.04 (dd, J = 7.8, 2.4 Hz, 1H), 4.67 (s, 2H), 4.33 (s, 2H), 3.87 (s, J = 3.6 Hz, 3H), 3.50 (s, 2H).	DMSO	412.4 (M+1)		Method C	99	Method AP
1632		2 HCl	¹ H NMR (300 MHz, DMSO) δ 9.00 (d, J = 6.5 Hz, 2H), 8.78 (d, J = 6.5 Hz, 2H), 8.46 - 8.32 (m, 2H), 8.15 (d, J = 8.4 Hz, 1H), 7.53 - 7.30 (m, 3H), 7.02 (d, J = 7.8 Hz, 1H), 4.34 (s, J = 2.4 Hz, 3H), 3.86 (s, J = 2.5 Hz, 3H).	DMSO	344.4 (M+1)		Method C	99	Method AP

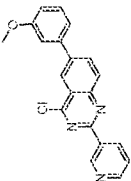
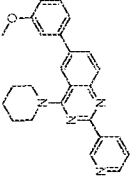
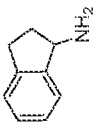
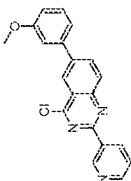
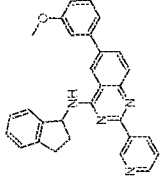
(continued)

Number	1633		Salt	¹ H-NMR ¹ H NMR (300 MHz, DMSO) δ 9.55 (dd, J = 2.1, 0.8 Hz, 1H), 8.76 - 8.62 (m, 3H), 8.60 (d, J = 1.7 Hz, 1H), 8.24-8.10 (m, 2H), 8.02 (s, 2H), 7.83 (d, J = 8.7 Hz, 1H), 7.56 - 7.46 (m, 1H), 6.98 (d, J = 9.2 Hz, 1H), 3.91 (s, 3H).	¹ H NMR Solvent	DMSO	LCMS	330.1 (M+1)	Retention Time	LCMS Protocol	Purity Percent	Method for Coupling
Number	1634		Starting Material R ³		Salt Type	¹ H NMR ¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.87 (s, 3H), 4.78 (d, 2H, J = 5.4 Hz), 6.58 (d, 1H, J = 1.1 Hz), 7.00-7.75 (m, 8H), 7.85 (d, 1H, J = 8.6 Hz), 8.64-8.97 (m, 4H), 9.64 (s, 1H)	¹ H NMR Solvent	DMSO	Purity percent	>98	Method of Coupling	Method AQ3, F5, G2 (reflux)

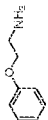
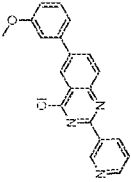
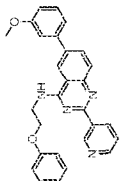
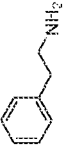
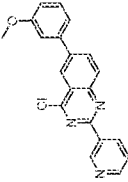
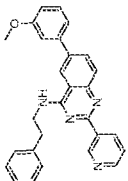
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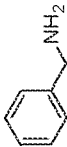
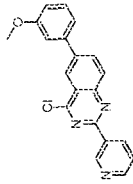
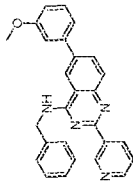
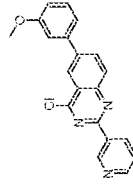
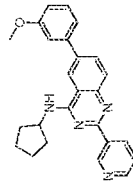
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1635					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.84–3.95 (m, 11H), 7.01 (d, 1H, <i>J</i> = 8.1 Hz), 7.32–7.46 (m, 3H), 7.56 (dd, 1H, <i>J</i> = 6.1, 8.1 Hz), 7.97 (d, 1H, <i>J</i> = 8.2 Hz), 8.16–8.76 (m, 4H), 9.61 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1636					¹ H NMR (DMSO- <i>d</i> ₆) ppm 2.23 (m, 2H), 3.85 (s, 3H), 4.71 (br s, 4H), 6.98 (dd, 1H, <i>J</i> = 2.1, 8.0 Hz), 7.31–7.55 (m, 4H), 7.87 (d, 1H, <i>J</i> = 8.2 Hz), 8.12–8.74 (m, 4H), 9.59 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

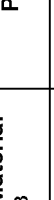
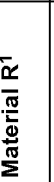
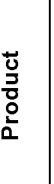
(continued)

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1637					¹ H NMR (DMSO-d ₆) ppm 1.77 (br s, 6H), 3.86–3.89 (br m, 7H), 6.99 (dd, 1H, J = 1.4, 8.6 Hz), 7.31–7.57 (m, 4H), 7.93 (d, 1H, J = 1.4, 8.6 Hz), 8.11–8.16 (m, 2H), 8.69–8.76 (m, 2H), 9.61 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1638					¹ H NMR (DMSO-d ₆) ppm 2.18–2.22 (m, 1H), 3.00–3.09 (m, 1H), 3.11–3.22 (m, 2H), 3.31–3.34 (m, 1H), 3.86 (s, 3H), 6.29–6.31 (m, 1H), 7.17–8.82 (m, 14H), 9.63 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

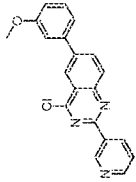
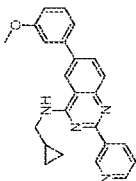
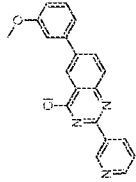
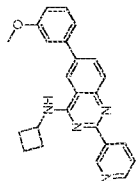
(continued)

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1639					¹ H NMR (DMSO-d ₆) ppm 3.87 (s, 3H), 4.10-4.13 (m, 2H), 4.35-4.38 (m, 2H), 6.92-7.46 (m, 10H), 7.85 (d, 1H, J = 8.6 Hz), 8.16 (d, 1H, J = 8.6 Hz), 8.66-8.75 (m, 4H), 9.61 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1640					¹ H NMR (DMSO-d ₆) ppm 3.07 (t, 2H, J = 6.9 Hz), 3.87 (s, 3H), 3.92-3.96 (m, 2H), 7.00-7.58 (m, 10H), 7.84 (d, 1H, J = 8.6 Hz), 8.14 (d, 1H, J = 8.6 Hz), 8.59-8.78 (m, 4H), 9.64 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

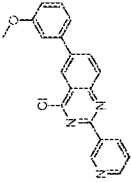
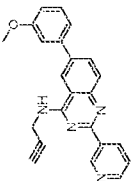
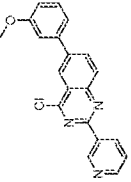
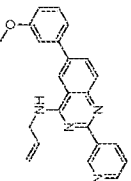
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1641					¹ H NMR (DMSO-d ₆) ppm 3.87 (s, 3H), 4.95 (d, 2H, J = 5.6 Hz), 7.01-7.51 (m, 10H), 7.85 (d, 1H, J = 8.6 Hz), 8.16 (d, 1H, J = 8.6 Hz), 8.65-8.73 (m, 4H), 9.56 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1642					¹ H NMR (DMSO-d ₆) ppm 1.66-1.781 (m, 6H), 2.16-2.19 (m, 2H), 3.87 (s, 3H), 4.78 (br s, 1H), 7.00-7.55 (m, 5H), 7.83 (d, 1H, J = 8.6 Hz), 8.12 (d, 1H, J = 8.6 Hz), 8.29 (s, 1H), 8.67-8.77 (m, 3H), 9.62 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

(continued)						
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent
1643				HCl	¹ H NMR (DMSO-d ₆) ppm 0.81 (t, 3H, J = 7.4 Hz), 1.47-1.54 (m, 2H), 2.00-2.05 (m, 2H), 3.32-3.37 (m, 4H), 3.86-3.91 (m, 5H), 7.03-7.80 (m, 5H), 8.16 (d, 1H, J = 8.7 Hz), 8.34 (d, 1H, J = 8.7 Hz), 8.86-8.96 (m, 3H), 9.61 (s, 1H), 10.03 (s, 1H)	DMSO
					>98	Method AQ3, F5, G2 (reflux)

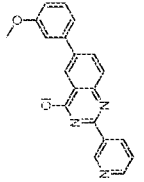
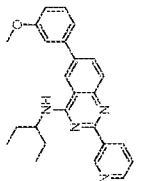
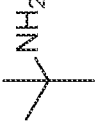
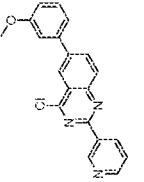
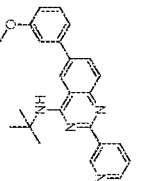
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Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1644					¹ H NMR (DMSO-d ₆) ppm 0.38-0.54 (m, 4H), 1.30-1.32 (m, 1H), 3.58-3.62 (m, 2H), 3.87 (s, 3H), 7.00-7.54 (m, 5H), 7.86 (d, 1H, J = 8.6 Hz), 8.14 (d, 1H, J = 8.6 Hz), 8.64-8.76 (m, 4H), 9.62 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1645					¹ H NMR (DMSO-d ₆) ppm 1.83-1.88 (m, 2H), 2.21-2.27 (m, 2H), 2.49 (br s, 2H), 3.87 (s, 3H), 4.88-4.91 (m, 1H), 7.03-7.57 (m, 5H), 7.83 (d, 1H, J = 8.6 Hz), 8.13 (d, 1H, J = 8.6 Hz), 8.62-8.77 (m, 4H), 9.63 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

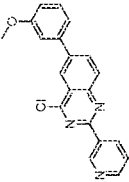
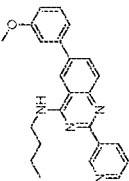
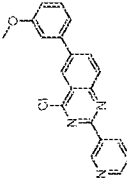
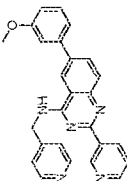
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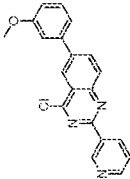
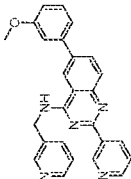
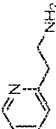
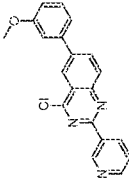
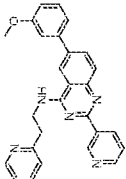
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1646					¹ H NMR (DMSO-d ₆) ppm 3.21 (s, 1H), 3.88 (s, 3H), 4.52 (dd, 2H, J=2.2, 5.2 Hz), 7.01-7.58 (m, 5H), 7.88 (d, 1H, J = 8.7 Hz), 8.19 (d, 1H, J = 8.6 Hz), 8.64-9.01 (m, 4H), 9.66 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1647					¹ H NMR (DMSO-d ₆) ppm 3.87 (s, 3H), 4.37 (br s, 2H), 5.19 (dd, 1H, J = 1.5, 10.2 Hz), 5.31 (dd, 1H, J = 1.5, 17.2 Hz), 6.01-6.22 (m, 1H), 7.00-7.56 (m, 5H), 7.85 (d, 1H, J = 8.6 Hz), 8.16 (d, 1H, J = 8.6 Hz), 8.67-8.82 (m, 4H), 9.61 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

(continued)

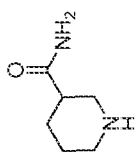
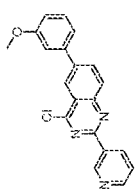
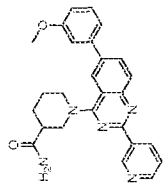
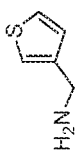
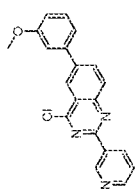
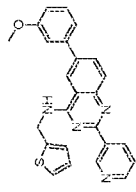
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1648					¹ H NMR (DMSO-d ₆) ppm 0.94–0.99 (t, 6H, J = 7.4 Hz), 1.68–1.80 (m, 4H), 3.87 (s, 3H), 4.49–4.52 (m, 1H), 7.01–7.47 (m, 5H), 7.83 (d, 1H, J = 8.6 Hz), 8.11–8.15 (m, 2H), 8.67–8.75 (m, 3H), 9.63 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1649					¹ H NMR (DMSO-d ₆) ppm 1.68 (s, 9H), 3.87 (s, 3H), 7.01–7.70 (m, 6H), 7.82 (d, 1H, J = 8.6 Hz), 8.11 (d, 1H, J = 8.6 Hz), 8.67–8.74 (m, 3H), 9.60 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

(continued)

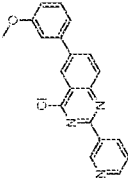
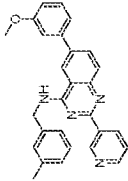
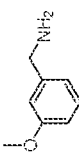
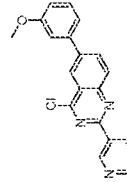
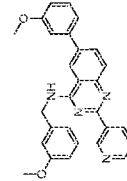
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1650					¹ H NMR (DMSO-d ₆) ppm 0.97 (t, 3H, J = 7.2 Hz), 1.44-1.48 (m, 2H), 1.74-1.78 (m, 2H), 3.71-3.75 (m, 2H), 3.89 (s, 3H), 7.00-7.56 (m, 5H), 7.83 (d, 1H, J = 8.6 Hz), 8.15 (d, 1H, J = 8.6 Hz), 8.59-8.76 (m, 4H), 9.62 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1651					¹ H NMR (DMSO-d ₆) ppm 3.87 (s, 3H), 4.98 (d, 2H, J = 5.6 Hz), 7.02-7.46 (m, 7H), 7.87 (d, 1H, J = 8.7 Hz), 8.51 (dd, 1H, J = 1.5, 4.5 Hz), 8.50-8.71 (m, 5H), 9.26 (s, 1H), 9.48 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1652					¹ H NMR (DMSO-d ₆) ppm 3.87 (s, 3H), 4.98 (d, 2H, J = 5.5 Hz), 7.00-7.46 (m, 6H), 7.86-7.90 (m, 2H), 8.16 (d, 1H, J = 1.7 Hz), 8.46 (d, 1H, J = 1.7 Hz), 8.66-8.76 (m, 4H), 9.22 (s, 1H), 9.56 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1653					¹ H NMR (DMSO-d ₆) ppm 3.22-3.27 (m, 2H), 3.87 (s, 3H), 4.04-4.11 (m, 2H), 7.02-7.46 (m, 8H), 7.83 (d, 1H, J = 8.6 Hz), 8.13 (d, 1H, J = 1.6 Hz), 8.67-8.78 (m, 5H), 9.64 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

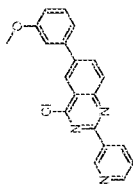
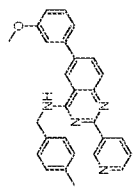
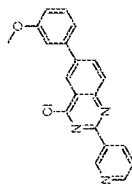
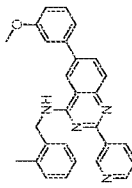
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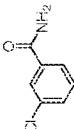
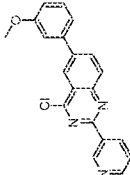
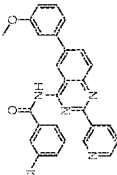
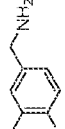
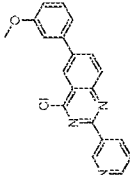
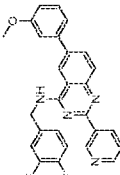
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1654					¹ H NMR (DMSO-d ₆) ppm 1.76–1.88 (m, 4H), 2.66 (br s, 1H), 3.37–3.48 (m, 2H), 3.86 (s, 3H), 4.39–4.47 (m, 2H), 6.95–7.57 (m, 7H), 7.95 (d, 1H, J = 8.7 Hz), 8.15–8.19 (m, 2H), 8.69–8.76 (m, 2H), 9.61 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1655					¹ H NMR (DMSO-d ₆) ppm 3.87 (s, 3H), 4.97 (d, 2H, J = 5.6 Hz), 7.00–7.50 (m, 8H), 7.85 (d, 1H, J = 8.7 Hz), 8.16 (s, 1H), 8.66–8.77 (m, 3H), 9.11 (s, 1H), 9.61 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

(continued)

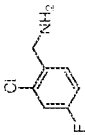
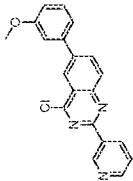
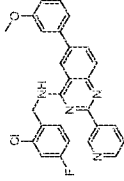
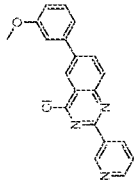
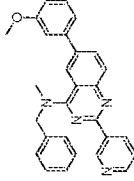
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1656					¹ H NMR (DMSO- <i>d</i> ₆) ppm 2.28 (s, 3H), 3.87 (s, 3H), 4.91 (d, 2H, <i>J</i> = 5.5 Hz), 7.00-7.52 (m, 9H), 7.86 (d, 1H, <i>J</i> = 8.6 Hz), 8.17 (dd, 1H, <i>J</i> = 8.6, 1.8 Hz), 8.66-9.16 (m, 3H), 9.18-9.16 (m, 1H), 9.57 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1657					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.71 (s, 3H), 3.87 (s, 3H), 4.91 (d, 2H, <i>J</i> = 5.7 Hz), 6.82-7.52 (m, 9H), 7.86 (d, 1H, <i>J</i> = 8.7 Hz), 8.17 (dd, 1H, <i>J</i> = 8.7, 1.8 Hz), 8.66-9.15 (m, 3H), 9.17-9.19 (m, 1H), 9.57 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

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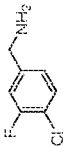
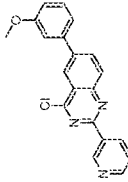
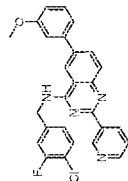
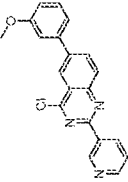
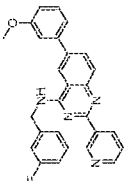
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1658					¹ H NMR (DMSO- <i>d</i> ₆) ppm 2.25 (s, 3H), 3.87 (s, 3H), 4.91 (d, 2H, J = 5.7 Hz), 7.00-7.53 (m, 9H), 7.86 (d, 1H, J = 8.6 Hz), 8.17 (dd, 1H, J = 8.6, 1.8 Hz), 8.66-9.15 (m, 3H), 9.17-9.19 (m, 1H), 9.57 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1659					¹ H NMR (DMSO- <i>d</i> ₆) ppm 2.44 (s, 3H), 3.86 (s, 3H), 4.95 (d, 2H, J = 5.7 Hz), 7.01-7.53 (m, 9H), 7.86 (d, 1H, J = 8.6 Hz), 8.17 (dd, 1H, J = 8.6, 1.8 Hz), 8.65-9.02 (m, 3H), 9.50-9.52 (m, 1H), 9.54 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1660					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.87 (s, 3H), 7.03 (s, 1H), 7.44-7.94 (m, 10H), 8.07-8.70 (m, 4H), 9.41 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1661					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.87 (s, 3H), 4.91 (d, 2H, J = 5.5 Hz), 7.00-7.04 (m, 1H), 7.04-7.55 (m, 7H), 7.87 (d, 1H, J = 8.6 Hz), 8.18 (dd, 1H, J = 8.8, 1.8 Hz), 8.66-9.18 (m, 3H), 9.18-9.21 (m, 1H), 9.55 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

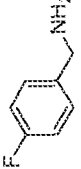
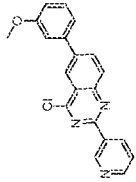
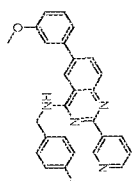
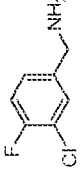
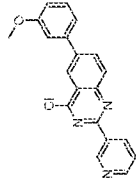
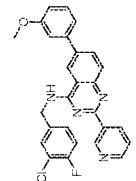
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Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1662					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.87 (s, 3H), 4.99 (d, 2H, J = 5.5 Hz), 7.01-7.05 (m, 2H), 7.15-7.56 (m, 6H), 7.88 (d, 1H, J = 8.6 Hz), 8.20 (dd, 1H, J = 8.6, 1.8 Hz), 8.64-9.18 (m, 3H), 9.19-9.21 (m, 1H), 9.50 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1663					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.49 (s, 3H), 3.74 (s, 3H), 5.15 (s, 2H), 6.90 (d, J = 8.2 Hz, 1H), 7.04-7.29 (m, 2H), 7.33-7.54 (m, 7H), 7.93 (d, 1H, J = 8.6 Hz), 8.15 (dd, 1H, J = 8.6, 1.8 Hz), 8.21 (s, 1H), 8.68-8.75 (m, 2H), 9.59 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

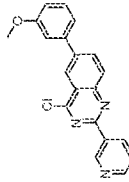
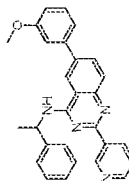
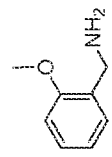
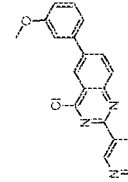
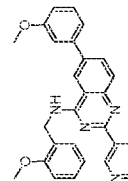
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Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1664					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.87 (s, 3H), 4.94 (d, 2H, J = 5.5 Hz), 7.01-7.03 (m, 1H), 7.35-7.57 (m, 7H), 7.88 (d, 1H, J = 8.6 Hz), 8.20 (dd, 1H, J = 8.6, 1.8 Hz), 8.66-9.22 (m, 3H), 9.19-9.21 (m, 1H), 9.50 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1665					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.87 (s, 3H), 4.96 (d, 2H, J = 5.5 Hz), 7.00-7.03 (m, 2H), 7.33-7.52 (m, 7H), 7.88 (d, 1H, J = 8.6 Hz), 8.19 (dd, 1H, J = 8.6, 1.8 Hz), 8.65-9.20 (m, 3H), 9.19-9.21 (m, 1H), 9.54 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

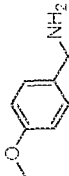
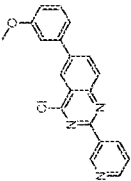
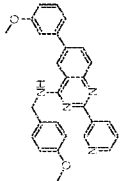
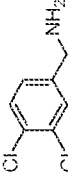
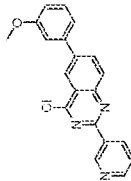
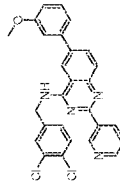
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Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1666					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.87 (s, 3H), 4.93 (d, 2H, J = 5.5 Hz), 7.00-7.19 (m, 3H), 7.41-7.55 (m, 6H), 7.88 (d, 1H, J = 8.6 Hz), 8.17 (dd, 1H, J = 8.6, 1.8 Hz), 8.65-9.18 (m, 3H), 9.19-9.21 (m, 1H), 9.57 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1667					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.87 (s, 3H), 4.92 (d, 2H, J = 5.5 Hz), 7.01-7.03 (d, 1H, J = 7.0 Hz), 7.37-7.53 (m, 6H), 7.72 (d, 1H, J = 8.6 Hz), 8.18 (dd, 1H, J = 8.6, 1.8 Hz), 8.66-9.19 (m, 3H), 9.20-9.23 (m, 1H), 9.56 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

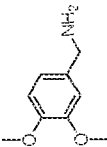
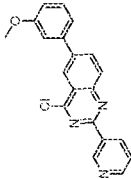
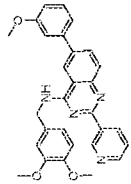
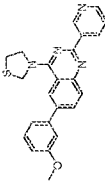
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Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1668					¹ H NMR (DMSO- <i>d</i> ₆) ppm 1.68 (d, 3H, <i>J</i> = 7.0 Hz), 3.87 (s, 3H), 4.92 (d, 2H, <i>J</i> = 5.5 Hz), 5.75-5.78 (m, 1H), 7.02 (d, 1H, <i>J</i> = 7.0 Hz), 7.32-7.56 (m, 9H), 7.84 (d, 1H, <i>J</i> = 8.6 Hz), 8.15 (dd, 1H, <i>J</i> = 8.6, 1.8 Hz), 8.64-8.84 (m, 4H), 9.51 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1669					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.70 (s, 3H), 3.86 (s, 3H), 4.92 (d, <i>J</i> = 5.1 Hz, 2H), 6.89-7.01 (m, 3H), 7.41-7.54 (m, 6H), 7.85 (d, <i>J</i> = 8.6 Hz, 1H), 7.85 (d, <i>J</i> = 8.6 Hz, 1H), 8.67-8.75 (m, 3H), 9.19 (s, 1H), 9.56 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

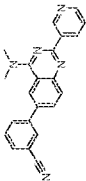
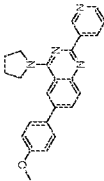
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Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1670					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.87 (s, 3H), 3.91 (s, 3H), 4.92 (d, J = 5.0 Hz, 2H), 6.87-7.52 (m, 9H), 7.85 (d, J = 8.5 Hz, 1H), 8.17 (d, J = 8.5 Hz, 1H), 8.64-8.73 (m, 3H), 9.05 (s, 1H), 9.52 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1671					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.87 (s, 3H), 4.93 (d, J = 5.6 Hz, 2H), 7.02 (s, 1H), 7.41-7.61 (m, 7H), 7.83 (d, J = 8.9 Hz, 1H), 8.18 (d, J = 8.9 Hz, 1H), 8.66-8.71 (m, 3H), 9.21 (s, 1H), 9.54 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)

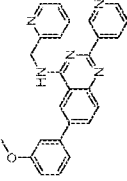
(continued)

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1672					¹ H NMR (DMSO- <i>d</i> ₆) ppm 3.69 (s, 6H), 3.87 (s, 3H), 4.93 (d, <i>J</i> = 5.6 Hz, 2H), 6.90 (s, 1H), 7.00-7.54 (m, 7H), 7.85 (d, <i>J</i> = 8.6 Hz, 1H), 8.17 (d, <i>J</i> = 8.6 Hz, 1H), 8.66-8.78 (m, 3H), 9.14 (s, 1H), 9.63 (s, 1H)	DMSO	>98	Method AQ3, F5, G2 (reflux)
1673						DMSO	95	Method AQ3, AP

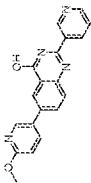
(continued)

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1674				2HCl	¹ H-NMR (300 MHz, DMSO): δ 9.71 (d, J = 1.6 Hz, 1H), 9.16 (d, J = 8.2 Hz, 1H), 8.98 (dd, J = 5.1, 1.5 Hz, 1H), 8.58 (s, 1H), 8.48–8.34 (m, 3H), 8.20 (d, J = 8.0 Hz, 1H), 7.98–7.87 (m, 2H), 7.75 (t, J = 7.8 Hz, 1H), 3.73 (s, 6H).	DMSO	95	Method AQ3, AP
1675					¹ H-NMR (300 MHz, DMSO): δ 9.59 (s, 1H), 8.78–8.62 (m, 2H), 8.38 (s, 1H), 8.05 (dd, J = 8.6, 1.8 Hz, 1H), 7.84 (d, J = 8.7 Hz, 1H), 7.73 (d, J = 8.7 Hz, 2H), 7.52 (dd, J = 7.8, 4.7 Hz, 1H), 7.07 (d, J = 8.8 Hz, 2H), 4.04 (s, 4H), 3.82 (s, 3H), 2.02 (s, 4H).	DMSO	95	Method AQ3, AP

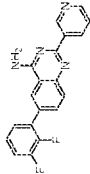
(continued)

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1676				3HCl	¹ H-NMR (300 MHz, DMSO): δ 10.40 (s, 1H), 9.48 (d, J = 2.0 Hz, 1H), 8.95 (s, 2H), 8.88 (d, J = 5.2 Hz, 1H), 8.72 (d, J = 4.3 Hz, 1H), 8.35 (d, J = 8.9 Hz, 1H), 8.18 (s, 1H), 8.08 (d, J = 8.8 Hz, 1H), 7.94 - 7.81 (m, 2H), 7.64 (s, 1H), 7.54 - 7.45 (m, 3H), 7.06 - 7.00 (m, 1H), 5.25 (d, J = 5.1 Hz, 2H), 3.88 (s, 3H).	DMSO	95	Method AQ3, AP

(continued)

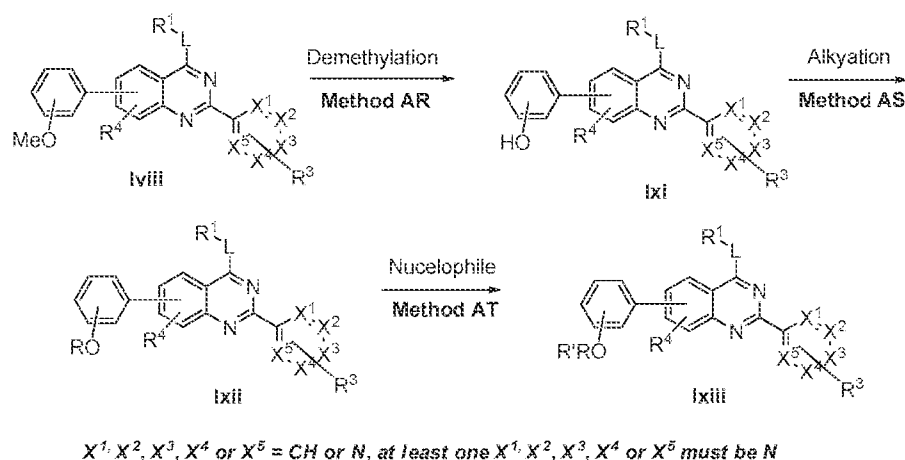
Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1677					¹ H-NMR (300 MHz, DMSO): δ 12.83 (s, 1H), 9.32 (d, J = 2.2 Hz, 1H), 8.77 (dd, J = 4.8, 1.5 Hz, 1H), 8.63 (d, J = 2.5 Hz, 1H), 8.56-8.49 (m, 1H), 8.37 (d, J = 2.2 Hz, 1H), 8.23-8.14 (m, 2H), 7.85 (d, J = 8.5 Hz, 1H), 7.60 (dd, J = 8.0, 4.8 Hz, 1H), 6.96 (d, J = 8.7 Hz, 1H), 3.92 (s, 3H).	DMSO	95	Method AQ3, AP

(continued)

Number	Starting Material R ¹	Starting Material R ³	Product	Salt Type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling
1678	NH3			2HCl	¹ H NMR (300 MHz, DMSO) δ 9.79 (brs, 1H), 9.60 (s, 2H), 9.07 (d, <i>J</i> = 7.4 Hz, 1H), 9.00 (d, <i>J</i> = 4.5 Hz, 1H), 8.80 (s, 1H), 8.35 (d, <i>J</i> = 8.7 Hz, 1H), 8.26 (d, <i>J</i> = 8.6 Hz, 1H), 8.02 - 7.88 (m, 1H), 7.67 - 7.48 (m, 2H), 7.49 - 7.37 (m, 1H).	DMSO	>98	Method AQ3, F5, G2 (reflux)

Scheme 78: General route for the synthesis of compounds with general formula Ixiii

[0504]



Method AT for alkylation:

[0505]

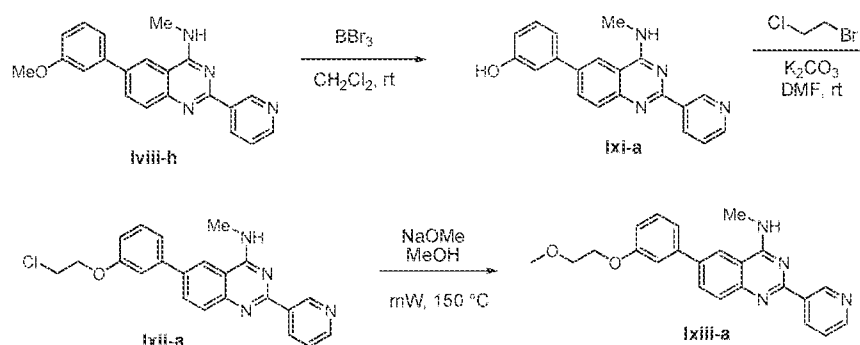
Method AT1: NaOMe/MeOH/microwave/150°C

Method AT2: MeOH/microwave/150°C

Method AT3: DEPEA/KI/DMF/microwave/150°C

Scheme 79: Representative synthesis of compounds of formula Ixiii (see Scheme 76)

[0506]



Method AR: 3-(4-(methylamino)-2-(pyridin-3-yl)quinazolin-6-yl)phenol (Ixi-a)

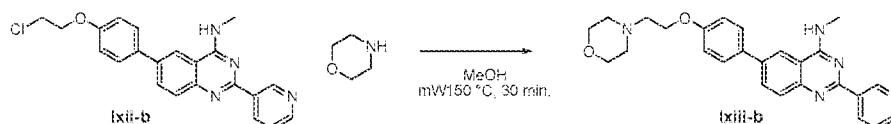
[0507] To a suspension of 6-(3-methoxyphenyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine (prepared in a similar method described for 6-(3-methoxyphenyl)-2-(pyridine-3-yl)-4-(pyrrolidin-1-yl)quinazoline using Scheme 74, substituting N-methylamine for pyrrolidine) (1.00 g, 2.9 mmol) in CH_2Cl_2 (15 mL) and boron tribromide 1M solution in dichloromethane (8.76 mL, 8.76 mmol) was added at 0 °C. The reaction mixture was stirred overnight at room temperature after which it was carefully poured into a vigorously stirring mixture of ice and saturated aqueous solution of $NaHCO_3$. The resultant solid was collected by filtration, dried and then dissolved in a mixture of K_2CO_3 (2 g) and methanol (50 mL). The solution was then acidified using aqueous NH_4Cl solution. (50 mL) and the precipitate which formed was collected by filtration and dried to give 0.95 g of 3-(4-(methylamino)-2-(pyridin-3-yl)quinazolin-6-yl)phenol as pale yellow solid (99 %). LCMS m/z = 329 (M + 1) (Method D) (retention time = 1.30 min). 1H NMR (300 MHz, DMSO) δ 9.63 (s, 1H), 9.60 (s, 1H), 8.77 (d, J = 8.0 Hz, 1H), 8.74 - 8.59 (m, 2H), 8.54 (s, 1H), 8.05 (d, J = 8.7 Hz, 1H), 7.83 (d, J = 8.7 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 7.38 - 7.12 (m, 3H), 6.82 (d, J = 7.8 Hz, 1H), 3.17 (d, J = 4.3 Hz, 3H).

[0508] Method AS: 6-(3-(2-chloroethoxy)phenyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine (Ixii-a) A suspension of 3-(4-(methylamino)-2-(pyridin-3-yl)quinazolin-6-yl)phenol (0.30 g, 0.914 mmol), 1-bromo-2-chloroethane (0.38 ml, 4.57 mmol), and potassium carbonate (0.38 g, 2.74 mmol) in DMF (10 mL) was stirred for 2 days at room temperature. Water (10 mL) and ethyl acetate (10 mL) were added to the mixture and extracted. The organic layer was separated and concentrated *in vacuo* to leave a solid, which was collected by filtration and washed with hexane and dried to give 0.30 g of 6-(3-(2-chloroethoxy)phenyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine as brown solid (83 %). The product was used without further purification.

[0509] Method AT1: 6-(3-(2-methoxyethoxy)phenyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine (Ixiii-a) A solution of 6-(3-(2-chloroethoxy)phenyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine (70 mg, 0.18 mmol) and sodium methoxide (97 mg, 1.8 mmol) in methanol (3 mL) was placed in a microwave reaction vial. The mixture was heated under microwave irradiation conditions at 150 °C for 30 minutes after which the solvent was removed *in vacuo*. The crude product was obtained, which was purified by column chromatography on basic silica gel (eluted with hexane/ethyl acetate 3:1→1:4) to give 20 mg of 6-(3-(2-methoxyethoxy)phenyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine as an off-white powder (29 %). LCMS *m/z* = 387 (*M* + 1) (Method D) (retention time = 1.49 min). ¹H NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.77 (d, *J* = 8.2 Hz, 1H), 8.73 - 8.51 (m, 3H), 8.16 (d, *J* = 8.7 Hz, 1H), 7.84 (d, *J* = 8.3 Hz, 1H), 7.60 - 7.32 (m, 4H), 7.01 (s, 1H), 4.21 (s, 2H), 3.70 (s, 2H), 3.33 (s, 3H), 3.19 (s, 3H).

Method AT2: N-Methyl-6-(4-(2-morpholinoethoxy)phenyl)-2-(pyridin-3-yl)quinazolin-4-amine (Ixiii-b)

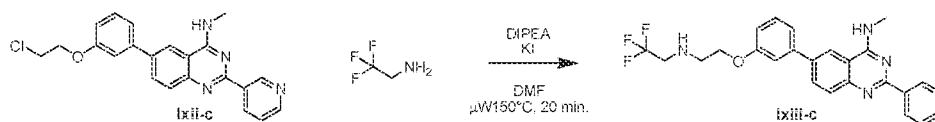
[0510]



[0511] In a 10 mL microwave vial 6-(4-(2-chloroethoxy)phenyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine (70.0 mg, 0.179 mmol) and morpholine (0.155 ml, 1.791 mmol) were added in methanol (3 mL) to give a yellow suspension. The vial was irradiated at 150 °C in the microwave for 30 min. The volatiles were evaporated *in vacuo*. Water (10 mL) was added to the reaction mixture and extracted with ethyl acetate (2 x 10 mL). The organic layers were combined and washed with brine (1 x 20 mL), then dried over MgSO₄, filtered and concentrated. The residue was purified by column chromatography on basic silica gel (eluted with hexane/ethyl acetate 3:2 to 0:1). The product was obtained as the parent and converted to the HCl salt by addition of 4 M HCl-dioxane, then crystallized from EtOH-H₂O to give 55 mg of N-methyl-6-(4-(2-morpholinoethoxy)phenyl)-2-(pyridin-3-yl)quinazolin-4-amine as a yellow powder (60 % yield). LCMS *m/z* = 442 (*M* + 1) (Method D) (retention time = 1.12 min). ¹H NMR (300 MHz, DMSO) δ 11.31 (s, 1H), 10.41 (s, 1H), 9.63 (s, 1H), 9.10 - 8.79 (m, 3H), 8.34 (d, *J* = 8.2 Hz, 1H), 8.28 - 8.12 (m, 1H), 7.92 (d, *J* = 8.0 Hz, 2H), 7.87 - 7.74 (m, 1H), 7.18 (d, *J* = 8.0 Hz, 2H), 4.60 - 4.46 (m, 2H), 4.06 - 3.91 (m, 2H), 3.91 - 3.74 (m, 2H), 3.67 - 3.41 (m, 4H), 3.38 - 3.09 (m, *J* = 10.1 Hz, 5H).

Method AT3: N-methyl-2-(pyridin-3-yl)-6-(3-(2-(2,2,2-trifluoroethylamino)ethoxy)phenyl)quinazolin-4-amine (Ixiii-c)

[0512]

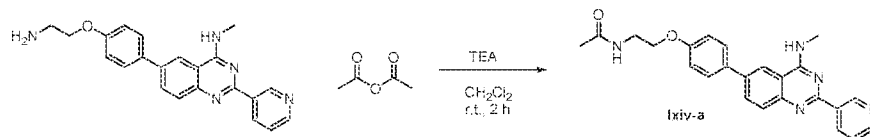


[0513] In a 10 mL microwave vial was added 6-(3-(2-chloroethoxy)phenyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine (50.0 mg, 0.128 mmol), 2,2,2-trifluoroethylamine (0.100 ml, 1.279 mmol), potassium iodide (42.5 mg, 0.256 mmol), and *N,N'*-diisopropylethylamine (0.045 ml, 0.256 mmol) in DMF (3 mL) to give a yellow suspension. The vial was irradiated at 150 °C in the microwave for 20 min. Water (10 mL) was added to the mixture and extracted with ethyl acetate (2 x 20 mL). The organic layers were combined and washed with water (1 x 20 mL) and brine (1 x 20 mL), dried over MgSO₄, filtered and concentrated. The residue was purified by column chromatography on silica gel (eluted with CH₂Cl₂/CH₂Cl₂-MeOH-NH₄OH=100:20:1 1:0 to 0:1). The product was converted to the HCl salt by addition of 4 M HCl-dioxane, then crystallization from IPA-H₂O to give 30 mg of N-methyl-2-(pyridin-3-yl)-6-(3-(2-(2,2,2-trifluoroethylamino)ethoxy)phenyl)quinazolin-4-amine as a yellow powder (60 % yield).

no)ethoxy)phenyl)quinazolin-4-amine as a yellow powder (42 %). LCMS m/z = 454 ($M + 1$) (Method C) (retention time = 2.26 min). ^1H NMR (300 MHz, DMSO) δ 10.73 - 10.36 (m, 1H), 9.65 (s, 1H), 9.13 - 8.98 (m, 2H), 8.93 (d, J = 5.0 Hz, 1H), 8.38 (d, J = 8.4 Hz, 1H), 8.24 (d, J = 7.7 Hz, 1H), 7.94 - 7.75 (m, 1H), 7.69 - 7.41 (m, 3H), 7.09 (d, J = 8.1 Hz, 1H), 4.61 - 4.43 (m, 2H), 4.28 - 4.07 (m, 2H), 3.59 - 3.39 (m, 2H), 3.30 (d, J = 2.8 Hz, 3H).

Scheme 80: Method AU: N-(2-(4-(4-(methylamino)-2-(pyridin-3-yl)quinazolin-6-yl)phenoxy)ethyl)acetamide (Ixiv-a)

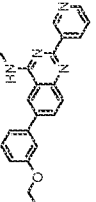
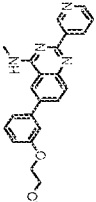
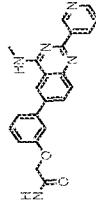
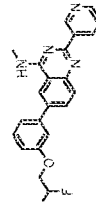
[0514]



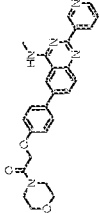
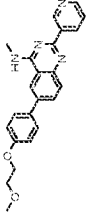
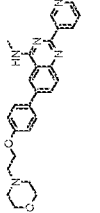
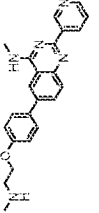
[0515] In a 50 mL round-bottomed flask was added 6-(4-(2-aminoethoxy)phenyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine (15.0 mg, 0.040 mmol) and triethylamine (0.017 ml, 0.121 mmol) in CH_2Cl_2 (5 mL) to give a yellow solution. Acetic anhydride (4.58 μl , 0.048 mmol) was added and the mixture was stirred for 2 h at room temperature. Water (10 mL) was added to the mixture and extracted with ethyl acetate (2 x 10 mL). The organic layers were combined and washed with brine (1 x 20 mL), dried over Mg_2SO_4 , filtered and concentrated. The residue was purified by column chromatography on silica gel (eluted with $\text{CH}_2\text{Cl}_2/\text{MeOH}$ 1:0 to 9:1). The product was converted to the HCl salt by addition of 4 M HCl-dioxane, then dissolved in a small amount of methanol, followed by ethyl acetate. The resulting solid was filtered and dried to give 10 mg of N-(2-(4-(4-(methylamino)-2-(pyridin-3-yl)quinazolin-6-yl)phenoxy)ethyl)acetamide as a HCl salt as a yellow solid in a 51% yield. LCMS m/z = 414 ($M + 1$) (Method C) (retention time = 1.28 min). ^1H NMR (300 MHz, DMSO) δ 10.26 (s, 1H), 9.62 (s, 1H), 9.06 - 8.90 (m, 2H), 8.81 (s, 1H), 8.34 (d, J = 9.0 Hz, 1H), 8.17 (d, J = 6.6 Hz, 2H), 7.86 (d, J = 8.0 Hz, 3H), 7.13 (d, J = 7.9 Hz, 2H), 4.06 (t, J = 5.4 Hz, 2H), 3.52 - 3.38 (m, 2H), 3.30 (d, J = 4.1 Hz, 3H), 1.84 (s, 3H).

[0516] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 78**, replacing 1-bromo-2-chloroethane with the corresponding alkyl halide.

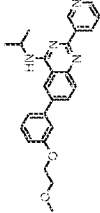
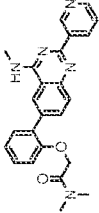
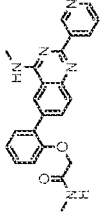
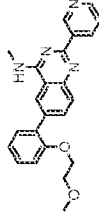
Table 26:

Number	PRODUCT	Salt	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1679		HCl	¹ H NMR (300 MHz, DMSO) δ 10.36 (s, 1H), 9.65 (s, 1H), 9.13 - 8.99 (m, 1H), 8.99 - 8.83 (m, 2H), 8.38 (d, J = 8.8 Hz, 1H), 8.30 - 8.16 (m, 1H), 7.94 - 7.80 (m, 1H), 7.55 - 7.35 (m, 3H), 7.09 - 6.97 (m, 1H), 4.16 (q, J = 7.0 Hz, 2H), 3.31 (d, J = 4.3 Hz, 3H), 1.37 (t, J = 6.9 Hz, 3H).	DMSO	357 (M+1)	1.62	Method D	100	Method AR/AS/AT1
1680			¹ H NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.77 (d, J = 8.2 Hz, 1H), 8.73 - 8.51 (m, 3H), 8.16 (d, J = 8.7 Hz, 1H), 7.84 (d, J = 8.3 Hz, 1H), 7.60 - 7.32 (m, 4H), 7.01 (s, 1H), 4.21 (s, 2H), 3.70 (s, 2H), 3.33 (s, 3H), 3.19 (s, 3H).	DMSO	387 (M+1)	1.49	Method D	100	Method AR/AS/AT1
1681			¹ H NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.78 (d, J = 7.9 Hz, 1H), 8.72 - 8.49 (m, 3H), 8.22 - 8.01 (m, 2H), 7.85 (d, J = 8.6 Hz, 1H), 7.62 - 7.36 (m, 4H), 7.01 (s, 1H), 4.59 (s, 2H), 3.19 (d, J = 4.0 Hz, 3H), 2.68 (d, J = 4.5 Hz, 3H).	DMSO	400 (M+1)	1.31	Method D	100	Method AR/W with KI
1682		2HCl	¹ H NMR (300 MHz, DMSO) δ 10.47 (s, 1H), 9.65 (d, J = 1.7 Hz, 1H), 9.05 (d, J = 7.9 Hz, 1H), 9.01 - 8.88 (m, 2H), 8.39 (m, 1H), 8.23 (d, J = 8.8 Hz, 1H), 7.87 (dd, J = 8.0, 5.3 Hz, 1H), 7.65 - 7.39 (m, 3H), 7.09 (m, 1H), 6.44 (m, 1H), 4.47 (d, J = 14.7, 3.4 Hz, 2H), 3.29 (d, J = 4.5 Hz, 3H).	DMSO	393.4 (M+1)		Method C (NH ₄ HCO ₃)	100	Method AR/AS (K ₂ CO ₃ , DMF-THF (1 : 1), 60 °C) then Method AT

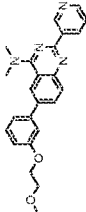
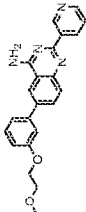
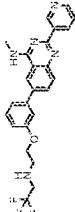
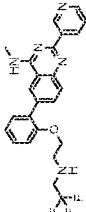
(continued)

Number	PRODUCT	Salt	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1683		NCl	¹ H NMR (300 MHz, DMSO) δ 10.12 (s, 1H), 9.60 (s, 1H), 9.03 - 8.87 (m, 2H), 8.77 (s, 1H), 8.32 (d, J = 8.5 Hz, 1H), 8.21 - 8.02 (m, 1H), 7.92 - 7.69 (m, 3H), 7.20 - 7.02 (m, 2H), 4.93 (s, 2H), 3.70 - 3.40 (m, 8H), 3.29 (d, J = 3.5 Hz, 3H).	DMSO	456 (M+1)	1.73	Method B (Ammonium formate)	100	Method AR/W with KI
1684		HCl	¹ H NMR (300 MHz, DMSO) δ 10.38 (s, 1H), 9.64 (s, 1H), 9.02 (d, J = 6.7 Hz, 1H), 8.93 (d, J = 4.9 Hz, 1H), 8.84 (s, 1H), 8.34 (d, J = 9.0 Hz, 1H), 8.22 (d, J = 8.7 Hz, 1H), 7.98 - 7.70 (m, 3H), 7.11 (d, J = 7.3 Hz, 2H), 4.26 - 4.05 (m, 2H), 3.78 - 3.57 (m, 2H), 3.45 - 3.19 (m, 6H).	DMSO	387 (M+1)	1.47	Method A (Formic acid)	100	Method AR/AS/AT1
1685		HCl	¹ H NMR (300 MHz, DMSO) δ 11.31 (s, 1H), 10.41 (s, 1H), 9.63 (s, 1H), 9.10 - 8.79 (m, 3H), 8.34 (d, J = 8.2 Hz, 1H), 8.28 - 8.12 (m, 1H), 7.92 (d, J = 8.0 Hz, 2H), 7.87 - 7.74 (m, 1H), 7.18 (d, J = 8.0 Hz, 2H), 4.60 - 4.46 (m, 2H), 4.06 - 3.91 (m, 2H), 3.91 - 3.74 (m, 2H), 3.67 - 3.41 (m, 4H), 3.38 - 3.09 (m, J = 10.1 Hz, 5H).	DMSO	442 (M+1)	1.22	Method A (Formic acid)	100	Method AR/AS/AT2
1686		HCl	¹ H NMR (300 MHz, DMSO) δ 10.32 - 9.70 (m, 1H), 9.61 (s, 1H), 9.13 - 8.70 (m, 5H), 8.29 (d, J = 8.5 Hz, 1H), 8.20 - 8.01 (m, 1H), 7.89 (d, J = 8.6 Hz, 2H), 7.83 - 7.67 (m, 1H), 7.16 (d, J = 8.8 Hz, 2H), 4.38 - 4.27 (m, 2H), 3.43 - 3.32 (m, 2H), 3.27 (d, J = 3.7 Hz, 3H), 2.63 (t, J = 4.8 Hz, 3H).	DMSO	386 (M+1)	1.46	Method B (Ammonium formate)	100	Method AR/AS/AT2

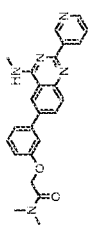
(continued)

Number	PRODUCT	Salt	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1687		HCl	¹ H NMR (300 MHz, DMSO) δ 9.95 (s, 1H), 9.62 (s, 1H), 9.13 - 8.86 (m, J = 5.2 Hz, 3H), 8.39 (d, J = 8.7 Hz, 1H), 8.26 (s, 1H), 7.97 - 7.80 (m, 1H), 7.61 - 7.37 (m, 3H), 7.14 - 6.98 (m, 1H), 5.01 - 4.81 (m, 1H), 4.31 - 4.13 (m, 2H), 3.79 - 3.63 (m, 2H), 1.42 (d, J = 6.5 Hz, 6H).	DMSO	415 (M+1)	1.65	Method A (Formic acid)	100	Method AR/AS/AT1
1688		HCl	¹ H NMR (300 MHz, DMSO) δ 10.12 (s, 1H), 9.63 (s, 1H), 9.10 - 8.88 (m, 2H), 8.72 (s, 1H), 8.44 (d, J = 8.3 Hz, 1H), 8.14 (d, J = 8.3 Hz, 1H), 7.85 (dd, J = 7.6, 4.8 Hz, 1H), 7.52 (d, J = 6.3 Hz, 1H), 7.39 (t, J = 7.9 Hz, 1H), 7.22 - 7.04 (m, 2H), 4.97 (s, 2H), 3.30 (d, J = 4.1 Hz, 3H), 2.97 (s, 3H), 2.85 (s, 3H).	DMSO	414 (M+1)	1.28	Method A (Formic acid)	100	Method AR/W with KI
1689		HCl	¹ H NMR (300 MHz, DMSO) δ 10.22 (s, 1H), 9.63 (s, 1H), 9.10 - 8.87 (m, 2H), 8.75 (s, 1H), 8.35 (d, J = 8.8 Hz, 1H), 8.15 (d, J = 8.7 Hz, 1H), 8.01 (d, J = 4.2 Hz, 1H), 7.93 - 7.77 (m, 1H), 7.51 (d, J = 7.3 Hz, 1H), 7.43 (t, J = 7.8 Hz, 1H), 7.16 (t, J = 7.5 Hz, 1H), 7.07 (d, J = 8.5 Hz, 1H), 4.54 (s, 2H), 3.29 (d, J = 3.9 Hz, 3H), 2.64 (d, J = 4.4 Hz, 3H).	DMSO	400 (M+1)	1.33	Method A (Formic acid)	100	Method AR/W with KI
1690		HCl	¹ H NMR (300 MHz, DMSO) δ 10.09 (s, 1H), 9.62 (s, 1H), 9.05 - 8.87 (m, 2H), 8.60 (s, 1H), 8.26 (d, J = 8.6 Hz, 1H), 8.13 (d, J = 8.8 Hz, 1H), 7.90 - 7.77 (m, 1H), 7.54 - 7.36 (m, 2H), 7.26 - 7.06 (m, 2H), 4.23 - 4.11 (m, 2H), 3.70 - 3.58 (m, 2H), 3.29 (d, J = 4.4 Hz, 3H), 3.24 (s, 3H).	DMSO	387 (M+1)	1.51	Method A (Formic acid)	100	Method AR/AS/AT1

(continued)

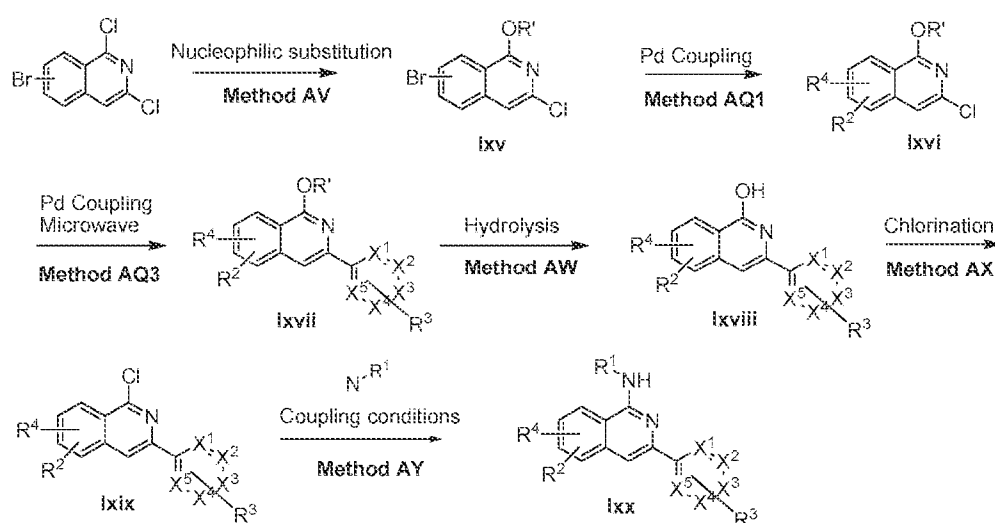
Number	PRODUCT	Salt	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1691		HCl	¹ H NMR (300 MHz, DMSO) δ 9.65 (s, 1H), 9.04 (d, J = 7.8 Hz, 1H), 8.94 (d, J = 5.0 Hz, 1H), 8.50 (s, 1H), 8.39-8.22 (m, 2H), 7.97-7.80 (m, 1H), 7.53-7.31 (m, 3H), 7.03 (d, J = 6.8 Hz, 1H), 4.26-4.13 (m, 2H), 3.82-3.59 (m, 8H), 3.32 (s, 3H).	DMSO	401 (M+1)	1.47	Method A (Formic acid)	100	Method AR/AS/AT1
1692		HCl	¹ H NMR (300 MHz, DMSO) δ 9.83-9.26 (m, 3H), 9.00-8.76 (m, 3H), 8.40 (d, J = 8.8 Hz, 1H), 8.18 (d, J = 8.6 Hz, 1H), 7.89-7.72 (m, 1H), 7.58-7.36 (m, 3H), 7.04 (d, J = 3.4 Hz, 1H), 4.30-4.18 (m, 2H), 3.79-3.66 (m, 2H), 3.33 (s, 3H).	DMSO	373 (M+1)	1.42	Method A (Formic acid)	100	Method AR/AS/AT1
1693		HCl	¹ H NMR (300 MHz, DMSO) δ 10.73-10.36 (m, 1H), 9.65 (s, 1H), 9.13-8.98 (m, 2H), 8.93 (d, J = 5.0 Hz, 1H), 8.38 (d, J = 8.4 Hz, 1H), 8.24 (d, J = 7.7 Hz, 1H), 7.94-7.75 (m, 1H), 7.69-7.41 (m, 3H), 7.09 (d, J = 8.1 Hz, 1H), 4.61-4.43 (m, 2H), 4.28-4.07 (m, 2H), 3.59-3.39 (m, 2H), 3.30 (d, J = 2.8 Hz, 3H).	DMSO	454 (M+1)	2.26	Method B (Ammonium formate)	100	Method AR/AS/AT3
1694		HCl	¹ H NMR (300 MHz, DMSO) δ 10.35 (s, 1H), 9.67 (s, 1H), 9.04 (d, J = 7.3 Hz, 1H), 8.94 (d, J = 5.0 Hz, 1H), 8.78 (s, 1H), 8.24 (s, 2H), 7.96-7.77 (m, 1H), 7.60-7.40 (m, 2H), 7.33-7.10 (m, 2H), 4.50-4.36 (m, 2H), 4.05-3.85 (m, 2H), 3.53-3.37 (m, 2H), 3.28 (d, J = 3.6 Hz, 3H).	DMSO	454 (M+1)	2.11	Method B (Ammonium formate)	100	Method AR/AS/AT3

(continued)

Number	PRODUCT	Salt	¹ H-NMR	¹ H-NMR Solvent	LCMS	Retention Time	LCMS Protocol	Purity percent	Method for Coupling
1695			¹ H NMR (300 MHz, DMSO) δ 9.64 (s, 1H), 8.77 (d, J = 8.0 Hz, 1H), 8.73 - 8.50 (m, 3H), 8.14 (d, J = 8.7 Hz, 1H), 7.84 (d, J = 8.7 Hz, 1H), 7.54 (dd, J = 7.9, 4.8 Hz, 1H), 7.48 - 7.34 (m, 3H), 6.96 (d, J = 3.2 Hz, 1H), 4.91 (s, 2H), 3.18 (d, J = 4.3 Hz, 3H), 3.03 (s, 3H), 2.86 (s, 3H).	DMSO	414 (M+1)	1.30	Method A (Formic acid)	100	Method AR/W with KI

Scheme 81: General route for the synthesis of compounds with general formula Ixx

[0517]



(14 mg, 0.02 mmol, 0.05 eq) under N₂ atmosphere. The sealed tube was irradiated in the CEM microwave at 130°C for 1h. After the reaction was completed, the volatiles were removed *in vacuo* and the residue was purified with chromatography on silica gel (hexane/ ethyl acetate 50:1) to give 200 mg of crude **lxvii-a** as a brown oil, which was used directly in the next step without purification. LCMS m/z = 419.0 (M + 1) (Method B) (retention time = 2.25 min).

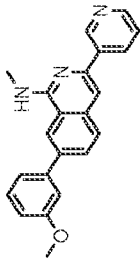
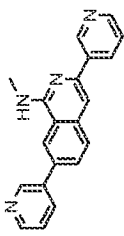
[0522] Method AW: 7-(3-Methoxyphenyl)-3-(pyridin-3-yl)isoquinolin-1-ol (lxviii-a) A mixture of 1-(benzyloxy)-7-(3-methoxyphenyl)-3-(pyridin-3-yl)isoquinoline (200 mg, 0.48 mmol), and concentrated HCl (10 mL) in ethanol (20 mL) was stirred at room temperature overnight. After the reaction was completed, the mixture was filtered and concentrated to afford 130 mg of **lxviii-a** as a yellow solid (yield 90% for two steps). LCMS m/z = 329.0 (M + 1) (Method B) (retention time = 1.66 min).

[0523] Method AX: 1-Chloro-7-(3-methoxyphenyl)-3-(pyridin-3-yl)isoquinoline (lxix-a) To a mixture of 7-(3-methoxyphenyl)-3-(pyridin-3-yl)isoquinolin-1-ol (130 mg, 0.395 mmol) in phenylphosphonic dichloride (5 mL) was stirred at 120°C overnight. After the reaction was completed, the mixture was added to ice-water slowly. The pH was adjusted to 7 by slow addition of NH₃ H₂O at 0°C. Then the mixture was extracted with dichloromethane (100 mL x 3). The combined organic layer were dried over MgSO₄ filtered and concentrated *in vacuo* to give 130 mg of crude **lxix-a** as a white solid, which was used directly in the next step without purification. LCMS m/z = 346.9 (M + 1) (Method B) (retention time = 1.80 min).

[0524] Method AY: 7-(3-Methoxyphenyl)-N-methyl-3-(pyridin-3-yl)isoquinolin-1-amine (lxx-a) A mixture of 1-chloro-7-(3-methoxyphenyl)-3-(pyridin-3-yl)isoquinoline (90 mg, 0.26 mmol), methylamine hydrochloride (200 mg, 2.96 mmol, 10.0 eq) and Et₃N (0.5 mL) was added to a sealed tube with i-AmOH (2 mL). The sealed tube was irradiated in the CEM microwave at 160 °C for 1 h. After the reaction was completed, the i-AmOH was removed *in vacuo* and the residue was dissolved in 10 mL of water and dichloromethane, the mixture was extracted with dichloromethane (3x 50 mL). The combined organic layers were dried over MgSO₄, filtered and concentrated *in vacuo* and the residue was washed with MeOH to give 5 mg of **lxx-a** as a brown solid (yield 6%). LCMS m/z = 342.1 (M+1) (Method B) (retention time = 2.02 min). ¹H NMR (400 MHz, DMSO-d₆): δ 9.40 (d, *J* = 0.9 Hz, 1H), 8.67 - 8.47 (m, 3H), 8.08 - 7.95 (m, 1H), 7.89 - 7.84 (m, 2H), 7.66 (s, 1H), 7.57 - 7.49 (m, 1H), 7.46 - 7.42 (m, 3H), 7.08 - 6.95 (m, 1H), 3.88 (s, 3H), 3.14 (d, *J* = 4.3 Hz, 3H).

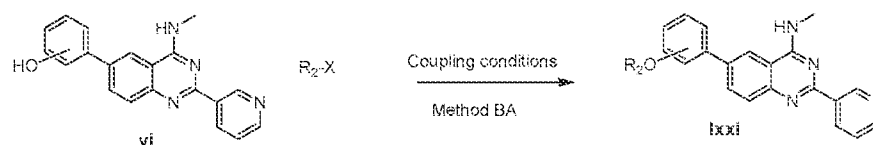
[0525] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 81**, replacing with the appropriate isoquinoline and aniline.

Table 27:

Number	PRODUCT	Salt type	Molecular Mass	¹ H-NMR	¹ H-NMR Solvent	LCMS	LCMS Protocol	Purity percent	Method for coupling
1697			341.41	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.40 (d, J = 0.9 Hz, 1H), 8.67 - 8.47 (m, 3H), 8.08 - 7.95 (m, 1H), 7.89 - 7.84 (m, 2H), 7.66 (s, 1H), 7.57 - 7.49 (m, 1H), 7.46 - 7.42 (m, 3H), 7.08 - 6.95 (m, 1H), 3.88 (s, 3H), 3.14 (d, J = 4.3 Hz, 3H).	DMSO	342.1 (M+1)	Method B (NH ₄ HCO ₃)	95	Method AX, H1
1698			312.37	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.40 (d, J = 1.8 Hz, 1H), 9.12 (d, J = 2.1 Hz, 1H), 8.66 (s, 1H), 8.60 (m, 2H), 8.54 (dt, J = 8.0, 1.9 Hz, 1H), 8.29 - 8.23 (m, 1H), 8.08 (dd, J = 8.5, 1.6 Hz, 1H), 7.92 (d, J = 8.5 Hz, 1H), 7.87 (d, J = 4.5 Hz, 1H), 7.68 (s, 1H), 7.56 (dd, J = 7.8, 4.8 Hz, 1H), 7.51 (dd, J = 7.8, 4.8 Hz, 1H), 3.14 (d, J = 4.4 Hz, 3H).	DMSO	312.9 (M+1)	Method B (NH ₄ HCO ₃)	95	Method AX, H1

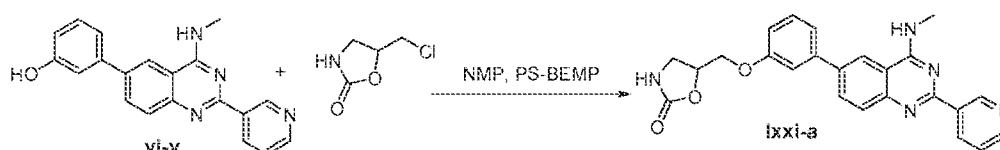
Scheme 83: General route for the synthesis of compounds with general formula lxxi

[0526]



Scheme 84: Representative synthesis of compounds of formula lxxi-a (see Scheme 81)

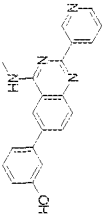
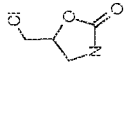
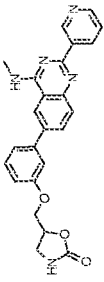
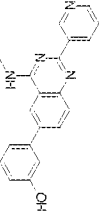
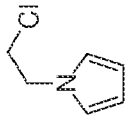
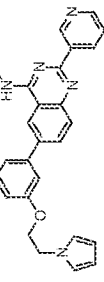
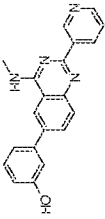
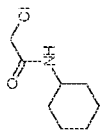
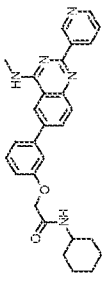
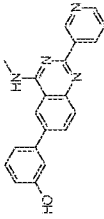
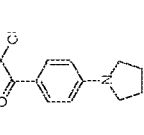
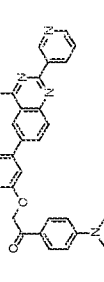
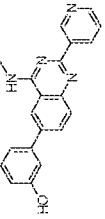
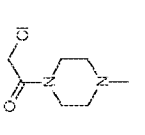
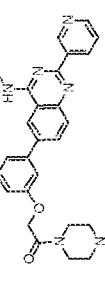
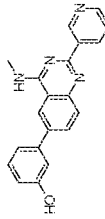
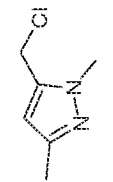
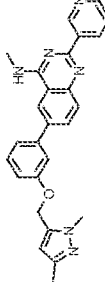
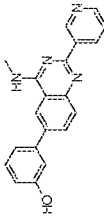
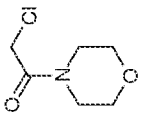
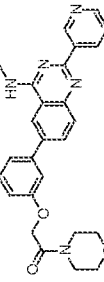
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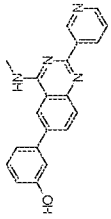
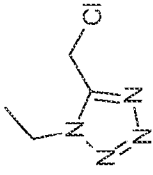
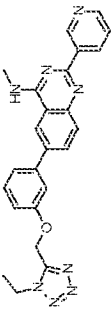
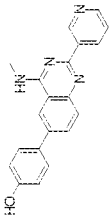
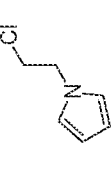
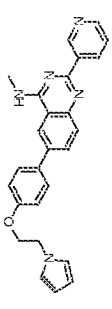
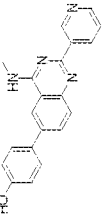
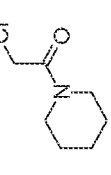
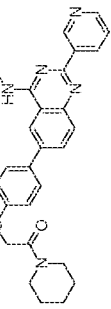
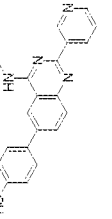
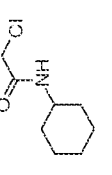
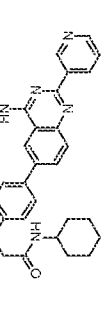
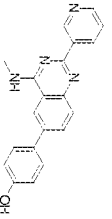
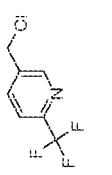
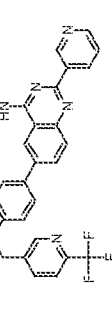
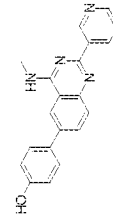
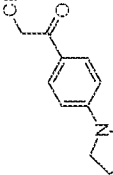
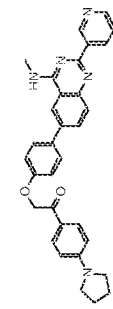
[0528] Method BA: Synthesis of 5-[3-(4-Methylamino-2-pyridin-3-yl-quinazolin-6-yl)-phenoxy-methyl]-oxazolidin-2-one (lxxi-a) To 5-(chloromethyl)oxazolidin-2-one (45 μ mol) was added the solution of 3-(4-(methylamino)-2-(pyridin-3-yl)quinazolin-6-yl)phenol (30 μ mol) in NMP (200 μ L). PS-BEMP (90 μ mol) was added to the vials by resin dispenser. After the reaction mixture was heated at 90°C for 12 h, the residue was diluted with methanol and purified by **PREP-HPLC Condition D**. The target fraction was lyophilized to afford the titled compound whose structure was finally confirmed by LCMS using LCMS **Method E**.

[0529] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 80**, replacing 5-(chloromethyl)oxazolidin-2-one with the appropriate alkyl halide.

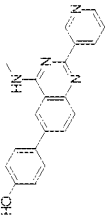
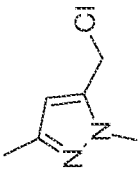
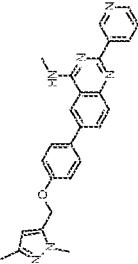
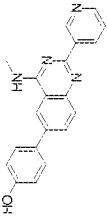
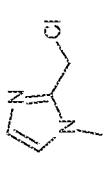
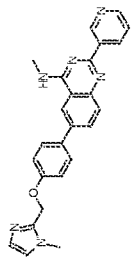
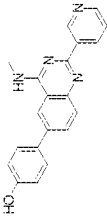
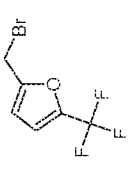
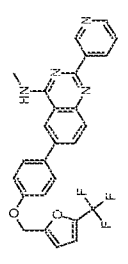
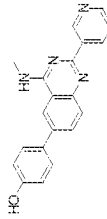
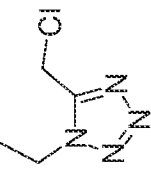
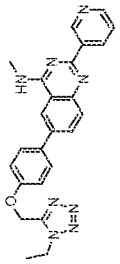
Table 28:

Number	Starting Material 1	Starting Material 2	Product	Salt Type	Exact Mass	Mass Found (M+1)	Purity (%)
1699					427	428	98
1700					421	422	98
1701				TFA	467	468	98
1702					515	516	98
1703				2TFA	468	469	98
1704				TFA	436	437	98
1705					455	456	98

(continued)

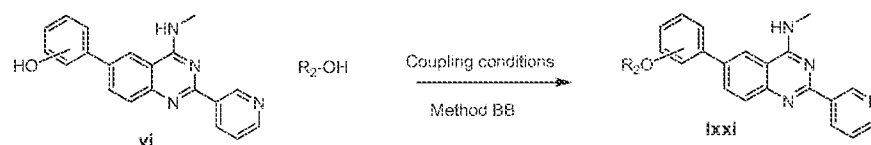
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1706					438	439	98
1707					421	422	98
1708					453	454	98
1709				TFA	467	468	98
1710					487	488	98
1711				TFA	515	516	98

(continued)

Number	Starting Material 1	Starting Material 2	Product	Salt Type	Exact Mass	Mass Found (M+1)	Purity (%)
1712				TFA	436	437	98
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1714				TFA	476	477	98
1715				2TFA	438	439	98

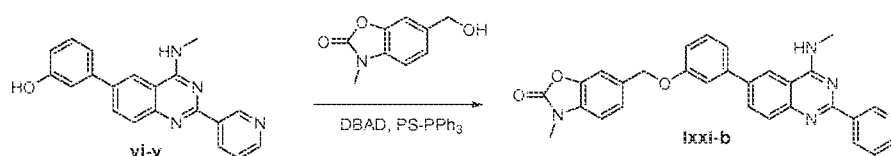
Scheme 85: General route for the synthesis of compounds with general formula lxxi

[0530]



Scheme 86: Representative synthesis of compounds of formula lxxi (see Scheme 85)

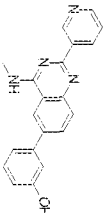
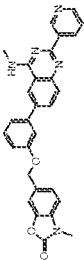
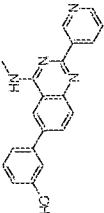
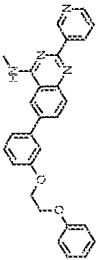
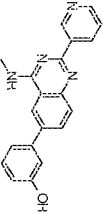
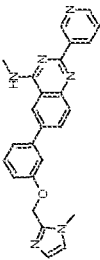
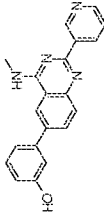
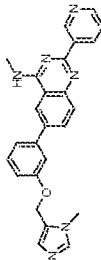
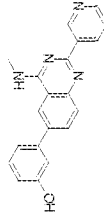
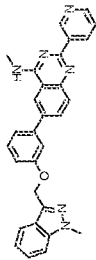
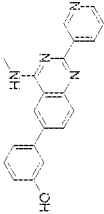
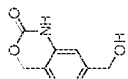
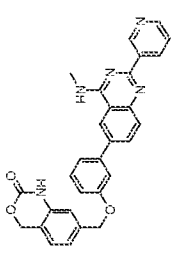
[0531]



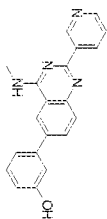
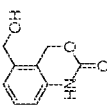
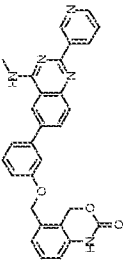
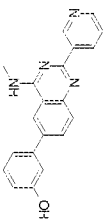
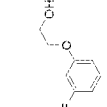
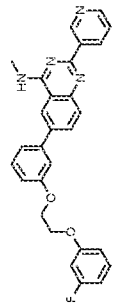
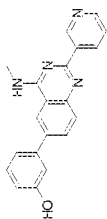
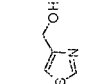
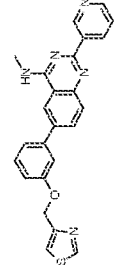
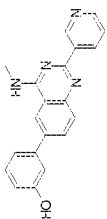
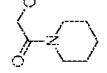
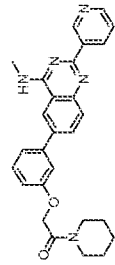
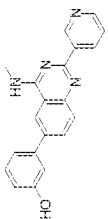
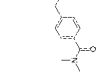
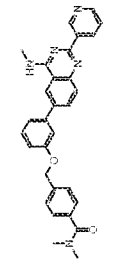
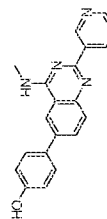
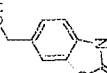
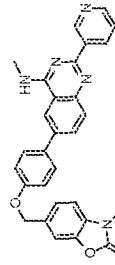
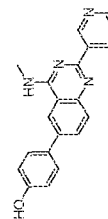
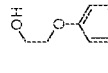
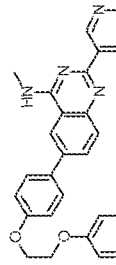
[0532] Method BB: Synthesis of 3-methyl-6-((3-(4-(methoxycarbonyl)-2-methylphenyl)propan-1-yloxy)methyl)benzo[d]oxazol-2(3H)-one (lxxi-b) To 6-(hydroxymethyl)-3-methylbenzo[d]oxazol-2(3H)-one (45 μ mol) was added the solution of 3-(4-(methoxycarbonyl)-2-methylphenyl)propan-1-ol (30 μ mol) in THF (400 μ L). After PS-triphenylphosphine (60 μ mol) was added, the solution of DBAD (di-*tert*-butyl azodicarboxylate, 66 μ mol) in THF was dispensed to the vials. The mixture was heated at 50°C for 8 h. After the solvent was removed, the residue was diluted with methanol and purified by Mass triggered **PREP-HPLC Condition D**. The target fraction was lyophilized to afford the titled compound whose structure was finally confirmed by LCMS using LCMS **Method E**.

[0533] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 85**, replacing 6-(hydroxymethyl)-3-methylbenzo[d]oxazol-2(3H)-one with the appropriate alkyl alcohol.

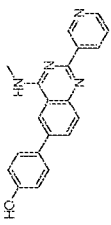
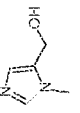
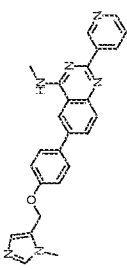
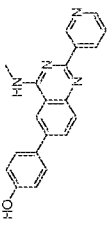
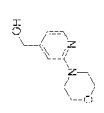
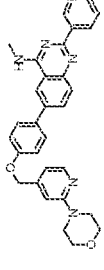
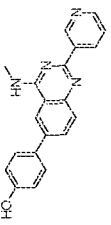
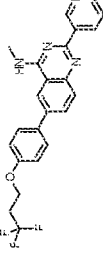
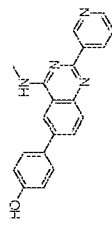
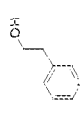
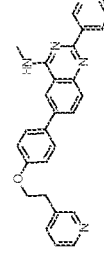
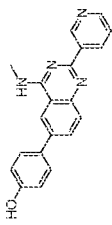
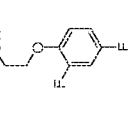
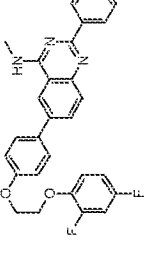
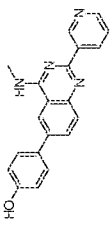
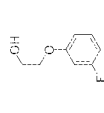
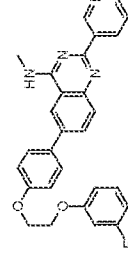
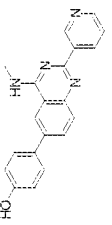
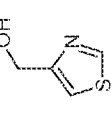
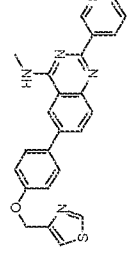
Table 29:

Number	Starting Material 1	Starting Material 2	Product	Salt Type	Exact Mass	Mass Found (M+1)	Purity (%)
1716				TFA	489	490	100
1717				TFA	448	449	100
1718				TFA	422	423	100
1719				TFA	422	423	100
1720				TFA	472	473	100
1721				TFA	489	490	100

(continued)

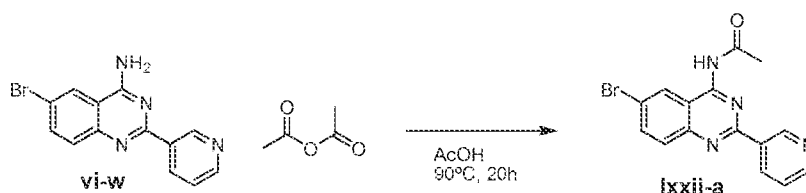
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1722				TFA	489	490	100
1723				TFA	466	467	100
1724				TFA	425	426	94
1725				TFA	453	454	100
1726				TFA	489	490	100
1727				TFA	489	490	100
1728				TFA	448	449	100

(continued)

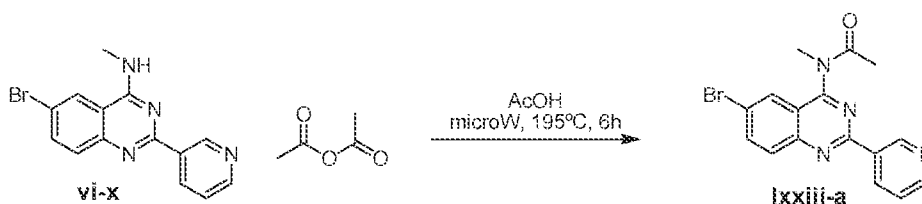
Number	Starting Material 1	Starting Material 2	Product	Salt Type	Exact Mass	Mass Found (M+1)	Purity (%)
1729				TFA	422	423	100
1730				TFA	504	505	100
1731				TFA	424	425	100
1732				TFA	433	434	100
1733				TFA	484	485	100
1734				TFA	466	467	100
1735				TFA	425	426	100

(continued)

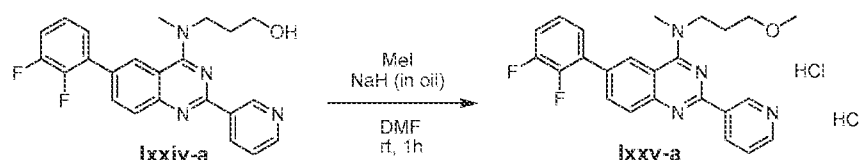
Number	Starting Material 1	Starting Material 2	Product	Salt Type	Exact Mass	Mass Found (M+1)	Purity (%)
1736				TFA	464	465	95
1737				TFA	466	467	100
1738				TFA	489	490	100

Scheme 87: Synthesis of N-(6-bromo-2-(pyridin-3-yl)quinazolin-4-yl)acetamide with formula Ixxii (Compound 1739)**[0534]**

[0535] Synthesis of N-(6-bromo-2-(pyridin-3-yl)quinazolin-4-yl)acetamide (Ixxii-a) A 100 mL round bottom flask was fitted with a reflux condenser and charged with 6-bromo-2-(pyridin-3-yl)quinazolin-4-amine (2.45 g, 8.14 mmol), acetic anhydride (5.81 g, 57.0 mmol) and acetic acid (30 mL). The reaction mixture was stirred at 90 °C for 20 min and then cooled to room temperature. A precipitate formed during the reaction and was collected by filtration and washed with water (100 mL). The solid was dried at 60 °C to give N-(6-bromo-2-(pyridin-3-yl)quinazolin-4-yl)acetamide as a white powder (1.83 g, 66%). ¹H NMR (300 MHz, CDCl₃) δ 9.72 (s, 1H), 8.84 - 8.69 (m, 3H), 8.22 (s, 1H), 8.02 - 7.90 (m, 2H), 7.53 - 7.41 (m, 1H), 2.83 (s, 3H).

Scheme 88: Synthesis of N-(6-bromo-2-(pyridin-3-yl)quinazolin-4-yl)-N-methylacetamide with formula Ixxiii (Compound 1740)**[0536]**

[0537] Synthesis of N-(6-bromo-2-(pyridin-3-yl)quinazolin-4-yl)-N-methylacetamide (Ixxiii-a) To a solution of 6-bromo-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine (1.20 g, 3.81 mmol) in acetic acid (10 mL) was added acetic anhydride (1.94 g, 19.0 mmol) and stirred at 195 °C using microwave for 6h. The reaction mixture was checked by LC-MS, no starting material was observed, ice was added into the reaction. The precipitate was collected by filtration and washed with water. The product was dried at 60 °C to give N-(6-bromo-2-(pyridin-3-yl)quinazolin-4-yl)-N-methylacetamide (711 mg, 52%) as a light brown powder. ¹H NMR (300 MHz, CDCl₃) δ 9.76 (dd, J = 2.2, 0.8 Hz, 1H), 8.85 - 8.79 (m, 1H), 8.76 (dd, J = 4.8, 1.7 Hz, 1H), 8.09 (dd, J = 1.8, 0.8 Hz, 1H), 8.05 - 8.01 (m, 2H), 7.46 (ddd, J = 8.0, 4.8, 0.8 Hz, 1H), 3.52 (s, 3H), 2.11 (s, 3H).

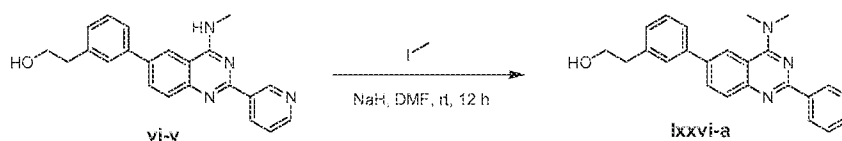
Scheme 89: Synthesis of compounds of formula Ixxv**[0538]****Method BC : Synthesis of 6-(2,3-difluorophenyl)-N-(3-methoxypropyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride (Ixxv-a) (Compound 1742)**

[0539] To a solution of the hydroxyl derivative (520 mg, 1.27 mmol) in DMF (20 mL) was added methyl iodide (262

mg, 1.84 mmol) and NaH (55% in oil; 69 mg, 1.9 mmol) at 0 °C. The reaction mixture was stirred at room temperature for 1 h and the ice water was added into the reaction mixture. The resulting solution was extracted with ethyl acetate (3 times), dried over Na₂SO₄, concentrated and purified by column chromatography (NH-silica gel, 50% hexane/50% ethyl acetate). The resulting product was dissolved in isopropyl alcohol and 1N-HCl (5 mL) was added and a precipitate formed which was collected by filtration and dried at 60 °C to give 6-(2,3-difluorophenyl)-N-(3-methoxypropyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride as an orange powder (130 mg, 21%). ¹H NMR (300 MHz, DMSO) δ 9.66 (d, *J* = 1.6 Hz, 1H), 9.09 (d, *J* = 8.1 Hz, 1H), 8.96 (dd, *J* = 5.1, 1.5 Hz, 1H), 8.51 (s, 1H), 8.33 (d, *J* = 8.7 Hz, 1H), 8.20 (d, *J* = 8.8 Hz, 1H), 7.90 (dd, *J* = 8.1, 5.1 Hz, 1H), 7.66 - 7.47 (m, 2H), 7.47 - 7.32 (m, 1H), 4.22 - 4.07 (m, 2H), 3.70 (s, 3H), 3.48 (t, *J* = 5.8 Hz, 2H), 3.22 (s, 3H), 2.19 - 2.01 (m, 2H).

Scheme 90: Synthesis of 2-(3-(4-(dimethylamino)-2-(pyridin-3-yl)quinazolin-6-yl)phenyl)ethanol of formula Ixxvi (Compound 1743)

[0540]

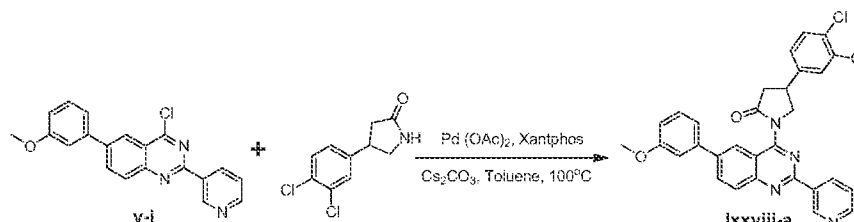


[0541] Method BD: 2-(3-(4-(dimethylamino)-2-(pyridin-3-yl)quinazolin-6-yl)phenyl)ethanol, 2HCl (Ixxvi-a) To a reaction vial containing 2-(3-(4-(methylamino)-2-(pyridin-3-yl)quinazolin-6-yl)phenyl)ethanol (89.0 mg, 0.25 mmol) in DMF (1 mL) was added 60% sodium hydride (13 mg, 0.325 mmol) and iodomethane (0.02 mL, 0.325 mmol). The reaction mixture was allowed to stir at ambient temperature for 12 h. Water (20 mL) was added to the reaction mixture, and the crude product was extracted with ethyl acetate (4 x 15 mL). The crude material was purified via ISCO (silica, 4 g column, 95% CH₂Cl₂ - 5% MeOH - 0.1% NH₄OH) to give the product as an off-white solid. The free base was then converted to the HCl salt to yield the final product as a yellow solid (24.6 mg, 0.055 mmol, 22%). LC-MS *m/z* = 371.5 (*M*+1) (retention time = 1.86) ¹H NMR (300 MHz, DMSO) δ 9.61 (d, *J* = 2.1 Hz, 1H), 8.99 (d, *J* = 7.5 Hz, 1H), 8.93 (dd, *J* = 5.0, 1.5 Hz, 1H), 8.48 (d, *J* = 1.0 Hz, 1H), 8.28 (dd, *J* = 17.4, 9.4 Hz, 2H), 7.84 (dd, *J* = 7.7, 5.1 Hz, 1H), 7.69 - 7.60 (m, 2H), 7.44 (t, *J* = 7.5 Hz, 1H), 7.30 (d, *J* = 7.3 Hz, 1H), 3.78 - 3.61 (m, 9H), 2.82 (t, *J* = 7.0 Hz, 2H).

[0542] Synthesis of 6-(2,4-difluorophenyl)-N-(3-methoxypropyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride (Ixxvii-a) (Compound 1744) 6-(2,4-difluorophenyl)-N-(3-methoxypropyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride was synthesized in a similar manner to that described for 6-(2,3-difluorophenyl)-N-(3-methoxypropyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine dihydrochloride substituting the appropriate hydroxyl derivative. 6-(2,4-difluorophenyl)-N-(3-methoxypropyl)-N-methyl-2-(pyridin-3-yl)quinazolin-4-amine was obtained as the dihydrochloride salt ¹H NMR (300 MHz, DMSO) δ 9.69 (d, *J* = 2.0 Hz, 1H), 9.14 (d, *J* = 8.3 Hz, 1H), 8.97 (dd, *J* = 5.1, 1.2 Hz, 1H), 8.45 (s, 1H), 8.39 (d, *J* = 8.6 Hz, 1H), 8.17 (d, *J* = 8.8 Hz, 1H), 7.92 (dd, *J* = 8.0, 5.2 Hz, 1H), 7.86 - 7.73 (m, 1H), 7.57 - 7.38 (m, 1H), 7.36 - 7.23 (m, 1H), 4.19 - 4.04 (m, 2H), 3.71 (s, 3H), 3.48 (t, *J* = 5.7 Hz, 2H), 3.23 (s, 3H), 2.21 - 1.96 (m, 2H).

Scheme 91: Synthesis of compounds of formula Ixxviii

[0543]

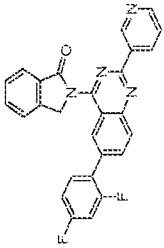
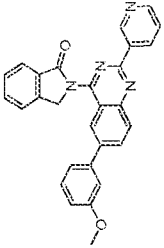
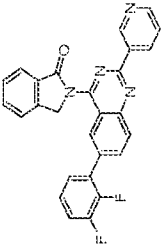
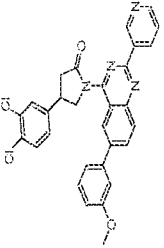
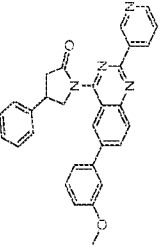


[0544] Method BE: 4-(3,4-dichlorophenyl)-1-(6-(3-methoxyphenyl)-2-(pyridin-3-yl)quinazolin-4-yl)pyrrolidin-2-one (Ixxviii-a) To a mixture of 4-chloro-6-(3-methoxyphenyl)-2-(pyridin-3-yl)quinazoline (40 mg, 0.12 mmol), 4-(3,4-dichlorophenyl)pyrrolidin-2-one (100 mg, 0.437 mmol) and Cs₂CO₃ (42 mg, 0.127 mmol) in dry toluene (6 mL) was added Pd(OAc)₂ (3 mg, 0.01 mmol) and Xantphos (10 mg, 0.02 mmol) under a nitrogen atmosphere. The resulting

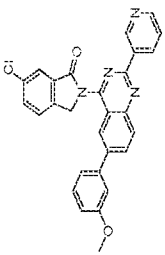
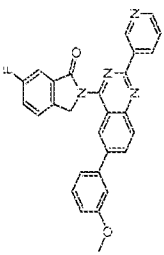
mixture was stirred at 100 °C for 12h. After cooling, the mixture was filtered through a pad of celite. The residue was purified by silica gel chromatography, eluted with petroleum ether/ethyl acetate (5:4) to give the desired product as a yellow solid. 14 mg of the desired product was obtained in a 22.5% yield, LCMS: retention time = 1.802 min, $[MH]^+ = 541.0, 543.0$. 1H -NMR (400 MHz, DMSO- d_6): δ 9.73 (s, 1H), 9.05 - 9.04 (m, 1H), 8.86 - 8.85 (m, 1H), 8.42 - 8.38 (m, 2H), 8.19 (d, $J = 8.8$ Hz, 1H), 7.89 (d, $J = 1.6$ Hz, 1H), 7.82 - 7.81 (m, 1H), 7.69 (d, $J = 8.0$ Hz, 1H), 7.58 (dd, $J = 8.0, 1.6$ Hz, 1H), 7.50 (t, $J = 8.0$ Hz, 1H), 7.41 - 7.36 (m, 2H), 7.07 (dd, $J = 8.0, 2.0$ Hz, 1H), 4.57 - 4.46 (m, 2H), 4.06 - 4.03 (m, 1H), 3.20 - 3.01 (m, 2H), 3.88 (s, 3H).

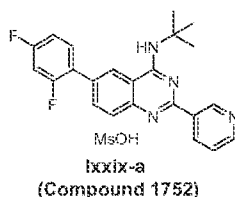
[0545] The compounds in the following table were prepared in a manner analogous to that described in **Scheme 91**, replacing 4-(3,4-dichlorophenyl)pyrrolidin-2-one with the appropriate amide

Table 30:

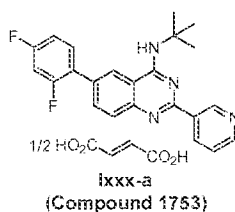
Number	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
1745			¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.75 (s, 1H), 8.90 (d, J = 8.0 Hz, 1H), 8.78 (d, J = 4.0 Hz, 1H), 8.35 (s, 1H), 8.28 - 8.19 (m, 2H), 7.92 (d, J = 7.6 Hz, 1H), 7.82 (d, J = 5.0 Hz, 2H), 7.73 - 7.70 (m, 1H), 7.68 - 7.61 (m, 2H), 7.52 - 7.42 (m, 1H), 7.29 (t, J = 8.4 Hz, 1H), 5.61 (s, 2H).	DMSO	95	Method BE	451.0 (M+1)	Method B (NH ₄ HCO ₃)
1746			¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.74 (s, 1H), 8.89 (d, J = 8.0 Hz, 1H), 8.77 (d, J = 3.8 Hz, 1H), 8.45 (s, 1H), 8.39 (d, J = 8.7 Hz, 1H), 8.18 (d, J = 8.7 Hz, 1H), 7.94 (d, J = 7.6 Hz, 1H), 7.83 (s, 2H), 7.72 - 7.56 (m, 2H), 7.52 - 7.27 (m, 3H), 7.03 (d, J = 7.9 Hz, 1H), 5.60 (s, 2H), 3.85 (s, 3H).	DMSO	95	Method BE	445.1 (M+1)	Method B (NH ₄ HCO ₃)
1747			¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.76 (s, 1H), 8.91 (d, J = 6.4 Hz, 1H), 8.79 (s, 1H), 8.41 (s, 1H), 8.30 - 8.23 (m, 2H), 7.92 (d, J = 8.0 Hz, 1H), 7.83 (s, 2H), 7.65 (d, J = 3.6 Hz, 2H), 7.55 - 7.49 (m, 2H), 7.40 (s, 1H), 5.61 (s, 2H).	DMSO	95	Method BE	451.0, 452.0 (M+1)	Method B (NH ₄ HCO ₃)
1748		2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.73 (s, 1H), 9.05 - 9.04 (m, 1H), 8.86 - 8.85 (m, 1H), 8.42 - 8.38 (m, 2H), 8.19 (d, J = 8.8 Hz, 1H), 7.89 (d, J = 1.6 Hz, 1H), 7.82 - 7.81 (m, 1H), 7.69 (d, J = 8.0 Hz, 1H), 7.58 (dd, J = 8.0, 1.6 Hz, 1H), 7.50 (t, J = 8.0 Hz, 1H), 7.41 - 7.36 (m, 2H), 7.07 (dd, J = 8.0, 2.0 Hz, 1H), 4.57 - 4.46 (m, 2H), 4.06 - 4.03 (m, 1H), 3.20 - 3.01 (m, 2H), 3.88 (s, 3H).	DMSO	95	Method BE	541.0, 543.0 (M+1)	Method B (NH ₄ HCO ₃)
1749		2HCl	¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.68 (s, 1H), 8.83 (d, J = 8.0 Hz, 1H), 8.76 (s, 1H), 8.39 - 8.37 (m, 2H), 8.17 (d, J = 10.0 Hz, 1H), 7.63 - 7.31 (m, 9H), 7.06 (dd, J = 8.4, 2.0 Hz, 1H), 4.56 - 4.45 (m, 2H), 4.03 - 3.99 (m, 1H), 3.88 (s, 3H), 3.19 - 2.97 (m, 2H).	DMSO	95	Method BE	472.9 (M+1)	Method B (NH ₄ HCO ₃)

(continued)

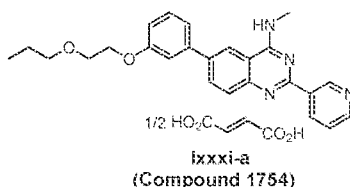
Number	Product	Salt type	¹ H NMR	¹ H NMR Solvent	Purity percent	Method of Coupling	LCMS	LCMS Method
1750			¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.74 (d, J = 8.4 Hz, 1H), 8.89 (dt, J = 8.4, 1.6 Hz, 1H), 8.78 (dd, J = 4.4, 1.2 Hz, 1H), 8.46 (d, J = 1.6 Hz, 1H), 8.41 (dd, J = 8.8, 1.6 Hz, 1H), 8.20 (d, J = 8.4 Hz, 1H), 7.97 (s, 1H), 7.88 (s, 2H), 7.65 - 7.62 (m, 1H), 7.48 - 7.44 (m, 1H), 7.41 - 7.37 (m, 2H), 7.04 (dd, J = 8.0, 1.6 Hz, 1H), 5.59 (s, 2H), 3.86 (s, 3H).	DMSO	95	Method BE	479.0 (M+1)	Method B (NH ₄ HCO ₃)
1751			¹ H-NMR (400 MHz, DMSO-d ₆): δ 9.74 (s, 1H), 8.89 (d, J = 8.4 Hz, 1H), 8.78 (d, J = 4.4 Hz, 1H), 8.46 (s, 1H), 8.41 (d, J = 9.2 Hz, 1H), 8.20 (d, J = 8.8 Hz, 1H), 7.91 - 7.87 (m, 1H), 7.77 - 7.62 (m, 3H), 7.48 - 7.37 (m, 3H), 7.03 (d, J = 6.4 Hz, 1H), 5.58 (s, 2H), 3.86 (s, 3H).	DMSO	95	Method BE	463.1 (M+1)	Method B (NH ₄ HCO ₃)

Scheme 91: Synthesis of N-tert-butyl-6-(2,4-difluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine methanesulfonate (Ixxix-a) (Compound 1752)**[0546]**

[0547] N-tert-butyl-6-(2,4-difluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine (1.51 g, 3.87 mmol) was dissolved in $\text{CH}_2\text{Cl}_2/\text{MeOH}$ (20 mL/20 mL). Methanesulfonic acid (0.251 mL, 3.87 mmol) was added to the solution. The volatiles were evaporated *in vacuo*. The resultant residue was crystallized from EtOH (30 mL) to give 1.35 g of N-tert-butyl-6-(2,4-difluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine methanesulfonate as a light yellow powder (72 %). LCMS m/z = 391 (M + 1) (Method D) (Retention time = 1.91 min). ^1H NMR (300 MHz, DMSO) δ 9.52 (s, 1H), 9.01 - 8.83 (m, 2H), 8.75 (s, 1H), 8.58 (s, 1H), 8.09 (d, J = 7.2 Hz, 1H), 8.03 - 7.85 (m, 2H), 7.83 - 7.68 (m, 1H), 7.59 - 7.43 (m, 1H), 7.33 (t, J = 7.9 Hz, 1H), 2.31 (s, 3H), 1.66 (s, 9H).

Scheme 92: Synthesis of N-tert-butyl-6-(2,4-difluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine fumarate (Ixxx-a) (Compound 1753)**[0548]**

[0549] N-tert-butyl-6-(2,4-difluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine (1.22 g, 3.12 mmol) was dissolved in $\text{CH}_2\text{Cl}_2/\text{MeOH}$ (20 mL/20 mL). Fumaric acid (0.363 g, 3.12 mmol) was added to the solution. The mixture was sonicated until fumaric acid was dissolved. Then, the volatiles were evaporated *in vacuo*. The resultant solid was washed with MeOH and dried to give 1.28 g of N-tert-butyl-6-(2,4-difluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine 1/2 fumarate as a light yellow powder (91 %). LCMS m/z = 391 (M + 1) (Method D) (Retention time = 1.95 min). ^1H NMR (300 MHz, DMSO) δ 13.14 (s, 1H), 9.59 (s, 1H), 8.82 - 8.65 (m, 2H), 8.57 (s, 1H), 7.89 (d, J = 8.7 Hz, 1H), 7.82 (d, J = 8.7 Hz, 1H), 7.79 - 7.62 (m, 2H), 7.61 - 7.52 (m, 1H), 7.51 - 7.38 (m, 1H), 7.28 (t, J = 8.6 Hz, 1H), 6.61 (s, 1H), 1.64 (s, 9H).

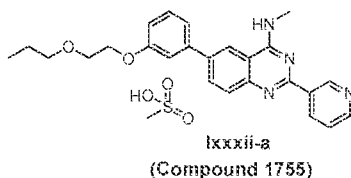
Scheme 93: Synthesis of N-methyl-6-(3-(2-propoxyethoxy)phenyl)-2-(pyridin-3-yl)quinazolin-4-amine fumarate (Ixxxi-a) (Compound 1754)**[0550]**

[0551] N-methyl-6-(3-(2-propoxyethoxy)phenyl)-2-(pyridin-3-yl)quinazolin-4-amine fumarate was synthesized in a similar manner to that described N-tert-butyl-6-(2,4-difluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine fumarate. LCMS m/z = 415.5 (M + 1) (Method C (NH_4HCO_3)) (Retention time = 2.43 min). ^1H NMR (300 MHz, CD_3OD) δ 9.57 (s, 1H), 8.84

(m, 1H), 8.63 (s, 1H), 8.35 (m, 1H), 8.07 (m, 1H), 7.88 (d, $J = 8.7$ Hz, 1H), 7.58 (m, 1H), 7.48 - 7.24 (m, 4H), 7.00 (m, 1H), 6.74 (s, 1H), 4.36 - 4.05 (m, 2H), 4.00 - 3.70 (m, 2H), 3.53 (t, $J = 6.6$ Hz, 2H), 3.29 - 3.13 (m, 3H), 1.75 - 1.51 (m, 2H), 0.95 (t, $J = 7.4$ Hz, 3H).

Scheme 94: Synthesis of N-methyl-6-(3-(2-propoxyethoxy)phenyl)-2-(pyridin-3-yl)quinazolin-4-amine methanesulfonate (Ixxxii-a) (Compound 1755)

[0552]



[0553] N-methyl-6-(3-(2-propoxyethoxy)phenyl)-2-(pyridin-3-yl)quinazolin-4-amine methanesulfonate was synthesized in a similar manner to that described N-tert-butyl-6-(2,4-difluorophenyl)-2-(pyridin-3-yl)quinazolin-4-amine methanesulfonate. LCMS $m/z = 415.5$ ($M + 1$) (Method C (NH_4HCO_3)) (Retention time = 2.43 min). ^1H NMR (300 MHz, CD_3OD) δ 9.49 (s, 1H), 9.00 - 8.72 (m, 1H), 8.53 (s, 1H), 8.29 (m, 1H), 7.99 (d, $J = 8.7$ Hz, 1H), 7.80 (dd, $J = 8.0, 5.1$ Hz, 1H), 7.52 - 7.25 (m, 3H), 7.02 (m, 1H), 4.38-4.06 (m, 2H), 3.98-3.72 (m, 2H), 3.61 - 3.48 (m, 2H), 3.42 (d, $J = 0.6$ Hz, 3H), 3.34 - 3.26 (m, 2H), 2.71 (s, 3H), 1.80-1.47 (m, 2H), 0.96 (t, $J = 7.4$ Hz, 3H).

Biological Testing:

STEP46 biochemical assays

[0554] Serial dilutions of compounds were performed in 100% DMSO and 1 μL of compounds were dispensed into 384-well black polystyrene plates (Corning, NY). Compounds were incubated with 24 μL of buffer containing 50 mM Hepes, 1 mM DTT, 0.02% Brij35, 1 ng/well purified STEP46 enzyme for 30 min at room temperature. The reaction was initiated by addition of 25 μL of DiFMUP (6, 8-difluoro-4-methylumbelliferyl phosphate) (Invitrogen, CA) with a final concentration of 10 μM and incubated at 27 $^\circ\text{C}$ for 90 min. Final DMSO concentration is 2%. Plates were read with fluorescence intensity at excitation/emission of 360/460nm using a PheraStar plate reader (BMG Labtech, NC).

Data analysis

[0555] Data were expressed as percentage (%) inhibition of enzyme activity. 0% inhibition is defined as the RFUs (relative fluorescence units) in the absence of compounds and 100% inhibition is defined as RFUs in the absence of STEP46 enzyme. IC_{50} values of compounds with inhibitory activity against STEP46 were determined by GraphPad Prism (version 4.03) using four parameter logistic equation. Some compounds act as activators. For compounds showing STEP46 enzymatic activation, data are represented as percentage of inhibition but with negative values at three representative concentrations (25, 50 and 100 μM).

[0556] Compounds **1-1760** show either inhibition or activation > 50% at 100 μM , 50 or 25 μM .

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Number	STEP IC50 (μ M)		Number	STEP IC50 (μ M)		Number	STEP IC50 (μ M)		Number	STEP IC50 (μ M)	
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296	++		329	+++		392	++		428	++	
297	+		330	++		393	++		429	+++	
300	+		332	+++		396	++		430	+++	
303	++		333	+++		399	+++		431	+++	
306	+		334	++		400	++		433	++	
309	++		335	+++		403	++		434	++	
312	+++		336	+++		407	++		435	+++	
313	+++		337	+++		412	+++		436	++	
314	++		339	++		415	++		437	++	
315	++		341	++		416	+++		439	+++	
316	+++		342	++		418	++		440	++	
317	++		343	+		419	+++		441	++	
318	++		344	++		421	+		442	+++	
320	++		346	++		423	++		443	+++	
322	+		348	++		424	++		444	+++	
325	+++		352	++		425	++		445	+++	
326	++		390	++		426	+		446	++	

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Number	STEP IC50 (μ M)	Number	STEP IC50 (μ M)	Number	STEP IC50 (μ M)	Number	STEP IC50 (μ M)
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448	+++	471	++	643	+++	664	++
449	++	472	++	644	++	665	++
451	+++	473	++	645	++	666	+++
452	+++	474	++	647	++	667	+++
453	+++	629	++	648	++	668	+++
454	++	630	++	649	++	669	+++
455	++	631	++	650	+++	670	+++
456	+++	632	+++	651	+++	671	++
457	++	633	++	652	+++	672	++
458	+++	634	++	653	+++	673	++
459	+++	635	+++	655	+++	674	++
460	++	636	+	656	+++	675	+++
461	++	637	++	657	+++	676	+++
463	++	638	++	658	++	677	+
466	++	639	+++	660	+++	678	++
467	++	640	++	661	+++	679	+++
468	+++	641	+++	662	+++	680	++

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Number	STEP IC50 (μ M)	Number	STEP IC50 (μ M)	Number	STEP IC50 (μ M)	Number	STEP IC50 (μ M)
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682	++	700	+++	901	++	1151	++
683	++	701	++	923	++	1152	++
684	++	702	+++	932	++	1153	++
685	+++	703	+++	933	++	1155	++
686	+++	704	+++	934	+++	1157	++
687	+++	705	+++	962	+++	1158	++
688	+++	706	+++	1124	+++	1159	++
689	+++	707	+++	1125	++	1160	++
690	++	708	++	1126	++	1161	++
691	+++	709	+++	1140	+++	1165	+++
692	+++	846	++	1141	+++	1167	++
693	+++	847	+++	1142	++	1168	++
694	+++	884	+	1143	++	1171	++
695	+++	888	+	1144	++	1172	++
696	+++	889	++	1145	++	1173	+
697	++	898	+	1146	++	1176	++
698	+++	899	++	1148	++	1178	++

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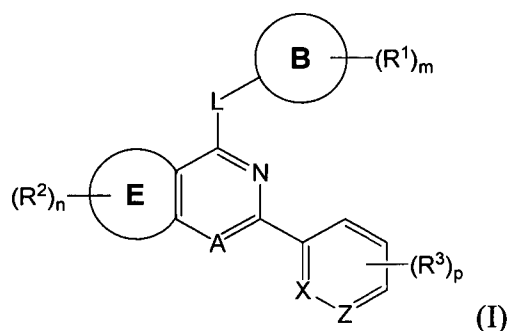
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1182	++	1206	++	1228	++	1328	+
1183	++	1208	+	1229	++	1330	+
1184	++	1211	+	1230	++	1332	++
1185	++	1212	++	1234	++	1333	+
1186	++	1213	++	1237	++	1334	+
1187	+++	1214	++	1238	++	1335	++
1188	++	1215	+++	1239	++	1336	++
1191	++	1216	+++	1248	++	1337	++
1192	++	1217	++	1279	++	1339	++
1193	++	1218	++	1280	++	1340	++
1194	+++	1219	++	1281	+++	1341	++
1198	++	1220	+	1282	++	1346	++
1200	++	1221	+	1283	+	1348	+
1201	++	1222	++	1284	++	1350	++
1202	++	1223	+++	1285	+	1351	++
1203	++	1225	++	1286	++	1352	+

Number	STEP IC ₅₀ (μ M)	Number	STEP IC ₅₀ (μ M)	Number	STEP IC ₅₀ (μ M)	Number	STEP IC ₅₀ (μ M)
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1354	++	1375	++	1411	+	1449	+
1356	++	1376	++	1412	++	1450	++
1357	++	1380	++	1413	++	1451	++
1358	++	1381	++	1414	++	1452	+++
1359	++	1382	+++	1415	+	1453	++
1360	++	1391	+	1416	++	1454	+
1361	+	1395	++	1419	++	1455	++
1362	++	1396	++	1420	++	1456	++
1363	++	1400	++	1422	++	1458	+
1364	++	1401	++	1423	+	1467	++
1365	++	1402	++	1424	+	1497	++
1366	++	1403	+	1425	++	1516	+
1369	++	1404	+	1430	++	1601	++
1370	++	1406	++	1437	++	1611	++
1371	++	1407	++	1445	++	1680	++
1372	++	1408	++	1446	++		
1373	++	1409	++	1447	+		
Key							
+	IC ₅₀ > 10 μ M						
++	IC ₅₀ 1-10 μ M						
+++	IC ₅₀ < 1 μ M						

[0557] Having thus described several aspects of at least one embodiment of this invention, it is to be appreciated various alterations, modifications, and improvements will readily occur to those skilled in the art. Such alterations, modifications, and improvements are intended to be part of this disclosure, and are intended to be within the scope of the invention. Accordingly, the foregoing description and drawings are by way of example only.

Claims

1. A compound of formula (I):



or a salt thereof,
wherein:

A is CH or N;

B is phenyl, dihydroindenyl, dihydrobenzoxazinyl, dihydrobenzodioxinyl, chromenyl, tetrahydroisoquinolyl, tetrahydroquinolyl, dihydroquinolyl, quinolyl, tetrahydroquinazolinyl, indolyl, dihydrobenzothiazolyl, dihydrobenzimidazolyl, dihydrobenzoxazolyl, isoindolyl, benzofuryl, benzothienyl, benzodioxolyl, indolyl, indazolyl, benzoimidazolyl, benzotriazolyl, benzisoxazolyl, benzoxazolyl, benzothiazolyl, benzoxadiazolyl, tetrahydrobenzothienyl, thienyl, thiazolyl, imidazolyl, pyrazolyl, thiadiazolyl or pyridyl;

m is 1, 2, 3, or 4;

E is phenyl, thienyl or pyrrolyl;

when E is phenyl, n is 1, or 2; and when E is thienyl or pyrrolyl, n is 0, or 1;

L is NR⁵;

one of X and Z is N and the other is CH;

p is 0, 1, or 2;

R¹ is C₁-C₈ alkyl, halo, haloalkyl, haloalkoxy, alkoxyalkyl, -CN, oxo, -NO₂, -C(Y)NR^bR^{b'}, -NR^cC(Y)R^{c'}, -SO₂NR^bR^{b'}, -OR^d, or -S(O)_qR^f;

R² is C₁-C₈ alkyl, C₃-C₈ cycloalkyl, thiomorpholinylalkyl, pyrrolidinylalkyl, morpholinylalkyl, haloalkyl, haloalkoxy, aminoalkyl, alkylaminoalkyl, dialkylaminoalkyl, hydroxyalkyl, alkoxyalkyl, silyloxyalkyl, -CN, -NO₂, -C(O)OR^a, -C(Y)NR^bR^{b'}, -NR^bR^{b'}, -OR^d, -C(Y)R^e or -S(O)_qR^f;

R³ is C₁-C₈ alkyl, halo, haloalkyl, -NR^bR^{b'} or -OR^d

R⁵ is hydrogen;

Y is O or S;

q is 1 or 2 and

each R^a, R^b, R^{b'}, R^c, R^{c'}, R^d, R^e and R^f is independently hydrogen, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₃-C₈ cycloalkyl, phenyl, dihydroindenyl, morpholinyl, tetrahydropyranyl, piperidyl, pyrrolidinyl, thiomorpholinyl, phenylalkyl, thienylalkyl, pyridylalkyl, tetrahydropyranylalkyl, dihydroindenylalkyl, tetrahydrofurylalkyl, hydroxyalkyl, morpholinylalkyl, pyrrolidinylalkyl, dialkylaminoalkyl, piperidylalkyl, benzodioxolilalkyl, dihydrobenzodioxinylalkyl, C₃-C₈ cycloalkylalkyl, haloalkyl or alkoxyalkyl.

2. The compound according to claim 1 or a salt thereof, wherein

A is N;

B is phenyl or pyridyl;

m is 1, 2 for 3;

E is phenyl;

n is 1, or 2; and

p is 0 or 1.

3. The compound according to claim 2 or a salt thereof, wherein

m is 1 or 2 ;

p is 0;

R¹ is C₁-C₈ alkyl, halo, haloalkyl, haloalkoxy, -CN, -NO₂, -C(Y)NR^bR^{b'}, -NR^cC(Y)R^{c'}, -SO₂NR^bR^{b'}, -OR^d, or -S(O)_qR^f;

and

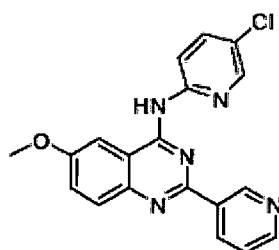
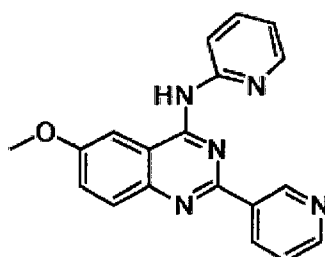
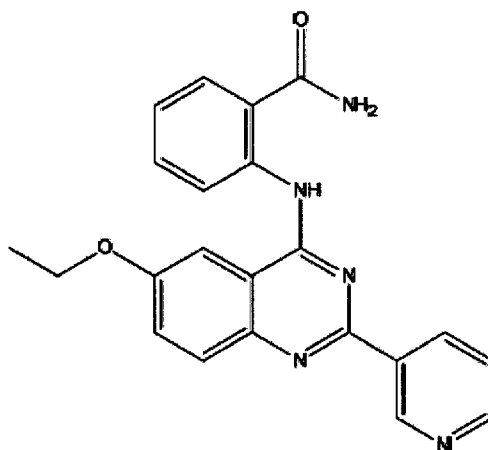
R² is -OR^d.

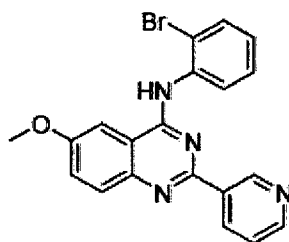
4. The compound according to claim 3 or a salt thereof, wherein

R^2 is $-OR^d$, in which R^d is C_1 - C_8 alkyl; and

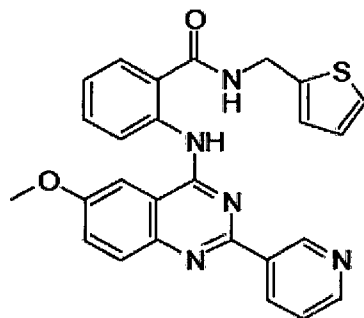
each R^b , $R^{b'}$, R^c , $R^{c'}$, R^d and R^f is independently hydrogen, C_1 - C_8 alkyl, C_2 - C_8 alkenyl, C_2 - C_8 alkynyl, C_3 - C_8 cycloalkyl, phenyl, dihydroindenyl, tetrahydropyranyl, phenylalkyl, thienylalkyl, tetrahydropyranylalkyl, dihydroindenylalkyl, or haloalkyl.

- 5
5. The compound according to claim 4 or a salt thereof, wherein
 R^1 is C_1 - C_8 alkyl, halo, haloalkyl, haloalkoxy, $-C(Y)NR^bR^{b'}$ or $-OR^d$; and
each R^b , $R^{b'}$, R^d and R^f is independently hydrogen, C_1 - C_8 alkyl, C_3 - C_8 cycloalkyl or haloalkyl.
- 10
6. The compound according to claim 1 or a salt thereof, wherein the compound of formula (I) is a compound selected from the group consisting of

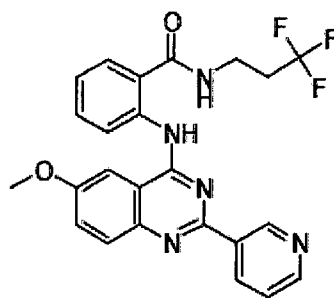




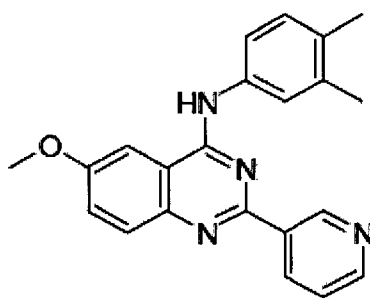
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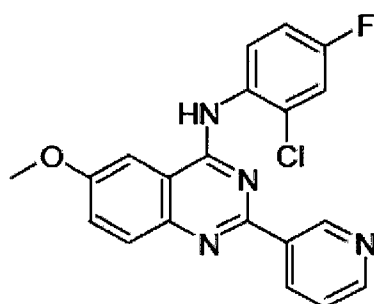
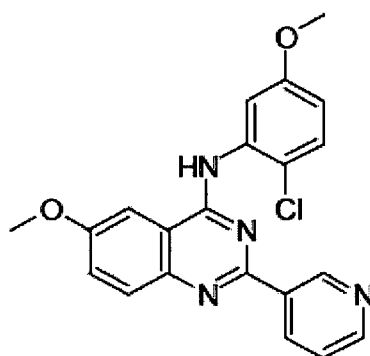
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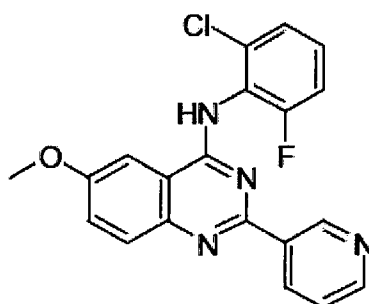
,



,



, and



7. A pharmaceutical composition comprising the compound according to any one of claims 1 to 6 or a salt thereof as an active ingredient and a pharmaceutically acceptable carrier.
8. The pharmaceutical composition according to claim 7 for use in preventing or treating central nervous system diseases.
9. The pharmaceutical composition according to claim 8 for use in treating or preventing central nervous system disorders selected from the group consisting of schizophrenia; refractory, intractable or chronic schizophrenia; emotional disturbance; psychotic disorder; mood disorder; bipolar I type disorder; bipolar II type disorder; depression; endogenous depression; major depression; melancholy and refractory depression; dysthymic disorder; cyclothymic disorder; panic attack; panic disorder; agoraphobia; social phobia; obsessive-compulsive disorder; post-traumatic stress disorder; generalized anxiety disorder; acute stress disorder; hysteria; somatization disorder; conversion disorder; pain disorder; hypochondriasis; factitious disorder; dissociative disorder; sexual dysfunction; sexual desire disorder; sexual arousal disorder; erectile dysfunction; anorexia nervosa; bulimia nervosa; sleep disorder; adjustment disorder; alcohol abuse; alcohol intoxication; drug addiction; stimulant intoxication; narcotism; anhedonia; iatrogenic anhedonia; anhedonia of a psychic or mental cause; anhedonia associated with depression; anhedonia associated with schizophrenia; delirium; cognitive impairment; cognitive impairment associated with Alzheimer's disease, Parkinson's disease and other neurodegenerative diseases; cognitive impairment caused by Alzheimer's disease; Parkinson's disease and associated neurodegenerative diseases; cognitive impairment of schizophrenia; cognitive impairment caused by refractory, intractable or chronic schizophrenia; vomiting; motion sickness; obesity; migraine;

pain (ache) ; mental retardation; autism disorder (autism); Tourette's disorder; tic disorder; attention-deficit/hyperactivity disorder; conduct disorder; and Down's syndrome.

10. A process for producing a pharmaceutical composition comprising mixing a compound or a salt thereof according to any one of claims 1 to 6 with a pharmaceutically acceptable carrier.

11. A compound or a salt thereof according to any one of claims 1 to 6 for use as a drug.

12. A compound or a salt thereof according to any one of claims 1 to 6 for use in treating schizophrenia or cognitive deficit.

13. A combination of a compound or a salt thereof according to any one of claims 1 to 6 with an additional therapeutic agent for use in treating a disorder that would benefit by the modulation of STEP.

14. The combination for use according to claim 13, wherein the additional therapeutic agent is an atypical antipsychotic.

15. The combination for use according to claim 13, wherein the additional therapeutic agent is selected from the group consisting of aripiprazole, clozapine, ziprasidone, risperidone, quetiapine, olanzapine, amisulpride, asenapine, iloperidone, melperone, paliperidone, perospirone, sertindole and sulpride.

16. The combination for use according to claim 13, wherein the additional therapeutic agent is a typical antipsychotic.

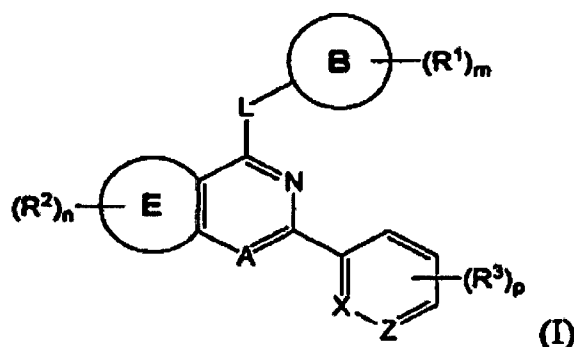
17. The combination for use according to claim 13, wherein the additional therapeutic agent is selected from the group consisting of haloperidol, molindone, loxapine, thioridazine, thiothixene, pimozide, fluphenazine, trifluoperazine, mesoridazine, chlorprothixene, chlorpromazine, perphenazine, triflupromazine and zuclopenthixol.

18. A kit comprising a composition comprising a compound or a salt thereof according to any one of claims 1 to 6 and an acceptable carrier.

19. A kit comprising a pharmaceutical composition comprising a compound or a salt thereof according to any one of claims 1 to 6 and a pharmaceutically acceptable carrier.

Patentansprüche

1. Verbindung der Formel (I):



oder ein Salz davon,
wobei:

A für CH oder N steht,

B für Phenyl, Dihydroindenyl, Dihydrobenzoxazinyl, Dihydrobenzodioxinyl, Chromenyl, Tetrahydroisochinolinyl, Tetrahydrochinolinyl, Dihydrochinolinyl, Chinolinyl, Tetrahydrochinazolinyl, Indolinyl, Dihydrobenzothiazolyl, Dihydrobenzimidazolyl, Dihydrobenzoxazolyl, Isoindolinyl, Benzofuryl, Benzothienyl, Benzodioxolyl, Indolyl, Indazolyl, Benzimidazolyl, Benzotriazolyl, Benzisoxazolyl, Benzoxazolyl, Benzothiazolyl, Benzoxadiazolyl, Tetrahydrobenzothienyl, Thienyl, Thiazolyl, Imidazolyl, Pyrazolyl, Thiadiazolyl oder Pyridyl steht,

m für 1, 2, 3 oder 4 steht,

E für Phenyl, Thienyl oder Pyrrolyl steht,

wenn E für Phenyl steht, n für 1 oder 2 steht, und wenn E für Thienyl oder Pyrrolyl steht, n für 0 oder 1 steht,

L für NR^5 steht,

einer der Reste X und Z für N und der andere für CH steht,

p für 0, 1 oder 2 steht,

R^1 für $\text{C}_1\text{-C}_8\text{-Alkyl}$, Halogen, Halogenalkyl, Halogenalkoxy, Alkoxyalkyl, -CN, Oxo, $-\text{NO}_2$, $-\text{C}(\text{Y})\text{NR}^b\text{R}^{b'}$, $-\text{NR}^c\text{C}(\text{Y})\text{R}^c$, $-\text{SO}_2\text{NR}^b\text{R}^{b'}$, $-\text{OR}^d$ oder $-\text{S}(\text{O})_q\text{R}^f$ steht,

R^2 für $\text{C}_1\text{-C}_8\text{-Alkyl}$, $\text{C}_3\text{-C}_8\text{-Cycloalkyl}$, Thiomorpholinylalkyl, Pyrrolidinylalkyl, Morpholinylalkyl, Halogenalkyl, Halogenalkoxy, Aminoalkyl, Alkylaminoalkyl, Dialkylaminoalkyl, Hydroxyalkyl, Alkoxyalkyl, Silyloxyalkyl, -CN, $-\text{NO}_2$, $-\text{C}(\text{O})\text{OR}^a$, $-\text{C}(\text{Y})\text{NR}^b\text{R}^{b'}$, $-\text{NR}^b\text{R}^{b'}$, $-\text{OR}^d$, $-\text{C}(\text{Y})\text{R}^e$ oder $-\text{S}(\text{O})_q\text{R}^f$,

R^3 für $\text{C}_1\text{-C}_8\text{-Alkyl}$, Halogen, Halogenalkyl, $-\text{NR}^b\text{R}^{b'}$ oder $-\text{OR}^d$ steht,

R^5 für Wasserstoff steht,

Y für O oder S steht,

q für 1 oder 2 steht und

R^a , R^b , $\text{R}^{b'}$, R^c , R^d , R^e und R^f jeweils unabhängig voneinander für Wasserstoff, $\text{C}_1\text{-C}_8\text{-Alkyl}$, $\text{C}_2\text{-C}_8\text{-Alkenyl}$, $\text{C}_2\text{-C}_8\text{-Alkinyl}$, $\text{C}_3\text{-C}_8\text{-Cycloalkyl}$, Phenyl, Dihydroindenyl, Morpholinyl, Tetrahydropyranyl, Piperidyl, Pyrrolidinyl, Thiomorpholinyl, Phenylalkyl, Thienylalkyl, Pyridylalkyl, Tetrahydropyranylalkyl, Dihydroindenylalkyl, Tetrahydrofurylalkyl, Hydroxyalkyl, Morpholinylalkyl, Pyrrolidinylalkyl, Dialkylaminoalkyl, Piperidylalkyl, Benzodioxolylalkyl, Dihydrobenzodioxinylalkyl, $\text{C}_3\text{-C}_8\text{-Cycloalkylalkyl}$, Halogenalkyl oder Alkoxyalkyl stehen.

2. Verbindung nach Anspruch 1 oder ein Salz davon, wobei

A für N steht,

B für Phenyl oder Pyridyl steht,

m für 1, 2 oder 3 steht,

E für Phenyl steht,

n für 1 oder 2 steht und

p für 0 oder 1 steht.

3. Verbindung nach Anspruch 2 oder ein Salz davon, wobei

m für 1 oder 2 steht,

p für 0 steht,

R^1 für $\text{C}_1\text{-C}_8\text{-Alkyl}$, Halogen, Halogenalkyl, Halogenalkoxy, -CN, $-\text{NO}_2$, $-\text{C}(\text{Y})\text{NR}^b\text{R}^{b'}$, $-\text{NR}^c\text{C}(\text{Y})\text{R}^c$, $-\text{SO}_2\text{NR}^b\text{R}^{b'}$, $-\text{OR}^d$ oder $-\text{S}(\text{O})_q\text{R}^f$ steht und

R^2 für $-\text{OR}^d$ steht.

4. Verbindung nach Anspruch 3 oder ein Salz davon, wobei

R^2 für $-\text{OR}^d$ steht, wobei R^d für $\text{C}_1\text{-C}_8\text{-Alkyl}$ steht, und

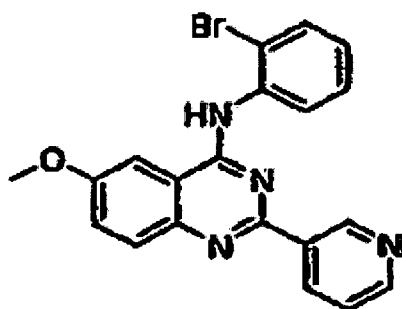
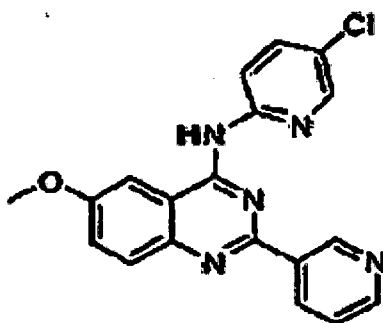
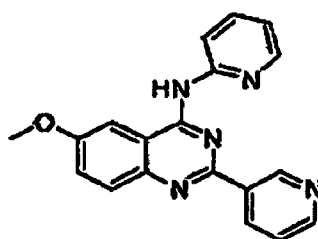
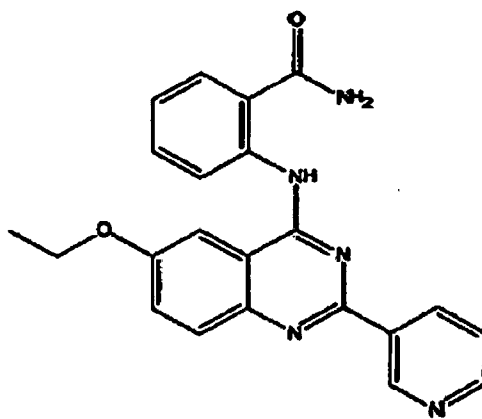
R^b , $\text{R}^{b'}$, R^c , R^d und R^f jeweils unabhängig voneinander für Wasserstoff, $\text{C}_1\text{-C}_8\text{-Alkyl}$, $\text{C}_2\text{-C}_8\text{-Alkenyl}$, $\text{C}_2\text{-C}_8\text{-Alkinyl}$, $\text{C}_3\text{-C}_8\text{-Cycloalkyl}$, Phenyl, Dihydroindenyl, Tetrahydropyranyl, Phenylalkyl, Thienylalkyl, Tetrahydropyranylalkyl, Dihydroindenylalkyl oder Halogenalkyl stehen.

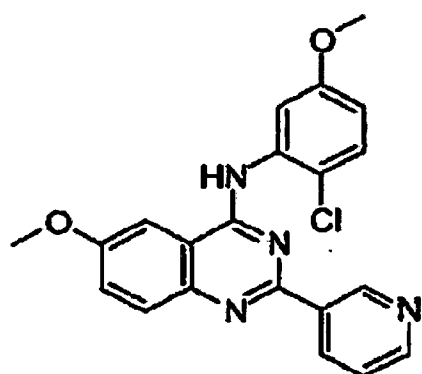
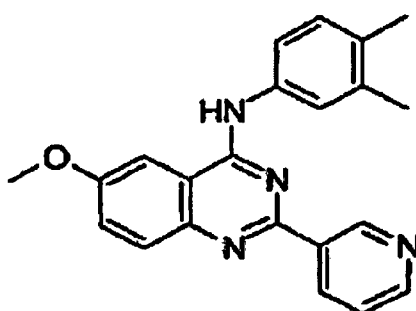
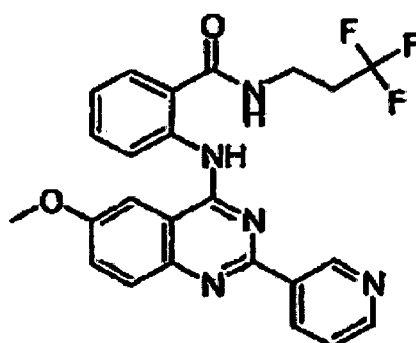
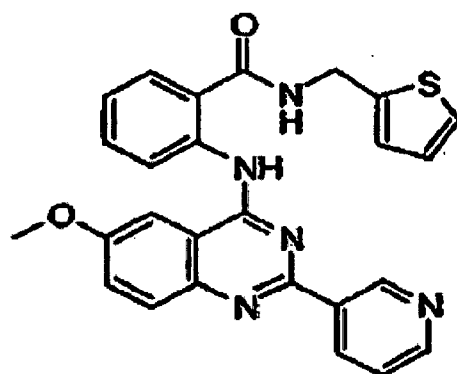
5. Verbindung nach Anspruch 4 oder ein Salz davon, wobei

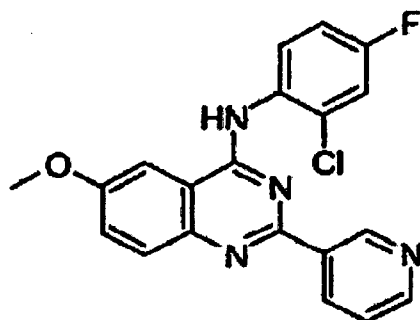
R^1 für $\text{C}_1\text{-C}_8\text{-Alkyl}$, Halogen, Halogenalkyl, Halogenalkoxy, $-\text{C}(\text{Y})\text{NR}^b\text{R}^{b'}$ oder $-\text{OR}^d$ steht und

R^b , $\text{R}^{b'}$, R^d und R^f jeweils unabhängig voneinander für Wasserstoff, $\text{C}_1\text{-C}_8\text{-Alkyl}$, $\text{C}_3\text{-C}_8\text{-Cycloalkyl}$ oder Halogenalkyl stehen.

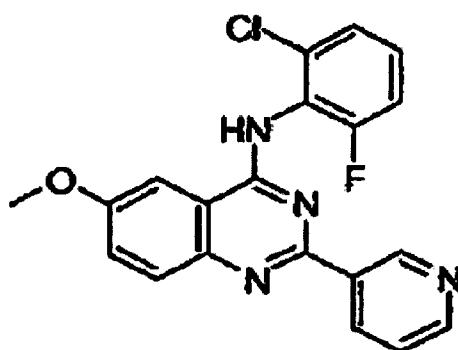
6. Verbindung nach Anspruch 1 oder ein Salz davon, wobei es sich bei der Verbindung der Formel (I) um eine Verbindung ausgewählt aus der Gruppe bestehend aus







und



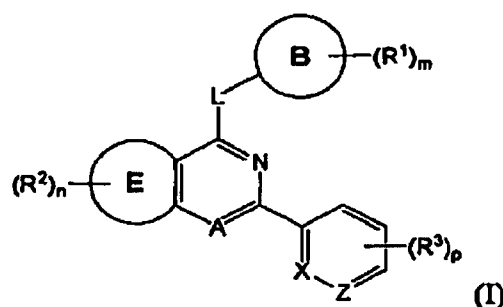
handelt.

7. Pharmazeutische Zusammensetzung, umfassend die Verbindung nach einem der Ansprüche 1 bis 6 oder ein Salz davon als Wirkstoff und einen pharmazeutisch unbedenklichen Träger.
8. Pharmazeutische Zusammensetzung nach Anspruch 7 zur Verwendung bei der Prävention oder Behandlung von Krankheiten des zentralen Nervensystems.
9. Pharmazeutische Zusammensetzung nach Anspruch 8 zur Verwendung bei der Behandlung oder Prävention von Erkrankungen des zentralen Nervensystems ausgewählt aus der Gruppe bestehend aus Schizophrenie, refraktärer, therapieresistenter oder chronischer Schizophrenie, emotioneller Störung, psychotischer Erkrankung, affektiver Störung, bipolarer Störung vom Typ I, bipolarer Störung vom Typ II, Depression, endogener Depression, schwerer Depression, Melancholie und refraktärer Depression, dysthymischer Erkrankung, cyclothymischer Erkrankung, Panikanfall, Panikerkrankung, Agoraphobie, sozialer Phobie, Zwangsneurose, posttraumatischer Belastungsstörung, generalisierter Angststörung, akuter Stressstörung, Hysterie, Somatisierungsstörung, Konversionsstörung, Schmerzstörung, Hypochondriasie, artifizieller Störung, dissoziativer Störung, sexueller Dysfunktion, sexueller Apenzenzstörung, sexueller Erregungsstörung, erektiler Dysfunktion, Anorexia nervosa, Bulimia nervosa, Schlafstörung, Anpassungsstörung, Alkoholmissbrauch, Alkoholrausch, Drogensucht, Stimulanzienrausch, Narkotismus, Anhängdonie, iatrogene Anhängdonie, Anhängdonie psychischer oder geistiger Ursache, mit Depression assoziierter Anhängdonie, mit Schizophrenie assoziierter Anhängdonie, Delirium, kognitiver Störung, mit Alzheimer-Krankheit, Parkinson-Krankheit und anderen neurodegenerativen Krankheiten assoziierter kognitiver Störung, durch Alzheimer-Krankheit verursachter kognitiver Störung, Parkinson-Krankheit und assoziierten neurodegenerativen Krankheiten, kognitiver Störung bei Schizophrenie, durch refraktäre, therapieresistente oder chronische Schizophrenie verursachter kognitiver Störung, Erbrechen, Kinetose, Obesitas, Migräne, Schmerzen, geistiger Retardierung, Autismusstörung (Autismus), Tourette-Syndrom, Tic-Störung, Aufmerksamkeitsdefizit-/Hyperaktivitätsstörung, Verhaltensstörung und Down-Syndrom.
10. Verfahren zur Herstellung einer pharmazeutischen Zusammensetzung, bei dem man eine Verbindung oder ein Salz davon nach einem der Ansprüche 1 bis 6 mit einem pharmazeutisch unbedenklichen Träger mischt.
11. Verbindung oder Salz davon nach einem der Ansprüche 1 bis 6 zur Verwendung als Arzneimittel.

12. Verbindung oder Salz davon nach einem der Ansprüche 1 bis 6 zur Verwendung bei der Behandlung von Schizophrenie oder kognitivem Defizit.
13. Kombination einer Verbindung oder eines Salzes davon nach einem der Ansprüche 1 bis 6 mit einem zusätzlichen Therapeutikum zur Verwendung bei der Behandlung einer Erkrankung, bei der die Modulation von STEP von Nutzen sein würde.
14. Kombination zur Verwendung nach Anspruch 13, wobei es sich bei dem zusätzlichen Therapeutikum um ein atypisches Antipsychotikum handelt.
15. Kombination zur Verwendung nach Anspruch 13, wobei das zusätzliche Therapeutikum aus der aus Aripiprazol, Clozapin, Ziprasidon, Risperidon, Quetiapin, Olanzapin, Amisulprid, Asenapin, Iloperidon, Melperon, Paliperidon, Perospiron, Sertindol und Sulprid bestehenden Gruppe ausgewählt ist.
16. Kombination zur Verwendung nach Anspruch 13, wobei es sich bei dem zusätzlichen Therapeutikum um ein typisches Antipsychotikum handelt.
17. Kombination zur Verwendung nach Anspruch 13, wobei das zusätzliche Therapeutikum aus der aus Haloperidol, Molindon, Loxapin, Thioridazin, Thiothixen, Pimozid, Fluphenazin, Trifluoperazin, Mesoridazin, Chlorprothixen, Chlorpromazin, Perphenazin, Triflupromazin und Zuclopenthixol bestehenden Gruppe ausgewählt ist.
18. Kit, umfassend eine Zusammensetzung umfassend eine Verbindung oder ein Salz davon nach einem der Ansprüche 1 bis 6 und einen unbedenklichen Träger.
19. Kit, umfassend eine pharmazeutische Zusammensetzung umfassend eine Verbindung oder ein Salz davon nach einem der Ansprüche 1 bis 6 und einen pharmazeutisch unbedenklichen Träger.

Revendications

1. Composé de formule (I)



ou un sel correspondant,
dans laquelle :

A représente CH ou N ;

B représente phényle, dihydro-indényle, dihydrobenzoxazinyle, dihydrobenzodioxinyle, chroményle, tétrahydro-isoquinoléyle, tétrahydroquinoléinyle, dihydroquinoléyle, quinoléyle, tétrahydroquinazolinyle, indolinyle, dihydrobenzothiazolyle, dihydrobenzimidazolyle, dihydrobenzoxazolyle, iso-indolinyle, benzofuryle, benzothiényne, benzodioxolyle, indolyle, indazolyle, benzo-imidazolyle, benzotriazolyle, benzisoxazolyle, benzoxazolyle, benzothiazolyle, benzoxadiazolyle, tétrahydrobenzothiényne, thiényne, thiazolyle, imidazolyle, pyrazolyle, thiadiazolyle ou pyridyle ;

m vaut 1, 2, 3 ou 4 ;

E représente phényle, thiényne ou pyrrolyle ; lorsque E représente phényle, n vaut 1 ou 2 ; et lorsque E représente thiényne ou pyrrolyle, n vaut 0 ou 1 ;

L représente NR⁵ ;

l'un parmi X et Z représente N et l'autre représente CH ;

p vaut 0, 1 ou 2 ;

R¹ représente C₁-C₈-alkyle, halogéno, halogénoalkyle, halogénoalcoxy, alcoxyalkyle, -CN, oxo, -NO₂, -C(Y)NR^bR^{b'}, -NR^cC(Y)R^{c'}, -SO₂NR^bR^{b'}, -OR^d ou -S(O)_qR^f ;

R² représente C₁-C₈-alkyle, C₃-C₈-cycloalkyle, thiomorpholinylalkyle, pyrrolidinylalkyle, morpholinylalkyle, halogénoalkyle, halogénoalcoxy, aminoalkyle, alkylaminoalkyle, dialkylaminoalkyle, hydroxyalkyle, alcoxyalkyle, silyloxyalkyle, -CN, -NO₂, -C(O)OR^a, -C(Y)NR^bR^{b'}, -N R^bR^{b'}, -OR^d, -C(Y)R^e ou -S(O)_qR^f ;

R³ représente C₁-C₈-alkyle, halogéno, halogénoalkyle, -NR^bR^{b'} ou -OR^d ;

R⁵ représente hydrogène ;

Y représente O ou S ;

q vaut 1 ou 2 et

chaque R^a, R^b, R^{b'}, R^c, R^{c'}, R^d, R^e et R^f représente, indépendamment, hydrogène, C₁-C₈-alkyle, C₂-C₈-alcényle, C₂-C₈-alcynyle, C₃-C₈-cycloalkyle, phényle, dihydro-indényle, morpholinyle, tétrahydropyrannyle, pipéridyle, pyrrolidinyle, thiomorpholinyle, phénylalkyle, thiénylalkyle, pyridylalkyle, tétrahydropyrannylalkyle, dihydro-indénylalkyle, tétrahydrofurylalkyle, hydroxyalkyle, morpholinylalkyle, pyrrolidinylalkyle, dialkylaminoalkyle, pipéridylalkyle, benzodioxolylalkyle, dihydrobenzodioxinylalkyle, C₃-C₈-cycloalkylalkyle, halogénoalkyle ou alcoxyalkyle.

2. Composé selon la revendication 1, ou un sel correspondant,

A représentant N ;

B représentant phényle ou pyridyle ;

m valant 1, 2 ou 3 ;

E représentant phényle ;

n valant 1 ou 2 ; et

p valant 0 ou 1.

3. Composé selon la revendication 2, ou un sel correspondant,

m valant 1 ou 2 ;

p valant 0 ;

R¹ représentant C₁-C₈-alkyle, halogéno, halogénoalkyle, halogénoalcoxy, -CN, -NO₂, -C(Y)NR^bR^{b'}, -NR^cC(Y)R^{c'}, -SO₂NR^bR^{b'}, -OR^d ou -S(O)_qR^f ; et

R² valant -OR^d.

4. Composé selon la revendication 3, ou un sel correspondant,

R² représentant -OR^d, dans lequel R^d représente C₁-C₈-alkyle ; et

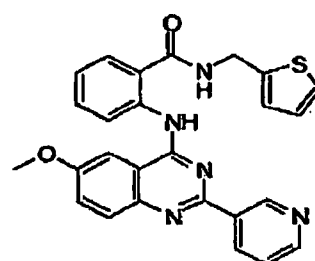
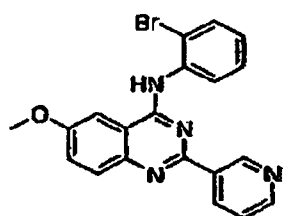
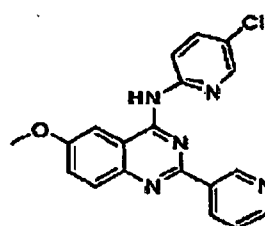
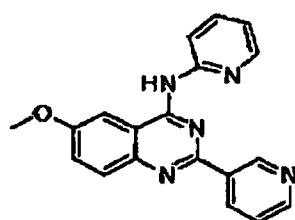
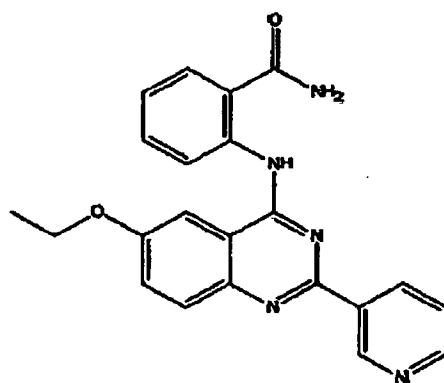
chaque R^b, R^{b'}, R^c, R^{c'}, R^d et R^f représentant, indépendamment, hydrogène, C₁-C₈-alkyle, C₂-C₈-alcényle, C₂-C₈-alcynyle, C₃-C₈-cycloalkyle, phényle, dihydro-indényle, tétrahydropyrannyle, phénylalkyle, thiénylalkyle, tétrahydropyrannylalkyle, dihydro-indénylalkyle ou halogénoalkyle.

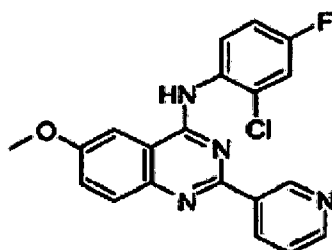
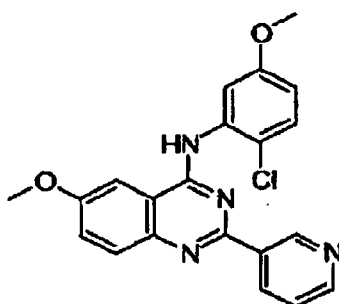
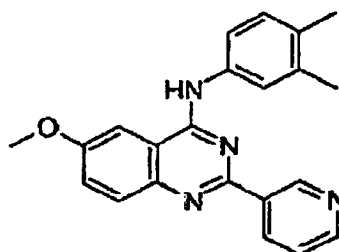
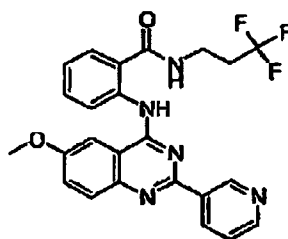
5. Composé selon la revendication 4, ou un sel correspondant,

R¹ représentant C₁-C₈-alkyle, halogéno, halogénoalkyle, halogénoalcoxy, -C(Y)NR^bR^{b'} ou -OR^d ; et

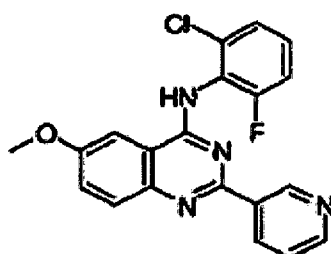
chaque R^b, R^{b'}, R^d et R^f représentant, indépendamment, hydrogène, C₁-C₈-alkyle, C₃-C₈-cycloalkyle ou halogénoalkyle.

6. Composé selon la revendication 1 ou un sel correspondant, le composé de formule (I) étant un composé choisi dans le groupe constitué par :





et



7. Composition pharmaceutique comprenant le composé selon l'une quelconque des revendications 1 à 6, ou un sel correspondant, comme principe actif et un support pharmaceutiquement acceptable.
8. Composition pharmaceutique selon la revendication 7 pour une utilisation dans la prévention ou le traitement de

maladies du système nerveux central.

- 5 9. Composition pharmaceutique selon la revendication 8 pour une utilisation dans le traitement ou la prévention de troubles du système nerveux central, choisis dans le groupe constitué par la schizophrénie ; la schizophrénie réfractaire, résistante ou chronique ; le trouble émotionnel ; le trouble psychotique ; le trouble de l'humeur ; le trouble bipolaire de type I ; le trouble bipolaire de type II ; la dépression ; la dépression endogène ; la dépression majeure ; la dépression mélancolique et réfractaire ; le trouble dysthymique ; le trouble cyclothymique ; l'attaque de panique ; le trouble panique ; l'agoraphobie ; la phobie sociale ; le trouble obsessionnel compulsif ; le trouble du stress post-traumatique ; le trouble de l'anxiété généralisée ; le trouble du stress aigu ; l'hystérie ; le trouble de somatisation ;

10 le trouble de conversion ; le trouble douloureux ; l'hypochondrie ; le trouble factice ; le trouble dissociatif ; la dysfonction sexuelle ; le trouble du désir sexuel ; le trouble de l'excitation sexuelle ; la dysfonction érectile ; l'anorexie mentale ; la boulimie ; le trouble du sommeil ; le trouble de l'adaptation ; l'abus d'alcool ; l'intoxication alcoolique ; la toxicomanie ; l'intoxication par un stimulant ; le narcotisme ; l'anhédonie ; l'anhédonie iatrogène ; l'anhédonie due à une cause psychique ou mentale ; l'anhédonie associée à une dépression ; l'anhédonie associée à la

15 schizophrénie ; le délire ; la déficience cognitive ; la déficience cognitive associée à la maladie d'Alzheimer, à la maladie de Parkinson et à d'autres maladies neurodégénératives ; la déficience cognitive provoquée par la maladie d'Alzheimer ; la maladie de Parkinson et les maladies neurodégénératives associées ; la déficience cognitive de la schizophrénie ; la déficience cognitive provoquée par une schizophrénie réfractaire, résistante ou chronique ; le vomissement ; le mal des transports ; l'obésité ; la migraine ; la douleur ; le retard mental ; le trouble de l'autisme (autisme) ; le trouble de Tourette ; le trouble de tic ; le trouble du déficit d'attention/d'hyperactivité ; le trouble des conduites ; et le syndrome de Down.
- 25 10. Procédé pour produire une composition pharmaceutique comprenant le mélange d'un composé, ou d'un sel correspondant, selon l'une quelconque des revendications 1 à 6 avec un support pharmaceutiquement acceptable.
11. Composé, ou sel correspondant, selon l'une quelconque des revendications 1 à 6 pour une utilisation comme médicament.
- 30 12. Composé, ou sel correspondant, selon l'une quelconque des revendications 1 à 6 pour une utilisation dans le traitement de la schizophrénie ou d'une déficience cognitive.
- 35 13. Combinaison d'un composé, ou d'un sel correspondant, selon l'une quelconque des revendications 1 à 6 avec un agent thérapeutique supplémentaire pour une utilisation dans le traitement d'un trouble qui tirerait un avantage de la modulation de la STEP.
14. Combinaison pour une utilisation selon la revendication 13, l'agent thérapeutique supplémentaire étant un antipsychotique atypique.
- 40 15. Combinaison pour une utilisation selon la revendication 13, l'agent thérapeutique supplémentaire étant choisi dans le groupe constitué par l'aripiprazole, la clozapine, la ziprasidone, la rispéridone, la quétiapine, l'olanzapine, l'amisulpride, l'asénapine, l'ilopéridone, la melpérone, la palipéridone, la pérospirone, la sertindole et le sulpride.
- 45 16. Combinaison pour une utilisation selon la revendication 13, l'agent thérapeutique supplémentaire étant un antipsychotique typique.
- 50 17. Combinaison pour une utilisation selon la revendication 13, l'agent thérapeutique supplémentaire étant choisi dans le groupe constitué par l'halopéridol, la molindone, la loxapine, la thioridazine, le thiothixène, le pimozide, la fluphénazine, la trifluopérazine, la mésoridazine, le chlorprothixène, la chlorpromazine, la perphénazine, la triflupromazine et le zuclopenthixol.
18. Kit comprenant une composition comprenant un composé, ou un sel correspondant, selon l'une quelconque des revendications 1 à 6 et un support acceptable.
- 55 19. Kit comprenant une composition pharmaceutique comprenant un composé, ou un sel correspondant, selon l'une quelconque des revendications 1 à 6 et un support pharmaceutiquement acceptable.

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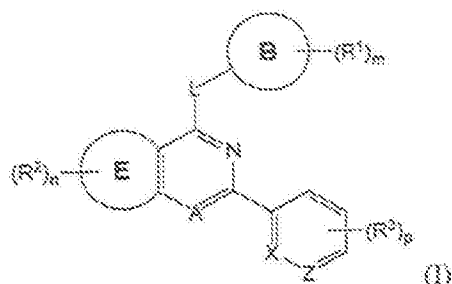
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TERÁPIÁS VEGYÜLETEK ÉS KAPCSOLÓDÓ ALKALMAZÁSI ELJÁRÁSOK

Szabadalmi igénypontok

1. (I) képletű vegyület:



vagy sója, ahol:

A jelentése CH vagy N;

B jelentése fenil, dihidroindenil, dihidrobenzoxazinil, dihidrobenzodioxinil, kromenil, tetrahidroizokinolil, tetrahidrokinolinil, dihidrokinolil, kinolil, tetrahidrokinazolinil, indolinil, dihidrobenzotiazolil, dihidrobenzimidazolil, dihidrobenzoxazolil, izoindolinil, benzofuril, benzotienil, benzodioxolil, indolil, indazolil, benzoimidazolil, benzotriazolil, benzizoxazolil, benzoxazolil, benzotiazolil, benzoxadiazolil, tetrahidrobenzotienil, tienil, tiazolil, imidazolil, pirazolil, tiadiazolil vagy piridil;

m jelentése 1, 2, 3, vagy 4;

E jelentése fenil, tienil vagy pirrolil;

ha E jelentése fenil, n értéke 1 vagy 2; és ha E jelentése tienil vagy pirrolil, n értéke 0 vagy 1;

L jelentése NR^1 ;

X és Z közül az egyiknek a jelentése N és a másiknak a jelentése CH;

p értéke 0, 1, vagy 2;

R^1 jelentése C_1-C_8 alkil, halo, haloalkil, haloalkoxi, alkoxialkil, -CN, oxo, $-NO_2$, $-C(Y)NR^bR^v$, $-NR^cC(Y)R^e$, $-SO_2NR^bR^v$, $-OR^d$, vagy $-S(O)_qR^f$;

R^2 jelentése C_1-C_8 alkil, C_3-C_8 cikloalkil, tiomorfolinilalkil, pirrolidinilalkil, morfolinilalkil, haloalkil, haloalkoxi, aminoalkil, alkilaminoalkil, dialkilaminoalkil, hidroxialkil, alkoxialkil, szililoxialkil, -CN, $-NO_2$, $-C(O)OR^a$, $-C(Y)NR^bR^v$, $-NR^bR^v$, $-OR^d$, $-C(Y)R^e$ vagy $-S(O)_qR^f$;

R^3 jelentése C_1-C_8 alkil, halo, haloalkil, $-NR^bR^v$ vagy $-OR^d$;

R^5 jelentése hidrogén;

Y jelentése O vagy S;

q értéke 1 vagy 2 és

R^a , R^b , R^v , R^c , R^d , R^e és R^f mindegyikének jelentése egymástól függetlenül hidrogén, C_1-C_8 alkil, C_2-C_8 alkenil, C_2-C_8 alkinil, C_3-C_8 cikloalkil, fenil, dihidroindenil, morfolinil, tetrahidropiranil, piperidil, pirrolidinil, tiomorfolinil, fenilalkil, tienilalkil, piridilalkil, tetrahidropiranilalkil, dihidroindenilalkil, tetrahidrofurilalkil, hidroxialkil, morfolinilalkil, pirrolidinilalkil, dialkilaminoalkil, piperidilalkil, benzodioxolilalkil, dihidrobenzodioxinilalkil, C_3-C_8 cikloalkilalkil, haloalkil vagy alkoxialkil.

2. Az 1. igénypont szerinti vegyület vagy sója, ahol

- A jelentése N;
 B jelentése fenil vagy piridil;
 m értéke 1, 2 vagy 3;
 E jelentése fenil;
 n értéke 1 vagy 2; and
 p értéke 0 vagy 1.

3. A 2. igénypont szerinti vegyület vagy sója, ahol

- m értéke 1 vagy 2;
 p értéke 0;
 R^1 jelentése C_1-C_8 alkil, halo, haloalkil, haloalkoxi, $-CN$, $-NO_2$, $-C(Y)NR^bR^v$, $-NR^cC(Y)R^v$, $-SO_2NR^bR^v$, $-OR^d$, vagy $-S(O)_pR^e$; és
 R^2 jelentése $-OR^d$.

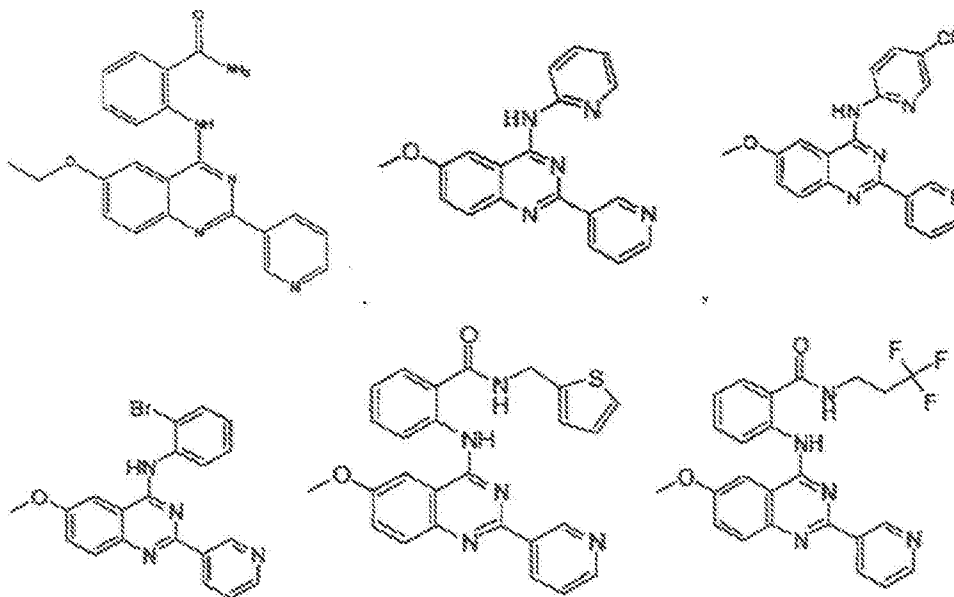
4. A 3. igénypont szerinti vegyület vagy sója, ahol

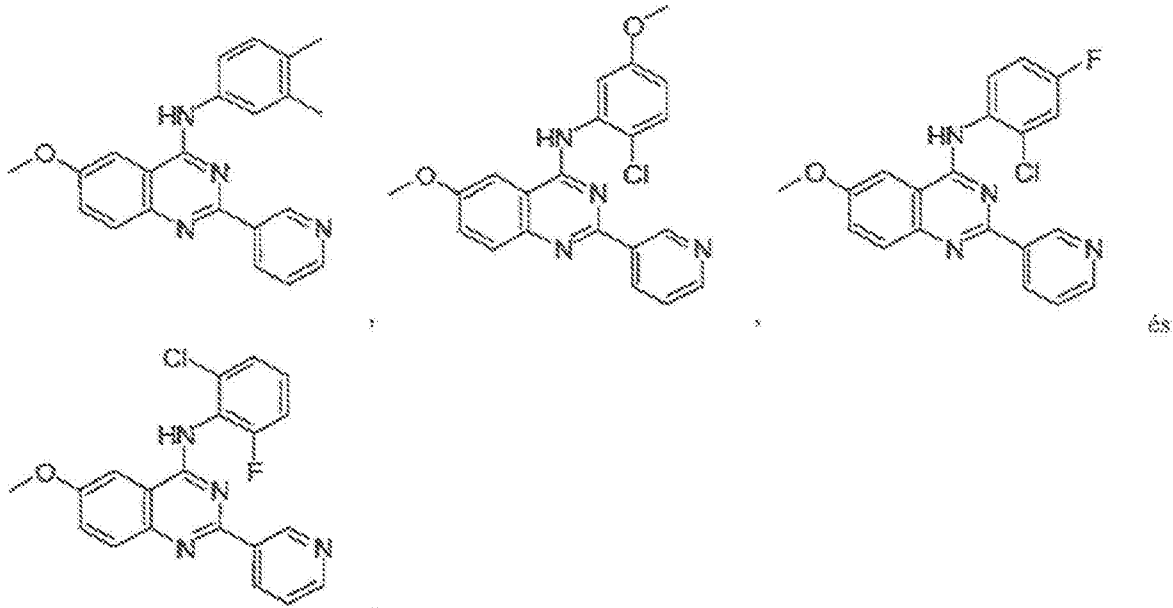
- R^2 jelentése $-OR^d$, ahol R^d jelentése C_1-C_8 alkil; és
 R^b , R^v , R^c , R^e , R^d és R^f mindegyikének jelentése egymástól függetlenül hidrogén, C_1-C_8 alkil, C_2-C_8 alkenil, C_2-C_8 alkinil, C_3-C_8 cikloalkil, fenil, dihidroindenil, tetrahidropiranil, fenilalkil, tienilalkil, tetrahidropiranilalkil, dihidroindenilalkil vagy haloalkil.

5. A 4. igénypont szerinti vegyület vagy sója, ahol

- R^1 jelentése C_1-C_8 alkil, halo, haloalkil, haloalkoxi, $-C(Y)NR^bR^v$ vagy $-OR^d$; és
 R^b , R^v , R^c és R^f mindegyikének jelentése egymástól függetlenül hidrogén, C_1-C_8 alkil, C_3-C_8 cikloalkil vagy haloalkil.

6. Az 1. igénypont szerinti vegyület vagy sója, ahol az (I) képletű vegyület a következők alkotta vegyületek közül választott:





7. Gyógyászati készítmény, amely az 1-6 igénypontok bármelyike szerinti készítményt vagy sóját tartalmazza hatóanyagként, valamint gyógyászatilag elfogadható hordozót tartalmaz.

8. A 7. igénypont szerinti gyógyászati készítmény központi idegrendszeri betegség megelőzésében vagy kezelésében történő alkalmazásra.

9. A 7. igénypont szerinti gyógyászati készítmény központi idegrendszeri betegség megelőzésében vagy kezelésében történő alkalmazásra, amely betegség a következők alkotta csoportból választott: skizofrénia; kezelésnek ellenálló (refrakter), makacs vagy krónikus skizofrénia; érzelmi zavar; pszichotikus zavar; hangulatzavar; I-es típusú bipoláris rendellenesség; II-es típusú bipoláris rendellenesség; depresszió; endogén depresszió; major depresszió; melankólia és kezelésnek ellenálló depresszió; disztímiás zavar; ciklotímiás zavar; pánikroham; pánikbetegség; agorafóbia; szociális fóbia; obszesszív-kompulzív betegség; poszt-traumás stressz-betegség; generalizált szorongásos zavar; akut stresszbetegség; hisztéria; szomatizációs zavar; konverziós betegség; fájdalombetegség; hypochondria; hamis (factitious) zavar; disszociatív betegség; szexuális diszfunkció; szexuális vágy rendellenessége; szexuális izgalmi rendellenesség; erekciós zavar; anorexia nervosa; bulimia nervosa; alvászavar; alkalmazkodási rendellenesség; alkohollal való visszaélés; alkoholemérgezés; drogfüggőség; stimulánsmérgezés; narkotizmus; anhedónia; iatrogén anhedónia; pszichikus vagy agyi eredetű anhedonia; depresszióval összefüggő anhedónia; skizofréniával összefüggő anhedónia; delírium; gondolkozási zavar; Alzheimer-betegséggel, Parkinson-betegséggel és más neurodegeneratív betegségekkel összefüggő kognitív zavar; Alzheimer-betegség, Parkinson-betegség és más neurodegeneratív betegségek okozta kognitív zavar, refrakter, makacs vagy krónikus skizofrénia okozta kognitív rendellenesség; hányás; mozgási betegség; elhízás; migrén; fájdalom; mentális retardáció; autizmus betegség (autizmus); Tourette-betegség; tik-betegség; figyelemzavar/hiperaktivitás betegség; viselkedési rendellenesség; és Down-szindróma.

10. Eljárás gyógyászati készítmény előállítására, amely tartalmazza az 1-6. igénypontok bármelyike szerinti vegyület vagy sója összekeverését gyógyászatilag elfogadható hordozóval.

11. Az 1-6. igénypontok bármelyike szerinti vegyület vagy sója gyógyszerként történő alkalmazásra.

12. Az 1-6. igénypontok bármelyike szerinti vegyület vagy sója skizofrénia vagy kognitív zavar kezelésében történő alkalmazásra.

13. Az 1-6. igénypontok bármelyike szerinti vegyület vagy sója további terápiás ágenssel alkotott kombinációja olyan betegség kezelésében történő alkalmazásra, amelyre STEP modulálása jótékony hatással bír.

14. A 13. igénypont szerinti kombináció az ott meghatározott alkalmazásra, ahol a további terápiás ágens atípusos antipszichotikum.

15. A 13. igénypont szerinti kombináció az ott meghatározott alkalmazásra, ahol a további terápiás ágens a következők alkotta csoportból választott: aripiprazol, klozapin, ziprazidon, risperidon, quetiapin, olanzapin, amiszulprid, asenapin, iloperidon, melperon, paliperidon, perospiron, szertindol és szulprid.

16. A 13. igénypont szerinti kombináció az ott meghatározott alkalmazásra, ahol a további terápiás ágens típusos antipszichotikum.

17. A 13. igénypont szerinti kombináció az ott meghatározott alkalmazásra, ahol a további terápiás ágens a következők alkotta csoportból választott: haloperidol, molindon, loxapin, tioridazin, tiotixén, pimozid, flufenazin, trifluoperazin, mezoridazin, klórprotixén, klórpromazin, perfenazin, triflupromazin és zuklopentixol.

18. Az 1-6. igénypontok bármelyike szerinti vegyületet vagy sóját tartalmazó készítményt és elfogadható hordozót tartalmazó készlet.

19. Az 1-6. igénypontok bármelyike szerinti vegyületet vagy sóját tartalmazó gyógyászati készítményt és gyógyászati lag elfogadható hordozót tartalmazó készlet.