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(54) Title: ARYL-SUBSTITUTED HETEROCYCLIC UREA MODULATORS OF FATTY ACID AMIDE HYDROLASE

(57) Abstract: Certain aryl-substituted heterocyclic urea compounds are described, which are useful as FAAH inhibitors. Such compounds may be used in pharmaceutical compositions and methods for the treatment of disease states, disorders, and conditions mediated by fatty acid amide hydrolase (FAAH) activity, such as anxiety, pain, inflammation, sleep disorders, eating disorders, energy metabolism disorders, and movement disorders (e.g., multiple sclerosis).



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5 **ARYL-SUBSTITUTED HETEROCYCLIC UREA MODULATORS OF FATTY
ACID AMIDE HYDROLASE**

Cross Reference to Related Application

This application claims the benefit of US provisional patent application
10 serial number 61/184,606, filed June 05, 2009.

Field of the Invention

The present invention relates to certain aryl-substituted heterocyclic
urea compounds, pharmaceutical compositions containing them, and methods
15 of using them for the treatment of disease states, disorders, and conditions
mediated by fatty acid amide hydrolase (FAAH) activity.

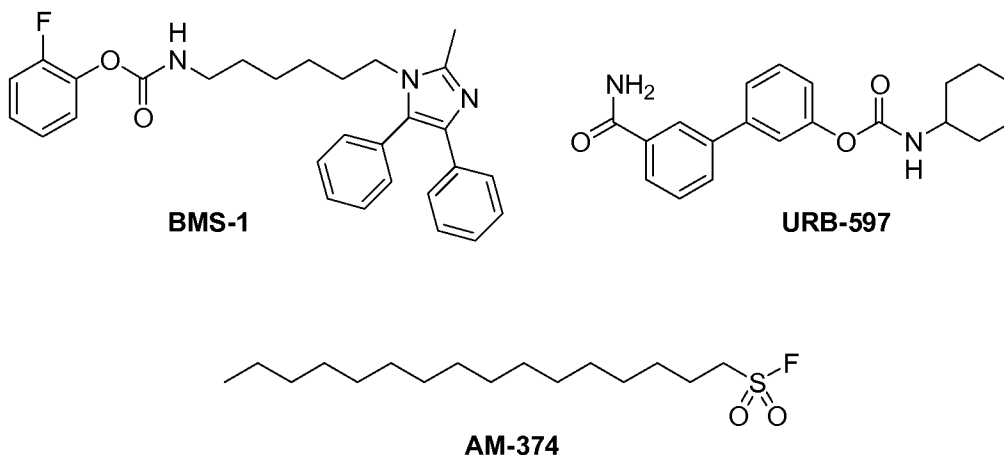
Background of the Invention

Medicinal benefits have been attributed to the cannabis plant for
20 centuries. The primary bioactive constituent of cannabis is Δ^9 -tetrahydro-
cannabinol (THC). The discovery of THC eventually led to the identification of
two endogenous cannabinoid receptors responsible for its pharmacological
actions, namely CB₁ and CB₂ (Goya, *Exp. Opin. Ther. Patents* 2000, 10,
1529). These discoveries not only established the site of action of THC, but
25 also inspired inquiries into the endogenous agonists of these receptors, or
“endocannabinoids”. The first endocannabinoid identified was the fatty acid
amide anandamide (AEA). AEA itself elicits many of the pharmacological
effects of exogenous cannabinoids (Piomelli, *Nat. Rev. Neurosci.* 2003, 4(11),
873).

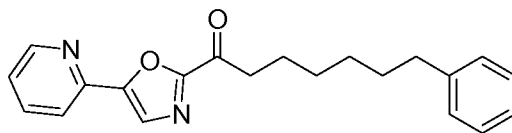
30 The catabolism of AEA is primarily attributable to the integral
membrane bound protein fatty acid amide hydrolase (FAAH), which
hydrolyzes AEA to arachidonic acid. FAAH was characterized in 1996 by
Cravatt and co-workers (Cravatt, *Nature* 1996, 384, 83). It was subsequently
determined that FAAH is additionally responsible for the catabolism of a large
35 number of important lipid signaling fatty acid amides including: another major
endocannabinoid, 2-arachidonoylglycerol (2-AG) (*Science* 1992, 258, 1946-
1949); the sleep-inducing substance, oleamide (OEA) (*Science* 1995, 268,

5 1506); the appetite-suppressing agent, N-oleoylethanolamine (Rodriguez de Fonesca, *Nature* 2001, 414, 209); and the anti-inflammatory agent, palmitoylethanolamide (PEA) (Lambert, *Curr. Med. Chem.* 2002, 9(6), 663).

Small-molecule inhibitors of FAAH should elevate the concentrations of these endogenous signaling lipids and thereby produce their associated
10 beneficial pharmacological effects. There have been some reports of the effects of various FAAH inhibitors in pre-clinical models. In particular, two carbamate-based inhibitors of FAAH were reported to have analgesic properties in animal models. In rats, BMS-1 (see WO 02/087569), which has the structure shown below, was reported to have an analgesic effect in the
15 Chung spinal nerve ligation model of neuropathic pain, and the Hargraves test of acute thermal nociception. URB-597 was reported to have efficacy in the zero plus maze model of anxiety in rats, as well as analgesic efficacy in the rat hot plate and formalin tests (Kathuria, *Nat. Med.* 2003, 9(1), 76). The sulfonylfluoride AM374 was also shown to significantly reduce spasticity in
20 chronic relapsing experimental autoimmune encephalomyelitis (CREAE) mice, an animal model of multiple sclerosis (Baker, *FASEB J.* 2001, 15(2), 300).



In addition, the oxazolopyridine ketone OL-135 is reported to be a
25 potent inhibitor of FAAH, and has been reported to have analgesic activity in both the hot plate and tail emersion tests of thermal nociception in rats (WO 04/033652).



OL-135

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Results of research on the effects of certain exogenous cannabinoids has elucidated that a FAAH inhibitor may be useful for treating various conditions, diseases, disorders, or symptoms. These include pain, nausea/emesis, anorexia, spasticity, movement disorders, epilepsy and glaucoma. To date, approved therapeutic uses for cannabinoids include the relief of chemotherapy-induced nausea and emesis among patients with cancer and appetite enhancement in patients with HIV/AIDs who experience anorexia as a result of wasting syndrome. Two products are commercially available in some countries for these indications, namely, dronabinol (Marinol[®]) and nabilone.

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Apart from the approved indications, a therapeutic field that has received much attention for cannabinoid use is analgesia, i.e., the treatment of pain. Five small randomized controlled trials showed that THC is superior to placebo, producing dose-related analgesia (Robson, *Br. J. Psychiatry* 2001, 178, 107-115). Atlantic Pharmaceuticals is reported to be developing a synthetic cannabinoid, CT-3, a 1,1-dimethyl heptyl derivative of the carboxylic metabolite of tetrahydrocannabinol, as an orally active analgesic and anti-inflammatory agent. A pilot phase II trial in chronic neuropathic pain with CT-3 was reportedly initiated in Germany in May 2002.

20

A number of individuals with locomotor activity-related diseases, such as multiple sclerosis have claimed a benefit from cannabis for both disease-related pain and spasticity, with support from small controlled trials (Croxford et al., *J. Neuroimmunol*, 2008, 193, 120-9; Svendsen, *Br. Med. J.* 2004, 329, 253). Likewise, various victims of spinal cord injuries, such as paraplegia, have reported that their painful spasms are alleviated after smoking marijuana. A report showing that cannabinoids appear to control spasticity and tremor in the CREAE model of multiple sclerosis demonstrated that these effects are mediated by CB₁ and CB₂ receptors (Baker, *Nature* 2000, 404, 84-87). Phase 3 clinical trials have been undertaken in multiple sclerosis and spinal cord injury patients with a narrow ratio mixture of

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5 tetrahydrocannabinol/cannabidiol (THC/CBD). It has been reported that
FAAH knockout mice consistently recover to a better clinical score than wild
type controls, and this improvement is not a result of anti-inflammatory
activity, but rather may reflect some neuroprotection or remyelination
10 promoting effect of lack of the enzyme (Webb et al, *Neurosci Lett.*, 2008, vol.
439, 106-110).

Reports of small-scale controlled trials to investigate other potential
commercial uses of cannabinoids have been made. Trials in volunteers have
been reported to have confirmed that oral, injected, and smoked cannabinoids
produced dose-related reductions in intraocular pressure (IOP) and therefore
15 may relieve glaucoma symptoms. Ophthalmologists have prescribed
cannabis for patients with glaucoma in whom other drugs have failed to
adequately control intraocular pressure (Robson, 2001, *supra*).

Inhibition of FAAH using a small-molecule inhibitor may be
advantageous compared to treatment with a direct-acting CB₁ agonist.
20 Administration of exogenous CB₁ agonists may produce a range of
responses, including reduced nociception, catalepsy, hypothermia, and
increased feeding behavior. These four in particular are termed the
“cannabinoid tetrad.” Experiments with FAAH *-/-* mice show reduced
responses in tests of nociception, but did not show catalepsy, hypothermia, or
25 increased feeding behavior (Cravatt, *Proc. Natl. Acad. Sci. USA* 2001, 98(16),
9371). Fasting caused levels of AEA to increase in rat limbic forebrain, but
not in other brain areas, providing evidence that stimulation of AEA
biosynthesis may be anatomically regionalized to targeted CNS pathways
(Kirkham, *Br. J. Pharmacol.* 2002, 136, 550). The finding that AEA increases
30 are localized within the brain, rather than systemic, suggests that FAAH
inhibition with a small molecule could enhance the actions of AEA and other
fatty acid amides in tissue regions where synthesis and release of these
signaling molecules is occurring in a given pathophysiological condition
(Piomelli, 2003, *supra*).

35 In addition to the effects of a FAAH inhibitor on AEA and other
endocannabinoids, inhibitors of FAAH’s catabolism of other lipid mediators
may be used in treating certain other therapeutic indications. For example,
PEA has demonstrated biological effects in animal models of inflammation

5 (Holt, et al. *Br. J. Pharmacol.* 2005, 146, 467-476), immunosuppression, analgesia, and neuroprotection (Ueda, *J. Biol. Chem.* 2001, 276(38), 35552). Oleamide, another substrate of FAAH, induces sleep (Boger, *Proc. Natl. Acad. Sci. USA* 2000, 97(10), 5044; Mendelson, *Neuropsychopharmacology* 2001, 25, S36). Inhibition of FAAH has also been implicated in cognition
10 (Varvel et al., *J. Pharmacol. Exp. Ther.* 2006, 317(1), 251-257) and depression (Gobbi et al., *Proc. Natl. Acad. Sci. USA* 2005, 102(51), 18620-18625).

Two additional indications for FAAH are supported by recent data indicating that FAAH substrate activated receptors are important in energy
15 metabolism, and in bone homeostasis (Overton et al., *Br. J. Pharmacol.* 2008, in press; and Plutzky, *Diab. Vasc. Dis. Res.* 2007, 4 Suppl 3, S12-4). It has been shown that the previously mentioned lipid signaling fatty acid amides catabolized by FAAH, oleoylethanolamide (OEA), is one of the most active agonists of the recently de-orphanised GPCR 119 (GPR119) (also termed
20 glucose dependent insulinotropic receptor). This receptor is expressed predominantly in the pancreas in humans and activation improves glucose homeostasis via glucose-dependent insulin release in pancreatic beta-cells. GPR119 agonists can suppress glucose excursions when administered during oral glucose tolerance tests, and OEA has also been shown independently to
25 regulate food intake and body weight gain when administered to rodents, indicating a probable benefit in energy metabolism disorders, such as insulin resistance and diabetes. The FAAH substrate palmitoylethanolamide (PEA) is an agonist at the PPAR α receptor. Evidence from surrogate markers in human studies with the PPAR α agonist fenofibrate is supportive of the
30 concept that PPAR α agonism offers the potential for inducing a coordinated PPAR α response that may improve dyslipidaemia, repress inflammation and limit atherosclerosis in patients with the metabolic syndrome or type 2 diabetes. The FAAH substrate anandamide (AEA) is an agonist at the PPAR γ receptor. Anandamide treatment induces 3T3-L1 differentiation into
35 adipocytes, as well as triglyceride droplet accumulation and expression of adiponectin (Bouaboula et al., *E. J. Pharmacol.* 2005, 517, 174-181). Low dose cannabinoid therapy has been shown to reduce atherosclerosis in mice,

5 further suggesting a therapeutic benefit of FAAH inhibition in dyslipidemia, liver steatosis, steatohepatitis, obesity, and metabolic syndrome (Steffens et al., *Nature*, 2005, 434, 782-6).

Osteoporosis is one of the most common degenerative diseases. It is characterized by reduced bone mineral density (BMD) with an increased risk
10 for bone fractures. CB₂-deficient mice have a markedly accelerated age-related trabecular bone loss and cortical expansion. A CB₂-selective agonism enhances endocortical osteoblast number and activity and restrains trabecular osteoclastogenesis and attenuates ovariectomy-induced bone loss (Ofek et al., *Proc. Natl. Acad. Sci. U.S.A.* 2006, 103, 696-701). There is a substantial
15 genetic contribution to BMD, although the genetic factors involved in the pathogenesis of human osteoporosis are largely unknown. The applicability to human BMD is suggested by genetic studies in which a significant association of single polymorphisms and haplotypes was found encompassing the CNR2 gene on human chromosome 1p36, demonstrating a role for the peripherally
20 expressed CB₂ receptor in the etiology of osteoporosis (Karsak et al., *Hum. Mol. Genet.*, 2005, 14, 3389-96).

Thus, small-molecule FAAH inhibitors should be useful in treating pain of various etiologies, anxiety, multiple sclerosis and other movement disorders, nausea/emesis, eating disorders, epilepsy, glaucoma,
25 inflammation, immunosuppression, neuroprotection, depression, cognition enhancement, and sleep disorders, and potentially with fewer side effects than treatment with an exogenous cannabinoid.

A number of heteroaryl-substituted ureas have been reported in various publications. Certain substituted thiophene ureas are described in US
30 Patent No. 6,881,741. Certain ureido-pyrazoles are described in US Patent No. 6,387,900. Certain benzothiazole amide derivatives are described in US Patent Publication US 2003/149036. Certain ureas are reported as prenyltransferase inhibitors in WO 2003/047569. Piperidiny l ureas are described as histamine H₃ receptor antagonists in US Patent No. 6,100,279.
35 Piperaziny l ureas are disclosed as calcitonin mimetics in US Patent Nos. 6,124,299 and 6,395,740. Certain piperidiny l ureas and piperaziny l ureas have been previously described as FAAH modulators in U.S. Pat. Appl. No. 12/126,389, filed May 23, 2008. Various ureas are reported as small-

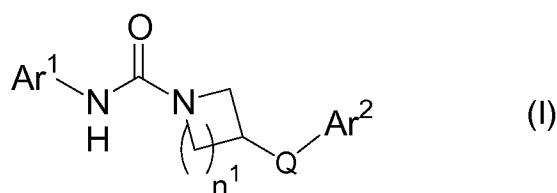
5 molecule FAAH modulators in US Patent Publication Nos. US 2006/173184 and US 2007/0004741, in Intl. Patent Appl. Nos. WO 2008/023720, WO 2008/047229, and WO 2008/024139, and by Cravatt et al. (Biochemistry 2007, 46(45), 13019. Ureas are described as modulators of other targets in U.S. Pat. Appl. Publ. US 2007/270433, and in Intl. Pat. Appl. Publ. Nos. WO
10 2007/096251 and WO 2006/085108.

However, there remains a desire for potent FAAH modulators with suitable pharmaceutical properties.

Summary of the Invention

Certain aryl-substituted heterocyclic urea derivatives have been found
15 to have FAAH-modulating activity. Thus, the invention is directed to the general and preferred embodiments defined, respectively, and by the independent and dependent claims appended hereto, which are incorporated by reference herein.

In one general aspect, the invention is directed to compounds of
20 Formula (I):



wherein

Q is $-(\text{CH}_2)_{1-3}-$, $-(\text{CH}_2)_{1-2}\text{O}-$, or $-(\text{CH}_2)_{1-2}\text{OCH}_2-$;

n^1 is 1 or 2, with the proviso that when n^1 is 1, then Q cannot be $-\text{CH}_2-$;

25 Ar^1 is a ring system selected from benzo[d]isoxazolyl, 1H-pyrrolo[2,3-b]pyridinyl, 2,1,3-benzoxadiazolyl, imidazo[1,2-a]pyridinyl, imidazo[1,2-b]pyridazinyl, isoquinolinyl, and pyridyl optionally substituted with triazolyl;

wherein each ring system is optionally substituted with halo;

30 Ar^2 is:

- (i) phenyl optionally substituted with one or two R^d moieties; wherein each R^d moiety is independently selected from $-\text{C}_{1-4}\text{alkyl}$, $-\text{OC}_{1-4}\text{alkyl}$, halo, $-\text{CF}_3$, $-\text{OCF}_3$, and $-\text{S}(\text{O})(\text{O})\text{C}_{1-4}\text{alkyl}$, or two adjacent R^d moieties taken together form $-\text{OCH}_2\text{O}-$ or $-\text{OCF}_2\text{O}-$;

5 (ii) phenyl substituted at the 3- or 4-position with $-L-Ar^3$, said phenyl optionally substituted with one additional R^d moiety, wherein:

L is $-O-(CH_2)_{0-1}-$ or a covalent bond;

Ar^3 is:

(a) phenyl optionally substituted with one or two R^d moieties; or

10 (b) pyridyl optionally substituted with one or two R^d moieties;

(iii) naphthyl;

(iv) 5,6,7,8-tetrahydro-naphthalenyl; or

(v) quinolinylyl optionally substituted with a halo.

The invention also relates to pharmaceutically acceptable salts of
15 compounds of Formula (I), pharmaceutically acceptable prodrugs of compounds of Formula (I), and pharmaceutically acceptable metabolites of compounds of Formula (I). In certain preferred embodiments, the compound of Formula (I) is a compound selected from those species described or exemplified in the detailed description below.

20 In a further general aspect, the invention relates to pharmaceutical compositions each comprising: (a) a therapeutically effective amount of at least one chemical entity selected from compounds of Formula (I), pharmaceutically acceptable salts of compounds of Formula (I), pharmaceutically acceptable prodrugs of compounds of Formula (I), and
25 pharmaceutically acceptable metabolites of compounds of Formula (I); and (b) a pharmaceutically acceptable excipient.

In another aspect, embodiments of the invention are useful as FAAH modulators. Thus, the invention is directed to a method for modulating FAAH activity, comprising exposing FAAH to a therapeutically effective amount of at
30 least one chemical entity selected from compounds of Formula (I), pharmaceutically acceptable salts of compounds of Formula (I), pharmaceutically acceptable prodrugs of compounds of Formula (I), and pharmaceutically active metabolites of compounds of Formula (I). Embodiments of this invention modulate FAAH activity.

35 In another general aspect, the invention is directed to a method of treating a subject suffering from or diagnosed with a disease, disorder, or medical condition (collectively, "indications") mediated by FAAH activity, comprising administering to the subject in need of such treatment a

5 therapeutically effective amount of a compound of Formula (I), a
pharmaceutically acceptable salt of a compound of Formula (I), a
pharmaceutically acceptable salt of a compound of Formula (I), and
pharmaceutically acceptable prodrug, or a pharmaceutically active metabolite
of a compound of Formula (I). In preferred embodiments of the inventive
10 method, the disease, disorder, or medical condition is selected from: anxiety,
depression, pain, sleep disorders, eating disorders, inflammation, multiple
sclerosis and other movement disorders, HIV wasting syndrome, closed head
injury, stroke, learning and memory disorders, Alzheimer's disease, epilepsy,
Tourette's syndrome, Niemann-Pick disease, Parkinson's disease,
15 Huntington's chorea, optic neuritis, autoimmune uveitis, symptoms of drug or
alcohol withdrawal, nausea, emesis, sexual dysfunction, post-traumatic stress
disorder, cerebral vasospasm, glaucoma, irritable bowel syndrome,
inflammatory bowel disease, immunosuppression, itch, gastroesophageal
reflux disease, paralytic ileus, secretory diarrhea, gastric ulcer, rheumatoid
20 arthritis, unwanted pregnancy, hypertension, cancer, hepatitis, allergic airway
disease, auto-immune diabetes, intractable pruritis, neuroinflammation,
diabetes, metabolic syndrome, and osteoporosis.

Additional embodiments, features, and advantages of the invention will be
apparent from the following detailed description and through practice of the
25 invention.

Detailed Description of Invention and Its Preferred Embodiments

The invention may be more fully appreciated by reference to the
following detailed description, including the following glossary of terms and
the concluding examples. For the sake of brevity, the disclosures of the
30 publications, including patents, cited in this specification are herein
incorporated by reference.

As used herein, the terms "including", "containing" and "comprising" are
used in their open, non-limiting sense.

The term "alkyl" refers to a straight- or branched-chain alkyl group
35 having from 1 to 12 carbon atoms in the chain. Such groups may contain
saturated or unsaturated carbon atoms within the chain. Examples of alkyl
groups include methyl (Me, which also may be structurally depicted by /

5 A structural formula given herein is intended to represent compounds having structures depicted by the formula as well as equivalent variations or forms. For example, compounds encompassed by Formula (I) may have asymmetric centers and therefore exist in different enantiomeric forms. All optical isomers and stereoisomers of the compounds of the general formula, and mixtures thereof, are considered within the scope of the formula. Thus, a general formula given herein is intended to represent a racemate, one or more enantiomeric forms, one or more diastereomeric forms, one or more atropisomeric forms, and mixtures thereof. Furthermore, certain structures may exist as geometric isomers (i.e., *cis* and *trans* isomers), as tautomers (e.g. pyrazole, benzimidazole, tetrazole, or benzotriazole tautomers), or as atropisomers, which are intended to be represented by the structural formula. Additionally, a formula given herein is intended to embrace hydrates, solvates, and polymorphs of such compounds, and mixtures thereof, even if such forms are not listed explicitly.

20 To provide a more concise description, some of the quantitative expressions given herein are not qualified with the term "about". It is understood that, whether the term "about" is used explicitly or not, every quantity given herein is meant to refer to the actual given value, and it is also meant to refer to the approximation to such given value that would reasonably be inferred based on the ordinary skill in the art, including equivalents and approximations due to the experimental and/or measurement conditions for such given value. Whenever a yield is given as a percentage, such yield refers to a mass of the entity for which the yield is given with respect to the maximum amount of the same entity that could be obtained under the particular stoichiometric conditions. Concentrations that are given as percentages refer to mass ratios, unless indicated differently.

35 Reference to a chemical entity herein stands for a reference to any one of: (a) the actually recited form of such chemical entity, and (b) any of the forms of such chemical entity in the medium in which the compound is being considered when named. For example, reference herein to a compound such as R-COOH, encompasses reference to any one of, for example, R-COOH(s), R-COOH(sol), and R-COO-(sol). In this example, R-COOH(s) refers to the solid compound, as it could be for example in a tablet or some other solid

5 pharmaceutical composition or preparation; R-COOH(sol) refers to the
undissociated form of the compound in a solvent; and R-COO-(sol) refers to
the dissociated form of the compound in a solvent, such as the dissociated
form of the compound in an aqueous environment, whether such dissociated
form derives from R-COOH, from a salt thereof, or from any other entity that
10 yields R-COO- upon dissociation in the medium being considered. In another
example, an expression such as “exposing an entity to compound of formula
R-COOH” refers to the exposure of such entity to the form, or forms, of the
compound R-COOH that exists, or exist, in the medium in which such
exposure takes place. In this regard, if such entity is for example in an
15 aqueous environment, it is understood that the compound R-COOH is in such
same medium, and therefore the entity is being exposed to species such as
R-COOH(aq) and/or R-COO-(aq), where the subscript “(aq)” stands for
“aqueous” according to its conventional meaning in chemistry and
biochemistry. A carboxylic acid functional group has been chosen in these
20 nomenclature examples; this choice is not intended, however, as a limitation
but it is merely an illustration. It is understood that analogous examples can
be provided in terms of other functional groups, including but not limited to
hydroxyl, basic nitrogen members, such as those in amines, and any other
group that interacts or transforms according to known manners in the medium
25 that contains the compound. Such interactions and transformations include,
but are not limited to, dissociation, association, tautomerism, solvolysis,
including hydrolysis, solvation, including hydration, protonation, and
deprotonation. No further examples in this regard are provided herein
because these interactions and transformations in a given medium are known
30 by any one of ordinary skill in the art.

Any structural formula given herein is also intended to represent
unlabeled forms as well as isotopically labeled forms of the compounds.
Isotopically labeled compounds have structures depicted by the formulas
given herein except that one or more atoms are replaced by an atom having a
35 selected atomic mass or mass number. Examples of isotopes that can be
incorporated into compounds of the invention include isotopes of hydrogen,
carbon, nitrogen, oxygen, phosphorous, fluorine, and chlorine, such as ²H, ³H,
¹¹C, ¹³C, ¹⁴C, ¹⁵N, ¹⁸O, ¹⁷O, ³²P, ³³P, ³⁵S, ¹⁸F, ³⁶Cl, and ¹²⁵I, respectively. Such

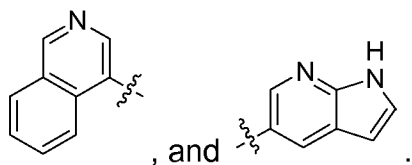
5 isotopically labeled compounds are useful in metabolic studies (preferably
with ^{14}C), reaction kinetic studies (with, for example ^2H or ^3H), detection or
imaging techniques [such as positron emission tomography (PET) or single-
photon emission computed tomography (SPECT), including drug or substrate
tissue distribution assays, or in radioactive treatment of patients. In particular,
10 an ^{18}F - or ^{11}C -labeled compound may be preferred for PET or SPECT studies.
Further, substitution with heavier isotopes such as deuterium (i.e., ^2H) may
afford certain therapeutic advantages resulting from greater metabolic
stability, for example increased *in vivo* half-life or reduced dosage
requirements. Isotopically labeled compounds of this invention and prodrugs
15 thereof can generally be prepared by carrying out the procedures disclosed in
the schemes or in the examples and preparations described below by
substituting a readily available isotopically labeled reagent for a non-
isotopically labeled reagent.

When referring to any formula given herein, the selection of a particular
20 moiety from a list of possible species for a specified variable is not intended to
define the moiety for the variable appearing elsewhere. In other words, where
a formula variable appears more than once, the choice of the species from a
specified list is independent of the choice of the species for the same variable
elsewhere in the formula, unless stated otherwise.

25 According to the foregoing interpretive considerations on assignments
and nomenclature, it is understood that explicit reference herein to a set
implies, where chemically meaningful and unless indicated otherwise,
independent reference to embodiments of such set, and reference to each
and every one of the possible embodiments of subsets of the set referred to
30 explicitly.

In some embodiments of Formula (I), Ar^1 is a ring system selected from
benzo[d]isoxazol-3-yl, 2,1,3-benzoxadiazol-4-yl, 4-chloropyridin-3-yl,
imidazo[1,2-a]pyridin-3-yl, imidazo[1,2-a]pyridin-5-yl, imidazo[1,2-a]pyridin-6-

5 yl, imidazo[1,2-b]pyridazin-3-yl, pyridin-3-yl, 6-[1,2,3]triazol-2-yl-pyridin-3-yl,



In some embodiments of Formula (I), Q is $-\text{CH}_2\text{CH}_2\text{O}-$. In further embodiments, Q is $-\text{CH}_2\text{O}-$.

In some embodiments of Formula (I), Ar^2 is phenyl substituted at the 3-
 10 or 4- position with $-\text{L}-\text{Ar}^3$. In further embodiments, Ar^2 is phenyl substituted at the 3- or 4- position with $-\text{L}-\text{Ar}^3$, L is $-\text{O}-$, and Ar^3 is phenyl optionally substituted with one or two R^d moieties. In other embodiments, Ar^2 is phenyl substituted at the 3- or 4- position with $-\text{L}-\text{Ar}^3$, L is $-\text{O}-$, and Ar^3 is pyridyl optionally substituted with one or two R^d moieties. In further embodiments,
 15 Ar^2 is phenyl substituted at the 3- or 4- position with $-\text{L}-\text{Ar}^3$ and Q is $-\text{CH}_2\text{CH}_2\text{O}-$. In further embodiments, Ar^2 is phenyl substituted at the 3- or 4- position with $-\text{L}-\text{Ar}^3$ and Q is $-\text{CH}_2\text{O}-$.

In some embodiments of Formula (I), Ar^2 is phenyl optionally substituted with one or two R^d . In further embodiments, Ar^2 is phenyl
 20 optionally substituted with one or two R^d and Q is $-\text{CH}_2\text{CH}_2\text{O}-$.

In some embodiments of Formula (I), n is 1. In further embodiments, n is 1 and Ar^1 is pyridin-3-yl optionally substituted with halo. In further
 embodiments, n is 1, Ar^1 is pyridin-3-yl optionally substituted with halo, and Ar^2 is phenyl substituted at the 3- or 4- position with $-\text{L}-\text{Ar}^3$. In further
 25 embodiments, n is 1, Ar^1 is pyridin-3-yl optionally substituted with halo, Ar^2 is phenyl substituted at the 3- or 4- position with $-\text{L}-\text{Ar}^3$, and Q is $-\text{CH}_2\text{CH}_2\text{O}-$. In further embodiments, n is 1, Ar^1 is pyridin-3-yl optionally substituted with halo, and Ar^2 is phenyl optionally substituted with one or two R^d .

In some embodiments of Formula (I), n is 2. In further embodiments, n
 30 is 2 and Ar^1 is pyridin-3-yl optionally substituted with halo. In further embodiments, n is 2, Ar^1 is pyridin-3-yl optionally substituted with halo, and Ar^2 is phenyl substituted at the 3- or 4- position with $-\text{L}-\text{Ar}^3$. In further embodiments, n is 2, Ar^1 is pyridin-3-yl optionally substituted with halo, Ar^2 is phenyl substituted at the 3- or 4- position with $-\text{L}-\text{Ar}^3$, and Q is $-\text{CH}_2\text{CH}_2\text{O}-$.

- 5 In further embodiments, n is 2, Ar¹ is pyridin-3-yl optionally substituted with halo, and Ar² is phenyl optionally substituted with one or two R^d.

The invention includes also pharmaceutically acceptable salts of the compounds represented by Formula (I), preferably of those described below and of the specific compounds exemplified herein, and methods using such
10 salts.

A "pharmaceutically acceptable salt" is intended to mean a salt of a free acid or base of a compound represented by Formula (I) that is non-toxic, biologically tolerable, or otherwise biologically suitable for administration to the subject. See, generally, S.M. Berge, et al., "Pharmaceutical Salts", J.
15 Pharm. Sci., 1977, 66:1-19, and Handbook of Pharmaceutical Salts, Properties, Selection, and Use, Stahl and Wermuth, Eds., Wiley-VCH and VHCA, Zurich, 2002.

Preferred pharmaceutically acceptable salts are those that are pharmacologically effective and suitable for contact with the tissues of
20 patients without undue toxicity, irritation, or allergic response. A compound of Formula (I) may possess a sufficiently acidic group, a sufficiently basic group, or both types of functional groups, and accordingly react with a number of inorganic or organic bases, and inorganic and organic acids, to form a pharmaceutically acceptable salt. Examples of pharmaceutically acceptable
25 salts include sulfates, pyrosulfates, bisulfates, sulfites, bisulfites, phosphates, monohydrogen-phosphates, dihydrogenphosphates, metaphosphates, pyrophosphates, chlorides, bromides, iodides, acetates, propionates, decanoates, caprylates, acrylates, formates, isobutyrate, caproates, heptanoates, propiolates, oxalates, malonates, succinates, suberates,
30 sebacates, fumarates, maleates, butyne-1,4-dioates, hexyne-1,6-dioates, benzoates, chlorobenzoates, methylbenzoates, dinitrobenzoates, hydroxybenzoates, methoxybenzoates, phthalates, sulfonates, xylenesulfonates, phenylacetates, phenylpropionates, phenylbutyrates, citrates, lactates, γ -hydroxybutyrates, glycolates, tartrates, methane-
35 sulfonates, propanesulfonates, naphthalene-1-sulfonates, naphthalene-2-sulfonates, and mandelates.

If the compound of Formula (I) contains a basic nitrogen, the desired pharmaceutically acceptable salt may be prepared by any suitable method

5 available in the art, for example, treatment of the free base with an inorganic acid, such as hydrochloric acid, hydrobromic acid, sulfuric acid, sulfamic acid, nitric acid, boric acid, phosphoric acid, and the like, or with an organic acid, such as acetic acid, phenylacetic acid, propionic acid, stearic acid, lactic acid, ascorbic acid, maleic acid, hydroxymaleic acid, isethionic acid, succinic acid, 10 valeric acid, fumaric acid, malonic acid, pyruvic acid, oxalic acid, glycolic acid, salicylic acid, oleic acid, palmitic acid, lauric acid, a pyranosidyl acid, such as glucuronic acid or galacturonic acid, an alpha-hydroxy acid, such as mandelic acid, citric acid, or tartaric acid, an amino acid, such as aspartic acid or glutamic acid, an aromatic acid, such as benzoic acid, 2-acetoxybenzoic acid, 15 naphthoic acid, or cinnamic acid, a sulfonic acid, such as laurylsulfonic acid, p-toluenesulfonic acid, methanesulfonic acid, ethanesulfonic acid, any compatible mixture of acids such as those given as examples herein, and any other acid and mixture thereof that are regarded as equivalents or acceptable substitutes in light of the ordinary level of skill in this technology.

20 If the compound of Formula (I) is an acid, such as a carboxylic acid or sulfonic acid, the desired pharmaceutically acceptable salt may be prepared by any suitable method, for example, treatment of the free acid with an inorganic or organic base, such as an amine (primary, secondary or tertiary), an alkali metal hydroxide, alkaline earth metal hydroxide, any compatible 25 mixture of bases such as those given as examples herein, and any other base and mixture thereof that are regarded as equivalents or acceptable substitutes in light of the ordinary level of skill in this technology. Illustrative examples of suitable salts include organic salts derived from amino acids, such as glycine and arginine, ammonia, carbonates, bicarbonates, primary, secondary, and 30 tertiary amines, and cyclic amines, such as benzylamines, pyrrolidines, piperidine, morpholine, and piperazine, and inorganic salts derived from sodium, calcium, potassium, magnesium, manganese, iron, copper, zinc, aluminum, and lithium.

The invention also relates to pharmaceutically acceptable prodrugs of 35 the compounds of Formula (I), and treatment methods employing such pharmaceutically acceptable prodrugs. The term "prodrug" means a precursor of a designated compound that, following administration to a subject, yields the compound in vivo via a chemical or physiological process

5 such as solvolysis or enzymatic cleavage, or under physiological conditions (e.g., a prodrug on being brought to physiological pH is converted to the compound of Formula (I)). A "pharmaceutically acceptable prodrug" is a prodrug that is non-toxic, biologically tolerable, and otherwise biologically suitable for administration to the subject. Illustrative procedures for the selection and preparation of suitable prodrug derivatives are described, for example, in "Design of Prodrugs", ed. H. Bundgaard, Elsevier, 1985.

Examples of prodrugs include compounds having an amino acid residue, or a polypeptide chain of two or more (e.g., two, three or four) amino acid residues, covalently joined through an amide or ester bond to a free amino, hydroxy, or carboxylic acid group of a compound of Formula (I). Examples of amino acid residues include the twenty naturally occurring amino acids, commonly designated by three letter symbols, as well as 4-hydroxyproline, hydroxylysine, demosine, isodemosine, 3-methylhistidine, norvalin, beta-alanine, gamma-aminobutyric acid, citrulline homocysteine, homoserine, ornithine and methionine sulfone.

Additional types of prodrugs may be produced, for instance, by derivatizing free carboxyl groups of structures of Formula (I) as amides or alkyl esters. Examples of amides include those derived from ammonia, primary C₁₋₆alkyl amines and secondary di(C₁₋₆alkyl) amines. Secondary amines include 5- or 6-membered heterocycloalkyl or heteroaryl ring moieties. Examples of amides include those that are derived from ammonia, C₁₋₃alkyl primary amines, and di(C₁₋₂alkyl)amines. Examples of esters of the invention include C₁₋₇alkyl, C₅₋₇cycloalkyl, phenyl, and phenyl(C₁₋₆alkyl) esters. Preferred esters include methyl esters. Prodrugs may also be prepared by derivatizing free hydroxy groups using groups including hemisuccinates, phosphate esters, dimethylaminoacetates, and phosphoryloxymethyloxycarbonyls, following procedures such as those outlined in Fleisher et al., Adv. Drug Delivery Rev. 1996, 19, 115-130. Carbamate derivatives of hydroxy and amino groups may also yield prodrugs. Carbonate derivatives, sulfonate esters, and sulfate esters of hydroxy groups may also provide prodrugs. Derivatization of hydroxy groups as (acyloxy)methyl and (acyloxy)ethyl ethers, wherein the acyl group may be an alkyl ester, optionally substituted with one or more ether, amine, or carboxylic acid functionalities, or where the acyl group is an amino

5 acid ester as described above, is also useful to yield prodrugs. Prodrugs of this type may be prepared as described in Robinson et al., J. Med. Chem. 1996, 39, 10-18. Free amines can also be derivatized as amides, sulfonamides or phosphoramidates. All of these prodrug moieties may incorporate groups including ether, amine, and carboxylic acid functionalities.

10 The present invention also relates to pharmaceutically active metabolites of compounds of Formula (I), and uses of such metabolites in the methods of the invention. A "pharmaceutically active metabolite" means a pharmacologically active product of metabolism in the body of a compound of
15 Formula (I) or salt thereof. Prodrugs and active metabolites of a compound may be determined using routine techniques known or available in the art. See, e.g., Bertolini et al., J. Med. Chem. 1997, 40, 2011-2016; Shan et al., J. Pharm. Sci. 1997, 86 (7), 765-767; Bagshawe, Drug Dev. Res. 1995, 34, 220-230; Bodor, Adv. Drug Res. 1984, 13, 255-331; Bundgaard, Design of
20 Prodrugs (Elsevier Press, 1985); and Larsen, Design and Application of Prodrugs, Drug Design and Development (Krogsgaard-Larsen et al., eds., Harwood Academic Publishers, 1991).

The compounds of Formula (I), and their pharmaceutically acceptable salts, pharmaceutically acceptable prodrugs, and pharmaceutically active
25 metabolites (collectively, "active agents") of the present invention are useful as FAAH inhibitors in the methods of the invention. The active agents may be used in the inventive methods for the treatment of medical conditions, diseases, or disorders mediated through inhibition or modulation of FAAH, such as those described herein. Active agents according to the invention may
30 therefore be used as an analgesic, anti-depressant, cognition enhancer, neuroprotectant, sedative, appetite stimulant, or contraceptive.

Thus, the active agents may be used to treat subjects diagnosed with or suffering from a disease, disorder, or condition mediated through FAAH activity. The term "treat" or "treating" as used herein is intended to refer to
35 administration of an active agent or composition of the invention to a subject for the purpose of effecting a therapeutic or prophylactic benefit through modulation of FAAH activity. Treating includes reversing, ameliorating, alleviating, inhibiting the progress of, lessening the severity of, or preventing a

5 disease, disorder, or condition, or one or more symptoms of such disease,
disorder or condition mediated through modulation of FAAH activity. The term
"subject" refers to a mammalian patient in need of such treatment, such as a
human. "Modulators" include both inhibitors and activators, where "inhibitors"
10 refer to compounds that decrease, prevent, inactivate, desensitize or down-
regulate FAAH expression or activity, and "activators" are compounds that
increase, activate, facilitate, sensitize, or up-regulate FAAH expression or
activity.

Accordingly, the invention relates to methods of using the active agents
described herein to treat subjects diagnosed with or suffering from a disease,
15 disorder, or condition mediated through FAAH activity, such as: anxiety, pain,
sleep disorders, eating disorders, inflammation, movement disorders (e.g.,
multiple sclerosis), energy metabolism (e.g. insulin resistance, diabetes,
dyslipidemia, liver steatosis, steatohepatitis, obesity, and metabolic
syndrome) and bone homeostasis (e.g. osteoporosis).

20 In certain preferred embodiments, active agents may be used in
methods to treat a FAAH mediated disease, disorder, or medical condition
where the disease, disorder, or medical condition is selected from the group
consisting of anxiety, depression, pain, sleep disorders, eating disorders,
inflammation, multiple sclerosis and other movement disorders, HIV wasting
25 syndrome, closed head injury, stroke, learning and memory disorders,
Alzheimer's disease, epilepsy, Tourette's syndrome, epilepsy, Niemann-Pick
disease, Parkinson's disease, Huntington's chorea, optic neuritis, autoimmune
uveitis, symptoms of drug withdrawal, nausea, emesis, sexual dysfunction,
post-traumatic stress disorder, cerebral vasospasm, glaucoma, irritable bowel
30 syndrome, inflammatory bowel disease, immunosuppression,
gastroesophageal reflux disease, paralytic ileus, secretory diarrhea, gastric
ulcer, rheumatoid arthritis, unwanted pregnancy, hypertension, cancer,
hepatitis, allergic airway disease, autoimmune diabetes, intractable pruritis,
neuroinflammation, diabetes, metabolic syndrome and osteoporosis. In
35 certain preferred embodiments, the disease, disorder, or medical condition is
pain or inflammation. In further embodiments, the disease, disorder, or
medical condition is anxiety, a sleep disorder, an eating disorder, or a
movement disorder. In further embodiments, the disease, disorder, or

5 medical condition is multiple sclerosis. In further embodiments, the disease, disorder, or medical condition is energy metabolism or bone homeostasis.

Symptoms or disease states are intended to be included within the scope of "medical conditions, disorders, or diseases." For example, pain may be associated with various diseases, disorders, or conditions, and may
10 include various etiologies. Illustrative types of pain treatable with a FAAH-modulating agent, in one example herein a FAAH-inhibiting agent, according to the invention include cancer pain, postoperative pain, GI tract pain, spinal cord injury pain, visceral hyperalgesia, thalamic pain, headache (including stress headache and migraine), low back pain, neck pain, musculoskeletal
15 pain, peripheral neuropathic pain, central neuropathic pain, neurodegenerative disorder related pain, and menstrual pain. HIV wasting syndrome includes associated symptoms such as appetite loss and nausea. Parkinson's disease includes, for example, levodopa-induced dyskinesia. Treatment of multiple sclerosis may include treatment of symptoms such as spasticity, neurogenic
20 pain, central pain, or bladder dysfunction. Symptoms of drug withdrawal may be caused by, for example, addiction to opiates or nicotine. Nausea or emesis may be due to chemotherapy, postoperative, or opioid related causes. Treatment of sexual dysfunction may include improving libido or delaying ejaculation. Treatment of cancer may include treatment of glioma. Sleep
25 disorders include, for example, sleep apnea, insomnia, and disorders calling for treatment with an agent having a sedative or narcotic-type effect. Eating disorders include, for example, anorexia or appetite loss associated with a disease such as cancer or HIV infection/AIDS.

In treatment methods according to the invention, an effective amount of
30 at least one active agent according to the invention is administered to a subject suffering from or diagnosed as having such a disease, disorder, or condition. A "therapeutically effective amount" or "effective amount" means an amount or dose of a FAAH-modulating agent sufficient to generally bring about a therapeutic benefit in patients in need of treatment for a disease,
35 disorder, or condition mediated by FAAH activity. Effective amounts or doses of the active agents of the present invention may be ascertained by routine methods such as modeling, dose escalation studies or clinical trials, and by taking into consideration routine factors, e.g., the mode or route of

5 administration or drug delivery, the pharmacokinetics of the agent, the severity and course of the disease, disorder, or condition, the subject's previous or ongoing therapy, the subject's health status and response to drugs, and the judgment of the treating physician. An exemplary dose is in the range of from about 0.0001 to about 200 mg of active agent per kg of
10 subject's body weight per day, preferably about 0.001 to 100 mg/kg/day, or about 0.01 to 35 mg/kg/day, or about 0.1 to 10 mg/kg daily in single or divided dosage units (e.g., BID, TID, QID). For a 70-kg human, an illustrative range for a suitable dosage amount is from about 0.05 to about 7 g/day, or about 0.2 to about 5 g/day. Once improvement of the patient's disease, disorder, or
15 condition has occurred, the dose may be adjusted for maintenance treatment. For example, the dosage or the frequency of administration, or both, may be reduced as a function of the symptoms, to a level at which the desired therapeutic effect is maintained. Of course, if symptoms have been alleviated to an appropriate level, treatment may cease. Patients may, however, require
20 intermittent treatment on a long-term basis upon any recurrence of symptoms.

Once improvement of the patient's disease, disorder, or condition has occurred, the dose may be adjusted for preventative or maintenance treatment. For example, the dosage or the frequency of administration, or both, may be reduced as a function of the symptoms, to a level at which the
25 desired therapeutic or prophylactic effect is maintained. Of course, if symptoms have been alleviated to an appropriate level, treatment may cease. Patients may, however, require intermittent treatment on a long-term basis upon any recurrence of symptoms.

In addition, the active agents of the invention may be used in
30 combination with additional active ingredients in the treatment of the above conditions. The additional active ingredients may be coadministered separately with an active agent of Formula (I) or included with such an agent in a pharmaceutical composition according to the invention. In an exemplary embodiment, additional active ingredients are those that are known or
35 discovered to be effective in the treatment of conditions, disorders, or diseases mediated by FAAH activity, such as another FAAH modulator or a compound active against another target associated with the particular condition, disorder, or disease. The combination may serve to increase

5 efficacy (e.g., by including in the combination a compound potentiating the
potency or effectiveness of an active agent according to the invention),
decrease one or more side effects, or decrease the required dose of the
active agent according to the invention. In one illustrative embodiment, a
composition according to the invention may contain one or more additional
10 active ingredients selected from opioids, non-steroidal anti-inflammatory drugs
(NSAIDs) (e.g., ibuprofen, cyclooxygenase-2 (COX-2) inhibitors, and
naproxen), gabapentin, pregabalin, tramadol, acetaminophen, and aspirin.

The active agents of the invention are used, alone or in combination
with one or more additional active ingredients, to formulate pharmaceutical
15 compositions of the invention. A pharmaceutical composition of the invention
comprises: (a) an effective amount of at least one active agent in accordance
with the invention; and (b) a pharmaceutically acceptable excipient.

When referring to modulating the target receptor, an "effective amount"
means an amount sufficient to affect the activity of such receptor. Measuring
20 the activity of the target receptor may be performed by routine analytical
methods. Target receptor modulation is useful in a variety of settings,
including assays.

A "pharmaceutically acceptable excipient" refers to a substance that is
non-toxic, biologically tolerable, and otherwise biologically suitable for
25 administration to a subject, such as an inert substance, added to a
pharmacological composition or otherwise used as a vehicle, carrier, or
diluent to facilitate administration of a agent and that is compatible therewith.
Examples of excipients include calcium carbonate, calcium phosphate,
various sugars and types of starch, cellulose derivatives, gelatin, vegetable
30 oils, and polyethylene glycols.

Delivery forms of the pharmaceutical compositions containing one or
more dosage units of the active agents may be prepared using suitable
pharmaceutical excipients and compounding techniques known or that
become available to those skilled in the art. The compositions may be
35 administered in the inventive methods by a suitable route of delivery, e.g.,
oral, parenteral, rectal, topical, or ocular routes, or by inhalation.

The preparation may be in the form of tablets, capsules, sachets,
dragees, powders, granules, lozenges, powders for reconstitution, liquid

5 preparations, or suppositories. Preferably, the compositions are formulated for intravenous infusion, topical administration, or oral administration.

For oral administration, the active agents of the invention can be provided in the form of tablets or capsules, or as a solution, emulsion, or suspension. To prepare the oral compositions, the active agents may be
10 formulated to yield a dosage of, e.g., from about 5 mg to 5 g daily, or from about 50 mg to 5 g daily, in single or divided doses. For example, a total daily dosage of about 5 mg to 5 g daily may be accomplished by dosing once, twice, three, or four times per day.

Oral tablets may include the active ingredient(s) mixed with compatible
15 pharmaceutically acceptable excipients such as diluents, disintegrating agents, binding agents, lubricating agents, sweetening agents, flavoring agents, coloring agents and preservative agents. Suitable inert fillers include sodium and calcium carbonate, sodium and calcium phosphate, lactose, starch, sugar, glucose, methyl cellulose, magnesium stearate, mannitol,
20 sorbitol, and the like. Exemplary liquid oral excipients include ethanol, glycerol, water, and the like. Starch, polyvinyl-pyrrolidone (PVP), sodium starch glycolate, microcrystalline cellulose, and alginic acid are exemplary disintegrating agents. Binding agents may include starch and gelatin. The lubricating agent, if present, may be magnesium stearate, stearic acid or talc.
25 If desired, the tablets may be coated with a material such as glyceryl monostearate or glyceryl distearate to delay absorption in the gastrointestinal tract, or may be coated with an enteric coating.

Capsules for oral administration include hard and soft gelatin capsules. To prepare hard gelatin capsules, active ingredient(s) may be mixed with a
30 solid, semi-solid, or liquid diluent. Soft gelatin capsules may be prepared by mixing the active ingredient with water, an oil such as peanut oil or olive oil, liquid paraffin, a mixture of mono and di-glycerides of short chain fatty acids, polyethylene glycol 400, or propylene glycol.

Liquids for oral administration may be in the form of suspensions,
35 solutions, emulsions or syrups or may be lyophilized or presented as a dry product for reconstitution with water or other suitable vehicle before use. Such liquid compositions may optionally contain: pharmaceutically-acceptable excipients such as suspending agents (for example, sorbitol,

5 methyl cellulose, sodium alginate, gelatin, hydroxyethylcellulose,
carboxymethylcellulose, aluminum stearate gel and the like); non-aqueous
vehicles, e.g., oil (for example, almond oil or fractionated coconut oil),
propylene glycol, ethyl alcohol, or water; preservatives (for example, methyl or
propyl p-hydroxybenzoate or sorbic acid); wetting agents such as lecithin;
10 and, if desired, flavoring or coloring agents.

The active agents of this invention may also be administered by non-oral routes. For example, compositions may be formulated for rectal administration as a suppository. For parenteral use, including intravenous, intramuscular, intraperitoneal, or subcutaneous routes, the agents of the
15 invention may be provided in sterile aqueous solutions or suspensions, buffered to an appropriate pH and isotonicity or in parenterally acceptable oil. Suitable aqueous vehicles include Ringer's solution and isotonic sodium chloride. Such forms may be presented in unit-dose form such as ampules or disposable injection devices, in multi-dose forms such as vials from which the
20 appropriate dose may be withdrawn, or in a solid form or pre-concentrate that can be used to prepare an injectable formulation. Illustrative infusion doses range from about 1 to 1000 $\mu\text{g}/\text{kg}/\text{minute}$ of agent admixed with a pharmaceutical carrier over a period ranging from several minutes to several days.

25 For topical administration, the agents may be mixed with a pharmaceutical carrier at a concentration of about 0.1% to about 10% of drug to vehicle. Another mode of administering the agents of the invention may utilize a patch formulation to affect transdermal delivery.

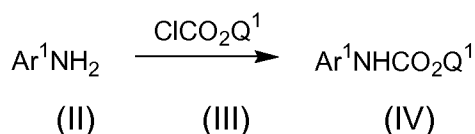
Active agents may alternatively be administered in methods of this
30 invention by inhalation, via the nasal or oral routes, e.g., in a spray formulation also containing a suitable carrier.

Exemplary active agents useful in methods of the invention will now be described by reference to illustrative synthetic schemes for their general preparation below and the specific examples that follow. Artisans will
35 recognize that, to obtain the various compounds herein, starting materials may be suitably selected so that the ultimately desired substituents will be carried through the reaction scheme with or without protection as appropriate to yield the desired product. Alternatively, it may be necessary or desirable to

- 5 employ, in the place of the ultimately desired substituent, a suitable group that may be carried through the reaction scheme and replaced as appropriate with the desired substituent. Unless otherwise specified, the variables are as defined above in reference to Formula (I).

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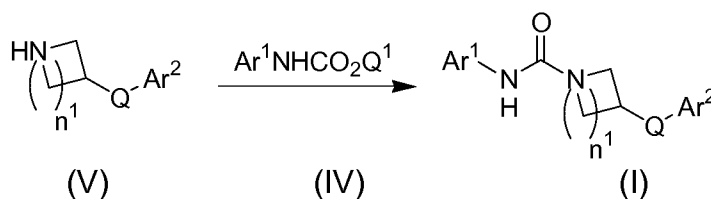
SCHEME A



- Referring to Scheme A, a carbamate of formula (IV) may be obtained by reacting a compound of formula (II) with a compound of formula (III), in which Q¹ represents a substituted or unsubstituted phenyl group, in a solvent such as dimethylformamide, acetonitrile or tetrahydrofuran, with or without a base such as pyridine, triethylamine, or diisopropylethylamine, at a temperature from about 0 °C to about 80 °C. Preferably, Q¹ is phenyl, and the reaction occurs with or without a base such as pyridine, in acetonitrile, at room temperature. In certain embodiments, the reaction occurs with 1.0 equivalent of a compound of formula (III) in the presence of 1.0 to 1.5 equivalents of base and 1.0 equivalent of a compound of formula (II). In other embodiments, the reaction occurs with 0.5 equivalent of a compound of formula (III) and 1.0 equivalent of a compound of formula (II) in the absence of base.

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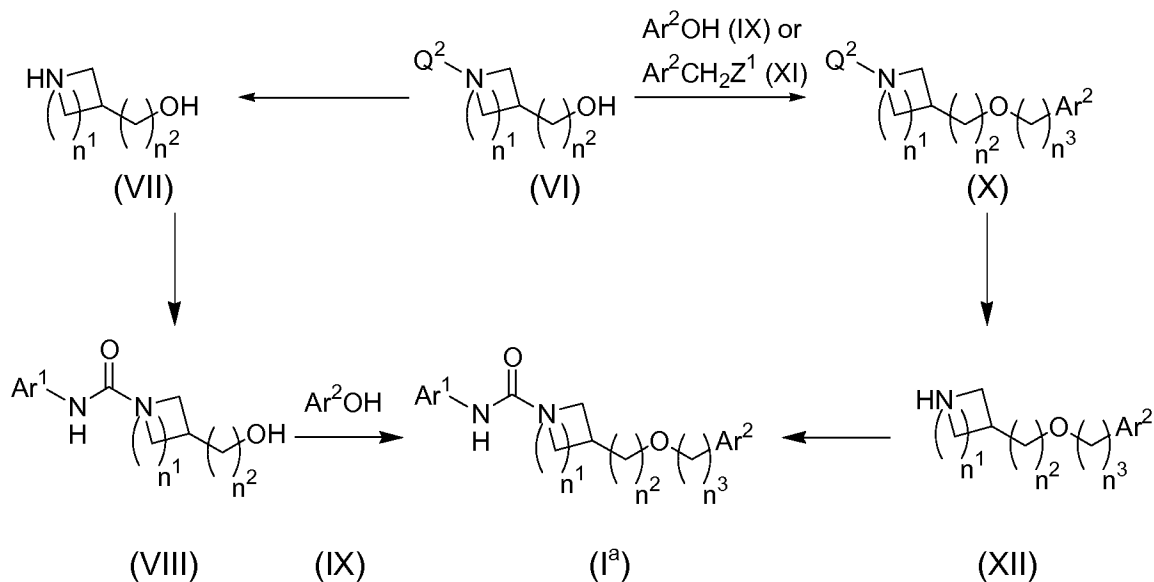
SCHEME B



5 Referring to Scheme B, a compound of formula (I) is obtained by reacting a compound of Formula (V) with a compound of formula (IV) in a solvent such as dimethylformamide or acetonitrile at a temperature from about rt to about 120 °C. Preferably, Q¹ is phenyl, and the reaction is performed in acetonitrile with heating from about rt to about 50 °C.

10

SCHEME C



15

Referring to Scheme C, compounds of formula (I^a) where n² is 1 or 2 and n³ is 0 or 1 are prepared from compounds of formula (VI). One skilled in the art would recognize that compounds of Formula (I^a) represent a subset of compounds of formula (I). Substituent Q² is a suitable amino protecting group compatible with the transformations in Scheme C. Preferably, Q² is tert-butylloxycarbonyl (BOC). An amine of formula (VII) is obtained by deprotecting a compound of formula (VI) under suitable Q² deprotection conditions. When Q² is BOC, removal may be preferably affected with HCl, trifluoroacetic acid (TFA) or formic acid in a solvent such as diethyl ether (Et₂O), DCM, or 1,4-dioxane. Alternatively, BOC removal may be effected in neat TFA or formic acid. A compound of formula (VIII) is obtained by reacting a compound of formula (VII) with a compound of formula (IV).

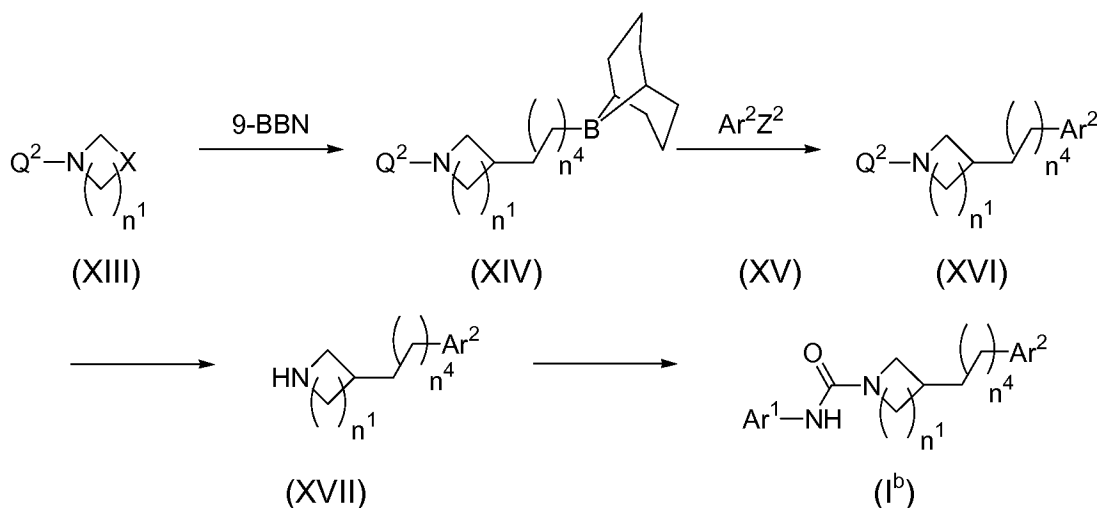
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Compounds of formula (VIII) are reacted with an alcohol of formula (IX) under Mitsunobu reaction conditions in a solvent such as tetrahydrofuran (THF), 1,2-dichloroethane (DCE), DCM, or Et₂O at a temperature from about

5 0 °C to 80 °C to produce a compound of Formula (I^a), where n³ is 0. In preferred embodiments, diisopropyl azodicarboxylate and polymer supported triphenyl phosphine in THF is employed at rt.

In alternative embodiments, compounds of formula (X), where n³ is 0, are obtained by reacting a compound of formula (VI) with a phenol of formula
 10 (IX) under Mitsunobu reaction conditions in a solvent such as tetrahydrofuran (THF), 1,2-dichloroethane (DCE), DCM, or Et₂O at a temperature from about 0 °C to 80 °C. Preferably, diisopropyl azodicarboxylate and polymer supported triphenyl phosphine in THF is employed at rt. In alternative
 15 embodiments, compounds of formula (X), where n³ is 1, are obtained by reacting a compound of formula (VI) with a benzylating agent of formula (XI), where Z¹ is I, Br, or Cl, under basic conditions in a solvent such as tetrahydrofuran (THF), dimethylformamide (DMF), or dimethylsulfoxide (DMSO) at a temperature from about 0 °C to 100 °C. Preferably, sodium
 20 hydride and Ar²CH₂Br in THF is employed at rt with tetrabutylammonium iodide as a catalyst. Deprotection of Q² from a compound of formula (X) under general deprotection conditions provides compounds of formula (XII). Where Q² is BOC, deprotection was achieved following the steps previously described above. Compounds of formula (I^a) are obtained by reacting a
 25 compound of formula (XII) with either a compound of formula (IV) or with a compound Ar¹NH₂ in the presence of di-(N-succinimidyl) carbonate.

Scheme D



30

Referring to Scheme D, compounds of formula (I^b), where n¹ is 1 or 2 and n⁴ is 0 or 1, are prepared from compounds of formula (XIII), where X is

5 $>C=CH_2$ or $>CHCH=CH_2$ utilizing a suitable protecting group Q^2 compatible with subsequent transformations. One skilled in the art would recognize that compounds of Formula (I^b) represent a subset of compounds of formula (I). In preferred embodiments Q^2 is BOC. A compound of formula (XIV) is obtained by reacting a compound of formula (XIII) under hydroboration conditions,
10 preferably using 9-borabicyclo[3.3.1]nonane as the borylating agent in a solvent such as THF at a temperature from about rt to about 80 °C. Compounds of formula (XVI) are obtained via palladium-mediated cross-coupling with reagents of formula (XV), where Z^2 is Br or I, with boronic esters of formula (XIV). In certain embodiments, boronic esters of formula (XIV) are treated with compounds of formula (XV) in the presence of a base such as
15 K_3PO_4 , K_2CO_3 , or KF in a suitable polar solvent such as CH_3CN , 1,2-dimethoxyethane (DME), tetrahydrofuran (THF), water, or a mixture thereof, at a temperature from about 50 °C to about 180 °C using conventional heating or a microwave reactor. An amine of formula (XVII) is obtained by
20 deprotecting a compound of formula (XVI) with a reagent under suitable Q^2 deprotection conditions. When Q^2 is BOC, deprotection may be affected using the methods described previously in Scheme C. A compound of formula (I^b) is obtained by reacting a compound of formula (XVII) with a compound of formula (IV). In alternative embodiments, a compound of
25 formula (I^b) is obtained by reacting a compound of formula (XVII) with a compound of formula Ar^1NH_2 in the presence of di-(N-succinimidyl)carbonate.

Compounds of formula (I) may be converted to their corresponding salts by applying general techniques described in the art. For example, a
30 compound of formula (I) may be treated with trifluoroacetic acid, HCl, or citric acid in a solvent such as Et_2O , 1,4-dioxane, DCM, THF, or MeOH to provide the corresponding salt forms.

Compounds prepared according to the schemes described above may be obtained as single enantiomers, diastereomers, or regioisomers, by enantio-, diastereo-, or regio-specific synthesis, or by resolution. Compounds
35 prepared according to the schemes above may alternatively be obtained as racemic (1:1) or non-racemic (not 1:1) mixtures or as mixtures of diastereomers or regioisomers. Where racemic and non-racemic mixtures of

5 enantiomers are obtained, single enantiomers may be isolated using conventional separation methods, such as chiral chromatography, recrystallization, diastereomeric salt formation, derivatization into diastereomeric adducts, biotransformation, or enzymatic transformation. Where regioisomeric or diastereomeric mixtures are obtained, single isomers
10 may be separated using conventional methods such as chromatography or crystallization.

The following specific examples are provided to further illustrate the invention and various preferred embodiments.

EXAMPLES

15 In preparing the examples listed below, the following general experimental and analytical methods were used.

Reaction mixtures were stirred under a nitrogen atmosphere at room temperature (rt) unless otherwise noted. Where solutions or mixtures are concentrated, they are typically concentrated under reduced pressure using a
20 rotary evaporator. Where solutions are dried, they are typically dried over a drying agent such as MgSO₄ or Na₂SO₄, unless otherwise noted.

Microwave reactions were carried out in either a CEM Discover or a Biotage Initiator™ Microwave at specified temperatures.

25 Normal-phase flash column chromatography (FCC) was performed on pre-packed Isco silica gel columns using 2 N NH₃ in methanol/CH₂Cl₂ or EtOAc/hexanes as eluents.

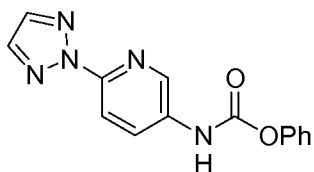
30 Reversed-Phase High Performance Liquid Chromatography (HPLC) was performed using: Shimadzu instrument with a Phenomenex Gemini column 5 μm C18 (150 x 21.2 mm) or Waters Xterra RP18 OBD column 5 μm (100 x 30 mm), a gradient of 95:5 to 0:100 water (0.05% TFA)/CH₃CN (0.05% TFA), a flow rate of 80 mL/min, and detection at 254 nM.

Mass spectra were obtained on an Agilent series 1100 MSD using electrospray ionization (ESI) in positive mode unless otherwise indicated.

35 NMR spectra were obtained on either a Bruker model DPX400 (400 MHz), DPX500 (500 MHz) or DRX600 (600 MHz) spectrometer. The format of the ¹H NMR data below is: chemical shift in ppm down field of the tetramethylsilane reference (multiplicity, coupling constant *J* in Hz, integration).

- 5 Chemical names were generated using ChemDraw Ultra 6.0.2 (CambridgeSoft Corp., Cambridge, MA) or ACD/Name Version 9 (Advanced Chemistry Development, Toronto, Ontario, Canada).

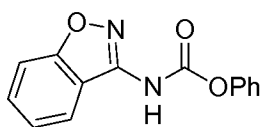
Intermediate 1: (6-[1,2,3]Triazol-2-yl-pyridin-3-yl)-carbamic acid phenyl ester.



- To a solution consisting of 6-[1,2,3]triazol-2-yl-pyridin-3-ylamine (1.00 g, 6.21 mmol) in CH₃CN (10 mL) was added phenyl chloroformate (0.389 mL, 3.10 mmol) dropwise at rt. After 16 h, the reaction mixture was diluted with EtOAc (30 mL) and washed with saturated aq. NaCl. The organic layer was separated, dried (Na₂SO₄), and concentrated. The crude residue was purified (FCC) to give the title compound as a white solid (0.808 g, 93%). MS (ESI⁺): calcd. for C₁₄H₁₁N₅O₂, 281.09; m/z found, 282.1 (M+H)⁺.
- 15

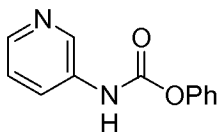
- Intermediates 2 to 10 were prepared using methods analogous to those described for intermediate 1, using the appropriate starting material.
- 20

Intermediate 2: Benzo[d]isoxazol-3-yl-carbamic acid phenyl ester.

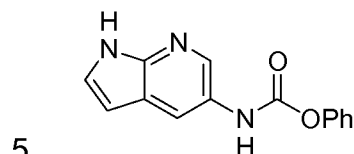


- MS (ESI⁺): calcd. for C₁₄H₁₀N₂O₃, 254.07; m/z found, 255.1 (M+H)⁺.
- 25

Intermediate 3: Pyridin-3-yl-carbamic acid phenyl ester.

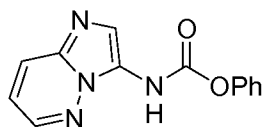


- MS (ESI⁺): calcd. for C₁₂H₁₀N₂O₂, 214.07; m/z found, 215.1 (M+H)⁺.
- 30 Intermediate 4: (1H-Pyrrolo[2,3-b]pyridin-5-yl)-carbamic acid phenyl ester.



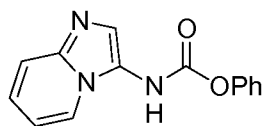
MS (ESI⁺): calcd. for C₁₄H₁₁N₃O₂, 253.09; m/z found, 254.1 (M+H)⁺.

Intermediate 5: Imidazo[1,2-b]pyridazin-3-yl-carbamic acid phenyl ester.



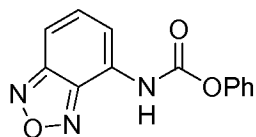
MS (ESI⁺): calcd. for C₁₃H₁₀N₄O₂, 254.08; m/z found, 255.1 (M+H)⁺.

Intermediate 6: Imidazo[1,2-a]pyridin-3-yl-carbamic acid phenyl ester.



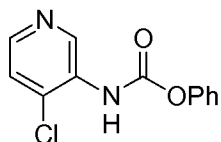
MS (ESI⁺): calcd. for C₁₄H₁₁N₃O₂, 253.09; m/z found, 254.1 (M+H)⁺.

Intermediate 7: N-2,1,3-Benzoxadiazol-4-yl-carbamic acid phenyl ester.



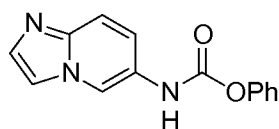
MS (ESI⁺): calcd. for C₁₃H₉N₃O₃, 255.06; m/z found, 256.1 (M+H)⁺.

Intermediate 8: (4-Chloro-pyridin-3-yl)-carbamic acid phenyl ester.

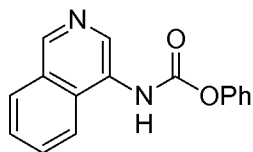


MS (ESI⁺): calcd. for C₁₂H₉ClN₂O₂, 248.04; m/z found, 249.1 (M+H)⁺.

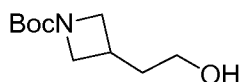
Intermediate 9: Imidazo[1,2-a]pyridin-6-yl-carbamic acid phenyl ester.



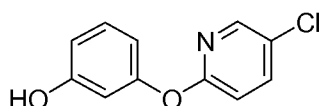
5

Intermediate 10: Isoquinolin-4-yl-carbamic acid phenyl ester.

MS (ESI⁺): calcd. for C₁₆H₁₂N₂O₂, 264.09; found, 265.1 (M+H)⁺.

10 Intermediate 11: 3-(2-Hydroxy-ethyl)-azetidine-1-carboxylic acid tert-butyl ester.

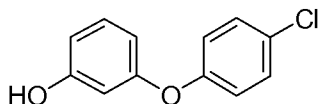
A solution consisting of 3-carboxymethyl-azetidine-1-carboxylic acid tert-butyl ester (10.0 g, 46.5 mmol) and THF (15 mL) was cooled to 0 °C and treated with a 1 M solution of borane in THF (70 mL, 70 mmol), then slowly warmed to rt. After stirring for 18 h, the reaction mixture was quenched with 2 N NaOH (100 mL), diluted with H₂O (100 mL), and extracted with Et₂O (3x200 mL). The organic layers were combined, dried (MgSO₄), and concentrated to give the title compound as a colorless oil (9.03 g, 97%). MS (ESI⁺): calcd. for C₁₀H₁₉NO₃, 201.14; m/z found, 224.2 (M+Na)⁺. ¹H NMR (d₆-DMSO): 4.37 (t, J = 5.1 Hz, 1H), 3.95-3.79 (m, 2H), 3.53-3.43 (m, 2H), 3.37 (dd, J = 11.5, 6.2 Hz, 2H), 2.62-2.52 (m, 1H), 1.66 (dd, J = 13.8, 6.4 Hz, 2H), 1.39-1.33 (m, 9H).

25 Intermediate 12: 3-(5-Chloro-pyridin-2-yloxy)-phenol.

To a solution consisting of 5-chloro-2-fluoropyridine (1.3 mL, 13 mmol) and resorcinol (2.1 g, 19 mmol) and DMSO (30 mL) was added Cs₂CO₃ (6.3 g, 19 mmol). The reaction vessel was heated at 40 °C for 22 hours then diluted with CH₂Cl₂ (60 mL) and washed with saturated aq. NH₄Cl (100 mL). The organic layer was dried (MgSO₄) and concentrated. The crude residue was purified (FCC) to give the title compound (1.0 g, 37%). MS (ESI⁺): calcd. for C₁₁H₈ClNO₂, 221.02; m/z found, 222.1 (M+H)⁺. ¹H NMR (CDCl₃): 8.14 (d,

- 5 $J = 2.6$ Hz, 1H), 7.65 (dd, $J = 8.7, 2.7$ Hz, 1H), 7.26-7.22 (m, 1H), 6.89 (dd, $J = 8.7, 0.5$ Hz, 1H), 6.69-6.65 (m, 2H), 6.60 (t, $J = 2.3$ Hz, 1H).

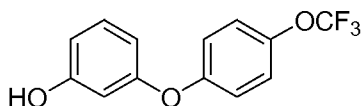
Intermediate 13: 3-(4-Chloro-phenoxy)-phenol.



- 10 A suspension of resorcinol (12.58 g, 114.2 mmol), 4-bromochlorobenzene (10.94 g, 57.14 mmol), dimethylglycine HCl (2.39 g, 17.1 mmol), and cesium carbonate (37.2 g, 114 mmol) in DMA (80 mL) was treated with CuI (1.09 g, 5.71 mmol) and heated at 90 °C for 16 h. The reaction mixture was cooled to room temperature, diluted with ethyl acetate
- 15 (500 mL) and extracted with 1 N NaOH (400 mL), and brine (200 mL). The organic layer was dried (MgSO₄) and concentrated. The residue was purified (FCC) to give the title compound as a yellow oil (7.6 g, 60%). MS (ESI⁺): calcd. for C₁₂H₉ClNO₂, 220.03; m/z found, 221.1 (M+H)⁺. ¹H NMR (d₆-DMSO): 9.63 (s, 1H), 7.43 (d, $J = 8.9$ Hz, 2H), 7.17 (t, $J = 8.1$ Hz, 1H), 7.03 (d, $J = 8.9$
- 20 Hz, 2H), 6.58-6.54 (m, 1H), 6.46-6.41 (m, 1H), 6.38 (t, $J = 2.3$ Hz, 1H).

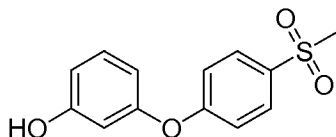
Intermediates 14 to 15 were prepared using methods analogous to those described for intermediate 13, using the appropriate starting material.

- 25 Intermediate 14: 3-(4-Trifluoromethoxy-phenoxy)-phenol.



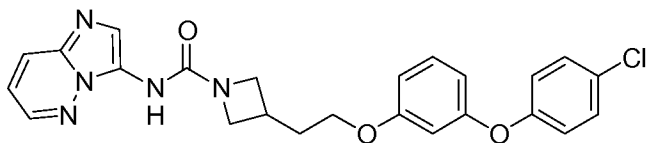
MS (ESI⁻): calcd. for C₁₃H₉F₃O₃, 270.05; m/z found, 269.0 (M-H)⁻.

Intermediate 15: 3-(4-Methanesulfonyl-phenoxy)-phenol.



- 30 MS (ESI⁻): calcd. for C₁₃H₁₂O₄S, 264.05; m/z found, 263.0 (M-H)⁻.

5 Example 1: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid imidazo[1,2-b]pyridazin-3-ylamide.



10 Step A: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid tert-butyl ester. A solution consisting of 3-(2-hydroxy-ethyl)-azetidine-1-carboxylic acid tert-butyl ester (4.39 g, 21.8 mmol), 3-(4-chloro-phenoxy)-phenol (4.81 g, 21.8 mmol) and THF (40 mL) was cooled at 0 °C, treated with polymer supported triphenyl phosphine (10.9 g, 32.7 mmol) then diisopropyl azodicarboxylate (6.62 g, 32.7 mmol) dropwise over 30 min. After stirring for 15 min at 0 °C, the reaction mixture was allowed to warm to room

15 temperature and stirred for 16 h. The reaction mixture was filtered, the filtrate evaporated, and the residue purified (FCC) to give the title compound as a clear oil (6.39 g, 73%). MS (ESI⁺): calcd. for C₂₂H₂₆ClNO₄, 403.16; m/z found, 426.2 (M+Na)⁺. ¹H NMR (d₆-DMSO): 7.43 (d, J = 9.0 Hz, 2H), 7.28 (t, J = 8.2 Hz, 1H), 7.03 (d, J = 9.0 Hz, 2H), 6.72 (dd, J = 8.3, 1.5 Hz, 1H), 6.59-6.55 (m, 2H), 3.98-3.84 (m, 4H), 3.59-3.50 (m, 2H), 2.70-2.58 (m, 1H), 2.02-1.89 (m, 2H), 1.36 (s, 9H).

20

Step B: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine, formic acid salt. 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid tert-butyl ester (545 mg, 1.35 mmol) was dissolved in formic acid (5.0

25 mL) and stirred for 4 h at room temperature. The formic acid was evaporated to provide the title compound as a yellow oil (460 mg, 100%). MS (ESI⁺): calcd. for C₁₇H₁₈ClNO₂, 303.10; m/z found, 304.1 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.41 (s, 1H), 7.43 (d, J = 8.8 Hz, 2H), 7.28 (t, J = 8.1 Hz, 1H), 7.03 (d, J = 8.8 Hz, 2H), 6.72 (dd, J = 8.2, 2.2 Hz, 1H), 6.60-6.53 (m, 2H), 3.94 (t, J = 6.2 Hz, 2H), 3.88 (t, J = 9.2 Hz, 2H), 3.67-3.59 (m, 2H), 2.95-2.85 (m, 1H), 2.03-1.96 (m, 2H).

30

Step C: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid imidazo[1,2-b]pyridazin-3-ylamide. To a solution consisting of 3-{2-[3-(4-chloro-phenoxy)-phenoxy]-ethyl}-azetidine formic acid salt (0.040 g, 0.13 mmol), TEA (0.018 mL, 0.13 mmol) and CH₃CN (1.0 mL) was added

35 imidazo[1,2-b]pyridazin-3-yl-carbamic acid phenyl ester (0.034 g, 0.13 mmol).

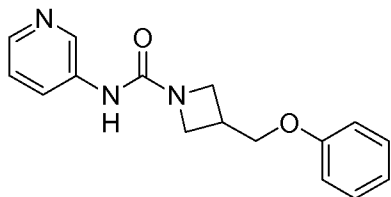
5 The reaction mixture was stirred at room temperature for 16 h, then diluted with EtOAc (40 mL) and washed with saturated aq. NaHCO₃ (40 mL). The organic layer was dried (Na₂SO₄) and concentrated. The crude residue was purified (FCC) to give the title compound (0.035 g, 57%). MS (ESI⁺): calcd. for C₂₄H₂₂ClN₅O₃, 463.14; m/z found, 464.1 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.55 (s, 1H), 8.52 (d, *J* = 4.4 Hz, 1H), 8.06 (d, *J* = 9.2 Hz, 1H), 7.64 (s, 1H), 7.43 (d, *J* = 7.3 Hz, 2H), 7.29 (t, *J* = 8.2 Hz, 1H), 7.19-7.15 (m, 1H), 7.04 (d, *J* = 7.4 Hz, 2H), 6.75 (d, *J* = 8.3 Hz, 1H), 6.61-6.55 (m, 2H), 4.09 (t, *J* = 8.1 Hz, 2H), 3.99 (t, *J* = 6.1 Hz, 2H), 3.77-3.66 (m, 2H), 2.83-2.69 (m, 1H), 2.10-2.01 (m, 2H).

15

Examples 2 to 48 were prepared using methods analogous to those described for Example 1, using the appropriate carbamate and phenol.

Example 2: 3-Phenoxymethyl-azetidone-1-carboxylic acid pyridin-3-ylamide, trifluoroacetate salt.

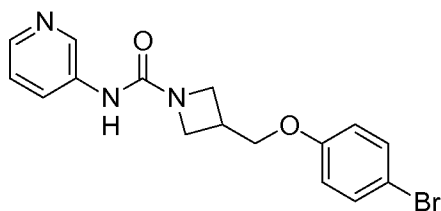
20



MS (ESI⁺): calcd. for C₁₆H₁₇N₃O₂, 283.13; m/z found, 284.2 (M+H)⁺. ¹H NMR (d₆-DMSO): 9.20 (s, 1H), 8.99 (d, *J* = 2.3 Hz, 1H), 8.39 (dd, *J* = 5.2, 1.0 Hz, 1H), 8.34-8.27 (m, 1H), 7.74 (dd, *J* = 8.6, 5.3 Hz, 1H), 7.35-7.26 (m, 2H), 7.00-6.91 (m, 3H), 4.19-4.11 (m, 4H), 3.86 (dd, *J* = 8.3, 5.5 Hz, 2H), 3.11-2.99 (m, 1H).

25

Example 3: 3-(4-Bromo-phenoxymethyl)-azetidone-1-carboxylic acid pyridin-3-ylamide.

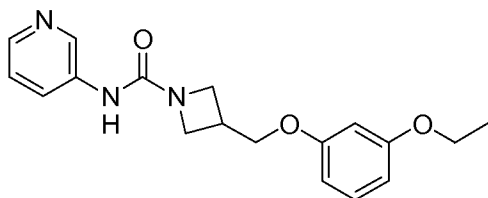


30

- 5 MS (ESI⁺): calcd. for C₁₆H₁₆BrN₃O₂, 361.04; m/z found, 362.1 (M+H)⁺.
¹H NMR (d₆-DMSO): 8.66 (d, *J* = 2.1 Hz, 1H), 8.62 (s, 1H), 8.14 (dd, *J* = 4.7, 1.5 Hz, 1H), 7.94-7.90 (m, 1H), 7.46 (d, *J* = 9.0 Hz, 2H), 7.26 (dd, *J* = 8.1, 4.4 Hz, 1H), 6.95 (d, *J* = 9.1 Hz, 2H), 4.16 (d, *J* = 6.6 Hz, 2H), 4.10 (t, *J* = 8.3 Hz, 2H), 3.80 (dd, *J* = 8.3, 5.4 Hz, 2H), 3.04-2.95 (m, 1H).

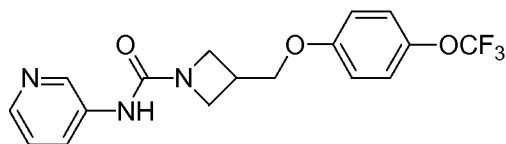
10

Example 4: 3-(3-Ethoxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide, trifluoroacetate salt.



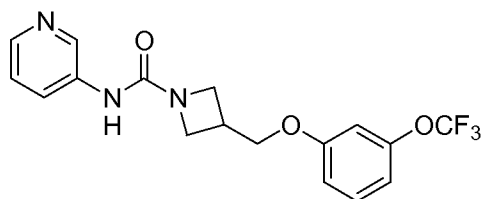
- 15 MS (ESI⁺): calcd. for C₁₈H₂₁N₃O₃, 327.16; m/z found, 328.2 (M+H)⁺. ¹H NMR (d₆-DMSO): 9.20 (s, 1H), 8.99 (d, *J* = 2.3 Hz, 1H), 8.39 (d, *J* = 4.5 Hz, 1H), 8.33-8.27 (m, 1H), 7.74 (dd, *J* = 8.6, 5.3 Hz, 1H), 7.17 (t, *J* = 8.3 Hz, 1H), 6.56-6.49 (m, 3H), 4.18-4.11 (m, 4H), 3.99 (q, *J* = 7.0 Hz, 2H), 3.85 (dd, *J* = 8.3, 5.5 Hz, 2H), 3.10-2.95 (m, 1H), 1.30 (t, *J* = 7.0 Hz, 3H).

- 20 Example 5: 3-(4-Trifluoromethoxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.



- 25 MS (ESI⁺): calcd. for C₁₇H₁₆F₃N₃O₃, 367.11; m/z found, 368.1 (M+H)⁺.
¹H NMR (d₆-DMSO): ¹H NMR (500 MHz, DMSO): 8.66 (d, *J* = 2.5 Hz, 1H), 8.62 (s, 1H), 8.14 (dd, *J* = 4.6, 1.5 Hz, 1H), 7.96-7.90 (m, 1H), 7.30 (d, *J* = 9.1 Hz, 2H), 7.26 (dd, *J* = 8.3, 4.7 Hz, 1H), 7.07 (d, *J* = 9.2 Hz, 2H), 4.19 (d, *J* = 6.6 Hz, 2H), 4.11 (t, *J* = 8.3 Hz, 2H), 3.82 (dd, *J* = 8.3, 5.4 Hz, 2H), 3.07-2.97 (m, 1H).

- 30 Example 6: 3-(3-Trifluoromethoxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.

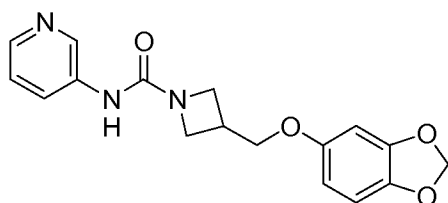


5

MS (ESI⁺): calcd. for C₁₇H₁₆F₃N₃O₃, 367.11; m/z found, 368.1 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.66 (d, *J* = 2.5 Hz, 1H), 8.62 (s, 1H), 8.14 (dd, *J* = 4.6, 1.4 Hz, 1H), 7.95-7.90 (m, 1H), 7.42 (t, *J* = 8.3 Hz, 1H), 7.26 (dd, *J* = 8.3, 4.7 Hz, 1H), 7.03 (dd, *J* = 8.3, 2.3 Hz, 1H), 6.97 (s, 1H), 6.95 (d, *J* = 9.1 Hz, 1H),
 10 4.22 (d, *J* = 6.7 Hz, 2H), 4.11 (t, *J* = 8.3 Hz, 2H), 3.82 (dd, *J* = 8.3, 5.4 Hz, 2H), 3.07-2.97 (m, 1H).

Example 7: 3-(Benzo[1,3]dioxol-5-yloxymethyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.

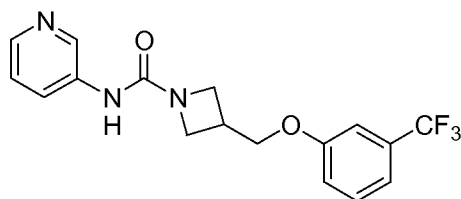


15

MS (ESI⁺): calcd. for C₁₇H₁₇N₃O₄, 327.12; m/z found, 328.2 (M+H)⁺. ¹H

NMR (d₆-DMSO): 8.66 (d, *J* = 2.1 Hz, 1H), 8.61 (s, 1H), 8.13 (dd, *J* = 4.6, 1.5 Hz, 1H), 7.94-7.90 (m, 1H), 7.26 (dd, *J* = 8.3, 4.7 Hz, 1H), 6.81 (d, *J* = 8.5 Hz, 1H), 6.66 (d, *J* = 2.5 Hz, 1H), 6.40 (dd, *J* = 8.5, 2.5 Hz, 1H), 5.96 (s, 2H), 4.12-
 20 4.05 (m, 4H), 3.78 (dd, *J* = 8.3, 5.4 Hz, 2H), 3.01-2.91 (m, 1H).

Example 8: 3-(3-Trifluoromethyl-phenoxy)methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide, trifluoroacetate salt.



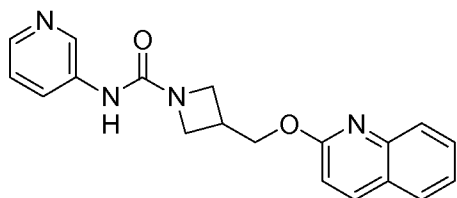
25

MS (ESI⁺): calcd. for C₁₇H₁₆F₃N₃O₂, 351.12; m/z found, 352.2 (M+H)⁺.

¹H NMR (d₆-DMSO): 9.29 (s, 1H), 9.04 (s, 1H), 8.42 (d, *J* = 4.9 Hz, 1H), 8.35 (d, *J* = 8.6 Hz, 1H), 7.80 (dd, *J* = 8.6, 5.3 Hz, 1H), 7.54 (t, *J* = 7.8 Hz, 1H), 7.33-7.26 (m, 3H), 4.27 (d, *J* = 6.5 Hz, 2H), 4.16 (t, *J* = 8.4 Hz, 2H), 3.93-3.83 (m, 2H), 3.12-3.00 (m, 1H).

5

Example 9: 3-(Quinolin-2-yloxymethyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.

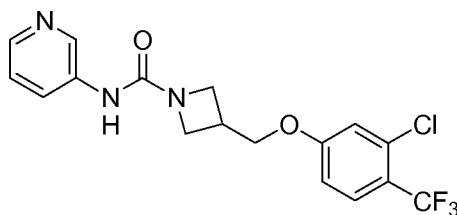


MS (ESI⁺): calcd. for C₁₉H₁₈N₄O₂, 334.14; m/z found, 335.2 (M+H)⁺. ¹H

10 NMR (d₆-DMSO): 8.68 (d, *J* = 2.3 Hz, 1H), 8.64 (s, 1H), 8.25 (d, *J* = 8.7 Hz, 1H), 8.14 (dd, *J* = 4.6, 1.2 Hz, 1H), 7.96-7.91 (m, 1H), 7.89 (dd, *J* = 8.0, 1.1 Hz, 1H), 7.78 (d, *J* = 8.4 Hz, 1H), 7.70-7.65 (m, 1H), 7.47-7.42 (m, 1H), 7.27 (dd, *J* = 8.3, 4.7 Hz, 1H), 7.04 (d, *J* = 8.8 Hz, 1H), 4.62 (d, *J* = 6.7 Hz, 2H), 4.14 (t, *J* = 8.4 Hz, 2H), 3.88 (dd, *J* = 8.3, 5.6 Hz, 2H), 3.14-3.04 (m, 1H).

15

Example 10: 3-(3-Chloro-4-trifluoromethyl-phenoxy)methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.

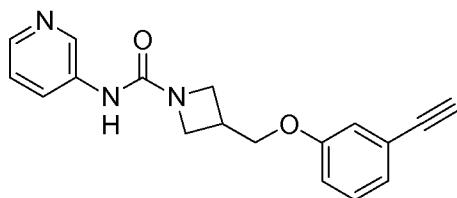


MS (ESI⁺): calcd. for C₁₇H₁₅ClF₃N₃O₂, 385.08; m/z found, 386.1

20 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.67 (d, *J* = 2.4 Hz, 1H), 8.63 (s, 1H), 8.14 (dd, *J* = 4.6, 1.3 Hz, 1H), 7.95-7.90 (m, 1H), 7.64 (d, *J* = 8.9 Hz, 1H), 7.39 (d, *J* = 3.0 Hz, 1H), 7.32 (dd, *J* = 8.9, 3.0 Hz, 1H), 7.26 (dd, *J* = 8.3, 4.7 Hz, 1H), 4.27 (d, *J* = 6.6 Hz, 2H), 4.11 (t, *J* = 8.3 Hz, 2H), 3.82 (dd, *J* = 8.3, 5.4 Hz, 2H), 3.07-2.97 (m, 1H).

25

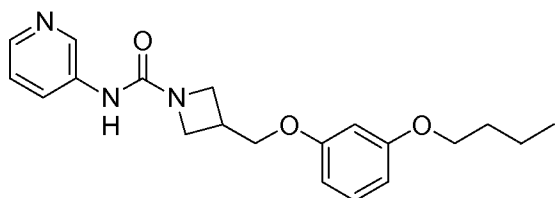
Example 11: 3-(3-Ethynyl-phenoxy)methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.



5 MS (ESI⁺): calcd. for C₁₈H₁₇N₃O₂, 307.13; m/z found, 308.2 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.66 (d, *J* = 2.3 Hz, 1H), 8.59 (s, 1H), 8.13 (dd, *J* = 4.7, 1.5 Hz, 1H), 7.96-7.88 (m, 1H), 7.30 (t, *J* = 8.1 Hz, 1H), 7.25 (dd, *J* = 8.3, 4.9 Hz, 1H), 7.09-6.99 (m, 3H), 4.19 (d, *J* = 6.7 Hz, 2H), 4.15 (s, 1H), 4.10 (t, *J* = 8.4 Hz, 2H), 3.81 (dd, *J* = 8.4, 5.4 Hz, 2H), 3.07-2.93 (m, 1H).

10

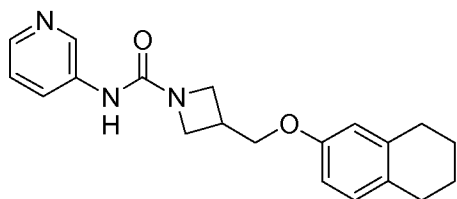
Example 12: 3-(3-Butoxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.



15 MS (ESI⁺): calcd. for C₂₀H₂₅N₃O₃, 355.19; m/z found, 356.2 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.66 (d, *J* = 2.4 Hz, 1H), 8.58 (s, 1H), 8.13 (dd, *J* = 4.6, 1.4 Hz, 1H), 7.95-7.89 (m, 1H), 7.25 (dd, *J* = 8.4, 4.6 Hz, 1H), 7.16 (dd, *J* = 8.8, 7.9 Hz, 1H), 6.56-6.48 (m, 3H), 4.14 (d, *J* = 6.6 Hz, 2H), 4.10 (t, *J* = 8.4 Hz, 2H), 3.93 (t, *J* = 6.5 Hz, 2H), 3.80 (dd, *J* = 8.4, 5.4 Hz, 2H), 3.05-2.93 (m, 1H), 1.73-1.60 (m, 2H), 1.50-1.34 (m, 2H), 0.93 (t, *J* = 7.4 Hz, 3H).

20

Example 13: 3-(5,6,7,8-Tetrahydro-naphthalen-2-yloxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.

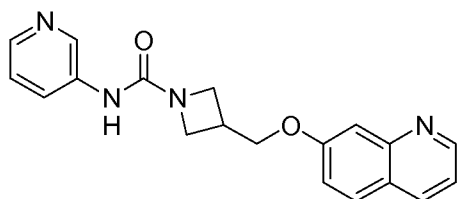


25 MS (ESI⁺): calcd. for C₂₀H₂₃N₃O₂, 337.18; m/z found, 338.2 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.66 (d, *J* = 2.0 Hz, 1H), 8.61 (s, 1H), 8.14 (dd, *J* = 4.6, 1.5 Hz, 1H), 7.95-7.90 (m, 1H), 7.26 (dd, *J* = 8.1, 4.8 Hz, 1H), 6.95 (d, *J* = 8.3 Hz, 1H), 6.69 (dd, *J* = 8.3, 2.7 Hz, 1H), 6.65 (d, *J* = 2.5 Hz, 1H), 4.12-4.06 (m, 4H), 3.79 (dd, *J* = 8.3, 5.4 Hz, 2H), 3.03-2.93 (m, 1H), 2.70-2.65 (m, 2H), 2.65-2.60 (m, 2H), 1.75-1.64 (m, 4H).

30

Example 14: 3-(Quinolin-7-yloxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.

5

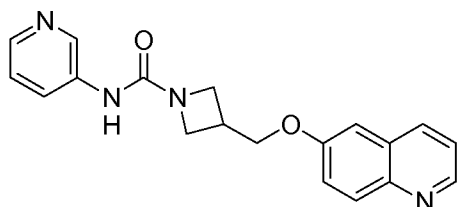


MS (ESI⁺): calcd. for C₁₉H₁₈N₄O₂, 334.14; m/z found, 335.2 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.83 (dd, *J* = 4.3, 1.7 Hz, 1H), 8.69 (s, 1H), 8.66 (s, 1H), 8.29 (dd, *J* = 8.2, 1.4 Hz, 1H), 8.15 (d, *J* = 4.0 Hz, 1H), 7.97-7.93 (m, 1H), 7.90 (d, *J* = 9.0 Hz, 1H), 7.46 (d, *J* = 2.4 Hz, 1H), 7.39 (dd, *J* = 8.2, 4.3 Hz, 1H), 7.31-7.26 (m, 2H), 4.36 (d, *J* = 6.6 Hz, 2H), 4.16 (t, *J* = 8.4 Hz, 2H), 3.89 (dd, *J* = 8.3, 5.4 Hz, 2H), 3.17-3.04 (m, 1H).

10

Example 15: 3-(Quinolin-6-yloxymethyl)-azetidine-1-carboxylic acid pyridin-3-ylamide, trifluoroacetate salt.

15

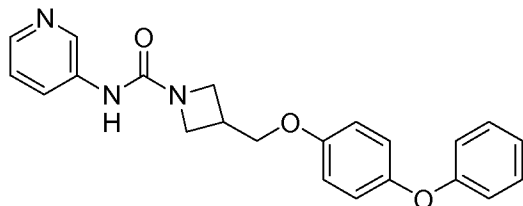


MS (ESI⁺): calcd. for C₁₉H₁₈N₄O₂, 334.14; m/z found, 335.2 (M+H)⁺. ¹H NMR (d₆-DMSO): 9.19 (s, 1H), 8.98 (s, 1H), 8.86 (dd, *J* = 4.4, 1.4 Hz, 1H), 8.46 (d, *J* = 8.1 Hz, 1H), 8.38 (d, *J* = 5.0 Hz, 1H), 8.28 (d, *J* = 8.6 Hz, 1H), 8.00 (d, *J* = 8.9 Hz, 1H), 7.72 (dd, *J* = 8.4, 5.3 Hz, 1H), 7.65 (dd, *J* = 8.4, 4.4 Hz, 1H), 7.59-7.50 (m, 2H), 4.36 (d, *J* = 6.5 Hz, 2H), 4.20 (t, *J* = 8.4 Hz, 2H), 3.92 (dd, *J* = 8.2, 5.5 Hz, 2H), 3.18-3.11 (m, 1H).

20

Example 16: 3-(4-Phenoxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.

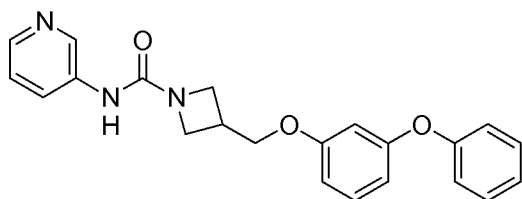
25



MS (ESI⁺): calcd. for C₂₂H₂₁N₃O₃, 375.16; m/z found, 376.2 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.67 (d, *J* = 2.2 Hz, 1H), 8.63 (s, 1H), 8.14 (dd, *J* = 4.7, 1.4 Hz, 1H), 7.96-7.89 (m, 1H), 7.38-7.32 (m, 2H), 7.26 (dd, *J* = 8.3, 4.7 Hz, 1H),

- 5 7.10-7.04 (m, 1H), 7.03-6.97 (m, 4H), 6.95-6.89 (m, 2H), 4.15 (d, $J = 6.6$ Hz, 2H), 4.11 (t, $J = 8.3$ Hz, 2H), 3.82 (dd, $J = 8.3, 5.4$ Hz, 2H), 3.07-2.97 (m, 1H).

Example 17: 3-(3-Phenoxy-phoxymethyl)-azetidone-1-carboxylic acid pyridin-3-ylamide.



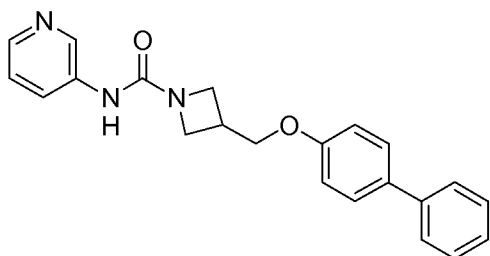
10

MS (ESI⁺): calcd. for C₂₂H₂₁N₃O₃, 375.16; m/z found, 376.2 (M+H)⁺. ¹H

NMR (d₆-DMSO): 8.66 (d, $J = 2.0$ Hz, 1H), 8.60 (s, 1H), 8.13 (dd, $J = 4.7, 1.5$ Hz, 1H), 7.95-7.88 (m, 1H), 7.40 (dd, $J = 8.6, 7.4$ Hz, 2H), 7.32-7.23 (m, 2H), 7.17-7.12 (m, 1H), 7.03 (dd, $J = 8.7, 1.0$ Hz, 2H), 6.76 (d, $J = 10.0$ Hz, 1H),

- 15 6.61 (t, $J = 2.3$ Hz, 1H), 6.56 (d, $J = 7.4$ Hz, 1H), 4.15 (d, $J = 6.7$ Hz, 2H), 4.09 (t, $J = 8.3$ Hz, 2H), 3.80 (dd, $J = 8.3, 5.5$ Hz, 2H), 3.04-2.95 (m, 1H).

Example 18: 3-(Biphenyl-4-yloxymethyl)-azetidone-1-carboxylic acid pyridin-3-ylamide.



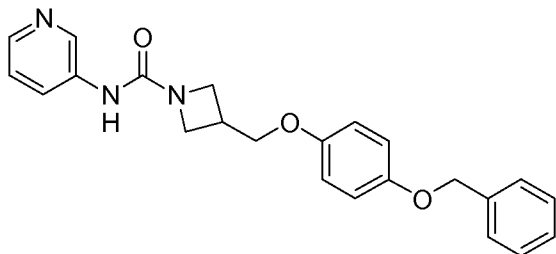
20

MS (ESI⁺): calcd. for C₂₂H₂₁N₃O₂, 359.16; m/z found, 360.2 (M+H)⁺. ¹H

NMR (d₆-DMSO): 8.67 (d, $J = 2.6$ Hz, 1H), 8.63 (s, 1H), 8.14 (dd, $J = 4.6, 1.4$ Hz, 1H), 7.93 (d, $J = 8.4$ Hz, 1H), 7.65-7.57 (m, 4H), 7.43 (t, $J = 7.7$ Hz, 2H), 7.31 (t, $J = 7.4$ Hz, 1H), 7.26 (dd, $J = 8.3, 4.7$ Hz, 1H), 7.07 (d, $J = 8.7$ Hz,

- 25 2H), 4.22 (d, $J = 6.6$ Hz, 2H), 4.13 (t, $J = 8.3$ Hz, 2H), 3.84 (dd, $J = 8.3, 5.4$ Hz, 2H), 3.09-3.00 (m, 1H).

Example 19: 3-(4-Benzyloxy-phoxymethyl)-azetidone-1-carboxylic acid pyridin-3-ylamide.



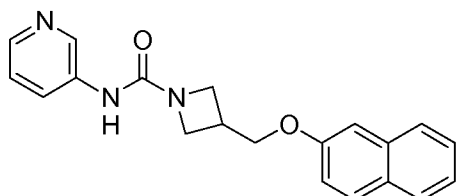
5

MS (ESI⁺): calcd. for C₂₃H₂₃N₃O₃, 389.17; m/z found, 390.2 (M+H)⁺. ¹H

NMR (d₆-DMSO): 8.66 (d, *J* = 2.2 Hz, 1H), 8.61 (s, 1H), 8.14 (d, *J* = 3.5 Hz, 1H), 7.95-7.90 (m, 1H), 7.45-7.41 (m, 2H), 7.40-7.36 (m, 2H), 7.35-7.29 (m, 1H), 7.26 (dd, *J* = 8.3, 4.7 Hz, 1H), 6.97-6.87 (m, 4H), 5.04 (s, 2H), 4.13-4.05 (m, 4H), 3.80 (dd, *J* = 8.3, 5.4 Hz, 2H), 3.04-2.90 (s, 1H).

10

Example 20: 3-(Naphthalen-2-yloxymethyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.



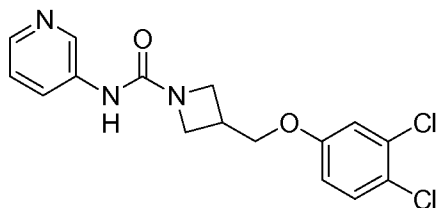
15

MS (ESI⁺): calcd. for C₂₀H₁₉N₃O₂, 333.15; m/z found, 334.2 (M+H)⁺. ¹H

NMR (d₆-DMSO): 8.68 (d, *J* = 2.5 Hz, 1H), 8.66 (s, 1H), 8.14 (dd, *J* = 4.6, 1.4 Hz, 1H), 7.96-7.92 (m, 1H), 7.86-7.79 (m, 3H), 7.46 (t, *J* = 7.5 Hz, 1H), 7.39 (d, *J* = 2.3 Hz, 1H), 7.35 (t, *J* = 7.5 Hz, 1H), 7.26 (dd, *J* = 8.3, 4.7 Hz, 1H), 7.19 (dd, *J* = 8.9, 2.5 Hz, 1H), 4.30 (d, *J* = 6.6 Hz, 2H), 4.15 (t, *J* = 8.3 Hz, 2H), 3.90-3.83 (m, 2H), 3.14-3.03 (m, 1H).

20

Example 21: 3-(3,4-Dichloro-phenoxy)methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.



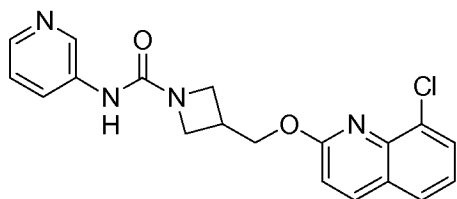
25

MS (ESI⁺): calcd. for C₁₆H₁₅Cl₂N₃O₂, 351.05; m/z found, 352.1 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.66 (d, *J* = 2.2 Hz, 1H), 8.62 (s, 1H), 8.14 (dd, *J* = 4.6, 1.4 Hz, 1H), 7.94-7.90 (m, 1H), 7.53 (d, *J* = 8.9 Hz, 1H), 7.29 (d, *J* = 2.9 Hz, 1H), 7.26 (dd, *J* = 8.1, 4.4 Hz, 1H), 7.00 (dd, *J* = 8.9, 2.9 Hz, 1H), 4.21 (d, *J* =

- 5 6.7 Hz, 2H), 4.10 (t, $J = 8.4$ Hz, 2H), 3.80 (dd, $J = 8.3, 5.4$ Hz, 2H), 3.05-2.95 (m, 1H).

Example 22: 3-(8-Chloro-quinolin-2-yloxymethyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.

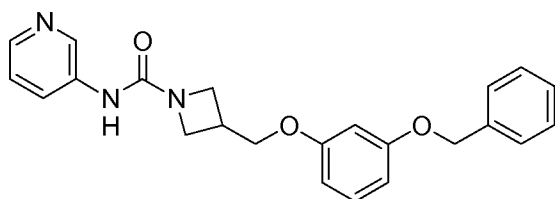


MS (ESI⁺): calcd. for C₁₉H₁₇ClN₄O₂, 368.10; m/z found, 369.1 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.66 (d, $J = 2.3$ Hz, 1H), 8.63 (s, 1H), 8.34 (d, $J = 8.9$ Hz, 1H), 8.14 (dd, $J = 4.6, 1.4$ Hz, 1H), 7.95-7.85 (m, 3H), 7.43 (t, $J = 7.8$ Hz, 1H), 7.26 (dd, $J = 8.2, 4.4$ Hz, 1H), 7.15 (d, $J = 8.8$ Hz, 1H), 4.70 (d, $J = 6.6$ Hz, 2H), 4.14 (t, $J = 8.4$ Hz, 2H), 3.90 (dd, $J = 8.3, 5.5$ Hz, 2H), 3.18-3.06 (m, 1H).

15

Example 23: 3-(3-Benzyloxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.

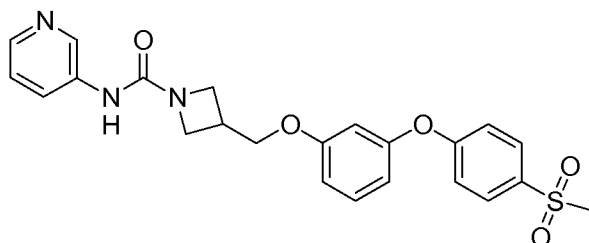


MS (ESI⁺): calcd. for C₂₃H₂₃N₃O₃, 389.17; m/z found, 390.2 (M+H)⁺. ¹H

NMR (d₆-DMSO): 8.67 (d, $J = 2.3$ Hz, 1H), 8.62 (s, 1H), 8.14 (d, $J = 3.4$ Hz, 1H), 7.96-7.89 (m, 1H), 7.46-7.42 (m, 2H), 7.42-7.36 (m, 2H), 7.35-7.30 (m, 1H), 7.26 (dd, $J = 8.3, 4.7$ Hz, 1H), 7.19 (t, $J = 8.1$ Hz, 1H), 6.64-6.58 (m, 2H), 6.58-6.52 (m, 1H), 5.08 (2H, s), 4.15 (d, $J = 6.7$ Hz, 2H), 4.10 (t, $J = 8.3$ Hz, 2H), 3.80 (dd, $J = 8.3, 5.4$ Hz, 2H), 3.04-2.94 (m, 1H).

25

Example 24: 3-[3-(4-Methanesulfonyl-phenoxy)-phenoxy-methyl]-azetidine-1-carboxylic acid pyridin-3-ylamide.



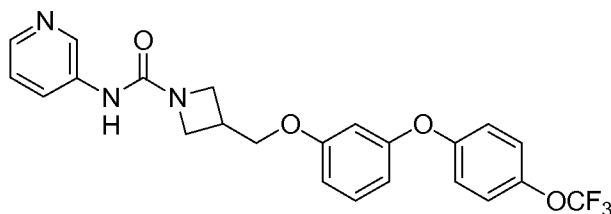
5

MS (ESI⁺): calcd. for C₂₃H₂₃N₃O₅S, 453.14; m/z found, 454.2 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.66 (d, *J* = 2.6 Hz, 1H), 8.62 (s, 1H), 8.13 (dd, *J* = 4.7, 1.4 Hz, 1H), 7.93-7.87 (m, 3H), 7.38 (t, *J* = 8.2 Hz, 1H), 7.25 (dd, *J* = 8.4, 4.7 Hz, 1H), 7.17 (d, *J* = 8.8 Hz, 2H), 6.89 (dd, *J* = 8.4, 2.4 Hz, 1H), 6.79 (t, *J* = 2.3 Hz, 1H), 6.72 (dd, *J* = 8.1, 2.2 Hz, 1H), 4.18 (d, *J* = 6.7 Hz, 2H), 4.10 (s, 2H), 3.81 (s, 2H), 3.19 (s, 3H), 3.07-2.94 (m, 1H).

10

Example 25: 3-[3-(4-Trifluoromethoxy-phenoxy)-phenoxy]methylazetidine-1-carboxylic acid pyridin-3-ylamide.



15

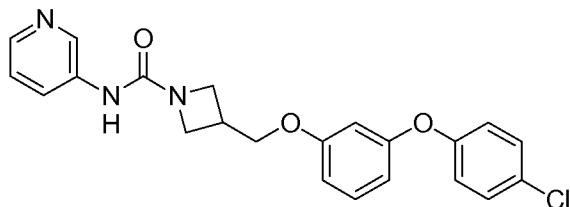
MS (ESI⁺): calcd. for C₂₃H₂₀F₃N₃O₄, 459.14; m/z found, 460.2 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.66 (d, *J* = 2.4 Hz, 1H), 8.61 (s, 1H), 8.13 (dd, *J* = 4.6, 1.4 Hz, 1H), 7.94-7.88 (m, 1H), 7.39 (d, *J* = 8.6 Hz, 2H), 7.32 (t, *J* = 8.2 Hz, 1H), 7.25 (dd, *J* = 8.4, 4.7 Hz, 1H), 7.11 (d, *J* = 9.1 Hz, 2H), 6.81 (dd, *J* = 8.2, 2.0 Hz, 1H), 6.69 (t, *J* = 2.3 Hz, 1H), 6.61 (dd, *J* = 8.1, 1.9 Hz, 1H), 4.16 (d, *J* = 6.7 Hz, 2H), 4.09 (t, *J* = 8.3 Hz, 2H), 3.80 (dd, *J* = 8.4, 5.5 Hz, 2H), 3.05-2.93 (m, 1H).

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Example 26: 3-[3-(4-Chloro-phenoxy)-phenoxy]methylazetidine-1-carboxylic acid pyridin-3-ylamide.

25



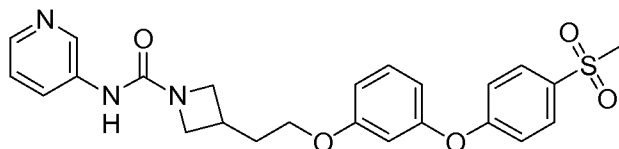
MS (ESI⁺): calcd. for C₂₂H₂₀ClN₃O₃, 409.12; m/z found, 410.2 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.66 (d, *J* = 2.5 Hz, 1H), 8.61 (s, 1H), 8.13 (dd, *J* = 4.6,

- 5 1.4 Hz, 1H), 7.94-7.89 (m, 1H), 7.43 (d, $J = 8.9$ Hz, 2H), 7.31 (t, $J = 8.2$ Hz, 1H), 7.26 (dd, $J = 8.3, 4.7$ Hz, 1H), 7.04 (d, $J = 8.9$ Hz, 2H), 6.79 (dd, $J = 8.3, 1.8$ Hz, 1H), 6.65 (t, $J = 2.3$ Hz, 1H), 6.59 (dd, $J = 8.1, 2.2$ Hz, 1H), 4.15 (d, $J = 6.7$ Hz, 2H), 4.09 (t, $J = 8.4$ Hz, 2H), 3.80 (dd, $J = 8.3, 5.4$ Hz, 2H), 3.03-2.95 (m, 1H).

10

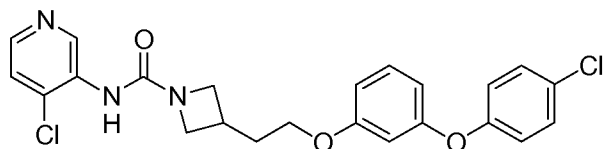
Example 27: 3-{2-[3-(4-Methanesulfonyl-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid pyridin-3-ylamide.



MS (ESI⁺): calcd. for C₂₄H₂₅N₃O₅S, 467.15; m/z found, 468.2 (M+H)⁺.

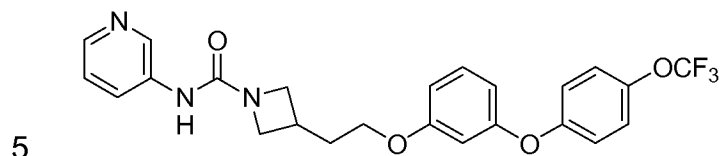
- 15 ¹H NMR (d₆-DMSO): 8.65 (d, $J = 2.4$ Hz, 1H), 8.56 (s, 1H), 8.12 (d, $J = 4.7$ Hz, 1H), 7.91-7.88 (m, 3H), 7.37 (t, $J = 8.2$ Hz, 1H), 7.25 (dd, $J = 8.3, 4.7$ Hz, 1H), 7.17 (d, $J = 8.8$ Hz, 2H), 6.85 (dd, $J = 8.2, 2.2$ Hz, 1H), 6.75-6.72 (m, 1H), 6.70 (dd, $J = 8.1$ Hz, 1H), 4.08 (t, $J = 8.2$ Hz, 2H), 4.00 (t, $J = 6.1$ Hz, 2H), 3.71 (dd, $J = 8.1, 5.9$ Hz, 2H), 3.19 (s, 3H), 2.79-2.68 (m, 1H), 2.08-1.97 (m, 2H).
- 20

Example 28: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.



- 25 MS (ESI⁺): calcd. for C₂₃H₂₁Cl₂N₃O₃, 457.10; m/z found, 458.1 (M+H)⁺.
¹H NMR (d₆-DMSO): 8.69 (s, 1H), 8.26 (d, $J = 5.2$ Hz, 1H), 8.15 (s, 1H), 7.55 (d, $J = 5.3$ Hz, 1H), 7.43 (d, $J = 8.8$ Hz, 2H), 7.29 (t, $J = 8.1$ Hz, 1H), 7.04 (d, $J = 8.8$ Hz, 2H), 6.74 (d, $J = 8.4$ Hz, 1H), 6.60 (s, 1H), 6.57 (d, $J = 10.1$ Hz, 1H), 4.09 (t, $J = 8.4$ Hz, 2H), 3.98 (t, $J = 6.0$ Hz, 2H), 3.72 (dd, $J = 7.6, 6.6$ Hz, 2H), 2.80-2.70 (m, 1H), 2.07-1.97 (m, 2H).
- 30

Example 29: 3-{2-[3-(4-Trifluoromethoxy-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid pyridin-3-ylamide.



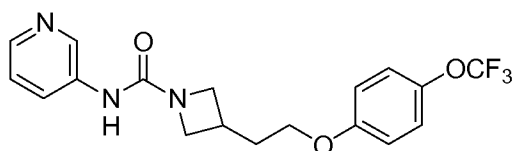
MS (ESI⁺): calcd. for C₂₄H₂₂F₃N₃O₄, 473.16; m/z found, 474.2 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.65 (d, *J* = 2.5 Hz, 1H), 8.56 (s, 1H), 8.13 (dd, *J* = 4.6, 1.3 Hz, 1H), 7.94-7.88 (m, 1H), 7.39 (d, *J* = 8.9 Hz, 2H), 7.31 (t, *J* = 8.2 Hz, 1H), 7.25 (dd, *J* = 8.3, 4.7 Hz, 1H), 7.11 (d, *J* = 9.0 Hz, 2H), 6.77 (dd, *J* = 8.3, 2.3 Hz, 1H), 6.66-6.62 (m, 1H), 6.59 (dd, *J* = 8.1, 2.2 Hz, 1H), 4.08 (t, *J* = 8.2 Hz, 2H), 3.99 (t, *J* = 6.1 Hz, 2H), 3.71 (dd, *J* = 8.1, 5.9 Hz, 2H), 2.79-2.67 (m, 1H), 2.05-1.98 (m, 2H).

10

Example 30: 3-[2-(4-Trifluoromethoxy-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide.

15



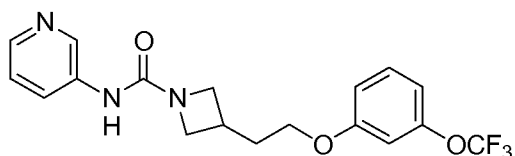
MS (ESI⁺): calcd. for C₁₈H₁₈F₃N₃O₃, 381.13; m/z found, 382.1 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.66 (d, *J* = 2.5 Hz, 1H), 8.57 (s, 1H), 8.13 (dd, *J* = 4.6, 1.4 Hz, 1H), 7.94-7.90 (m, 1H), 7.29 (d, *J* = 8.8 Hz, 2H), 7.25 (dd, *J* = 8.3, 4.7 Hz, 1H), 7.02 (d, *J* = 9.1 Hz, 2H), 4.09 (t, *J* = 8.2 Hz, 2H), 4.01 (t, *J* = 6.2 Hz, 2H), 3.72 (dd, *J* = 8.1, 5.9 Hz, 2H), 2.79-2.69 (m, 1H), 2.08-1.99 (m, 2H).

20

Example 31: 3-[2-(3-Trifluoromethoxy-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide.

25

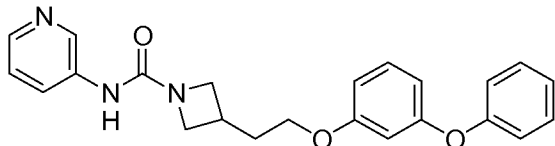


MS (ESI⁺): calcd. for C₁₈H₁₈F₃N₃O₃, 381.13; m/z found, 382.1 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.66 (d, *J* = 2.5 Hz, 1H), 8.57 (s, 1H), 8.13 (dd, *J* = 4.6, 1.4 Hz, 1H), 7.94-7.90 (m, 1H), 7.41 (t, *J* = 8.2 Hz, 1H), 7.25 (dd, *J* = 8.3, 4.7 Hz, 1H), 7.02-6.96 (m, 1H), 6.95-6.90 (m, 2H), 4.09 (t, *J* = 8.2 Hz, 2H), 4.03 (t, *J* = 6.2 Hz, 2H), 3.72 (dd, *J* = 8.1, 5.9 Hz, 2H), 2.80-2.69 (m, 1H), 2.04 (d, *J* = 7.2 Hz, 2H).

30

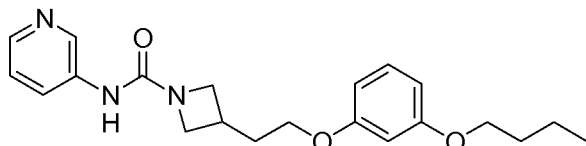
5 Example 32: 3-[2-(3-Phenoxy-phenoxy)-ethyl]-azetidone-1-carboxylic acid pyridin-3-ylamide.



MS (ESI⁺): calcd. for C₂₃H₂₃N₃O₃, 389.17; m/z found, 390.2 (M+H)⁺. ¹H

NMR (d₆-DMSO): 8.65 (d, *J* = 2.3 Hz, 1H), 8.56 (s, 1H), 8.13 (d, *J* = 4.6 Hz, 1H), 7.94-7.89 (m, 1H), 7.40 (t, *J* = 7.6 Hz, 2H), 7.30-7.22 (m, 2H), 7.15 (t, 1H), 7.02 (d, *J* = 8.6 Hz, 2H), 6.71 (d, *J* = 8.3 Hz, 1H), 6.58-6.51 (m, 2H), 4.08 (t, *J* = 8.2 Hz, 2H), 3.97 (t, *J* = 6.1 Hz, 2H), 3.74-3.66 (m, 2H), 2.78-2.67 (m, 1H), 2.05-1.95 (m, 2H).

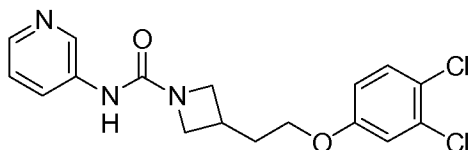
15 Example 33: 3-[2-(3-Butoxy-phenoxy)-ethyl]-azetidone-1-carboxylic acid pyridin-3-ylamide.



MS (ESI⁺): calcd. for C₂₁H₂₇N₃O₃, 369.21; m/z found, 370.2 (M+H)⁺. ¹H

NMR (d₆-DMSO): 8.66 (d, *J* = 2.3 Hz, 1H), 8.53 (s, 1H), 8.13 (dd, *J* = 4.6, 1.5 Hz, 1H), 7.95-7.88 (m, 1H), 7.24 (dd, *J* = 8.3, 4.7 Hz, 1H), 7.15 (t, *J* = 8.1 Hz, 1H), 6.52-6.44 (m, 3H), 4.09 (t, *J* = 8.2 Hz, 2H), 4.01-3.89 (m, 4H), 3.71 (dd, *J* = 8.2, 5.8 Hz, 2H), 2.85-2.67 (m, 1H), 2.10-1.93 (m, 2H), 1.74-1.57 (m, 2H), 1.50-1.33 (m, 2H), 0.93 (t, *J* = 7.4 Hz, 3H).

25 Example 34: 3-[2-(3,4-Dichloro-phenoxy)-ethyl]-azetidone-1-carboxylic acid pyridin-3-ylamide.



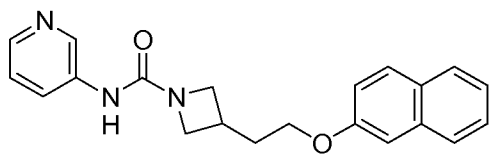
MS (ESI⁺): calcd. for C₁₇H₁₇Cl₂N₃O₂, 365.07; m/z found, 366.1 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.68 (d, *J* = 2.4 Hz, 1H), 8.58 (s, 1H), 8.15 (dd, *J* = 4.7, 1.1 Hz, 1H), 7.95 (d, *J* = 8.4 Hz, 1H), 7.52 (d, *J* = 8.9 Hz, 1H), 7.29 (dd, *J* = 8.3, 4.8 Hz, 1H), 7.23 (d, *J* = 2.9 Hz, 1H), 6.96 (dd, *J* = 8.9, 2.9 Hz, 1H), 4.08

- 5 (t, $J = 8.3$ Hz, 2H), 4.03 (t, $J = 6.2$ Hz, 2H), 3.71 (dd, $J = 8.2, 5.9$ Hz, 2H), 2.80-2.68 (m, 1H), 2.08-1.95 (m, 2H).

Example 35: 3-[2-(Naphthalen-2-yloxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide, trifluoroacetate salt.

10

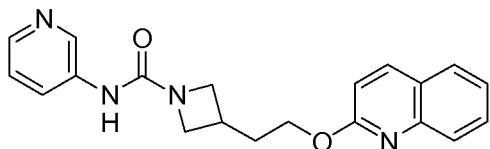


15

MS (ESI⁺): calcd. for C₂₁H₂₁N₃O₂, 347.16; m/z found, 348.2 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.98 (s, 1H), 8.91 (d, $J = 2.4$ Hz, 1H), 8.31 (d, $J = 5.0$ Hz, 1H), 8.20 (d, $J = 8.1$ Hz, 1H), 7.86-7.75 (m, 3H), 7.61 (dd, $J = 8.3, 5.3$ Hz, 1H), 7.45 (t, $J = 7.6$ Hz, 1H), 7.38-7.28 (m, 2H), 7.15 (dd, $J = 9.0, 2.5$ Hz, 1H), 4.21-4.07 (m, 4H), 3.83-3.73 (m, 2H), 2.88-2.75 (m, 1H), 2.17-2.07 (m, 2H).

Example 36: 3-[2-(Quinolin-2-yloxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide.

20

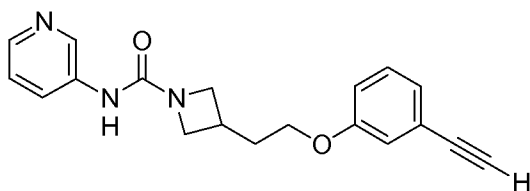


25

MS (ESI⁺): calcd. for C₂₀H₂₀N₄O₂, 348.16; m/z found, 349.2 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.66 (d, $J = 2.0$ Hz, 1H), 8.54 (s, 1H), 8.24 (d, $J = 8.7$ Hz, 1H), 8.12 (dd, $J = 4.6, 1.5$ Hz, 1H), 7.93-7.89 (m, 1H), 7.88 (dd, $J = 8.0, 1.2$ Hz, 1H), 7.77 (d, $J = 8.4$ Hz, 1H), 7.69-7.62 (m, 1H), 7.48-7.36 (m, 1H), 7.24 (dd, $J = 8.6, 4.4$ Hz, 1H), 7.00 (d, $J = 8.8$ Hz, 1H), 4.45 (t, $J = 6.3$ Hz, 2H), 4.11 (t, $J = 8.3$ Hz, 2H), 3.76-3.67 (m, 2H), 2.83-2.69 (m, 1H), 2.16-2.03 (m, 2H).

Example 37: 3-[2-(3-Ethynyl-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide, trifluoroacetate salt.

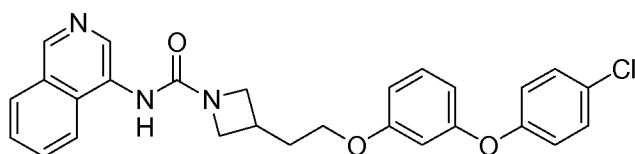
30



MS (ESI⁺): calcd. for C₁₉H₁₉N₃O₂, 321.15; m/z found, 322.2 (M+H)⁺. ¹H NMR (d₆-DMSO): 9.11 (s, 1H), 8.98 (d, $J = 2.3$ Hz, 1H), 8.37 (d, $J = 5.2$ Hz,

- 5 1H), 8.32-8.26 (m, 1H), 7.72 (dd, $J = 8.5, 5.3$ Hz, 1H), 7.29 (t, $J = 7.9$ Hz, 1H), 7.09-6.93 (m, 3H), 4.17-4.08 (m, 3H), 4.02 (t, $J = 6.1$ Hz, 2H), 3.76 (dd, $J = 8.3, 5.9$ Hz, 2H), 2.86-2.65 (m, 1H), 2.04 (dd, $J = 13.4, 6.3$ Hz, 2H).

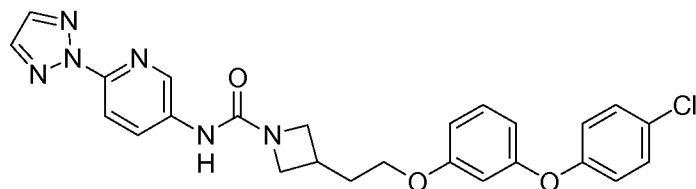
10 Example 38: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidone-1-carboxylic acid isoquinolin-4-ylamide.



MS (ESI⁺): calcd. for C₂₇H₂₄ClN₃O₃, 473.15; m/z found, 474.2 (M+H)⁺.

- 15 ¹H NMR (d₆-DMSO): 9.09 (s, 1H), 8.51 (d, $J = 12.9$ Hz, 2H), 8.11 (d, $J = 8.2$ Hz, 1H), 8.02 (d, $J = 8.0$ Hz, 1H), 7.78 (t, $J = 7.1$ Hz, 1H), 7.68 (t, $J = 7.1$ Hz, 1H), 7.43 (d, $J = 8.9$ Hz, 2H), 7.30 (t, $J = 8.2$ Hz, 1H), 7.04 (d, $J = 8.9$ Hz, 2H), 6.76 (dd, $J = 8.2, 1.9$ Hz, 1H), 6.62 (t, $J = 2.3$ Hz, 1H), 6.58 (dd, $J = 7.9, 2.0$ Hz, 1H), 4.15 (t, $J = 8.2$ Hz, 2H), 4.01 (t, $J = 6.2$ Hz, 2H), 3.78 (dd, $J = 8.0, 5.9$ Hz, 2H), 2.84-2.74 (m, 1H), 2.10-2.01 (m, 2H).

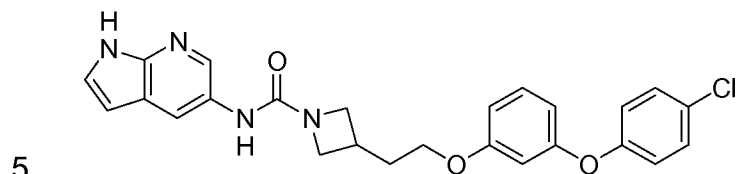
20 Example 39: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidone-1-carboxylic acid (6-[1,2,3]triazol-2-yl-pyridin-3-yl)-amide.



MS (ESI⁺): calcd. for C₂₅H₂₃ClN₆O₃, 490.15; m/z found, 491.1 (M+H)⁺.

- 25 ¹H NMR (d₆-DMSO): 8.82 (s, 1H), 8.65 (s, 1H), 8.20 (d, $J = 9.0$ Hz, 1H), 8.10 (s, 2H), 7.89 (d, $J = 9.0$ Hz, 1H), 7.43 (d, $J = 7.2$ Hz, 2H), 7.30 (t, $J = 7.5$ Hz, 1H), 7.04 (d, $J = 7.2$ Hz, 2H), 6.75 (d, $J = 8.2$ Hz, 1H), 6.62-6.60 (m, 1H), 6.58 (d, $J = 8.3$ Hz, 1H), 4.11 (t, $J = 8.1$ Hz, 2H), 3.99 (t, $J = 5.9$ Hz, 2H), 3.77-3.70 (m, 2H), 2.80-2.71 (m, 1H), 2.08-1.98 (m, 2H).

30 Example 40: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidone-1-carboxylic acid (1H-pyrrolo[2,3-b]pyridin-5-yl)-amide.



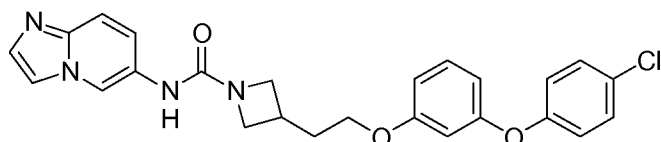
MS (ESI⁺): calcd. for C₂₅H₂₃ClN₄O₃, 462.15; m/z found, 463.2 (M+H)⁺.

¹H NMR (d₆-DMSO): 11.38 (s, 1H), 8.27 (s, 1H), 8.20 (d, *J* = 2.3 Hz, 1H), 8.01 (d, *J* = 2.2 Hz, 1H), 7.43 (d, *J* = 8.8 Hz, 2H), 7.40-7.35 (m, 1H), 7.30 (t, *J* = 8.2 Hz, 1H), 7.04 (d, *J* = 8.8 Hz, 2H), 6.75 (dd, *J* = 8.3, 2.3 Hz, 1H), 6.61 (t, *J* = 2.3 Hz, 1H), 6.59-6.55 (m, 1H), 6.36-6.32 (m, 1H), 4.06 (t, *J* = 8.2 Hz, 2H), 3.99 (t, *J* = 6.2 Hz, 2H), 3.69 (dd, *J* = 8.0, 5.8 Hz, 2H), 2.78-2.66 (m, 1H), 2.08-1.96 (m, 2H).

10

Example 41: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-
carboxylic acid imidazo[1,2-a]pyridin-6-ylamide.

15



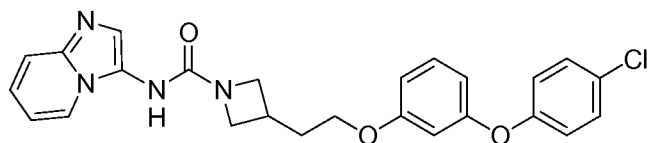
MS (ESI⁺): calcd. for C₂₅H₂₃ClN₄O₃, 462.15; m/z found, 463.2 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.88 (s, 1H), 8.37 (s, 1H), 7.91 (s, 1H), 7.48-7.40 (m, 4H), 7.30 (t, *J* = 8.3 Hz, 1H), 7.22 (d, *J* = 9.5 Hz, 1H), 7.04 (d, *J* = 7.7 Hz, 2H), 6.77-6.73 (m, 1H), 6.61-6.59 (m, 1H), 6.59-6.55 (m, 1H), 4.08 (t, *J* = 8.2 Hz, 2H), 3.98 (t, *J* = 6.3 Hz, 2H), 3.73-3.65 (m, 2H), 2.79-2.69 (m, 1H), 2.06-1.96 (m, 2H).

20

Example 42: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-
carboxylic acid imidazo[1,2-a]pyridin-3-ylamide.

25



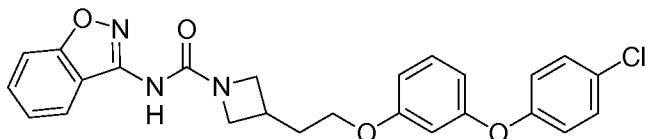
MS (ESI⁺): calcd. for C₂₅H₂₃ClN₄O₃, 462.15; m/z found, 463.2 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.44 (s, 1H), 8.04 (d, *J* = 7.0 Hz, 1H), 7.51 (d, *J* = 9.1 Hz, 1H), 7.45-7.40 (m, 2H), 7.34 (s, 1H), 7.30 (t, *J* = 8.2 Hz, 1H), 7.23-7.18 (m, 1H), 7.06-7.01 (m, 2H), 6.93-6.86 (m, 1H), 6.74 (d, *J* = 8.3 Hz, 1H), 6.62-6.53 (m, 2H), 4.08 (t, *J* = 8.4 Hz, 2H), 3.99 (t, *J* = 5.3 Hz, 2H), 3.72 (t, *J* = 6.5 Hz, 2H), 2.81-2.71 (m, 1H), 2.09-1.99 (m, 2H).

30

5

Example 43: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid benzo[d]isoxazol-3-ylamide.

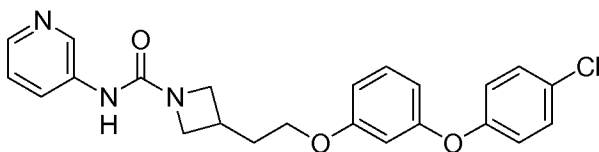


MS (ESI⁺): calcd. for C₂₅H₂₂ClN₃O₄, 463.13; m/z found, 464.1 (M+H)⁺.

10 ¹H NMR (d₆-DMSO): 9.70 (s, 1H), 7.99 (d, *J* = 8.1 Hz, 1H), 7.64-7.56 (m, 2H), 7.43 (d, *J* = 8.8 Hz, 2H), 7.32-7.26 (m, 2H), 7.04 (d, *J* = 8.8 Hz, 2H), 6.75 (dd, *J* = 8.3, 2.3 Hz, 1H), 6.61 (t, *J* = 2.3 Hz, 1H), 6.57 (dd, *J* = 8.0, 2.2 Hz, 1H), 4.15 (t, *J* = 8.0 Hz, 2H), 3.99 (t, *J* = 6.1 Hz, 2H), 3.79 (t, *J* = 5.8 Hz, 2H), 2.82-2.71 (m, 1H), 2.08-1.98 (m, 2H).

15

Example 44: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid pyridin-3-ylamide.

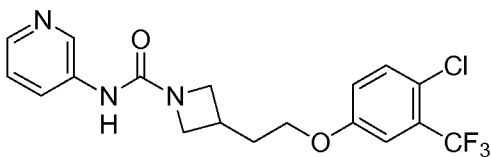


MS (ESI⁺): calcd. for C₂₃H₂₂ClN₃O₃, 423.13; m/z found, 424.2 (M+H)⁺.

20 ¹H NMR (d₆-DMSO): 8.65 (d, *J* = 2.4 Hz, 1H), 8.56 (s, 1H), 8.13 (dd, *J* = 4.6, 1.4 Hz, 1H), 7.93-7.88 (m, 1H), 7.43 (d, *J* = 8.9 Hz, 2H), 7.30 (t, *J* = 8.2 Hz, 1H), 7.25 (dd, *J* = 8.3, 4.6 Hz, 1H), 7.04 (d, *J* = 8.9 Hz, 2H), 6.78-6.71 (m, 1H), 6.63-6.55 (m, 2H), 4.08 (t, *J* = 8.2 Hz, 2H), 3.98 (t, *J* = 6.1 Hz, 2H), 3.73-3.65 (m, 2H), 2.78-2.66 (m, 1H), 2.05-1.96 (m, 2H).

25

Example 45: 3-[2-(4-Chloro-3-trifluoromethyl-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide.

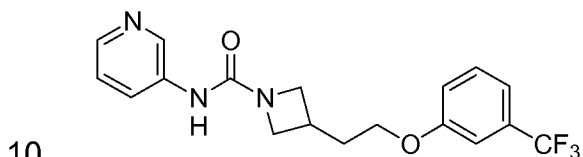


MS (ESI⁺): calcd. for C₁₈H₁₇ClF₃N₃O₂, 399.10; m/z found, 400.1

30 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.66 (d, *J* = 2.2 Hz, 1H), 8.57 (s, 1H), 8.13 (dd, *J* = 4.6, 1.4 Hz, 1H), 7.95-7.89 (m, 1H), 7.63 (d, *J* = 8.9 Hz, 1H), 7.33 (d, *J* = 2.9

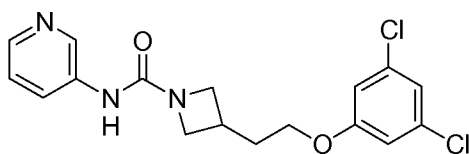
- 5 Hz, 1H), 7.29-7.23 (m, 2H), 4.13-4.04 (m, 4H), 3.72 (dd, $J = 8.1, 5.9$ Hz, 2H), 2.80-2.70 (m, 1H), 2.09-2.00 (m, 2H).

Example 46: 3-[2-(3-Trifluoromethyl-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide.



- MS (ESI⁺): calcd. for C₁₈H₁₈F₃N₃O₂, 365.14; m/z found, 366.2 (M+H)⁺.
¹H NMR (d₆-DMSO): 8.66 (d, $J = 2.2$ Hz, 1H), 8.56 (s, 1H), 8.13 (dd, $J = 4.6, 1.4$ Hz, 1H), 7.95-7.89 (m, 1H), 7.53 (t, $J = 8.0$ Hz, 1H), 7.31-7.19 (m, 4H), 4.14-4.03 (m, 4H), 3.73 (dd, $J = 8.1, 5.9$ Hz, 2H), 2.82-2.69 (m, 1H), 2.09-2.01
 15 (m, 2H).

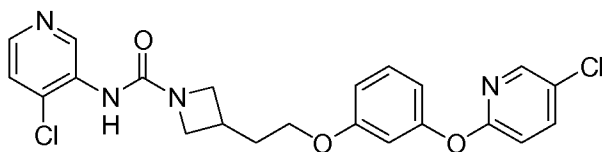
Example 47: 3-[2-(3,5-Dichloro-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide.



- MS (ESI⁺): calcd. for C₁₇H₁₇Cl₂N₃O₂, 365.07; m/z found, 366.1 (M+H)⁺.
¹H NMR (d₆-DMSO): 8.66 (d, $J = 2.2$ Hz, 1H), 8.56 (s, 1H), 8.13 (dd, $J = 4.6, 1.4$ Hz, 1H), 7.94-7.89 (m, 1H), 7.25 (dd, $J = 8.2, 4.6$ Hz, 1H), 7.15 (t, $J = 1.8$ Hz, 1H), 7.04 (d, $J = 1.8$ Hz, 2H), 4.11-4.01 (m, 4H), 3.71 (dd, $J = 8.2, 5.9$ Hz, 2H), 2.77-2.68 (m, 1H), 2.05-1.98 (m, 2H).

25

Example 48: 3-{2-[3-(5-Chloro-pyridin-2-yloxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.

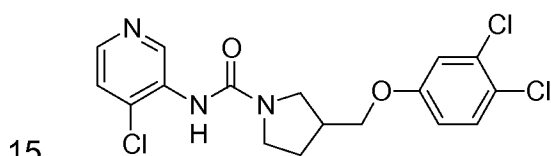


- MS (ESI⁺): calcd. for C₂₂H₂₀Cl₂N₄O₃, 458.09; m/z found, 459.1 (M+H)⁺.
¹H NMR (CDCl₃): 9.47 (s, 1H), 8.19 (d, $J = 5.2$ Hz, 1H), 8.16-8.12 (m, 1H), 7.65 (dd, $J = 8.7, 2.7$ Hz, 1H), 7.32-7.28 (m, 2H), 6.89 (dd, $J = 8.7, 0.4$ Hz,

- 5 1H), 6.76-6.70 (m, 2H), 6.65 (t, $J = 2.3$ Hz, 1H), 6.37 (s, 1H), 4.26 (t, $J = 8.2$ Hz, 2H), 4.00 (t, $J = 5.8$ Hz, 2H), 3.88 (dd, $J = 8.0, 5.7$ Hz, 2H), 2.99-2.87 (m, 1H), 2.15 (dd, $J = 13.3, 5.9$ Hz, 2H).

10 Examples 49 to 64 were prepared using methods analogous to those described for Example 1, using the appropriate carbamate, hydroxypyrrolidine-1-carboxylic acid tert-butyl ester and phenol.

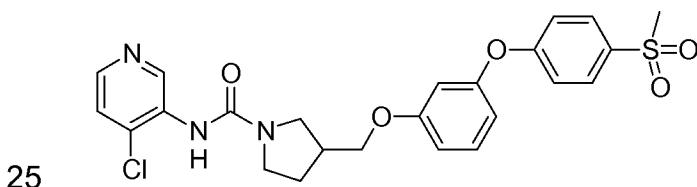
Example 49: N-(4-Chloropyridin-3-yl)-3-[(3,4-dichlorophenoxy)methyl]pyrrolidine-1-carboxamide.



MS (ESI): mass calcd. for $C_{17}H_{16}Cl_3N_3O_2$, 399.03; m/z found, 400.1 $[M+H]^+$. 1H NMR (CD_3OD): 8.79 (s, 1H), 8.24 (d, $J = 5.3$ Hz, 1H), 7.54 (d, $J = 5.3$ Hz, 1H), 7.40 (d, $J = 8.9$ Hz, 1H), 7.24-7.05 (m, 1H), 6.91 (dd, $J = 8.9, 2.9$ Hz, 1H), 4.14-4.04 (m, 1H), 4.04-3.95 (m, 1H), 3.82-3.72 (m, 1H), 3.72-3.63 (m, 1H), 3.63-3.48 (m, 1H), 3.48-3.37 (m, 1H), 2.88-2.73 (m, 1H), 2.30-2.17 (m, 1H), 2.03-1.90 (m, 1H).

20

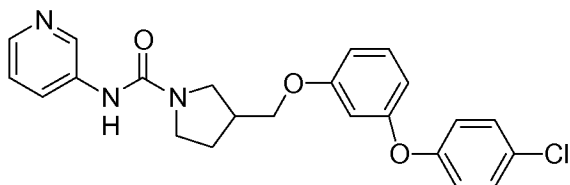
Example 50: N-(4-Chloropyridin-3-yl)-3-[(3-[4-(methylsulfonyl)phenoxy]phenoxy)methyl]pyrrolidine-1-carboxamide.



MS (ESI): mass calcd. for $C_{24}H_{24}ClN_3O_5S$, 501.11; m/z found, 502.1 $[M+H]^+$. 1H NMR (CD_3OD): 8.93-8.66 (m, 1H), 8.35-8.16 (m, 1H), 7.91 (d, $J = 8.8$ Hz, 2H), 7.62-7.44 (m, 1H), 7.44-7.25 (m, 1H), 7.24-7.06 (m, 2H), 6.94-6.79 (m, 1H), 6.78-6.60 (m, 2H), 4.16-3.95 (m, 2H), 3.83-3.63 (m, 2H), 3.64-3.51 (m, 2H), 3.16-3.04 (m, 3H), 2.91-2.72 (m, 1H), 2.33-2.13 (m, 1H), 2.04-1.76 (m, 1H).

30

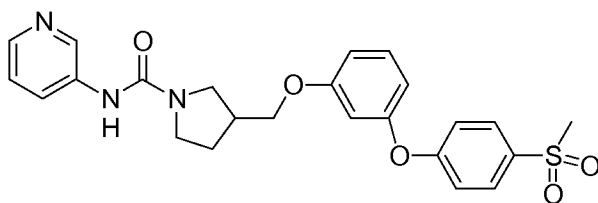
5 Example 51: 3-([3-(4-Chlorophenoxy)phenoxy]methyl)-N-pyridin-3-ylpyrrolidine-1-carboxamide.



MS (ESI): mass calcd. for C₂₃H₂₂ClN₃O₃, 423.13; m/z found, 424.2

[M+H]⁺. ¹H NMR (CD₃OD): 8.73-8.58 (m, 1H), 8.26-8.09 (m, 1H), 8.09-7.92 (m, 1H), 7.47-7.19 (m, 4H), 7.16-6.91 (m, 3H), 6.87-6.68 (m, 1H), 6.64-6.50 (m, 1H), 4.11-3.86 (m, 2H), 3.84-3.61 (m, 2H), 3.58-3.46 (m, 2H), 2.92-2.72 (m, 1H), 2.26-2.08 (m, 1H), 2.04-1.77 (m, 1H).

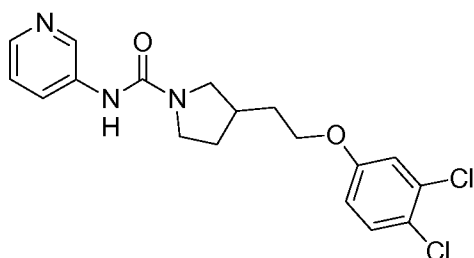
15 Example 52: 3-([3-[4-(Methylsulfonyl)phenoxy]phenoxy]methyl)-N-pyridin-3-ylpyrrolidine-1-carboxamide.



MS (ESI): mass calcd. for C₂₄H₂₅N₃O₅S, 467.15; m/z found, 468.2

[M+H]⁺. ¹H NMR (CD₃OD): 8.65 (d, *J* = 2.5 Hz, 1H), 8.18 (dd, *J* = 4.8, 1.4 Hz, 1H), 8.04-7.85 (m, 3H), 7.46-7.25 (m, 2H), 7.23-7.08 (m, 2H), 6.88 (dd, *J* = 7.9, 1.9 Hz, 1H), 6.82-6.60 (m, 2H), 4.14-3.95 (m, 2H), 3.82-3.63 (m, 2H), 3.59-3.37 (m, 2H), 3.18-3.08 (m, 3H), 2.88-2.75 (m, 1H), 2.29-2.15 (m, 1H), 2.02-1.87 (m, 1H).

25 Example 53: 3-[2-(3,4-Dichloro-phenoxy)-ethyl]-pyrrolidine-1-carboxylic acid pyridin-3-ylamide.



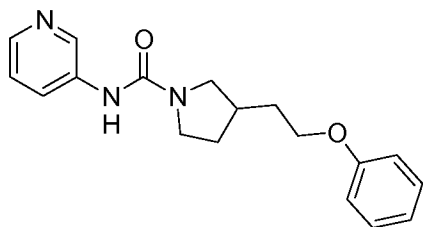
MS (ESI⁺): calcd. for C₁₈H₁₉Cl₂N₃O₂, 379.09; m/z found, 380.1 (M+H)⁺.

¹H NMR (CDCl₃): 8.44 (d, *J* = 2.5 Hz, 1H), 8.26 (dd, *J* = 4.7, 1.3 Hz, 1H), 8.06

- 5 (ddd, $J = 8.3, 2.5, 1.4$ Hz, 1H), 7.33 (d, $J = 8.9$ Hz, 1H), 7.23 (dd, $J = 8.3, 4.7$ Hz, 1H), 6.99 (d, $J = 2.8$ Hz, 1H), 6.75 (dd, $J = 8.9, 2.9$ Hz, 1H), 6.26 (s, 1H), 4.00 (t, $J = 6.1$ Hz, 2H), 3.79-3.71 (m, 1H), 3.67-3.58 (m, 1H), 3.45 (td, $J = 9.4, 7.1$ Hz, 1H), 3.14 (t, $J = 9.2$ Hz, 1H), 2.54-2.41 (m, 1H), 2.25-2.17 (m, 1H), 1.93 (dd, $J = 13.0, 6.3$ Hz, 2H), 1.79-1.68 (m, 1H).

10

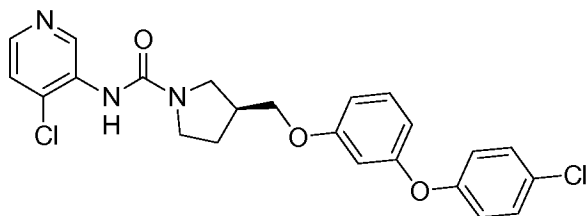
Example 54: 3-(2-Phenoxy-ethyl)-pyrrolidine-1-carboxylic acid pyridin-3-ylamide.



- MS (ESI⁺): calcd. for C₁₈H₂₁N₃O₂, 311.16; m/z found, 312.2 (M+H)⁺. ¹H NMR (CDCl₃): 8.44 (d, $J = 2.3$ Hz, 1H), 8.26 (dd, $J = 4.7, 1.4$ Hz, 1H), 8.07 (ddd, $J = 8.3, 2.6, 1.4$ Hz, 1H), 7.29 (dd, $J = 8.7, 7.4$ Hz, 2H), 7.23 (dd, $J = 8.4, 4.7$ Hz, 1H), 6.96 (t, $J = 7.4$ Hz, 1H), 6.90 (dd, $J = 8.7, 0.9$ Hz, 2H), 6.22 (s, 1H), 4.04 (t, $J = 6.1$ Hz, 2H), 3.79-3.73 (m, 1H), 3.67-3.60 (m, 1H), 3.45 (td, $J = 9.5, 7.0$ Hz, 1H), 3.15 (t, $J = 9.1$ Hz, 1H), 2.58-2.47 (m, 1H), 2.25-2.17 (m, 1H), 1.95 (dd, $J = 13.1, 6.2$ Hz, 2H), 1.78-1.68 (m, 1H).

20

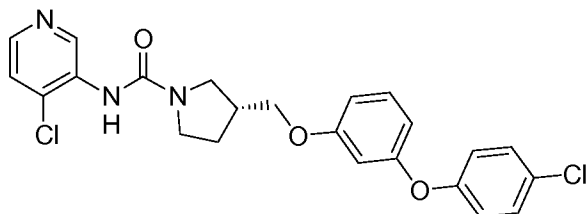
Example 55: (3S)-3-[3-(4-Chloro-phenoxy)-phoxymethyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.



- 25 MS (ESI⁺): calcd. for C₂₃H₂₁Cl₂N₃O₃, 457.10; m/z found, 458.1 (M+H)⁺. ¹H NMR (CDCl₃): 9.49 (s, 1H), 8.19 (d, $J = 5.2$, 1H), 7.31-7.28 (m, 3H), 7.24 (t, $J = 8.2$, 1H), 6.95 (d, $J = 8.9$, 2H), 6.68-6.65 (m, 2H), 6.61-6.58 (m, 1H), 6.54 (t, $J = 2.3$, 1H), 3.98 (dd, $J = 9.1, 6.2$, 1H), 3.93 (dd, $J = 9.2, 7.1$, 1H), 3.77 (dd, $J = 9.7, 7.7$, 1H), 3.72-3.65 (m, 1H), 3.60-3.54 (m, 1H), 3.43 (dd, $J = 9.9, 6.9$, 1H), 2.88-2.77 (m, 1H), 2.28-2.16 (m, 1H), 2.03-1.92 (m, 1H).

30

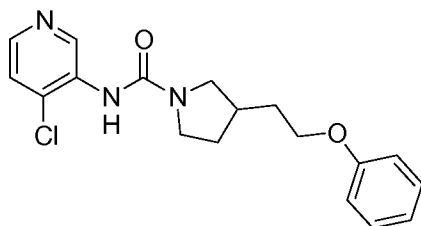
- 5 Example 56: (3R)-3-[3-(4-Chloro-phenoxy)-phenoxyethyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.



MS (ESI⁺): calcd. for C₂₃H₂₁Cl₂N₃O₃, 457.10; m/z found, 458.1 (M+H)⁺.

- 10 ¹H NMR (CDCl₃): 9.49 (s, 1H), 8.19 (d, J = 5.2, 1H), 7.31-7.27 (m, 3H), 7.24 (t, J = 8.2, 1H), 6.95 (d, J = 8.9, 2H), 6.68-6.64 (m, 2H), 6.61-6.58 (m, 1H), 6.54 (t, J = 2.3, 1H), 3.98 (dd, J = 9.1, 6.2, 1H), 3.93 (dd, J = 9.1, 7.1, 1H), 3.77 (dd, J = 9.6, 7.7, 1H), 3.71-3.65 (m, 1H), 3.60-3.54 (m, 1H), 3.43 (dd, J = 9.9, 6.9, 1H), 2.87-2.78 (m, 1H), 2.27-2.18 (m, 1H), 2.02-1.93 (m, 1H).

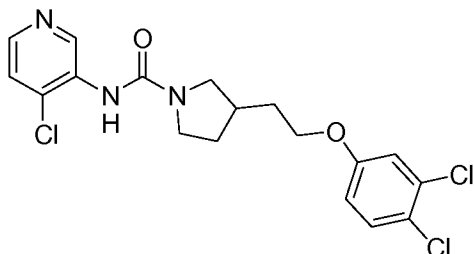
- 15 Example 57: 3-(2-Phenoxy-ethyl)-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.



MS (ESI⁺): calcd. for C₁₈H₂₀ClN₃O₂, 345.12; m/z found, 346.2 (M+H)⁺.

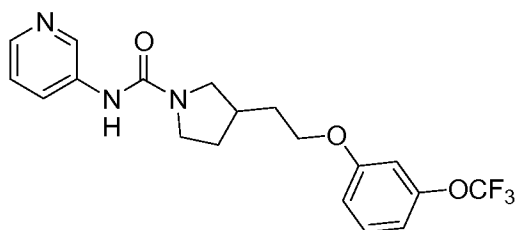
- ¹H NMR (CDCl₃): 9.51 (s, 1H), 8.18 (d, J = 5.2 Hz, 1H), 7.32-7.27 (m, 3H), 6.96 (t, J = 7.4 Hz, 1H), 6.90 (dd, J = 8.7, 0.9 Hz, 2H), 6.66 (s, 1H), 4.10-4.00 (m, 2H), 3.84-3.75 (m, 1H), 3.71-3.63 (m, 1H), 3.48 (td, J = 9.5, 7.0 Hz, 1H), 3.19 (t, J = 9.2 Hz, 1H), 2.59-2.47 (m, 1H), 2.28-2.19 (m, 1H), 1.96 (dd, J = 12.4, 6.4 Hz, 2H), 1.81-1.70 (m, 1H).

- 20 Example 58: 3-[2-(3,4-Dichloro-phenoxy)-ethyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.



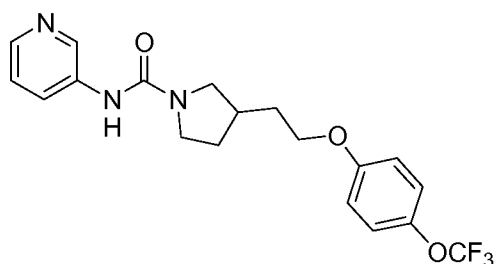
- 5 MS (ESI⁺): calcd. for C₁₈H₁₈Cl₃N₃O₂, 413.05; m/z found, 414.1 (M+H)⁺.
¹H NMR (CDCl₃): 9.50 (s, 1H), 8.19 (d, *J* = 5.2 Hz, 1H), 7.33 (d, *J* = 8.9 Hz,
 1H), 7.29 (d, *J* = 5.2 Hz, 1H), 6.99 (d, *J* = 2.8 Hz, 1H), 6.75 (dd, *J* = 8.9, 2.9
 Hz, 1H), 6.65 (s, 1H), 4.06-3.96 (m, 2H), 3.83-3.76 (m, 1H), 3.71-3.63 (m,
 1H), 3.48 (td, *J* = 9.4, 7.2 Hz, 1H), 3.18 (t, *J* = 9.2 Hz, 1H), 2.56-2.44 (m, 1H),
 10 2.27-2.19 (m, 1H), 1.95 (dd, *J* = 12.6, 6.3 Hz, 2H), 1.81-1.70 (m, 1H).

Example 59: 3-[2-(3-Trifluoromethoxy-phenoxy)-ethyl]-pyrrolidine-1-carboxylic acid pyridin-3-ylamide.



- 15 MS (ESI⁺): calcd. for C₁₉H₂₀F₃N₃O₃, 395.15; m/z found, 396.2 (M+H)⁺.
¹H NMR (CDCl₃): 8.44 (d, *J* = 2.3 Hz, 1H), 8.26 (dd, *J* = 4.7, 1.4 Hz, 1H), 8.06
 (ddd, *J* = 8.4, 2.6, 1.5 Hz, 1H), 7.29 (t, *J* = 8.3 Hz, 1H), 7.22 (dd, *J* = 8.4, 4.7
 Hz, 1H), 6.82 (dd, *J* = 8.3, 2.3 Hz, 2H), 6.76-6.73 (m, 1H), 6.23 (s, 1H), 4.03
 (t, *J* = 6.1 Hz, 2H), 3.77 (dd, *J* = 9.4, 7.6 Hz, 1H), 3.64 (td, *J* = 9.4, 2.8 Hz,
 20 1H), 3.46 (td, *J* = 9.4, 7.0 Hz, 1H), 3.19-3.12 (m, 1H), 2.57-2.44 (m, 1H), 2.26-
 2.16 (m, 1H), 1.95 (dd, *J* = 13.1, 6.2 Hz, 2H), 1.80-1.68 (m, 1H).

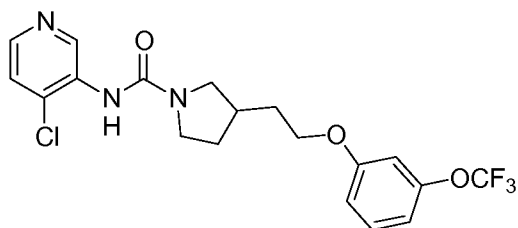
Example 60: 3-[2-(4-Trifluoromethoxy-phenoxy)-ethyl]-pyrrolidine-1-carboxylic acid pyridin-3-ylamide.



- 25 MS (ESI⁺): calcd. for C₁₉H₂₀F₃N₃O₃, 395.15; m/z found, 396.2 (M+H)⁺.
¹H NMR (CDCl₃): 8.44 (d, *J* = 2.3 Hz, 1H), 8.26 (dd, *J* = 4.7, 1.4 Hz, 1H), 8.07
 (ddd, *J* = 8.4, 2.6, 1.5 Hz, 1H), 7.23 (dd, *J* = 8.4, 4.7 Hz, 1H), 7.15 (dd, *J* =
 9.0, 0.7 Hz, 2H), 6.88 (d, *J* = 9.1 Hz, 2H), 6.18 (s, 1H), 4.03 (t, *J* = 6.1 Hz,
 30 2H), 3.77 (dd, *J* = 9.3, 7.7 Hz, 1H), 3.68-3.61 (m, 1H), 3.46 (td, *J* = 9.4, 7.0

- 5 Hz, 1H), 3.19-3.12 (m, 1H), 2.58-2.43 (m, 1H), 2.27-2.16 (m, 1H), 1.95 (dd, $J = 13.1, 6.2$ Hz, 2H), 1.80-1.67 (m, 1H).

Example 61: 3-[2-(3-Trifluoromethoxy-phenoxy)-ethyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.

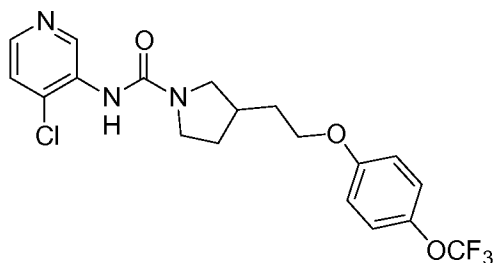


10

MS (ESI⁺): calcd. for C₁₉H₁₉ClF₃N₃O₃, 429.11; m/z found, 430.2

- (M+H)⁺. ¹H NMR (CDCl₃): 9.50 (s, 1H), 8.18 (d, $J = 5.2$ Hz, 1H), 7.31-7.26 (m, 2H), 6.82 (dd, $J = 8.3, 2.3$ Hz, 2H), 6.76-6.73 (m, 1H), 6.65 (s, 1H), 4.04 (t, $J = 6.1$ Hz, 2H), 3.83-3.76 (m, 1H), 3.71-3.63 (m, 1H), 3.49 (td, $J = 9.5, 7.0$ Hz, 1H), 3.23-3.14 (m, 1H), 2.59-2.44 (m, 1H), 2.30-2.18 (m, 1H), 1.96 (dd, $J = 12.8, 6.4$ Hz, 2H), 1.83-1.70 (m, 1H).
- 15

Example 62: 3-[2-(4-Trifluoromethoxy-phenoxy)-ethyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.

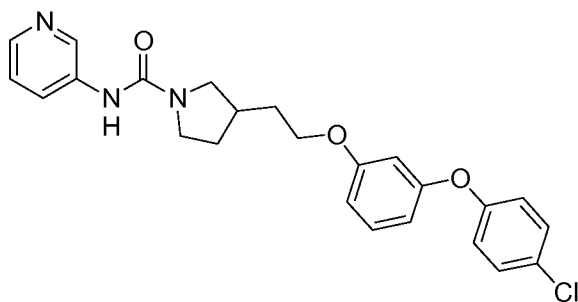


20

MS (ESI⁺): calcd. for C₁₉H₁₉ClF₃N₃O₃, 429.11; m/z found, 430.2

- (M+H)⁺. ¹H NMR (CDCl₃): 9.50 (s, 1H), 8.18 (d, $J = 5.2$ Hz, 1H), 7.28 (d, $J = 5.2$ Hz, 1H), 7.15 (d, $J = 8.6$ Hz, 2H), 6.88 (d, $J = 8.7$ Hz, 2H), 6.64 (s, 1H), 4.03 (t, $J = 5.8$ Hz, 2H), 3.80 (t, $J = 8.5$ Hz, 1H), 3.67 (t, $J = 7.9$ Hz, 1H), 3.48 (dd, $J = 16.9, 9.0$ Hz, 1H), 3.19 (t, $J = 9.1$ Hz, 1H), 2.59-2.43 (m, 1H), 2.29-2.17 (m, 1H), 2.01-1.90 (m, 2H), 1.83-1.69 (m, 1H).
- 25

Example 63: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-pyrrolidine-1-carboxylic acid pyridin-3-ylamide.

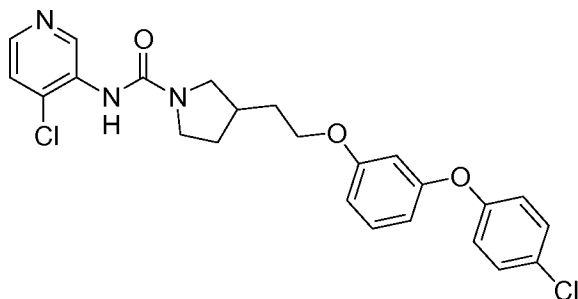


5

MS (ESI⁺): calcd. for C₂₄H₂₄ClN₃O₃, 437.15; m/z found, 438.2 (M+H)⁺.

¹H NMR (CDCl₃): 8.43 (d, *J* = 2.5 Hz, 1H), 8.26 (dd, *J* = 4.7, 1.3 Hz, 1H), 8.07 (ddd, *J* = 8.4, 2.5, 1.5 Hz, 1H), 7.29 (d, *J* = 8.9 Hz, 2H), 7.25-7.20 (m, 2H), 6.96 (d, *J* = 8.9 Hz, 2H), 6.66 (dd, *J* = 8.2, 1.8 Hz, 1H), 6.58 (dd, *J* = 8.1, 1.7 Hz, 1H), 6.54 (t, *J* = 2.3 Hz, 1H), 6.16 (s, 1H), 4.00 (t, *J* = 6.1 Hz, 2H), 3.76 (t, *J* = 8.4 Hz, 1H), 3.62 (d, *J* = 9.0 Hz, 1H), 3.45 (td, *J* = 9.4, 7.1 Hz, 1H), 3.14 (t, *J* = 9.1 Hz, 1H), 2.56-2.44 (m, 1H), 2.25-2.16 (m, 1H), 1.93 (dd, *J* = 13.0, 6.2, 2H), 1.78-1.68 (m, 1H).

15 Example 64: 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.

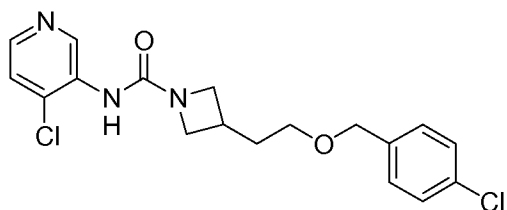


MS (ESI⁺): calcd. for C₂₄H₂₃Cl₂N₃O₃, 471.11; m/z found, 472.2 (M+H)⁺.

¹H NMR (CDCl₃): 9.50 (s, 1H), 8.18 (d, *J* = 5.2 Hz, 1H), 7.32-7.26 (m, 3H), 7.23 (t, *J* = 8.2 Hz, 1H), 6.98-6.93 (m, 2H), 6.68-6.62 (m, 2H), 6.58 (ddd, *J* = 8.1, 2.3, 0.8 Hz, 1H), 6.54 (dd, *J* = 3.5, 1.2 Hz, 1H), 4.01 (t, *J* = 6.1 Hz, 2H), 3.84-3.74 (m, 1H), 3.67 (td, *J* = 9.7, 2.4 Hz, 1H), 3.47 (td, *J* = 9.5, 7.0 Hz, 1H), 3.21-3.12 (m, 1H), 2.59-2.44 (m, 1H), 2.27-2.16 (m, 1H), 1.99-1.88 (m, 2H), 1.82-1.68 (m, 1H).

25

Example 65: 3-[2-(4-Chloro-benzyloxy)-ethyl]-azetidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.



5

Step A: 3-[2-(4-Chloro-benzyloxy)-ethyl]-azetidine-1-carboxylic acid tert-butyl ester. To a solution consisting of 3-(2-hydroxy-ethyl)-azetidine-1-carboxylic acid tert-butyl ester (0.250 g, 1.24 mmol) and THF (2 mL) at 0 °C was added sodium hydride (0.0459 g, 0.124 mmol). The reaction was stirred at 0 °C for 0.5 h, then treated with tetrabutylammonium iodide (0.0459 g, 0.124 mmol) and 4-chlorobenzyl bromide (0.383 g, 1.86 mmol), warmed to rt and stirred for 16 h. The reaction was quenched with saturated aqueous NH₄Cl (1.0 mL), diluted with ethyl acetate (40 mL) and extracted with 1 N NaOH (40 mL). The organic layer was dried (MgSO₄) and concentrated. The residue was purified (FCC) to give the title compound as a yellow oil (0.302 g, 74%). ¹H NMR (d₆-DMSO): 7.40 (d, *J* = 8.4 Hz, 2H), 7.32 (d, *J* = 8.4 Hz, 2H), 4.42 (s, 2H), 3.88 (t, *J* = 8.0 Hz, 2H), 3.53-3.44 (m, 2H), 3.40 (t, *J* = 6.2 Hz, 2H), 2.62-2.53 (m, 1H), 1.84-1.74 (m, 2H), 1.36 (s, 9H).

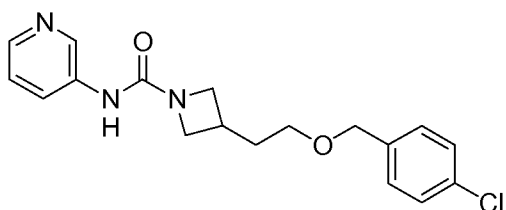
Step B: 3-[2-(4-chloro-benzyloxy)-ethyl]-azetidine, formic acid salt. The title compound was prepared using methods analogous to those described for Example 1 step B. MS (ESI⁺): calcd. for C₁₂H₁₆ClNO, 225.09; *m/z* found, 226.1 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.42 (s, 1H), 7.40 (d, *J* = 8.4 Hz, 2H), 7.32 (d, *J* = 8.4 Hz, 2H), 4.42 (s, 2H), 3.86 (dd, *J* = 10.2, 8.7 Hz, 2H), 3.63-3.54 (m, 2H), 3.41 (t, *J* = 6.1 Hz, 2H), 2.89-2.77 (m, 1H), 1.88-1.80 (m, 2H).

Step C: 3-[2-(4-Chloro-benzyloxy)-ethyl]-azetidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide. The title compound was prepared using methods analogous to those described for Example 1 step C. MS (ESI⁺): calcd. for C₁₈H₁₉Cl₂N₃O₂, 379.09; *m/z* found, 380.1 (M+H)⁺. ¹H NMR (d₆-DMSO): 8.70 (s, 1H), 8.25 (d, *J* = 5.2 Hz, 1H), 8.09 (s, 1H), 7.54 (d, *J* = 5.3 Hz, 1H), 7.41 (d, *J* = 8.5 Hz, 2H), 7.34 (d, *J* = 8.5 Hz, 2H), 4.45 (s, 2H), 4.06 (t, *J* = 8.2 Hz, 2H), 3.66 (dd, *J* = 8.1, 5.9 Hz, 2H), 3.45 (t, *J* = 6.2 Hz, 2H), 2.74-2.62 (m, 1H), 1.86 (dd, *J* = 13.6, 6.2 Hz, 2H).

30

- 5 Examples 66 to 70 were prepared using methods analogous to those described for Example 65, using the appropriate carbamate and benzyl bromide.

10 Example 66: 3-[2-(4-Chloro-benzyloxy)-ethyl]-azetidone-1-carboxylic acid pyridin-3-ylamide, trifluoroacetate salt.

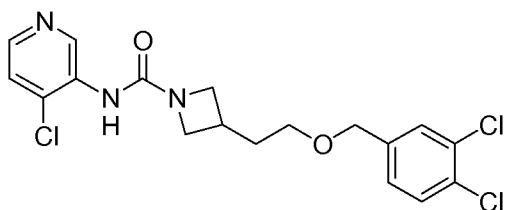


MS (ESI⁺): calcd. for C₁₈H₂₀ClN₃O₂, 345.12; m/z found, 346.2 (M+H)⁺.

¹H

- NMR (d₆-DMSO): 9.13 (s, 1H), 8.99 (d, *J* = 1.7 Hz, 1H), 8.39 (d, *J* = 5.0 Hz, 1H), 8.34-8.27 (m, 1H), 7.75 (dd, *J* = 8.6, 5.3 Hz, 1H), 7.41 (d, *J* = 8.5 Hz, 2H), 7.34 (d, *J* = 8.5 Hz, 2H), 4.45 (s, 2H), 4.09 (t, *J* = 8.3 Hz, 2H), 3.69 (dd, *J* = 8.3, 5.8 Hz, 2H), 3.45 (t, *J* = 6.1 Hz, 2H), 2.75-2.62 (m, 1H), 1.90-1.82 (m, 2H).
- 15

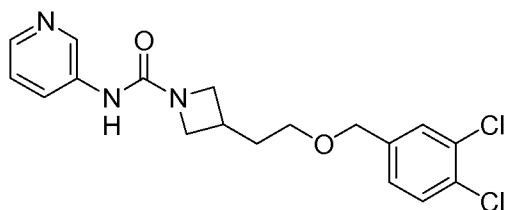
- 20 Example 67: 3-[2-(3,4-Dichloro-benzyloxy)-ethyl]-azetidone-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.



MS (ESI⁺): calcd. for C₁₈H₁₈Cl₃N₃O₂, 413.05; m/z found, 414.1 (M+H)⁺.

- ¹H NMR (d₆-DMSO): 8.70 (s, 1H), 8.25 (d, *J* = 5.3 Hz, 1H), 8.09 (s, 1H), 7.61 (d, *J* = 8.2 Hz, 1H), 7.56 (d, *J* = 1.8 Hz, 1H), 7.54 (d, *J* = 5.3 Hz, 1H), 7.32 (dd, *J* = 8.3, 1.9 Hz, 1H), 4.47 (s, 2H), 4.06 (t, *J* = 8.2 Hz, 2H), 3.66 (dd, *J* = 8.1, 5.9 Hz, 2H), 3.46 (t, *J* = 6.2 Hz, 2H), 2.75-2.64 (m, 1H), 1.92-1.83 (m, 2H).
- 25

- 30 Example 68: 3-[2-(3,4-Dichloro-benzyloxy)-ethyl]-azetidone-1-carboxylic acid pyridin-3-ylamide, trifluoroacetate salt.

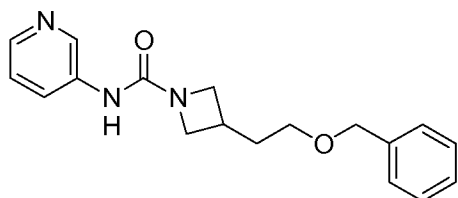


5

MS (ESI⁺): calcd. for C₁₉H₁₉Cl₂N₃O₂, 379.09; m/z found, 380.1 (M+H)⁺.

¹H NMR (d₆-DMSO): 9.05 (s, 1H), 8.96 (d, *J* = 2.3 Hz, 1H), 8.35 (dd, *J* = 5.2, 1.1 Hz, 1H), 8.28-8.23 (m, 1H), 7.70 (dd, *J* = 8.6, 5.3 Hz, 1H), 7.62 (d, *J* = 8.2 Hz, 1H), 7.56 (d, *J* = 1.9 Hz, 1H), 7.32 (dd, *J* = 8.2, 1.9 Hz, 1H), 4.47 (s, 2H),
 10 4.09 (t, *J* = 8.3 Hz, 2H), 3.68 (dd, *J* = 8.3, 5.9 Hz, 2H), 3.46 (t, *J* = 6.1 Hz, 2H), 2.76-2.65 (m, 1H), 1.88 (d, *J* = 7.3 Hz, 2H).

Example 69: 3-(2-Benzyloxy-ethyl)-azetidine-1-carboxylic acid pyridin-3-ylamide.

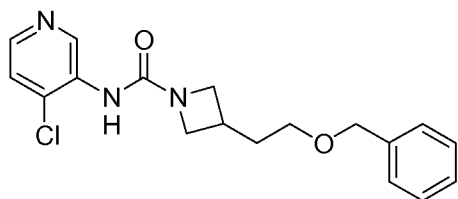


15

MS (ESI⁺): calcd. for C₁₈H₂₁N₃O₂, 311.16; m/z found, 312.2 (M+H)⁺. ¹H

NMR (d₆-DMSO): 8.65 (d, *J* = 2.5 Hz, 1H), 8.51 (s, 1H), 8.12 (dd, *J* = 4.7, 1.5 Hz, 1H), 7.93-7.89 (m, 1H), 7.39-7.26 (m, 5H), 7.24 (dd, *J* = 8.3, 4.6 Hz, 1H), 4.46 (s, 2H), 4.04 (t, *J* = 8.2 Hz, 2H), 3.64 (dd, *J* = 8.2, 5.8 Hz, 2H), 3.44 (t, *J* =
 20 6.2 Hz, 2H), 2.73-2.60 (m, 1H), 1.90-1.81 (m, 2H).

Example 70: 3-(2-Benzyloxy-ethyl)-azetidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.



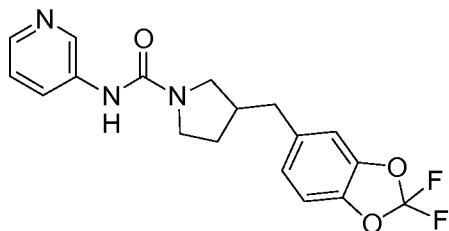
25

MS (ESI⁺): calcd. for C₁₈H₂₀ClN₃O₂, 345.12; m/z found, 346.2 (M+H)⁺.

¹H NMR (d₆-DMSO): 8.70 (s, 1H), 8.25 (d, *J* = 5.3 Hz, 1H), 8.09 (s, 1H), 7.53 (d, *J* = 5.3 Hz, 1H), 7.39-7.24 (m, 5H), 4.46 (s, 2H), 4.06 (t, *J* = 8.2 Hz, 2H),

- 5 3.66 (dd, $J = 8.1, 5.9$ Hz, 2H), 3.45 (t, $J = 6.2$ Hz, 2H), 2.75-2.62 (m, 1H), 1.91-1.82 (m, 2H).

Example 71: 3-(2,2-Difluoro-benzo[1,3]dioxol-5-ylmethyl)-pyrrolidine-1-carboxylic acid pyridin-3-ylamide.



Step A: 3-Methylene pyrrolidine-1-carboxylic acid *tert*-butyl ester. To a suspension consisting of methyl triphenylphosphonium bromide (18 g, 51 mmol) and THF (200 mL) at 0 °C was added *n*-butyl lithium (1.6 M solution in hexanes, 32 mL). The resulting orange solution was stirred at 0 °C for 5 min, then a solution of 3-oxo-pyrrolidine-1-carboxylic acid *tert*-butyl ester (9.0 g, 48 mmol) in THF (40 mL) was added via cannula. The reaction mixture was stirred at 0 °C for 90 min, then warmed to rt and stirring was continued for 1 h. The reaction mixture was then cooled to 0 °C, quenched with sat. NH₄Cl, and extracted with Et₂O. The organic layer was dried (MgSO₄) and concentrated. The crude residue was suspended in hot hexanes and filtered. The filtrate was concentrated and purified (FCC) to give the title compound (4.4 g, 50%).

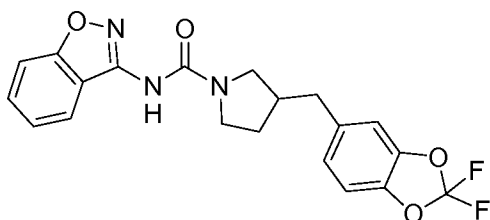
Step B: 3-(2,2-Difluoro-benzo[1,3]dioxol-5-ylmethyl)-pyrrolidine-1-carboxylic acid *tert*-butyl ester. A stream of N₂ was bubbled through neat 3-methylene pyrrolidine-1-carboxylic acid *tert*-butyl ester (792 mg, 4.33 mmol) for 15 min before charging the flask with 9-BBN (0.5 M in THF, 8.8 mL). The reaction mixture was heated at reflux for 2.5 h; then cooled to rt. The resultant mixture was then added, via cannula, to a preformed solution consisting of 5-bromo-2,2-difluoro-1,3-benzodioxole (1.01 g, 4.24 mmol), Pd(dppf)Cl₂•CH₂Cl₂ (92 mg), and potassium carbonate (760 mg, 5.50 mmol) in DMF/H₂O (10 mL/1 mL). The reaction mixture was heated at 60 °C for 18 h, cooled to rt and poured into water. The pH of the mixture was adjusted to 11 with NaOH (1 N), and extracted with EtOAc (3x). The organic layers were combined, dried (Na₂SO₄), and concentrated. The crude residue was purified (FCC) to give the title compound (1.4 g, 94%).

5 Step C: 3-(2,2-Difluoro-benzo[1,3]dioxol-5-ylmethyl)-pyrrolidine, hydrochloride salt. To a solution consisting of 3-(2,2-difluoro-benzo[1,3]dioxol-5-ylmethyl)-pyrrolidine-1-carboxylic acid tert-butyl ester (1.36 g, 3.98 mmol) and dichloromethane (20 mL) was added HCl (4 M solution in dioxane, 6 mL). The reaction mixture was stirred at rt for 18 h. The resultant
10 mixture was concentrated to give the title compound in quantitative yield.

Step D: 3-(2,2-Difluoro-benzo[1,3]dioxol-5-ylmethyl)-pyrrolidine-1-carboxylic acid pyridin-3-ylamide. To a solution consisting of 3-(2,2-difluoro-benzo[1,3]dioxol-5-ylmethyl)-pyrrolidine (111 mg, 0.399 mmol), triethylamine (0.1 mL, 0.72 mmol) and acetonitrile (2 mL) was added pyridin-3-yl-carbamic acid phenyl ester (90.1 mg, 0.421 mmol). The mixture was heated at 35 °C
15 for 18 h, cooled to rt and then concentrated. The crude residue was purified (FCC) to give the title compound (56 mg, 39%). MS (ESI): mass calcd. For C₁₈H₁₇F₂N₃O₃, 361.12; m/z found, 362.2 [M+H]⁺. ¹H NMR (CD₃OD): 8.63-8.59 (m, 1H), 8.16-8.13 (m, 1H), 7.97-7.92 (m, 1H), 7.35-7.30 (m, 1H), 7.16-7.09 (m, 2H), 7.06-7.02 (m, 1H), 3.66-3.53 (m, 2H), 3.48-3.38 (m, 1H), 3.19-3.11 (m, 1H), 2.85-2.72 (m, 2H), 2.62-2.50 (m, 1H), 2.11-2.02 (m, 1H), 1.78-1.66 (m, 1H).
20

 Examples 72 to 76 were prepared using methods analogous to those
25 described in Example 71.

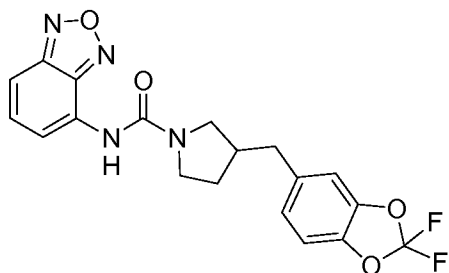
Example 72: 3-(2,2-Difluoro-benzo[1,3]dioxol-5-ylmethyl)-pyrrolidine-1-carboxylic acid benzo[d]isoxazol-3-ylamide.



30 MS (ESI): mass calcd. for C₂₀H₁₇F₂N₃O₄, 401.12; m/z found, 402.2 [M+H]⁺. ¹H NMR (CD₃OD): 7.93-7.88 (m, 1H), 7.61-7.48 (m, 2H), 7.33-7.26 (m, 1H), 7.17-7.08 (m, 2H), 7.07-7.02 (m, 1H), 3.74-3.60 (m, 2H), 3.55-3.45 (m, 1H), 3.26-3.16 (m, 1H), 2.86-2.75 (m, 2H), 2.65-2.52 (m, 1H), 2.15-2.03 (m, 1H), 1.81-1.68 (m, 1H).

5

Example 73: N-2,1,3-Benzoxadiazol-4-yl-3-[(2,2-difluoro-1,3-benzodioxol-5-yl)methyl]pyrrolidine-1-carboxamide.

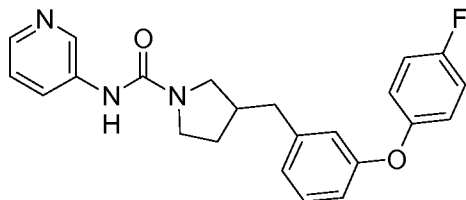


MS (ESI): mass calcd. for C₁₉H₁₆F₂N₄O₄, 402.11; m/z found, 403.2

10 [M+H]⁺. ¹H NMR (d₆-DMSO): 8.55 (s, 1H), 7.71-7.69 (m, 1H), 7.61-7.58 (m, 1H), 7.55-7.51 (m, 1H), 7.36 (d, J = 1.5 Hz, 1H), 7.34 (d, J = 8.2 Hz, 1H), 7.10 (dd, J = 8.2, 1.6 Hz, 1H), 3.66-3.54 (m, 2H), 3.45-3.37 (m, 1H), 3.18-3.09 (m, 1H), 2.75 (d, J = 7.5 Hz, 2H), 2.54-2.48 (m, 1H) (coincident with DMSO peak), 2.00-1.92 (m, 1H), 1.67-1.58 (m, 1H).

15

Example 74: 3-[3-(4-Fluoro-phenoxy)-benzyl]-pyrrolidine-1-carboxylic acid pyridin-3-ylamide.



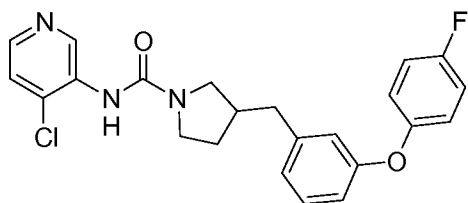
MS (ESI): mass calcd. for C₂₃H₂₂FN₃O₂, 391.17; m/z found, 392.2

20 [M+H]⁺. ¹H NMR (CD₃OD): 8.62-8.59 (m, 1H), 8.15 (dd, J = 4.8 Hz, 1.4, 1H), 7.94 (ddd, J = 8.4, 2.6, 1.5 Hz, 1H), 7.34-7.31 (m, 1H), 7.27 (t, J = 7.9 Hz, 1H), 7.10-7.05 (m, 2H), 7.01-6.97 (m, 3H), 6.88-6.86 (m, 1H), 6.81-6.78 (m, 1H), 3.64-3.54 (m, 2H), 3.46-3.40 (m, 1H), 3.15 (dd, J = 10.1, 8.0 Hz, 1H), 2.78-2.68 (m, 2H), 2.58-2.52 (m, 1H), 2.09-2.04 (m, 1H), 1.76-1.68 (m, 1H).

25

Example 75: 3-[3-(4-Fluoro-phenoxy)-benzyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.

5

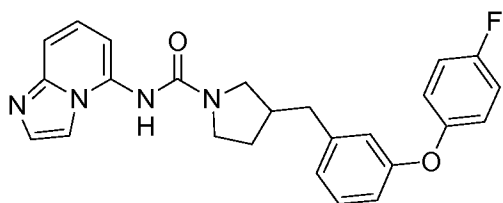


10

MS (ESI): mass calcd. for $C_{23}H_{21}ClFN_3O_2$, 425.13; m/z found, 426.2 $[M+H]^+$. 1H NMR (CD_3OD): 8.77 (s, 1H), 8.24-8.22 (m, 1H), 7.53 (d, $J = 5.4$ Hz, 1H), 7.28 (t, $J = 7.9$ Hz, 1H), 7.10-7.05 (m, 2H), 7.02-6.97 (m, 3H), 6.89-6.87 (m, 1H), 6.81-6.79 (m, 1H), 3.66-3.56 (m, 2H), 3.48-3.43 (m, 1H), 3.21-3.15 (m, 1H), 2.79-2.69 (m, 2H), 2.62-2.53 (m, 1H), 2.12-2.05 (m, 1H), 1.79-1.71 (m, 1H).

Example 76: 3-[3-(4-Fluoro-phenoxy)-benzyl]-pyrrolidine-1-carboxylic acid imidazo[1,2-a]pyridin-5-ylamide.

15

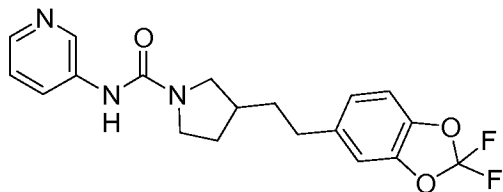


20

MS (ESI): mass calcd. for $C_{25}H_{23}FN_4O_2$, 430.18; m/z found, 431.2 $[M+H]^+$. 1H NMR (CD_3OD): 7.69-7.66 (m, 1H), 7.55 (d, $J = 1.3$ Hz, 1H), 7.45 (d, $J = 9.0$ Hz, 1H), 7.34-7.31 (m, 1H), 7.28 (t, $J = 7.9$ Hz, 1H), 7.10-7.05 (m, 2H), 7.03-6.97 (m, 3H), 6.91-6.88 (m, 1H), 6.82-6.79 (m, 2H), 3.71-3.60 (m, 2H), 3.53-3.45 (m, 1H), 3.24-3.16 (m, 1H), 2.84-2.71 (m, 2H), 2.65-2.53 (m, 1H), 2.15-2.05 (m, 1H), 1.83-1.71 (m, 1H).

Example 77: 3-[2-(2,2-Difluoro-benzo[1,3]dioxol-5-yl)-ethyl]-pyrrolidine-1-carboxylic acid pyridin-3-ylamide.

25



Step A: 3-Vinyl-pyrrolidine-1-carboxylic acid *tert*-butyl ester. To a suspension consisting of methyl triphenylphosphonium bromide (4.76 g, 13.3 mmol) and THF (20 mL) at 0 °C was added *n*-butyl lithium (1.6 M solution in hexanes, 8.3 mL). The resulting orange solution was stirred at 0 °C for 5 min.

5 A solution consisting of 3-formyl-pyrrolidine-1-carboxylic acid *tert*-butyl ester (2.50 g, 12.5 mmol) and THF (15 mL) was then added via cannula. The reaction mixture was stirred at 0 °C and was warmed to rt over 2.5 h. The reaction mixture was then cooled to 0 °C, quenched with sat. NH₄Cl, and extracted with Et₂O. The organic layer was combined, dried (MgSO₄) and
10 concentrated. The crude residue was suspended in hot hexanes and filtered. The filtrate was concentrated and purified (FCC) to give the title compound (1.7 g, 70%).

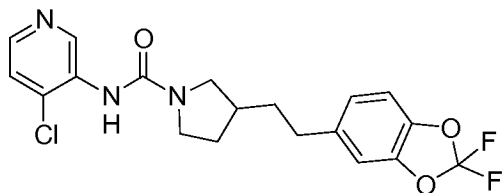
Step B: 3-[2-(2,2-Difluoro-benzo[1,3]dioxol-5-yl)-ethyl]-pyrrolidine-1-carboxylic acid *tert*-butyl ester. A stream of N₂ was bubbled through neat 3-vinyl-pyrrolidine-1-carboxylic acid *tert*-butyl ester (589 mg, 2.99 mmol) for 15
15 min before adding 9-BBN (0.5 M in THF, 6 mL). The reaction mixture was heated at reflux for 2 h, then cooled to rt. The reaction mixture was then added, via cannula, to a preformed solution consisting of 5-bromo-2,2-difluoro-1,3-benzodioxole (704 mg, 2.97 mmol), Pd(dppf)Cl₂•CH₂Cl₂ (69.7 mg,
20 0.0953 mmol), and potassium carbonate (559 mg, 4.04 mmol) in DMF/H₂O (10 mL/1 mL). The resultant mixture was heated at 60 °C for 18 h, cooled to rt, and poured into water. The pH of the mixture was adjusted to 11 with NaOH (1 N) and extracted with EtOAc (3x). The organic layers were combined, dried (Na₂SO₄), and concentrated. The crude residue was purified
25 (FCC) to give the title compound (1.0 g, 98%).

Step C: 3-[2-(2,2-Difluoro-benzo[1,3]dioxol-5-yl)-ethyl]-pyrrolidine, hydrochloride salt. Title compound was prepared using methods analogous to those described in Example 80, Step C.

Step D: 3-[2-(2,2-Difluoro-benzo[1,3]dioxol-5-yl)-ethyl]-pyrrolidine-1-carboxylic acid pyridin-3-ylamide. Title compound was synthesized using
30 procedures analogous to those described in Example 80, Step D. MS (ESI): mass calcd. for C₁₉H₁₉F₂N₃O₃, 375.14; m/z found, 376.2 [M+H]⁺. ¹H NMR (CD₃OD): 8.64-8.61 (m, 1H), 8.16-8.15 (m, 1H), 7.98-7.95 (m, 1H), 7.35-7.32 (m, 1H), 7.12-7.07 (m, 2H), 7.03-7.00 (m, 1H), 3.73-3.69 (m, 1H), 3.64-3.59
35 (m, 1H), 3.45-3.39 (m, 1H), 3.09 (t, *J* = 9.3 Hz, 1H), 2.75-2.70 (m, 2H), 2.28-2.22 (m, 1H), 2.18-2.12 (m, 1H), 1.80-1.74 (m, 2H), 1.71-1.63 (m, 1H).

5 Examples 78 to 79 were prepared using methods analogous to those described in Example 77.

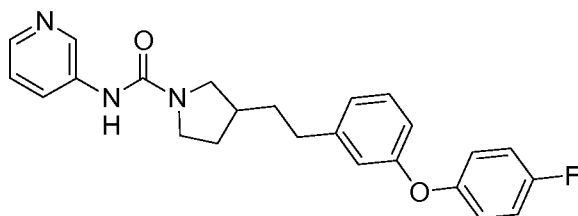
Example 78: 3-[2-(2,2-Difluoro-benzo[1,3]dioxol-5-yl)-ethyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide.



MS (ESI): mass calcd. for $C_{19}H_{18}ClF_2N_3O_3$, 409.10; m/z found, 410.1 $[M+H]^+$. 1H NMR (CD_3OD): 8.81 (s, 1H), 8.23 (d, $J = 5.3$ Hz, 1H), 7.54 (d, $J = 5.3$ Hz, 1H), 7.12-7.10 (m, 1H), 7.09 (d, $J = 8.2$ Hz, 1H), 7.03-7.01 (m, 1H), 3.72 (dd, $J = 9.9, 7.5$ Hz, 1H), 3.66-3.62 (m, 1H), 3.47-3.42 (m, 1H), 3.13-3.08 (m, 1H), 2.77-2.68 (m, 2H), 2.33-2.23 (m, 1H), 2.21-2.13 (m, 1H), 1.78 (q, $J = 7.7$ Hz, 2H), 1.73-1.66 (m, 1H).

15

Example 79: 3-[2-[3-(4-Fluoro-phenoxy)-phenyl]-ethyl]-pyrrolidine-1-carboxylic acid pyridin-3-ylamide.



MS (ESI): mass calcd. for $C_{24}H_{24}FN_3O_2$, 405.19; m/z found, 406.2 $[M+H]^+$. 1H NMR (CD_3OD): 8.64-8.61 (m, 1H), 8.15 (dd, $J = 4.8, 1.4$ Hz, 1H), 7.96 (ddd, $J = 8.4, 2.6, 1.5$ Hz, 1H), 7.35-7.31 (m, 1H), 7.26 (t, $J = 7.9$ Hz, 1H), 7.10-7.05 (m, 2H), 7.00-6.96 (m, 3H), 6.85-6.83 (m, 1H), 6.79-6.76 (m, 1H), 3.69 (dd, $J = 10.0, 7.4$ Hz, 1H), 3.63-3.58 (m, 1H), 3.43-3.37 (m, 1H), 3.07 (t, $J = 9.4$ Hz, 1H), 2.70-2.65 (m, 2H), 2.27-2.20 (m, 1H), 2.15-2.10 (m, 1H), 1.78-1.73 (m, 2H), 1.69-1.60 (m, 1H).

25

Biological Testing:

- 30
- Assay Method 1
 - A. Transfection of Cells with Human FAAH

5 A 10-cm tissue culture dish with a confluent monolayer of SK-N-MC cells was split 2 days (d) prior to transfection. Using sterile technique, the media was removed and the cells were detached from the dish by the addition of trypsin. One fifth of the cells were then placed onto a new 10-cm dish. Cells were grown in a 37 °C incubator with 5% CO₂ in Minimal Essential
10 Media Eagle with 10% Fetal Bovine Serum. After 2 d, cells were approximately 80% confluent. These cells were removed from the dish with trypsin and pelleted in a clinical centrifuge. The pellet was re-suspended in 400 µL complete media and transferred to an electroporation cuvette with a 0.4 cm gap between the electrodes. Supercoiled human FAAH cDNA (1 µg)
15 was added to the cells and mixed. The voltage for the electroporation was set at 0.25 kV, and the capacitance was set at 960 µF. After electroporation, the cells were diluted into complete media (10 mL) and plated onto four 10-cm dishes. Because of the variability in the efficiency of electroporation, four different concentrations of cells were plated. The ratios used were 1:20, 1:10,
20 and 1:5, with the remainder of the cells being added to the fourth dish. The cells were allowed to recover for 24 h before adding the selection media (complete media with 600 µg/mL G418). After 10 d, dishes were analyzed for surviving colonies of cells. Dishes with well-isolated colonies were used. Cells from individual colonies were isolated and tested. The clones that
25 showed the most FAAH activity, as measured by anandamide hydrolysis, were used for further study.

B. FAAH Assay

T84 frozen cell pellets or transfected SK-N-MC cells (contents of 1 x 15 cm culture dishes) were homogenized in 50 mL of FAAH assay buffer (125
30 mM Tris, 1mM EDTA, 0.2% Glycerol, 0.02% Triton X-100, 0.4 mM HEPES, pH 9). The assay mixture consisted of 50 µL of the cell homogenate, 10 µL of the test compound, and 40 µL of anandamide [1-³H-ethanolamine] (³H-AEA, Perkin-Elmer, 10.3 Ci/mmol), which was added last, for a final tracer concentration of 80 nM. The reaction mixture was incubated at rt for 1 h.
35 During the incubation, 96-well Multiscreen filter plates (catalog number MAFCNOB50; Millipore, Bedford, MA, USA) were loaded with 25 µL of activated charcoal (Multiscreen column loader, catalog number MACL09625,

5 Millipore) and washed once with 100 μ L of MeOH. Also during the incubation, 96-well DYNEX MicroLite plates (catalog number NL510410) were loaded with 100 μ L of MicroScint40 (catalog number 6013641, Packard Bioscience, Meriden, CT, USA). After the 1 h incubation, 60 μ L of the reaction mixture were transferred to the charcoal plates, which were then
10 assembled on top of the DYNEX plates using Centrifuge Alignment Frames (catalog number MACF09604, Millipore). The unbound labeled ethanolamine was centrifuged through to the bottom plate (5 min at 2000 rpm), which was preloaded with the scintillant, as described above. The plates were sealed and left at rt for 1 h before counting on a Hewlett Packard TopCount.

15 • Assay Method 2

A. Transfection of Cells with Rat FAAH

A 10-cm tissue culture dish with a confluent monolayer of SK-N-MC cells was split 2 days (d) prior to transfection. Using sterile technique, the media was removed and the cells were detached from the dish by the addition
20 of trypsin. One fifth of the cells were then placed onto a new 10-cm dish. Cells were grown in a 37 °C incubator with 5% CO₂ in Minimal Essential Media Eagle with 10% Fetal Bovine Serum. After 2 d, cells were approximately 80% confluent. These cells were removed from the dish with trypsin and pelleted in a clinical centrifuge. The pellet was re-suspended in
25 400 μ L complete media and transferred to an electroporation cuvette with a 0.4 cm gap between the electrodes. Supercoiled rat FAAH cDNA (1 μ g) was added to the cells and mixed. The voltage for the electroporation was set at 0.25 kV, and the capacitance was set at 960 μ F. After electroporation, the cells were diluted into complete media (10 mL) and plated onto four 10-cm
30 dishes. Because of the variability in the efficiency of electroporation, four different concentrations of cells were plated. The ratios used were 1:20, 1:10, and 1:5, with the remainder of the cells being added to the fourth dish. The cells were allowed to recover for 24 h before adding the selection media (complete media with 600 μ g/mL G418). After 10 d, dishes were analyzed for
35 surviving colonies of cells. Dishes with well-isolated colonies were used. Cells from individual colonies were isolated and tested. The clones that

5 showed the most FAAH activity, as measured by anandamide hydrolysis, were used for further study.

B. FAAH Assay

T84 frozen cell pellets or transfected SK-N-MC cells (contents of 1 x 15 cm culture dishes) were homogenized in 50 mL of FAAH assay buffer (125 mM Tris, 1 mM EDTA, 0.2% Glycerol, 0.02% Triton X-100, 0.4 mM HEPES, pH 9). The assay mixture consisted of 50 μ L of the cell homogenate, 10 μ L of the test compound, and 40 μ L of anandamide [1-³H-ethanolamine] (³H-AEA, Perkin-Elmer, 10.3 Ci/mmol), which was added last, for a final tracer concentration of 80 nM. The reaction mixture was incubated at rt for 1 h. During the incubation, 96-well Multiscreen filter plates (catalog number MAFCNOB50; Millipore, Bedford, MA, USA) were loaded with 25 μ L of activated charcoal (Multiscreen column loader, catalog number MACL09625, Millipore) and washed once with 100 μ L of MeOH. Also during the incubation, 96-well DYNEX MicroLite plates (catalog number NL510410) were loaded with 100 μ L of MicroScint40 (catalog number 6013641, Packard Bioscience, Meriden, CT, USA). After the 1 h incubation, 60 μ L of the reaction mixture were transferred to the charcoal plates, which were then assembled on top of the DYNEX plates using Centrifuge Alignment Frames (catalog number MACF09604, Millipore). The unbound labeled ethanolamine was centrifuged through to the bottom plate (5 min at 2000 rpm), which was preloaded with the scintillant, as described above. The plates were sealed and left at rt for 1 h before counting on a Hewlett Packard TopCount.

Results for compounds tested in these assays are summarized in Table 1, as an average of results obtained. Compounds were tested in the free base or trifluoroacetic acid salt form. Compounds tested in the trifluoroacetic acid salt form are indicated as such by a "*" designation after the example number. Where activity is shown as greater than (>) a particular value, the value is the solubility limit of the compound in the assay medium or the highest concentration tested in the assay.

35

5

Table 1

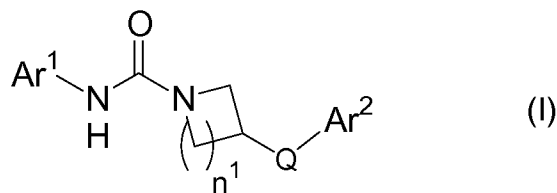
Ex.	Assay 1 IC ₅₀ (μM)	Assay 2 IC ₅₀ (μM)		Ex.	Assay 1 IC ₅₀ (μM)	Assay 2 IC ₅₀ (μM)
1	0.005	0.004		41	3.000	1.000
2*	>10	>10		42	0.030	0.024
3	>10	>10		43	0.004	0.001
4*	10.000	10.000		44	0.013	0.027
5	>10	3.000		45	0.700	0.043
6	1.100	0.175		46	1.600	0.615
7	>10	>10		47	1.000	0.250
8*	2.400	0.620		48	0.001	0.005
9	1.300	1.300		49	0.009	0.033
10	1.400	0.300		50	0.009	0.097
11	0.370	1.600		51	0.200	0.062
12	0.300	>10		52	0.100	1.000
13	2.500	1.700		53	5.999	10.000
14	>10	>10		54	>10	>10
15*	>10	>10		55	0.016	0.006
16	>10	>10		56	0.004	0.002
17	0.170	0.340		57	3.000	>10
18	>10	>10		58	0.300	0.700
19	>10	>10		59	>10	>10
20	1.000	1.600		60	>10	>10
21	1.000	0.530		61	1.600	1.600
22	1.300	0.500		62	1.300	10.000
23	0.300	0.940		63	0.600	0.630
24	0.058	0.380		64	0.063	0.073
25	0.050	0.230		65	0.275	0.739
26	0.120	0.200		66*	10.000	10.000
27	0.015	0.480		67	0.039	0.084
28	0.001	0.001		68*	0.849	1.225
29	0.053	0.400		69	10.000	>10
30	>10	10.000		70	0.936	10.000
31	3.000	0.900		71	>10	>10
32	0.360	0.430		72	>10	0.770
33	0.034	2.000		73	>10	10.000
34	0.030	0.054		74	1.200	1.100
35*	0.160	0.350		75	0.045	0.030
36	0.200	0.430		76	0.400	0.070
37	0.140	0.820		77	5.000	7.000
38	0.027	0.005		78	0.230	0.320
39	0.013	0.010		79	0.058	0.048
40	0.003	0.004				

While the invention has been illustrated by reference to exemplary and preferred embodiments, it will be understood that the invention is intended not to be limited to the foregoing detailed description, but to be defined by the

10 appended claims as properly construed under principles of patent law.

5 What is claimed is:

1. A composition of matter selected from compounds of Formula (I), pharmaceutically acceptable salts of compounds of Formula (I), and pharmaceutically acceptable prodrugs of compounds of Formula (I),



10 wherein

Q is $-(\text{CH}_2)_{1-3}-$, $-(\text{CH}_2)_{1-2}\text{O}-$, or $-(\text{CH}_2)_{1-2}\text{OCH}_2-$;

n^1 is 1 or 2, with the proviso that when n^1 is 1, then Q cannot be $-\text{CH}_2-$;

Ar^1 is a ring system selected from benzo[d]isoxazolyl, 1H-pyrrolo[2,3-
b]pyridinyl, 2,1,3-benzoxadiazolyl, imidazo[1,2-a]pyridinyl, imidazo[1,2-
15 b]pyridazinyl, isoquinolinyl, and pyridyl optionally substituted with triazolyl;

wherein each ring system is optionally substituted with halo;

Ar^2 is:

(i) phenyl optionally substituted with one or two R^d moieties;

20 wherein each R^d moiety is independently selected from $-\text{C}_{1-4}\text{alkyl}$, $-\text{OC}_{1-4}\text{alkyl}$, halo, $-\text{CF}_3$, $-\text{OCF}_3$, and $-\text{S}(\text{O})(\text{O})\text{C}_{1-4}\text{alkyl}$, or two adjacent R^d moieties taken together form $-\text{OCH}_2\text{O}-$ or $-\text{OCF}_2\text{O}-$;

(ii) phenyl substituted at the 3- or 4-position with $-\text{L}-\text{Ar}^3$, said phenyl optionally substituted with one additional R^d moiety, wherein:

25 L is $-\text{O}-(\text{CH}_2)_{0-1}-$ or a covalent bond;

Ar^3 is:

(b) phenyl optionally substituted with one or two R^d moieties; or

(b) pyridyl optionally substituted with one or two R^d moieties;

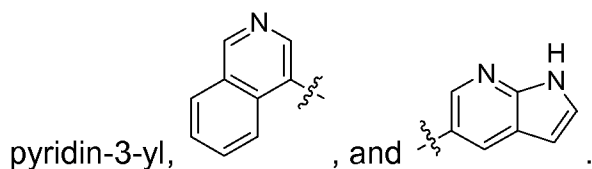
(iii) naphthyl;

30 (iv) 5,6,7,8-tetrahydro-naphthalenyl; or

(v) quinolinyl optionally substituted with a halo.

2. A composition of matter as in claim 1, wherein Ar^1 is a ring system selected from benzo[d]isoxazol-3-yl, 2,1,3-benzoxadiazol-4-yl, 4-chloropyridin-
35 3-yl, imidazo[1,2-a]pyridin-3-yl, imidazo[1,2-a]pyridin-5-yl, imidazo[1,2-

- 5 a]pyridin-6-yl, imidazo[1,2-b]pyridazin-3-yl, pyridin-3-yl, 6-[1,2,3]triazol-2-yl-



3. A composition of matter as in claim 1, wherein Q is -CH₂CH₂O-.
- 10 4. A composition of matter as in claim 1, wherein Q is -CH₂O-.
5. A composition of matter as in claim 1, wherein Ar² is phenyl substituted at the 3- or 4- position with -L-Ar³.
- 15 6. A composition of matter as in claim 5, wherein L is -O- and Ar³ is phenyl optionally substituted with one or two R^d moieties.
7. A composition of matter as in claim 5, wherein L is -O- and Ar³ is pyridyl optionally substituted with one or two R^d moieties.
- 20 8. A composition of matter as in claim 5, wherein Q is -CH₂CH₂O-.
9. A composition of matter as in claim 5, wherein Q is -CH₂O-.
- 25 10. A composition of matter as in claim 1, wherein Ar² is phenyl optionally substituted with one or two R^d.
11. A composition of matter as in claim 10, wherein Q is -CH₂CH₂O-.
- 30 12. A composition of matter as in claim 1, selected from the group consisting of:
 3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid imidazo[1,2-b]pyridazin-3-ylamide;
 3-Phenoxymethyl-azetidine-1-carboxylic acid pyridin-3-ylamide;
 3-(4-Bromo-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;

3-(3-Ethoxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(4-Trifluoromethoxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(3-Trifluoromethoxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(Benzo[1,3]dioxol-5-yloxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(3-Trifluoromethyl-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(Quinolin-2-yloxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(3-Chloro-4-trifluoromethyl-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(3-Ethynyl-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(3-Butoxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(5,6,7,8-Tetrahydro-naphthalen-2-yloxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(Quinolin-7-yloxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(Quinolin-6-yloxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(4-Phenoxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(3-Phenoxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(Biphenyl-4-yloxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(4-Benzyloxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(Naphthalen-2-yloxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(3,4-Dichloro-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(8-Chloro-quinolin-2-yloxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-(3-Benzyloxy-phenoxy-methyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[3-(4-Methanesulfonyl-phenoxy)-phenoxy-methyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[3-(4-Trifluoromethoxy-phenoxy)-phenoxy-methyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[3-(4-Chloro-phenoxy)-phenoxy-methyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-{2-[3-(4-Methanesulfonyl-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;
3-{2-[3-(4-Trifluoromethoxy-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid pyridin-3-ylamide;

3-[2-(4-Trifluoromethoxy-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[2-(3-Trifluoromethoxy-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[2-(3-Phenoxy-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[2-(3-Butoxy-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[2-(3,4-Dichloro-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[2-(Naphthalen-2-yloxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[2-(Quinolin-2-yloxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[2-(3-Ethynyl-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid isoquinolin-4-ylamide;
3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid (6-[1,2,3]triazol-2-yl-pyridin-3-yl)-amide;
3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid (1H-pyrrolo[2,3-b]pyridin-5-yl)-amide;
3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid imidazo[1,2-a]pyridin-6-ylamide;
3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid imidazo[1,2-a]pyridin-3-ylamide;
3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid benzo[d]isoxazol-3-ylamide;
3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[2-(4-Chloro-3-trifluoromethyl-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[2-(3-Trifluoromethyl-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-[2-(3,5-Dichloro-phenoxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;
3-{2-[3-(5-Chloro-pyridin-2-yloxy)-phenoxy]-ethyl}-azetidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;
N-(4-Chloropyridin-3-yl)-3-[(3,4-dichlorophenoxy)methyl]pyrrolidine-1-carboxamide;
N-(4-Chloropyridin-3-yl)-3-({3-[4-(methylsulfonyl)phenoxy]phenoxy}methyl)pyrrolidine-1-carboxamide;
3-{{3-(4-Chlorophenoxy)phenoxy}methyl}-N-pyridin-3-ylpyrrolidine-1-carboxamide;
3-{{3-[4-(Methylsulfonyl)phenoxy]phenoxy}methyl}-N-pyridin-3-ylpyrrolidine-1-

carboxamide;

3-[2-(3,4-Dichloro-phenoxy)-ethyl]-pyrrolidine-1-carboxylic acid pyridin-3-ylamide;

3-(2-Phenoxy-ethyl)-pyrrolidine-1-carboxylic acid pyridin-3-ylamide;

(3S)-3-[3-(4-Chloro-phenoxy)-phenoxyethyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;

(3R)-3-[3-(4-Chloro-phenoxy)-phenoxyethyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;

3-(2-Phenoxy-ethyl)-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;

3-[2-(3,4-Dichloro-phenoxy)-ethyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;

3-[2-(3-Trifluoromethoxy-phenoxy)-ethyl]-pyrrolidine-1-carboxylic acid pyridin-3-ylamide;

3-[2-(4-Trifluoromethoxy-phenoxy)-ethyl]-pyrrolidine-1-carboxylic acid pyridin-3-ylamide;

3-[2-(3-Trifluoromethoxy-phenoxy)-ethyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;

3-[2-(4-Trifluoromethoxy-phenoxy)-ethyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;

3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-pyrrolidine-1-carboxylic acid pyridin-3-ylamide;

3-{2-[3-(4-Chloro-phenoxy)-phenoxy]-ethyl}-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;

3-[2-(4-Chloro-benzyloxy)-ethyl]-azetidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;

3-[2-(4-Chloro-benzyloxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;

3-[2-(3,4-Dichloro-benzyloxy)-ethyl]-azetidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;

3-[2-(3,4-Dichloro-benzyloxy)-ethyl]-azetidine-1-carboxylic acid pyridin-3-ylamide;

3-(2-Benzyloxy-ethyl)-azetidine-1-carboxylic acid pyridin-3-ylamide;

3-(2-Benzyloxy-ethyl)-azetidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;

3-(2,2-Difluoro-benzo[1,3]dioxol-5-ylmethyl)-pyrrolidine-1-carboxylic acid pyridin-3-ylamide;

3-(2,2-Difluoro-benzo[1,3]dioxol-5-ylmethyl)-pyrrolidine-1-carboxylic acid

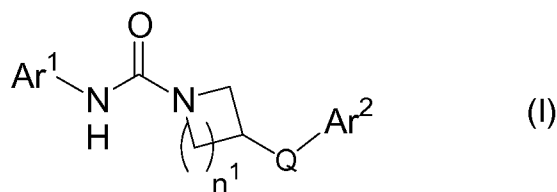
benzo[d]isoxazol-3-ylamide;

N-2,1,3-Benzoxadiazol-4-yl-3-[(2,2-difluoro-1,3-benzodioxol-5-yl)methyl]pyrrolidine-1-carboxamide;

- 3-[3-(4-Fluoro-phenoxy)-benzyl]-pyrrolidine-1-carboxylic acid pyridin-3-ylamide;
 3-[3-(4-Fluoro-phenoxy)-benzyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;
 3-[3-(4-Fluoro-phenoxy)-benzyl]-pyrrolidine-1-carboxylic acid imidazo[1,2-a]pyridin-5-ylamide;
 3-[2-(2,2-Difluoro-benzo[1,3]dioxol-5-yl)-ethyl]-pyrrolidine-1-carboxylic acid pyridin-3-ylamide;
 3-[2-(2,2-Difluoro-benzo[1,3]dioxol-5-yl)-ethyl]-pyrrolidine-1-carboxylic acid (4-chloro-pyridin-3-yl)-amide;
 3-{2-[3-(4-Fluoro-phenoxy)-phenyl]-ethyl}-pyrrolidine-1-carboxylic acid pyridin-3-ylamide;
 5 and pharmaceutically acceptable salts and prodrugs thereof.

13. A pharmaceutical composition comprising:

(a) a therapeutically effective amount of at least one chemical entity selected from the group consisting of compounds of Formula (I):



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wherein

Q is $-(CH_2)_{1-3}-$, $-(CH_2)_{1-2}O-$, or $-(CH_2)_{1-2}OCH_2-$;

n^1 is 1 or 2, with the proviso that when n^1 is 1, then Q cannot be $-CH_2-$;

Ar^1 is a ring system selected from benzo[d]isoxazolyl, 1H-pyrrolo[2,3-

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b]pyridinyl, 2,1,3-Benzoxadiazolyl, imidazo[1,2-a]pyridinyl, imidazo[1,2-b]pyridazinyl, isoquinolinyl, and pyridyl optionally substituted with triazolyl;

wherein each ring system is optionally substituted with halo;

Ar^2 is:

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- (i) phenyl optionally substituted with one or two R^d moieties; wherein each R^d moiety is independently selected from $-C_{1-4}alkyl$, $-OC_{1-4}alkyl$, halo, $-CF_3$, $-OCF_3$, and $-S(O)(O)C_{1-4}alkyl$, or two adjacent R^d moieties taken together form $-OCH_2O-$ or $-OCF_2O-$;

- (ii) phenyl substituted at the 3- or 4-position with $-L-Ar^3$, said phenyl optionally substituted with one additional R^d moiety, wherein:

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- 5 L is -O-(CH₂)₀₋₁- or a covalent bond;
Ar³ is:
- (c) phenyl optionally substituted with one or two R^d moieties; or
 - (b) pyridyl optionally substituted with one or two R^d moieties;
 - (iii) naphthyl;
- 10 (iv) 5,6,7,8-tetrahydro-naphthalenyl; or
(v) quinolinyl optionally substituted with a halo; and
pharmaceutically acceptable salts of compounds of Formula (I),
pharmaceutically acceptable prodrugs of compounds of Formula (I); and
(b) a pharmaceutically acceptable excipient.
- 15
14. A pharmaceutical composition according to claim 13, further comprising: an analgesic selected from the group consisting of opioids and non-steroidal anti-inflammatory drugs.
15. A pharmaceutical composition according to claim 13, further comprising: an additional active ingredient selected from the group consisting of aspirin, acetaminophen, opioids, ibuprofen, naproxen, COX-2 inhibitors, gabapentin, pregabalin, and tramadol.
16. A method for modulating FAAH activity, comprising exposing FAAH to an effective amount of at least one chemical entity as defined in claim 13.
- 20 17. A method for treating a subject suffering from or diagnosed with a disease, disorder, or medical condition mediated by FAAH activity, comprising administering to the subject in need of such treatment an effective amount of at least one chemical entity as defined in claim 13.
- 25 18. A method according to claim 17, wherein the disease, disorder, or medical condition is selected from the group consisting of: anxiety, depression, pain, sleep disorders, eating disorders, inflammation, movement disorders, HIV wasting syndrome, closed head injury, stroke, learning and memory disorders, Alzheimer's disease, epilepsy, Tourette's syndrome,
- 30 Niemann-Pick disease, Parkinson's disease, Huntington's chorea, optic

5 neuritis, autoimmune uveitis, drug withdrawal, nausea, emesis, sexual
dysfunction, post-traumatic stress disorder, cerebral vasospasm, glaucoma,
irritable bowel syndrome, inflammatory bowel disease, immunosuppression,
gastroesophageal reflux disease, paralytic ileus, secretory diarrhea, gastric
10 ulcer, rheumatoid arthritis, unwanted pregnancy, hypertension, cancer,
hepatitis, allergic airway disease, autoimmune diabetes, intractable pruritis,
neuroinflammation, diabetes, metabolic syndrome, and osteoporosis.

19. A method according to claim 17, wherein the disease, disorder, or
medical condition is pain or inflammation.

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20. A method according to claim 17, wherein the disease, disorder, or
medical condition is anxiety, a sleep disorder, an eating disorder, or a
movement disorder.

20 21. A method according to claim 17, wherein the disease, disorder, or
medical condition is multiple sclerosis.

22. A method according to claim 17, wherein the disease, disorder, or
medical condition is energy metabolism or bone homeostasis.

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INTERNATIONAL SEARCH REPORT

International application No
PCT/US2010/037388

A. CLASSIFICATION OF SUBJECT MATTER					
INV.	C07D401/12	C07D401/14	C07D413/12	C07D471/04	C07D487/04
	A61K31/422	A61K31/4245	A61K31/437	A61K31/4427	A61K31/4439
	A61K31/4725	A61K31/5025	A61P25/28	A61P29/00	A61P35/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
C07D A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, BEILSTEIN Data, BIOSIS, CHEM ABS Data, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2008/153752 A2 (JANSSEN PHARMACEUTICA NV [BE]; APODACA RICHARD [US]; BREITENBUCHER J G) 18 December 2008 (2008-12-18) claims; examples 15, 21, 22, 27	1-22
X	AHN K ET AL: "Discovery and Characterization of a Highly Selective FAAH Inhibitor that Reduces Inflammatory Pain" CHEMISTRY AND BIOLOGY, CURRENT BIOLOGY, LONDON, GB LNKD- DOI:10.1016/J.CHEMBIOL.2009.02.013, vol. 16, no. 4, 24 April 2009 (2009-04-24) , pages 411-420, XP026046192 ISSN: 1074-5521 [retrieved on 2009-04-23] table 2; compounds 5-13	1-22
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Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
20 July 2010	03/08/2010
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Authorized officer Stroeter, Thomas

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2010/037388

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 2008/075064 A1 (ASTRAZENECA AB [SE]; ASTRAZENECA UK LTD [GB]; BUTLIN ROGER JOHN [GB];) 26 June 2008 (2008-06-26) examples 10, 312, 344-346 -----	1-22
X,P	WANG J L ET AL: "Structure based design of novel irreversible FAAH inhibitors" BIOORGANIC & MEDICINAL CHEMISTRY LETTERS, PERGAMON, ELSEVIER SCIENCE, GB LNKD- DOI:10.1016/J.BMCL.2009.07.101, vol. 19, no. 20, 15 October 2009 (2009-10-15), pages 5970-5974, XP026640614 ISSN: 0960-894X [retrieved on 2009-07-25] compounds 6c,d -----	1-22

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No PCT/US2010/037388
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		CN 101686979 A	31-03-2010
		EP 2164493 A2	24-03-2010
		KR 20100017885 A	16-02-2010
		US 2009062294 A1	05-03-2009
WO 2008075064 A1	26-06-2008	AR 064496 A1	08-04-2009
		CL 38022007 A1	22-08-2008
		PE 18362008 A1	14-03-2009
		US 2009118332 A1	07-05-2009