



(86) Date de dépôt PCT/PCT Filing Date: 2014/11/07
(87) Date publication PCT/PCT Publication Date: 2015/05/14
(85) Entrée phase nationale/National Entry: 2016/05/03
(86) N° demande PCT/PCT Application No.: US 2014/064531
(87) N° publication PCT/PCT Publication No.: 2015/070007
(30) Priorité/Priority: 2013/11/08 (US61/901,689)

(51) Cl.Int./Int.Cl. *C07D 271/08* (2006.01),
A61K 31/4245 (2006.01), *A61P 35/00* (2006.01),
C07D 413/04 (2006.01)

(71) Demandeur/Applicant:
INCYTE HOLDINGS CORPORATION, US

(72) Inventeurs/Inventors:
TAO, MING, US;
FRIETZE, WILLIAM, US;
MELONI, DAVID J., US;
WENG, LINGKAI, US;
ZHOU, JIACHENG, US;
PAN, YONGCHUN, US

(74) Agent: SMART & BIGGAR

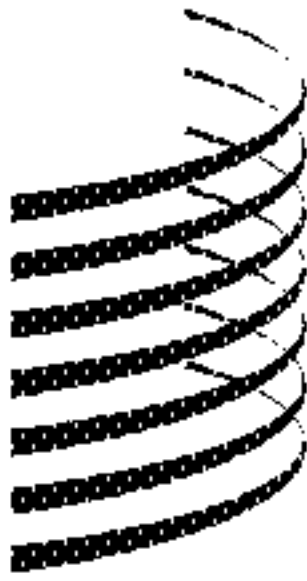
(54) Titre : PROCÉDE POUR LA SYNTHÈSE D'UN INHIBITEUR DE L'INDOLEAMINE 2,3-DIOXYGENASE
(54) Title: PROCESS FOR THE SYNTHESIS OF AN INDOLEAMINE 2,3-DIOXYGENASE INHIBITOR

(57) **Abrégé/Abstract:**

The present application is directed to processes and intermediates for making 4-({2-[(aminosulfonyl)amino]ethyl}amino)-N-(3-bromo-4-fluorophenyl)-N'-hydroxy-1,2,5-oxadiazole-3-carboximidamide, which is an inhibitor of indoleamine 2,3-dioxygenase, useful in the treatment of cancer and other disorders.



(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)
(19) World Intellectual Property Organization
International Bureau
(43) International Publication Date
14 May 2015 (14.05.2015)


WIPO | PCT


(10) International Publication Number
WO 2015/070007 A1

- (51) International Patent Classification:
C07D 271/08 (2006.01) A61K 31/4245 (2006.01)
C07D 413/04 (2006.01) A61P 35/00 (2006.01)

(21) International Application Number:
PCT/US2014/064531

(22) International Filing Date:
7 November 2014 (07.11.2014)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
61/901,689 8 November 2013 (08.11.2013) US

(71) Applicant: INCYTE CORPORATION [US/US]; 1801 Augustine Cut-Off, Wilmington, Delaware 19803 (US).

(72) Inventors: TAO, Ming; 1704 Brittany Drive, Maple Glen, Pennsylvania 19002 (US). FRIETZE, William; 900 Merrybell Lane, Kennett Square, Rhode Island 19348 (US). MELONI, David J.; 127 Megan Drive, Bear, Delaware 19701 (US). WENG, Lingkai; 28 Landmark Drive, Malvern, Pennsylvania 19355 (US). ZHOU, Jiacheng; 46 Bay Boulevard, Newark, Delaware 19702 (US). PAN, Yongchun; 11 Capano Drive, Apt. B6, Newark, Delaware 19702 (US).
- (74) Agent: GOODSON, Susanne H.; Fish & Richardson P.C., P.O. Box 1022, Minneapolis, Minnesota 55440-1022 (US).

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

Published:
— with international search report (Art. 21(3))

(54) Title: PROCESS FOR THE SYNTHESIS OF AN INDOLEAMINE 2,3-DIOXYGENASE INHIBITOR

(57) Abstract: The present application is directed to processes and intermediates for making 4-({2-[(aminosulfonyl)amino]ethyl}amino)-N-(3-bromo-4-fluorophenyl)-N'-hydroxy-1,2,5-oxadiazole-3-carboximidamide, which is an inhibitor of indoleamine 2,3-dioxygenase, useful in the treatment of cancer and other disorders.

PROCESS FOR THE SYNTHESIS OF AN INDOLEAMINE 2,3-DIOXYGENASE INHIBITOR

This application claims the benefit of priority of U.S. Prov. Appl. No. 61/901,689,
5 filed November 8, 2013, which is incorporated herein by reference in its entirety.

FIELD OF THE INVENTION

The present application relates to processes and intermediates for making 4-({2-
[(aminosulfonyl)amino]ethyl}amino)-N-(3-bromo-4-fluorophenyl)-N'-hydroxy-1,2,5-
10 oxadiazole-3-carboximidamide, which is an inhibitor of indoleamine 2,3-dioxygenase useful
in the treatment of cancer and other disorders.

BACKGROUND OF THE INVENTION

Tryptophan (Trp) is an essential amino acid required for the biosynthesis of proteins,
15 niacin and the neurotransmitter 5-hydroxytryptamine (serotonin). The enzyme indoleamine
2,3-dioxygenase (also known as INDO or IDO) catalyzes the first and rate limiting step in the
degradation of L-tryptophan to N-formyl-kynurenine. In human cells, a depletion of Trp
resulting from IDO activity is a prominent gamma interferon (IFN- γ) –inducible
antimicrobial effector mechanism. IFN- γ stimulation induces activation of IDO, which leads
20 to a depletion of Trp, thereby arresting the growth of Trp-dependent intracellular pathogens
such as *Toxoplasma gondii* and *Chlamydia trachomatis*. IDO activity also has an
antiproliferative effect on many tumor cells, and IDO induction has been observed *in vivo*
during rejection of allogeneic tumors, indicating a possible role for this enzyme in the tumor
rejection process (Daubener, *et al.*, 1999, *Adv. Exp. Med. Biol.*, 467: 517-24; Taylor, *et al.*,
25 1991, *FASEB J.*, 5: 2516-22).

It has been observed that HeLa cells co-cultured with peripheral blood lymphocytes
(PBLs) acquire an immuno-inhibitory phenotype through up-regulation of IDO activity. A
reduction in PBL proliferation upon treatment with interleukin-2 (IL2) was believed to result
from IDO released by the tumor cells in response to IFNG secretion by the PBLs. This effect
30 was reversed by treatment with 1-methyl-tryptophan (1MT), a specific IDO inhibitor. It was
proposed that IDO activity in tumor cells may serve to impair antitumor responses (Logan, *et al.*, 2002, *Immunology*, 105: 478-87).

Recently, an immunoregulatory role of Trp depletion has received much attention.
Several lines of evidence suggest that IDO is involved in induction of immune tolerance.

Studies of mammalian pregnancy, tumor resistance, chronic infections and autoimmune diseases have shown that cells expressing IDO can suppress T-cell responses and promote tolerance. Accelerated Trp catabolism has been observed in diseases and disorders associated with cellular immune activation, such as infection, malignancy, autoimmune diseases and
5 AIDS, as well as during pregnancy. For example, increased levels of IFNs and elevated levels of urinary Trp metabolites have been observed in autoimmune diseases; it has been postulated that systemic or local depletion of Trp occurring in autoimmune diseases may relate to the degeneration and wasting symptoms of these diseases. In support of this hypothesis, high levels of IDO were observed in cells isolated from the synovia of arthritic
10 joints. IFNs are also elevated in human immunodeficiency virus (HIV) patients and increasing IFN levels are associated with a worsening prognosis. Thus, it was proposed that IDO is induced chronically by HIV infection, and is further increased by opportunistic infections, and that the chronic loss of Trp initiates mechanisms responsible for cachexia, dementia and diarrhea and possibly immunosuppression of AIDS patients (Brown, *et al.*,
15 1991, *Adv. Exp. Med. Biol.*, 294: 425-35). To this end, it has recently been shown that IDO inhibition can enhance the levels of virus-specific T cells and, concomitantly, reduce the number of virally-infected macrophages in a mouse model of HIV (Portula *et al.*, 2005, *Blood*, 106: 2382-90).

IDO is believed to play a role in the immunosuppressive processes that prevent fetal
20 rejection in utero. More than 40 years ago, it was observed that, during pregnancy, the genetically disparate mammalian conceptus survives in spite of what would be predicted by tissue transplantation immunology (Medawar, 1953, *Symp. Soc. Exp. Biol.* 7: 320-38). Anatomic separation of mother and fetus and antigenic immaturity of the fetus cannot fully explain fetal allograft survival. Recent attention has focused on immunologic tolerance of the
25 mother. Because IDO is expressed by human syncytiotrophoblast cells and systemic tryptophan concentration falls during normal pregnancy, it was hypothesized that IDO expression at the maternal-fetal interface is necessary to prevent immunologic rejection of the fetal allografts. To test this hypothesis, pregnant mice (carrying syngeneic or allogeneic fetuses) were exposed to 1MT, and a rapid, T cell-induced rejection of all allogeneic concepti
30 was observed. Thus, by catabolizing tryptophan, the mammalian conceptus appears to suppresses T-cell activity and defends itself against rejection, and blocking tryptophan catabolism during murine pregnancy allows maternal T cells to provoke fetal allograft rejection (Munn, *et al.*, 1998, *Science*, 281: 1191-3).

Further evidence for a tumoral immune resistance mechanism based on tryptophan degradation by IDO comes from the observation that most human tumors constitutively express IDO, and that expression of IDO by immunogenic mouse tumor cells prevents their rejection by preimmunized mice. This effect is accompanied by a lack of accumulation of specific T cells at the tumor site and can be partly reverted by systemic treatment of mice with an inhibitor of IDO, in the absence of noticeable toxicity. Thus, it was suggested that the efficacy of therapeutic vaccination of cancer patients might be improved by concomitant administration of an IDO inhibitor (Uyttenhove *et al.*, 2003, *Nature Med.*, 9: 1269-74). It has also been shown that the IDO inhibitor, 1-MT, can synergize with chemotherapeutic agents to reduce tumor growth in mice, suggesting that IDO inhibition may also enhance the anti-tumor activity of conventional cytotoxic therapies (Muller *et al.*, 2005, *Nature Med.*, 11: 312-9).

One mechanism contributing to immunologic unresponsiveness toward tumors may be presentation of tumor antigens by tolerogenic host APCs. A subset of human IDO-expressing antigen-presenting cells (APCs) that coexpressed CD123 (IL3RA) and CCR6 and inhibited T-cell proliferation have also been described. Both mature and immature CD123-positive dendritic cells suppressed T-cell activity, and this IDO suppressive activity was blocked by 1MT (Munn, *et al.*, 2002, *Science*, 297: 1867-70). It has also been demonstrated that mouse tumor-draining lymph nodes (TDLNs) contain a subset of plasmacytoid dendritic cells (pDCs) that constitutively express immunosuppressive levels of IDO. Despite comprising only 0.5% of lymph node cells, *in vitro*, these pDCs potently suppressed T cell responses to antigens presented by the pDCs themselves and also, in a dominant fashion, suppressed T cell responses to third-party antigens presented by nonsuppressive APCs. Within the population of pDCs, the majority of the functional IDO-mediated suppressor activity segregated with a novel subset of pDCs coexpressing the B-lineage marker CD19. Thus, it was hypothesized that IDO-mediated suppression by pDCs in TDLNs creates a local microenvironment that is potently suppressive of host antitumor T cell responses (Munn, *et al.*, 2004, *J. Clin. Invest.*, 114(2): 280-90).

IDO degrades the indole moiety of tryptophan, serotonin and melatonin, and initiates the production of neuroactive and immunoregulatory metabolites, collectively known as kynurenines. By locally depleting tryptophan and increasing proapoptotic kynurenines, IDO expressed by dendritic cells (DCs) can greatly affect T-cell proliferation and survival. IDO induction in DCs could be a common mechanism of deletional tolerance driven by regulatory T cells. Because such tolerogenic responses can be expected to operate in a variety of physiopathological conditions, tryptophan metabolism and kynurenine production might

represent a crucial interface between the immune and nervous systems (Grohmann, *et al.*, 2003, *Trends Immunol.*, 24: 242-8). In states of persistent immune activation, availability of free serum Trp is diminished and, as a consequence of reduced serotonin production, serotonergic functions may also be affected (Wirleitner, *et al.*, 2003, *Curr. Med. Chem.*, 10: 1581-91).

Interestingly, administration of interferon- α has been observed to induce neuropsychiatric side effects, such as depressive symptoms and changes in cognitive function. Direct influence on serotonergic neurotransmission may contribute to these side effects. In addition, because IDO activation leads to reduced levels of tryptophan, the precursor of serotonin (5-HT), IDO may play a role in these neuropsychiatric side effects by reducing central 5-HT synthesis. Furthermore, kynurenine metabolites such as 3-hydroxy-kynurenine (3-OH-KYN) and quinolinic acid (QUIN) have toxic effects on brain function. 3-OH-KYN is able to produce oxidative stress by increasing the production of reactive oxygen species (ROS), and QUIN may produce overstimulation of hippocampal N-methyl-D-aspartate (NMDA) receptors, which leads to apoptosis and hippocampal atrophy. Both ROS overproduction and hippocampal atrophy caused by NMDA overstimulation have been associated with depression (Wichers and Maes, 2004, *J. Psychiatry Neurosci.*, 29: 11-17). Thus, IDO activity may play a role in depression.

Small molecule inhibitors of IDO are being developed to treat or prevent IDO-related diseases such as those described above. For example, oxadiazole and other heterocyclic IDO inhibitors are reported in US 2006/0258719 and US 2007/0185165. PCT Publication WO 99/29310 reports methods for altering T cell-mediated immunity comprising altering local extracellular concentrations of tryptophan and tryptophan metabolites, using an inhibitor of IDO such as 1-methyl-DL-tryptophan, p-(3-benzofuranyl)-DL-alanine, p-[3-benzo(b)thienyl]-DL-alanine, and 6-nitro-L-tryptophan (Munn, 1999). Reported in WO 03/087347, also published as European Patent 1501918, are methods of making antigen-presenting cells for enhancing or reducing T cell tolerance (Munn, 2003). Compounds having indoleamine-2,3-dioxygenase (IDO) inhibitory activity are further reported in WO 2004/094409; and U.S. Patent Application Publication No. 2004/0234623 is directed to methods of treating a subject with a cancer or an infection by the administration of an inhibitor of indoleamine-2,3-dioxygenase in combination with other therapeutic modalities.

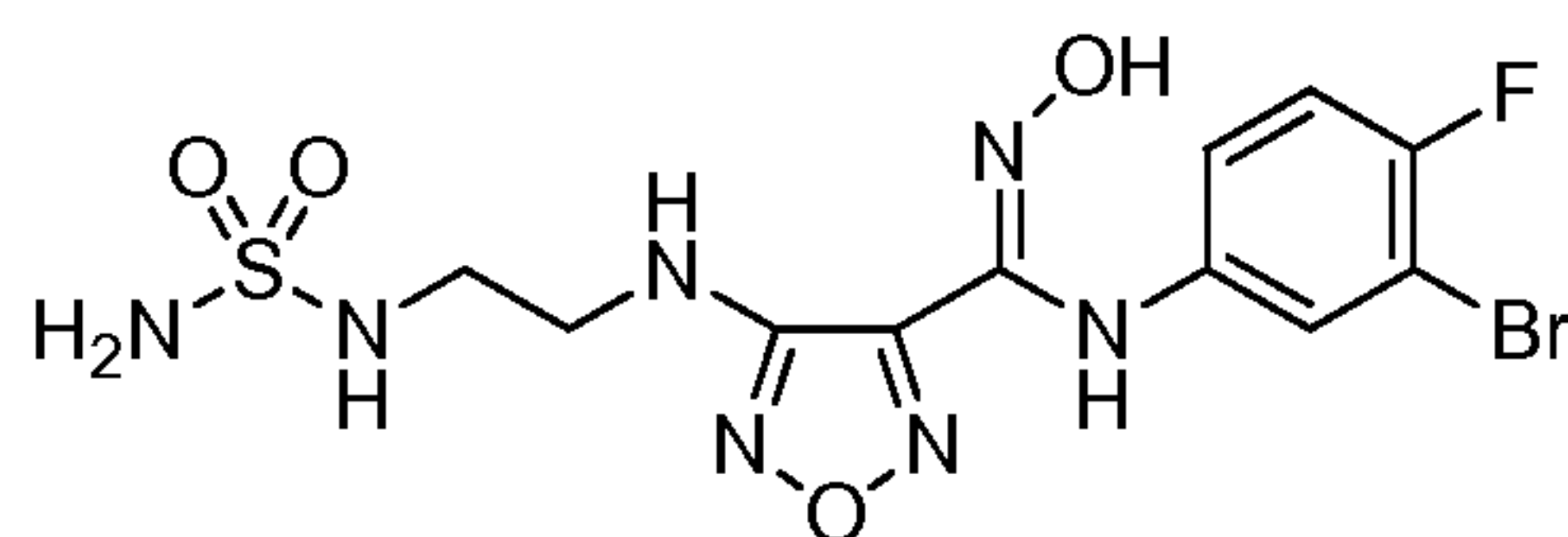
In light of the experimental data indicating a role for IDO in immunosuppression, tumor resistance and/or rejection, chronic infections, HIV-infection, AIDS (including its manifestations such as cachexia, dementia and diarrhea), autoimmune diseases or disorders

(such as rheumatoid arthritis), and immunologic tolerance and prevention of fetal rejection *in utero*, therapeutic agents aimed at suppression of tryptophan degradation by inhibiting IDO activity are desirable. Inhibitors of IDO can be used to activate T cells and therefore enhance T cell activation when the T cells are suppressed by pregnancy, malignancy or a virus such as HIV. Inhibition of IDO may also be an important treatment strategy for patients with neurological or neuropsychiatric diseases or disorders such as depression.

Due to the usefulness of IDO inhibitors, there is a need for development of new processes for making IDO inhibitors. This application is directed towards this need and others.

SUMMARY OF THE INVENTION

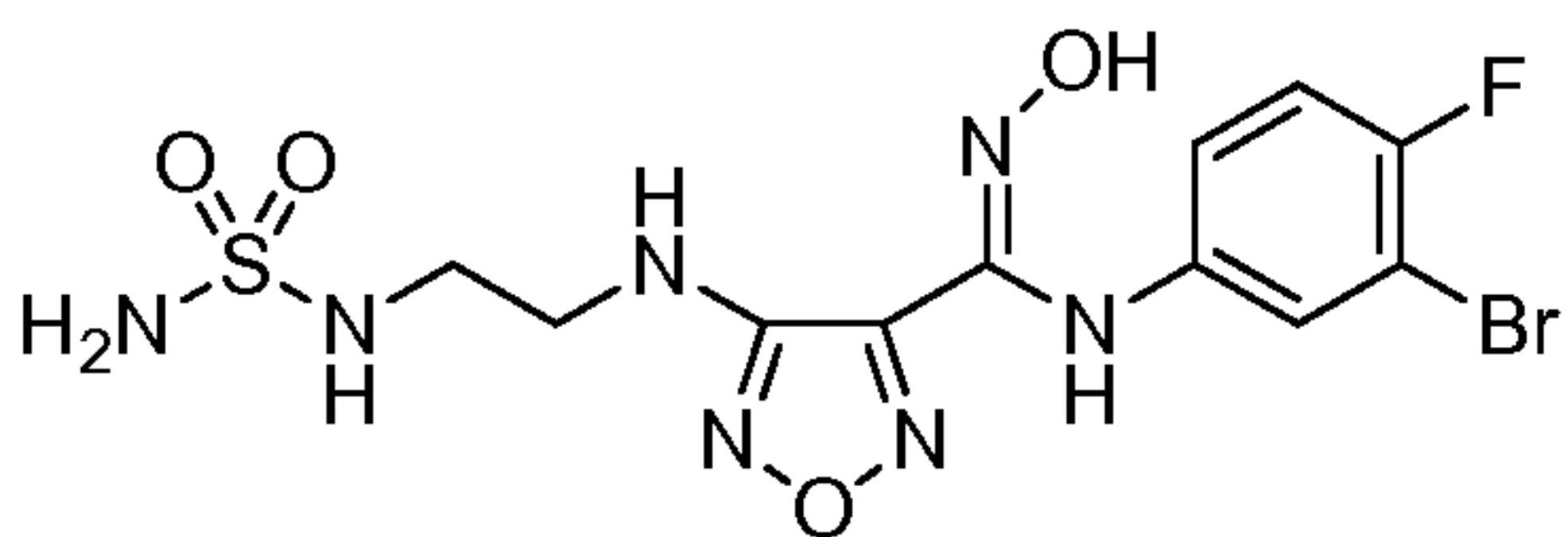
The compound 4-({2-[(aminosulfonyl)amino]ethyl} amino)-N-(3-bromo-4-fluorophenyl)-N'-hydroxy-1,2,5-oxadiazole-3-carboximidamide having Formula I:



I

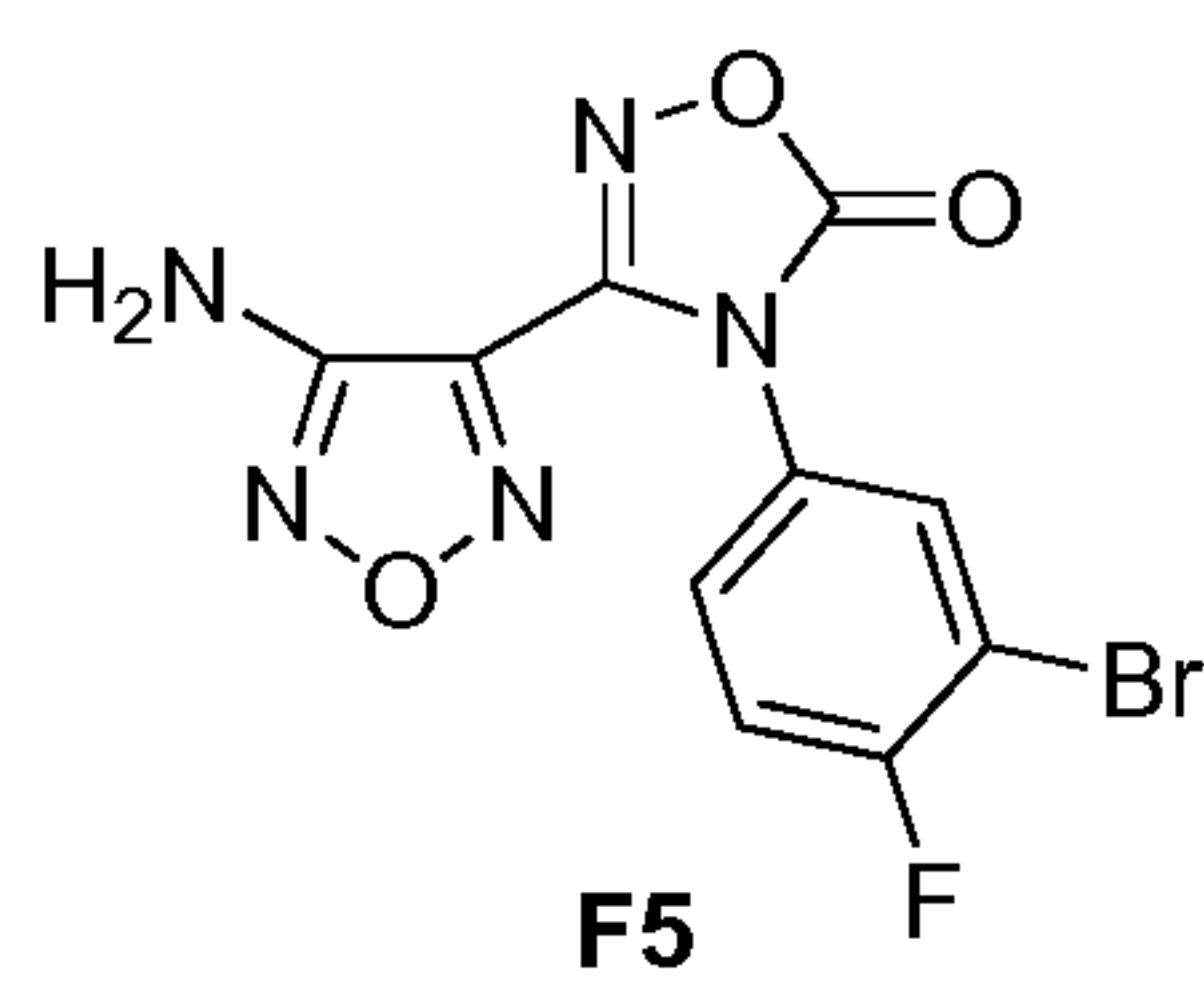
is an inhibitor of the enzyme indoleamine 2,3-dioxygenase (also known as IDO). The compound of Formula I, as well as its preparation and use, has been described in U.S. Patent No. 8,088,803, which is incorporated herein by reference in its entirety. The intermediates and processes provided herein help satisfy the ongoing need for the development of IDO inhibitors for the treatment of serious diseases.

The present application provides, *inter alia*, intermediates and processes for preparing a compound of Formula I:

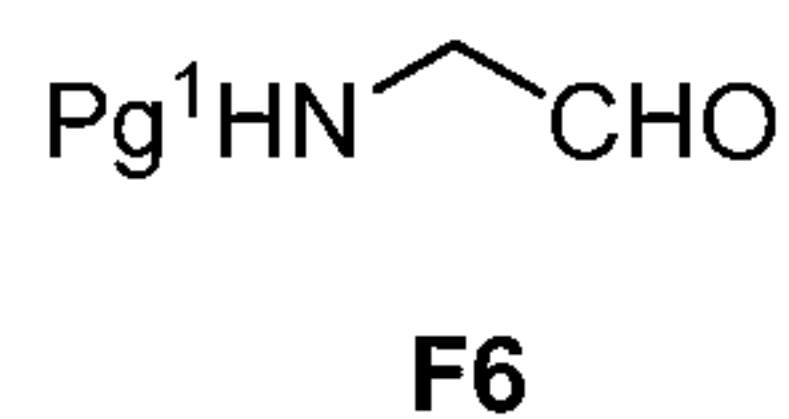


I

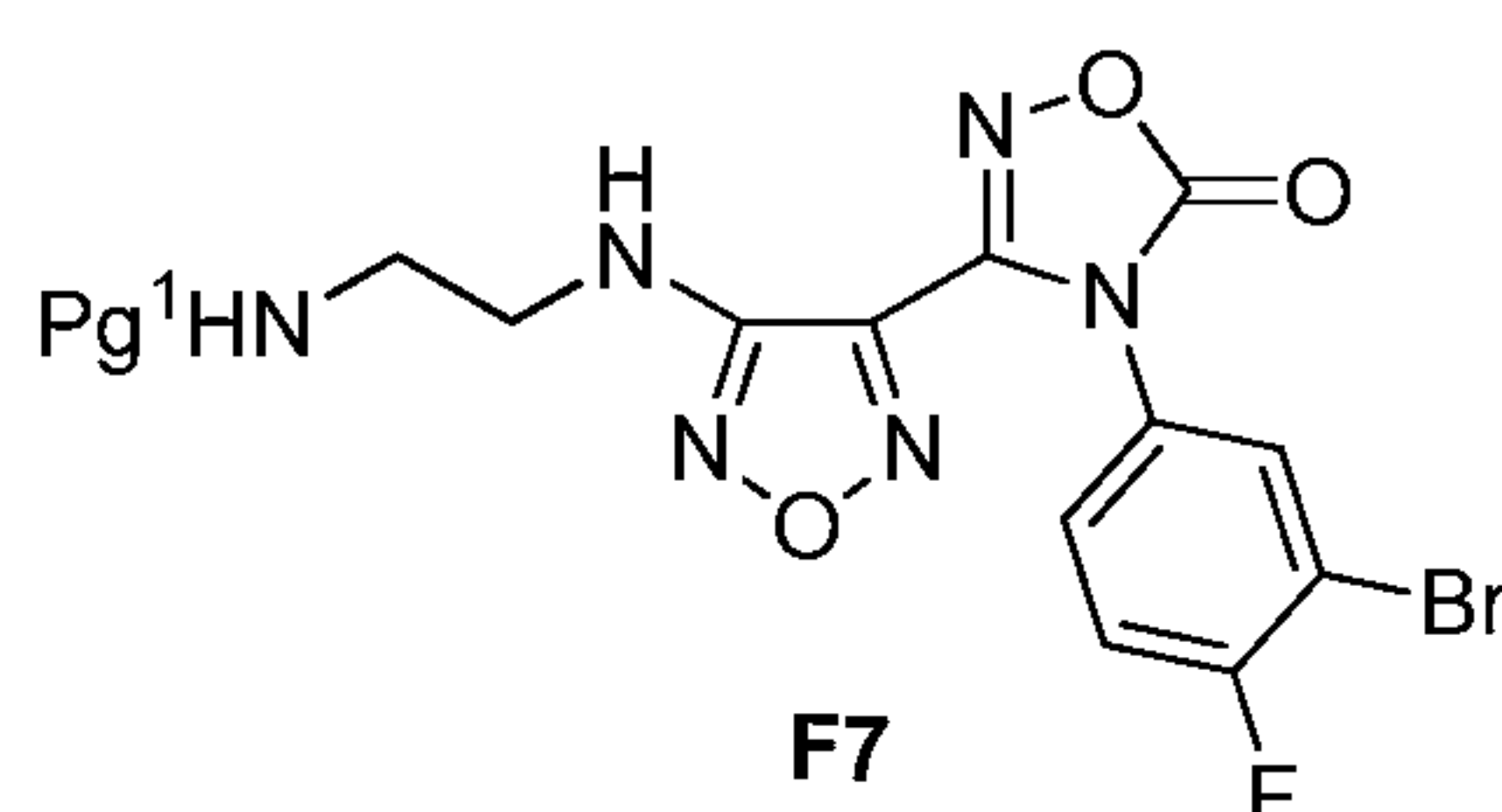
Accordingly, the present application provides a process comprising reacting a compound of Formula F5:



with an aldehyde of Formula F6:



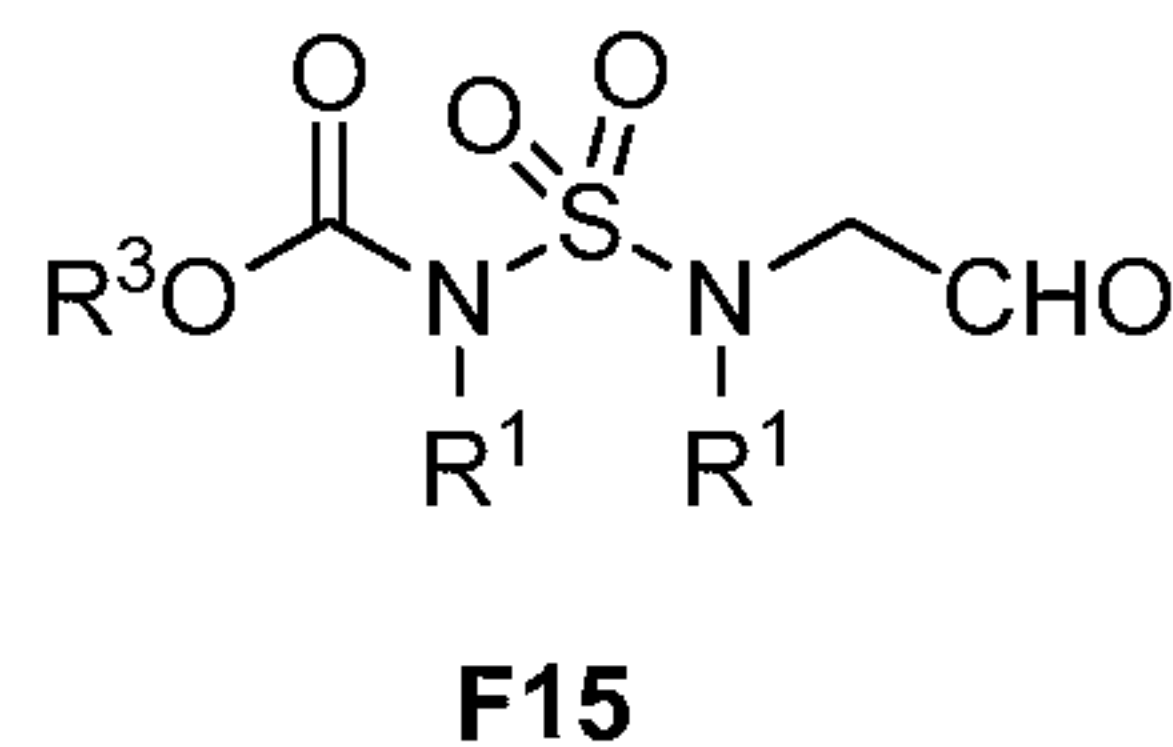
to afford a compound of Formula F7:



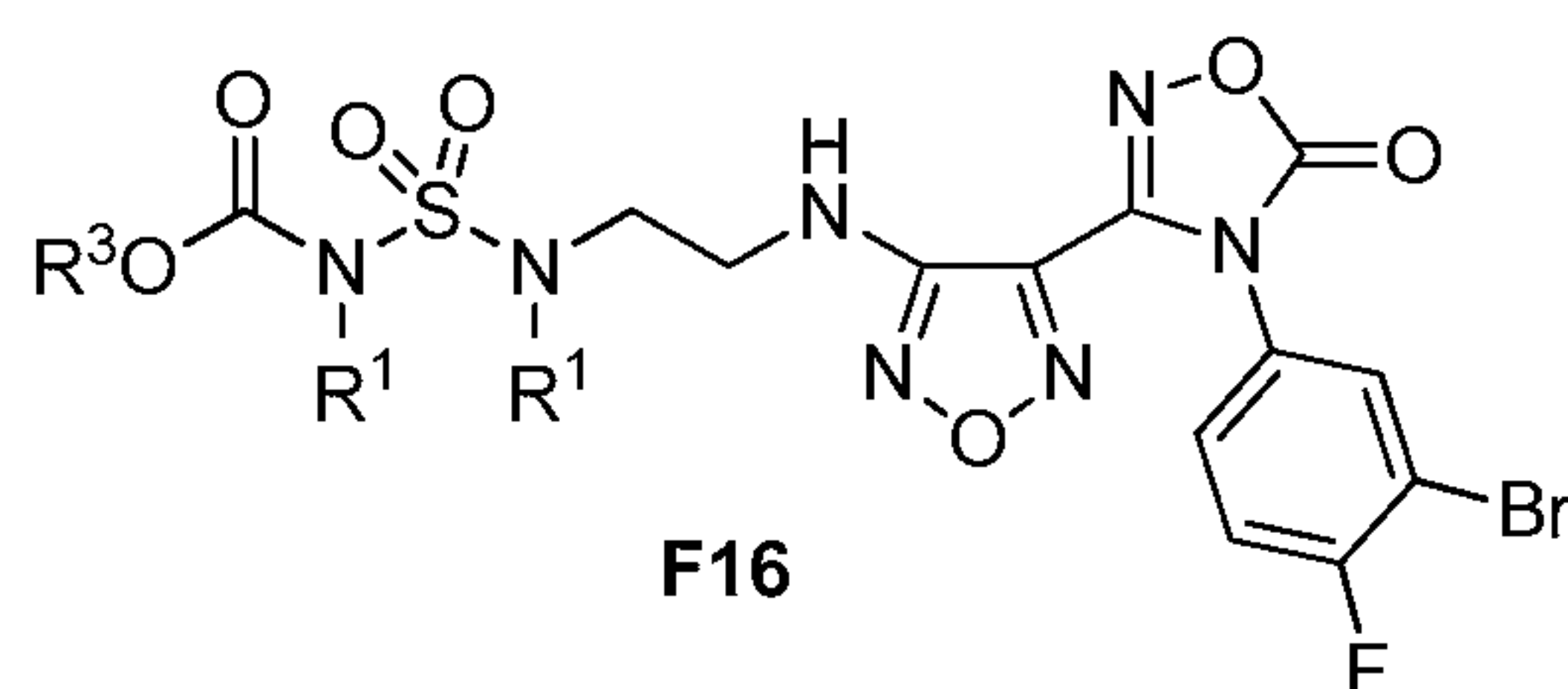
5

wherein Pg¹ is defined *infra*.

The present application further provides a process comprising reacting a compound of Formula F15:

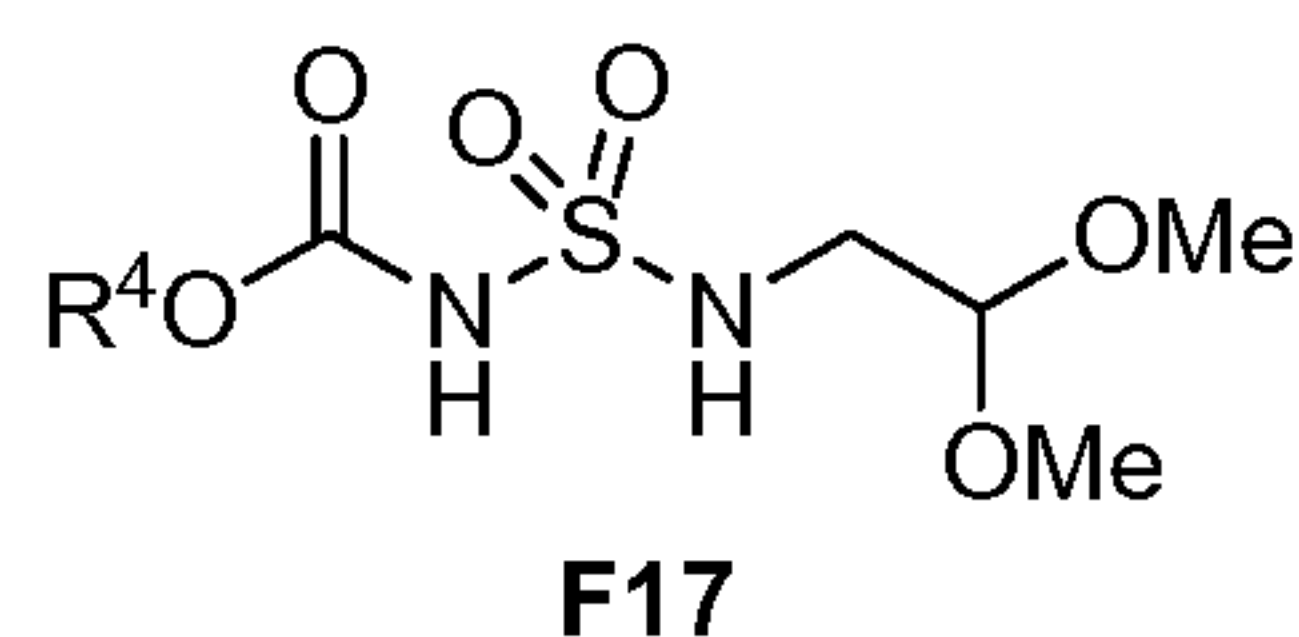


10 with a compound of Formula F5 to afford a compound of Formula F16:



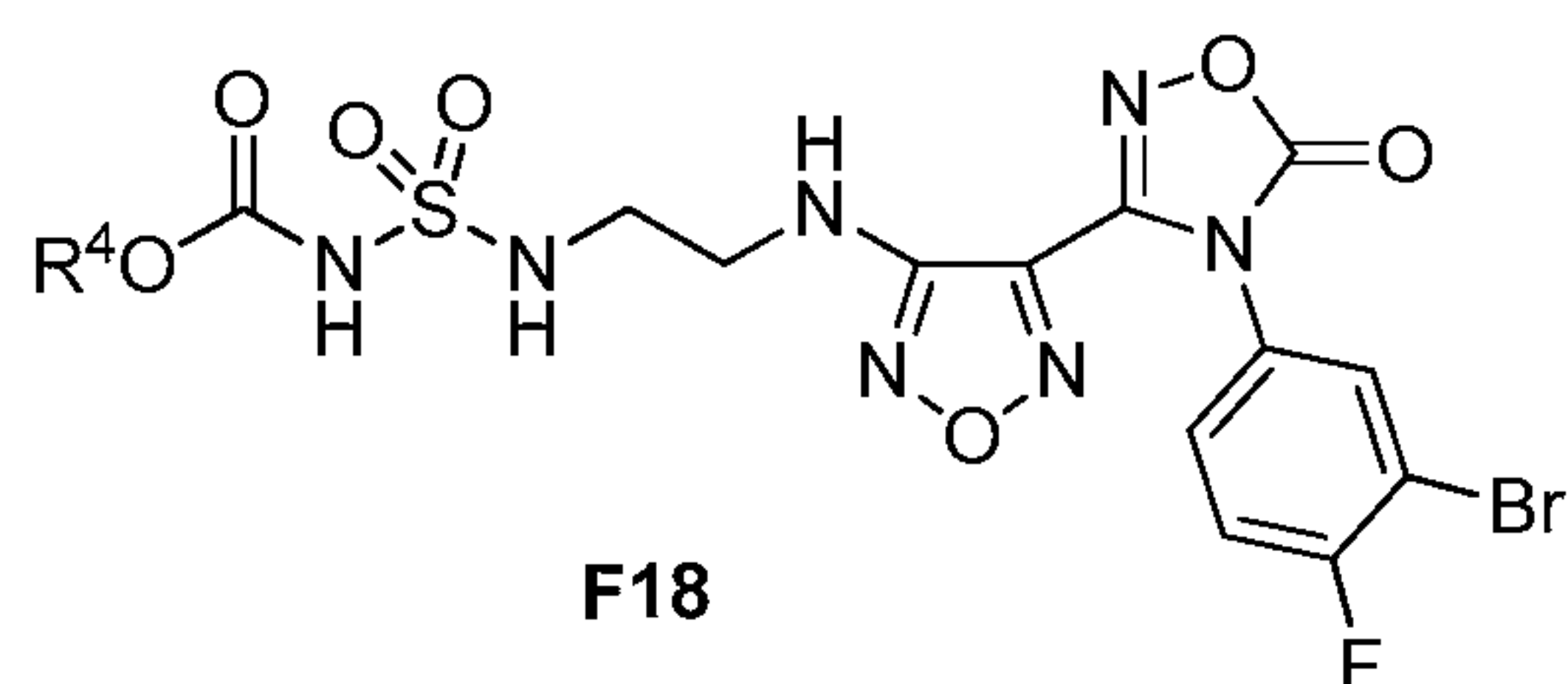
wherein R¹ and R³ are defined *infra*.

The present application further provides a process comprising reacting a compound of Formula F17:



15

with a compound of Formula F5 to afford a compound of Formula F18:



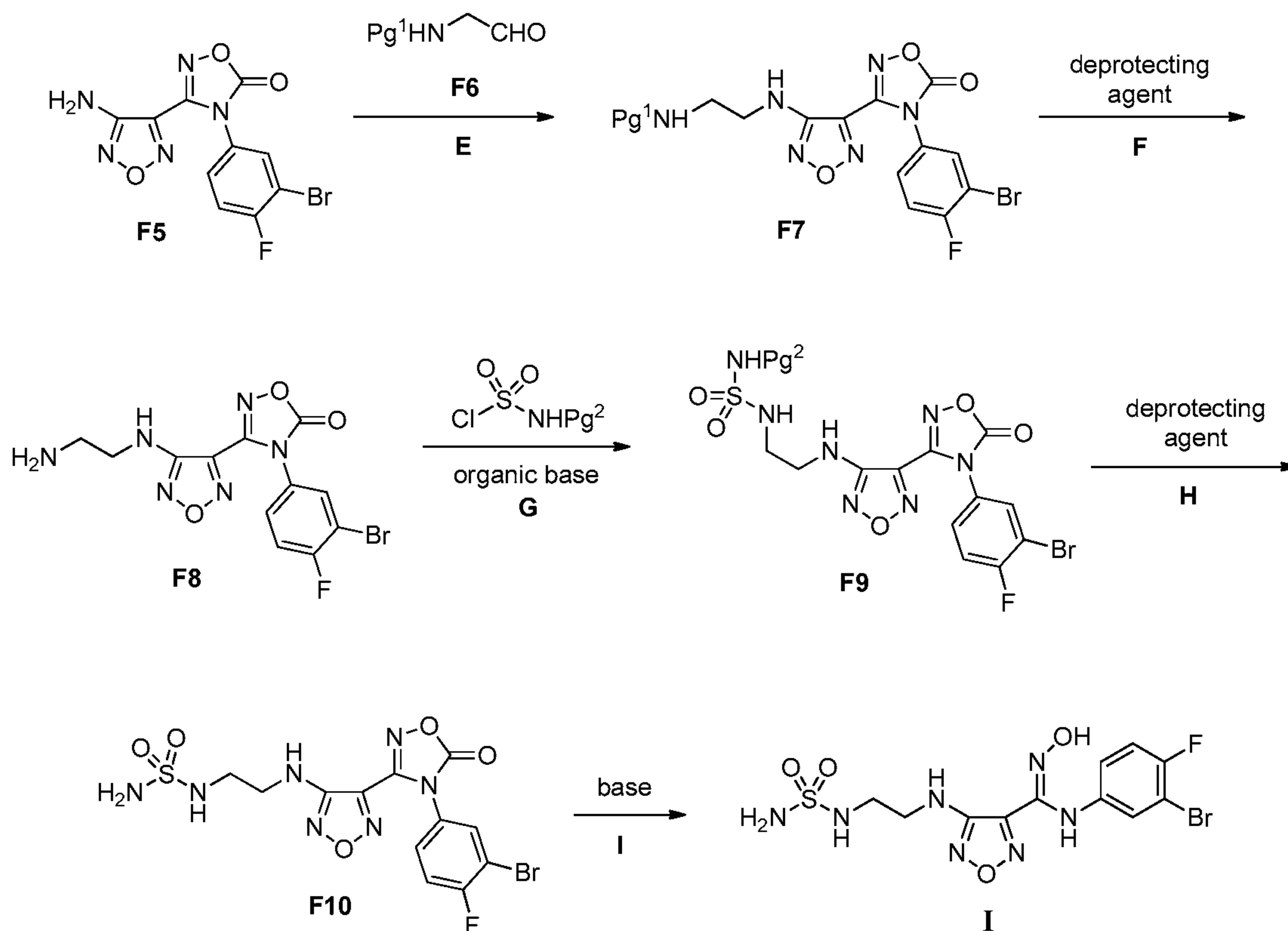
wherein R^4 is defined *infra*.

DETAILED DESCRIPTION

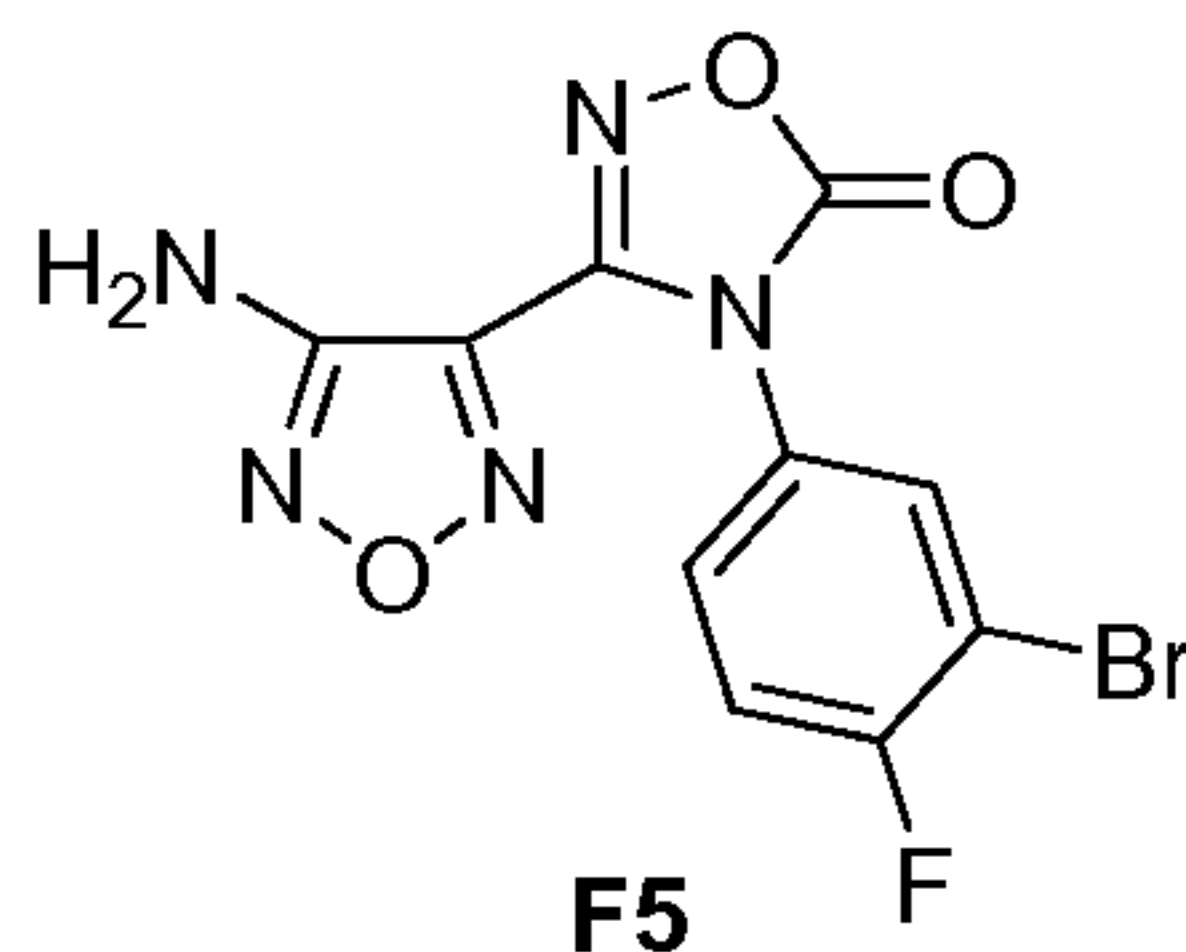
5 While certain of the processes steps are illustrated in the Schemes shown below, it is intended that the individual process steps may be claimed individually or in any combination (e.g., in Scheme I, steps E, F, G, H, and I may be claimed individually or in combination). It is not intended that the processes be limited to an overall process having each and every step in the Schemes below.

10 Accordingly, general scheme for the preparation of the compound of Formula I is described in Scheme 1.

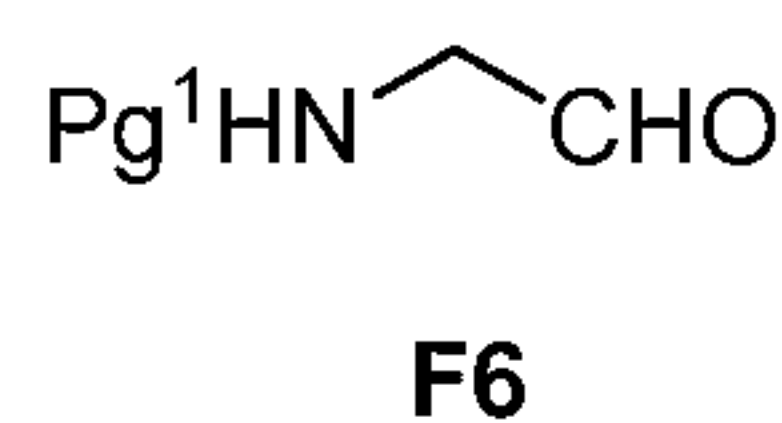
Scheme 1



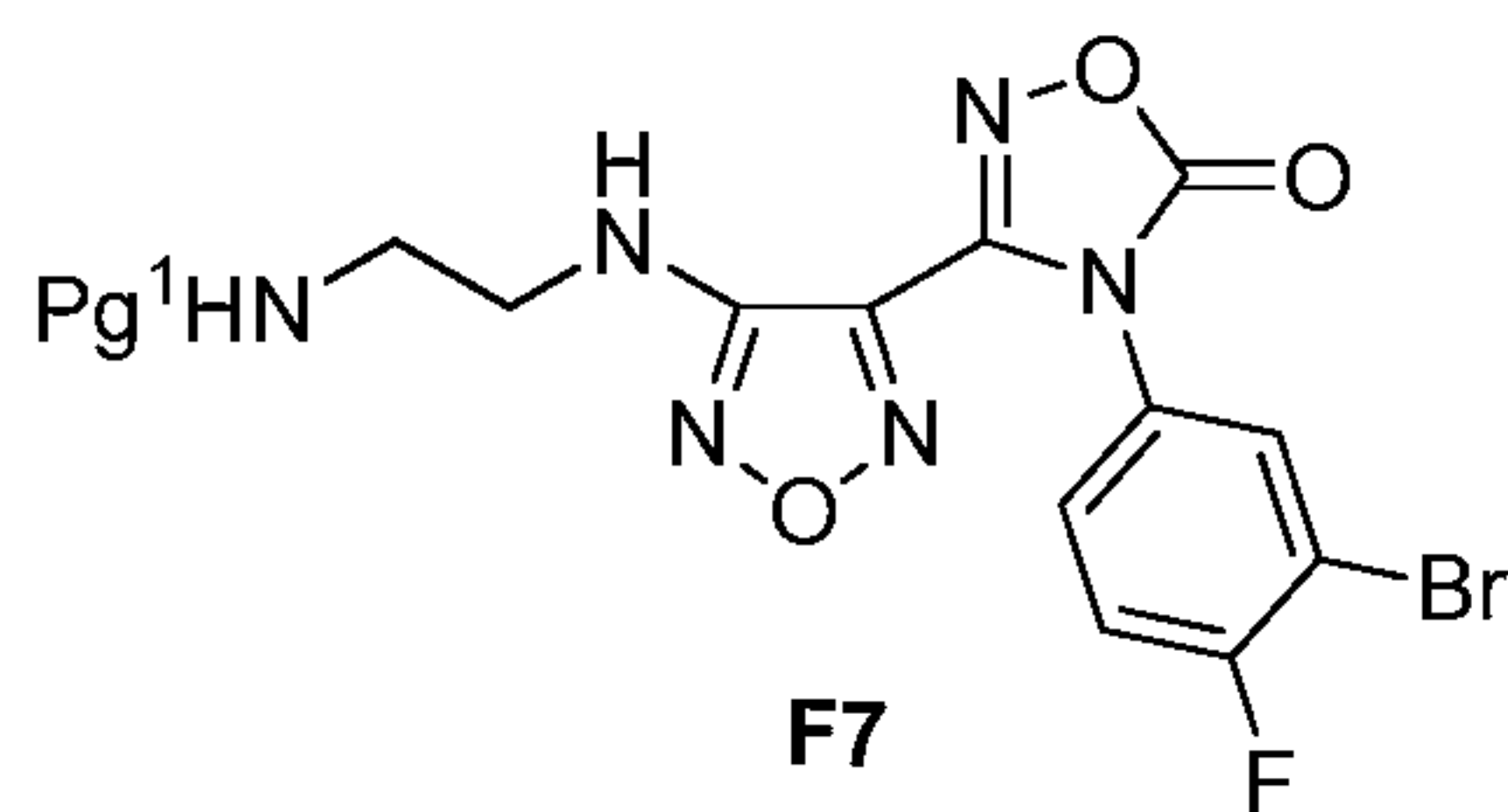
Accordingly, the present application provides a process comprising reacting a compound of Formula F5:



with an aldehyde of Formula F6:



wherein Pg¹ is an amino protecting group, to afford a compound of Formula F7:



Amino protecting groups Pg¹ may be used to prevent unwanted reactions of an amino group while performing a desired transformation. Amino protecting groups allow easy covalent attachment to a nitrogen atom as well as selective cleavage from the nitrogen atom. Suitable “amino protecting groups”, such as alkoxycarbonyl (such as ethoxycarbonyl, *tert*-butoxycarbonyl (Boc), benzyloxycarbonyl (Cbz), 9-fluorenylmethyloxycarbonyl (Fmoc), and the like), acyl (such as acetyl (Ac), benzoyl (Bz), and the like), sulfonyl (such as methanesulfonyl, trifluoromethanesulfonyl, and the like), arylalkyl (such as benzyl, 4-methoxybenzyl, diphenylmethyl, triphenylmethyl (trityl), and the like), alkenylalkyl (such as allyl, prenyl, and the like), diarylmethyleneyl (such as (C₆H₅)₂C=N, and the like), and silyl (such as *tert*-butyldimethylsilyl, triisopropylsilyl, and the like), are known to one skilled in the art. The chemistry of amino protecting groups can be found in Wuts and Greene, *Greene’s Protective Groups in Organic Synthesis*, 4th Ed., pp 696-926, John Wiley & Sons: New York, 2006, which is incorporated herein by reference in its entirety.

In some embodiments, Pg¹ is ethoxycarbonyl, *tert*-butoxycarbonyl, benzyloxycarbonyl, or 9-fluorenylmethyloxycarbonyl.

In some embodiments, Pg¹ is C₁₋₆ alkoxycarbonyl.

In some embodiments, Pg¹ is *tert*-butoxycarbonyl.

Appropriate solvents for Step E include, but are not limited to, methanol or tetrahydrofuran (THF), acetonitrile and the like. Halogenated hydrocarbon solvents (i.e., halogenated alkanes, such as dichloromethane, chloroform, dichloroethane or tetrachloroethane) can also be used.

5 In some embodiments, said reacting is performed in a solvent component comprising tetrahydrofuran. As used herein, a solvent component may refer to one solvent or a mixture of solvents. In some embodiments, the solvent component is an organic solvent. In some embodiments, said reacting is performed in a solvent component comprising a halogenated hydrocarbon solvent. In some embodiments, said halogenated hydrocarbon solvent is
10 dichloromethane.

In some embodiments, said reacting is performed in a solvent component comprising acetonitrile.

In some embodiments, said reacting is performed in a solvent component comprising dichloromethane and acetonitrile.

15 In some embodiments, said reacting is performed in the presence of a reducing agent.

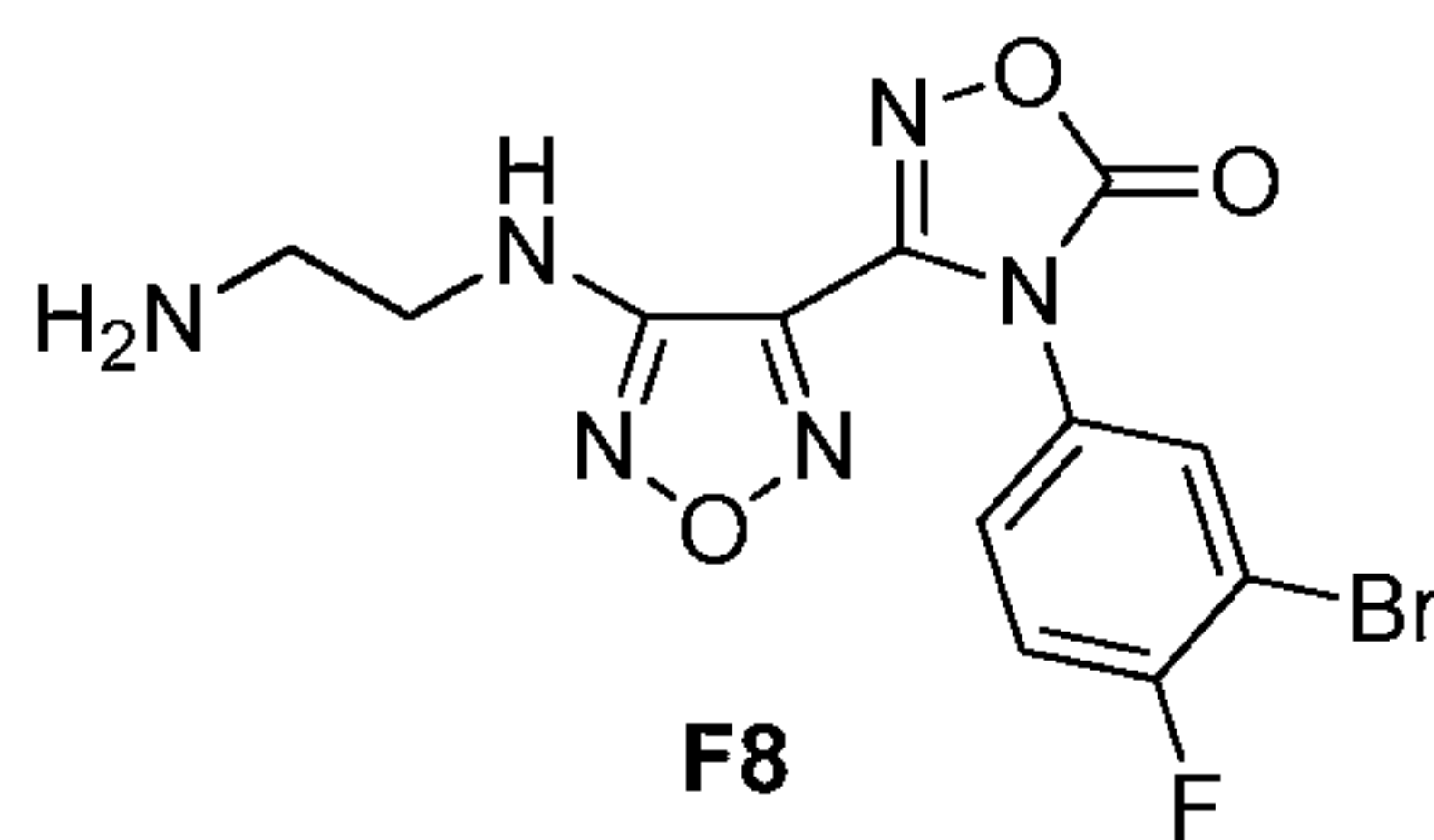
The reducing agent can be any compound capable of reducing an organic compound to a lower oxidation state. Reduction usually involves addition of hydrogen atoms or removal of oxygen atoms from a group. For example, aldehydes such as F6 can be reduced in the presence of an amine of Formula F5 (Step E, Scheme 1) by the addition of hydrogen,
20 either in the form of hydrogen gas (H_2) or using a hydride reagent (such as $NaB(OAc)_3H$, $NaBH_4$, $LiAlH_4$, and the like); using triphenylphosphine; or using a combination of sodium iodide, chlorotrimethylsilane, and methanol. In some embodiments, this step can be performed under acidic conditions in the presence of an acid (such as trifluoroacetic acid). In some embodiments, this step can be performed at a temperature from about $-15\text{ }^{\circ}C$ to about
25 $30\text{ }^{\circ}C$, e.g., from about $-15\text{ }^{\circ}C$ to about $0\text{ }^{\circ}C$, from about $-5\text{ }^{\circ}C$ to about $5\text{ }^{\circ}C$, from about $-5\text{ }^{\circ}C$ to about $0\text{ }^{\circ}C$, or from about $0\text{ }^{\circ}C$ to about $45\text{ }^{\circ}C$.

In some embodiments, said reducing agent is a borohydride reducing agent (e.g., $NaB(OAc)_3H$, $NaBH_4$, or other boron containing hydride reducing agent).

In some embodiments, said borohydride reducing agent is sodium
30 triacetoxyborohydride.

In some embodiments, said reacting is performed in the presence of trifluoroacetic acid.

In some embodiments, the process further comprises deprotecting said compound of Formula F7 to afford a compound of Formula F8:



Amino deprotecting agents useful for this Step F are known to those skilled in the art, such as those in Wuts and Greene (*supra*). In particular, the amino protecting groups described above can be conveniently removed using many available amino deprotecting agents that are specific to the various groups mentioned above without affecting other desired portions of the compound. The *tert*-butoxycarbonyl group can be removed (e.g., hydrolyzed) from the nitrogen atom, for example, by treatment with an acid (such as hydrochloric acid, trifluoroacetic acid, toluenesulfonic acid, and the like); a combination of reagents (e.g., mixture of acetyl chloride and methanol) known to generate an acid; or a Lewis acid (e.g., $\text{BF}_3 \cdot \text{Et}_2\text{O}$). The benzyloxycarbonyl group can be removed (e.g., hydrogenolyzed) from the nitrogen atom, for example, by treatment with hydrogen and a catalyst (such as palladium on carbon).

In some embodiments, the amino deprotecting agent is trifluoroacetic acid. In some embodiments, the amino deprotecting agent contains trifluoroacetic acid and >0.5% by volume of water, e.g., >1.0% by volume of water, >1.5% by volume of water, >2.0% by volume of water, from about 2% to about 10% by volume of water, from about 10% to about 20% by volume of water, or from about 20% to about 50% by volume of water. In some embodiments, the amino deprotecting agent can be a mixture of trifluoroacetic acid and water in a volumetric ratio of about 98:2. In some embodiments, the amino deprotecting agent can be hydrochloric acid, optionally in a solvent (e.g., water, THF, dioxane, ethyl acetate, etc.). In some embodiments, the solvent component is ethyl acetate. In some embodiments, the amino deprotecting agent can be hydrochloric acid optionally in a solvent such as an alcohol (such as isopropanol, methanol or ethanol). Halogenated hydrocarbon solvents (e.g., dichloromethane, chloroform, dichloroethane or tetrachloroethane) can also be used. In some embodiments, the molar ratio of hydrochloric acid and the compound of Formula F7 is about 6.0, about 5.0, about 4.0, about 3.0, about 2.0, about 1.0, or about 1.1. In some embodiments, Step F can be performed at a temperature from about -10 °C to about 60 °C, e.g., from about -10 °C to about 0 °C, from about 0 °C to about 25 °C, from about 25 °C to about 45 °C, or from about 45 °C to about 60 °C.

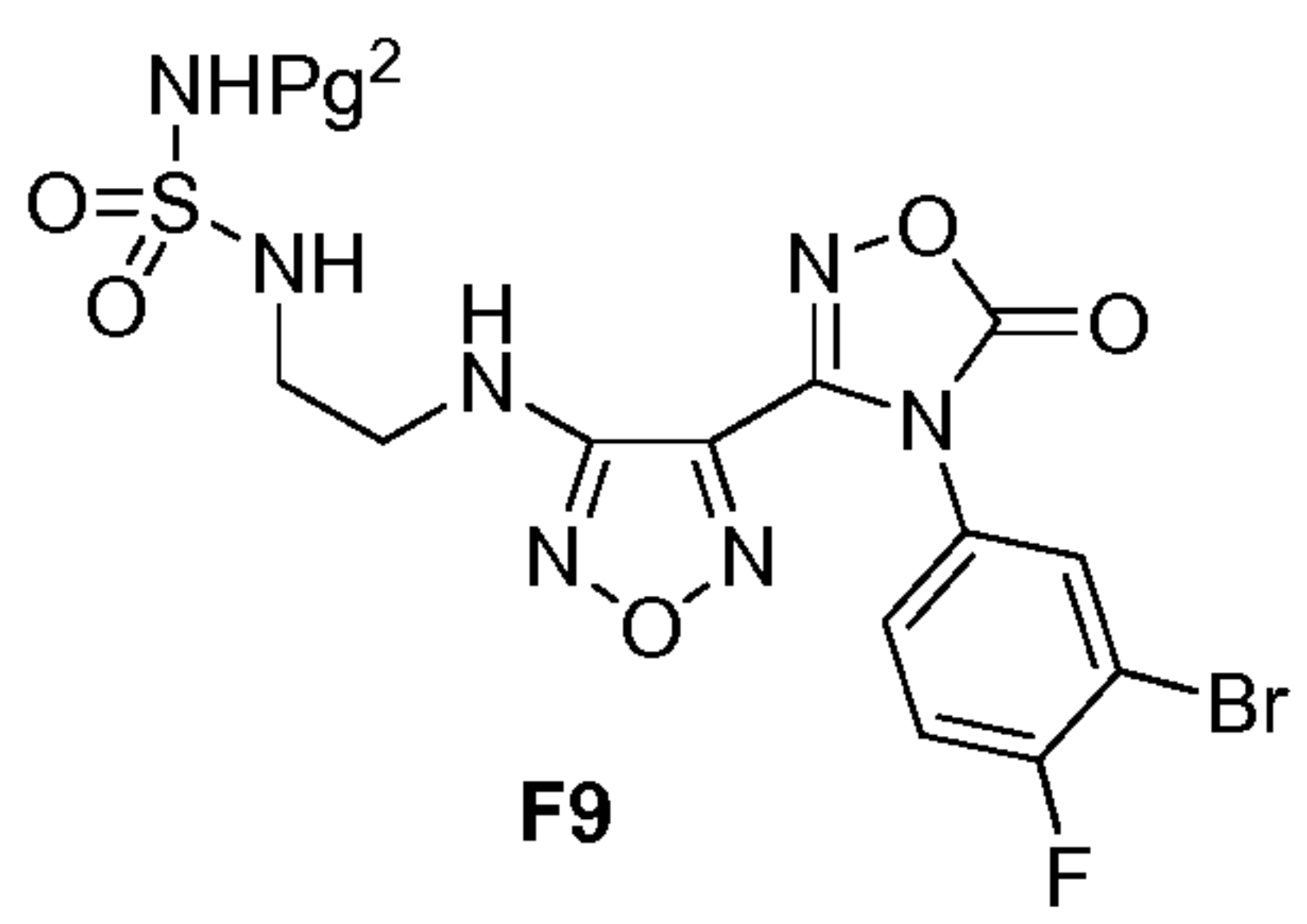
In some embodiments, said deprotecting comprises reacting the compound of Formula F7 with hydrochloric acid.

In some embodiments, said deprotecting comprises reacting the compound of Formula F7 with hydrochloric acid in a solvent component comprising isopropanol.

5 In some embodiments, said deprotecting comprises reacting the compound of Formula F7 with hydrochloric acid in a solvent component comprising a halogenated hydrocarbon solvent.

In some embodiments, said halogenated hydrocarbon solvent is dichloromethane.

10 In some embodiments, the invention further comprises reacting said compound of Formula F8, with $\text{Pg}^2\text{-NH-SO}_2\text{-X}$, in the presence of an organic base to afford a compound of Formula F9:



wherein:

Pg^2 is an amino protecting group; and

15 X is halo.

In some embodiments, $\text{Pg}^2\text{-NH-SO}_2\text{Cl}$ can be prepared and immediately used in the reaction with the compound of Formula F8. The protecting group Pg^2 could be selected from any of the protecting groups known in the art for protecting amines or sulfonamides (such as those described above for Pg^1). In some embodiments, Pg^2 can be an alkoxycarbonyl group
20 (such as *tert*-butoxycarbonyl).

Appropriate solvents include, but are not limited to, halogenated hydrocarbon solvents such as dichloromethane and the like. The organic base can be any base that serves to neutralize the HCl generated during the reaction of the compound of Formula F8 and the protected amino-sulfonyl chloride. The organic base can include acyclic tertiary amines such
25 as tri(C_{1-6})alkylamine (e.g., triethylamine, diisopropylethylamine (DIPEA) and the like), cyclic tertiary amines (e.g., N-methyl piperidine, 1,4-diazabicyclo[2.2.2]octane (DABCO) and the like). In some embodiments, the organic base can be triethylamine. In some embodiments, this step can be performed at a temperature from about $-15\text{ }^\circ\text{C}$ to about $60\text{ }^\circ\text{C}$,

e.g., from about -15 °C to about 0 °C, from about 0 °C to about 25 °C, from about 25 °C to about 45 °C, or from about 45 °C to about 60 °C.

In such embodiments, the $\text{Pg}^2\text{-NH-SO}_2\text{Cl}$ can be obtained by the reaction of an alcohol (such as, ethanol, *tert*-butyl alcohol and the like) with chlorosulfonyl isocyanate (ClS(O)₂NCO).

In some embodiments, Pg^2 is ethoxycarbonyl, *tert*-butoxycarbonyl, benzyloxycarbonyl, or 9-fluorenylmethyloxycarbonyl.

In some embodiments, Pg^2 is C₁₋₆ alkoxycarbonyl.

In some embodiments, Pg^2 is *tert*-butoxycarbonyl.

In some embodiments, said reacting is performed in a solvent component comprising a halogenated hydrocarbon solvent.

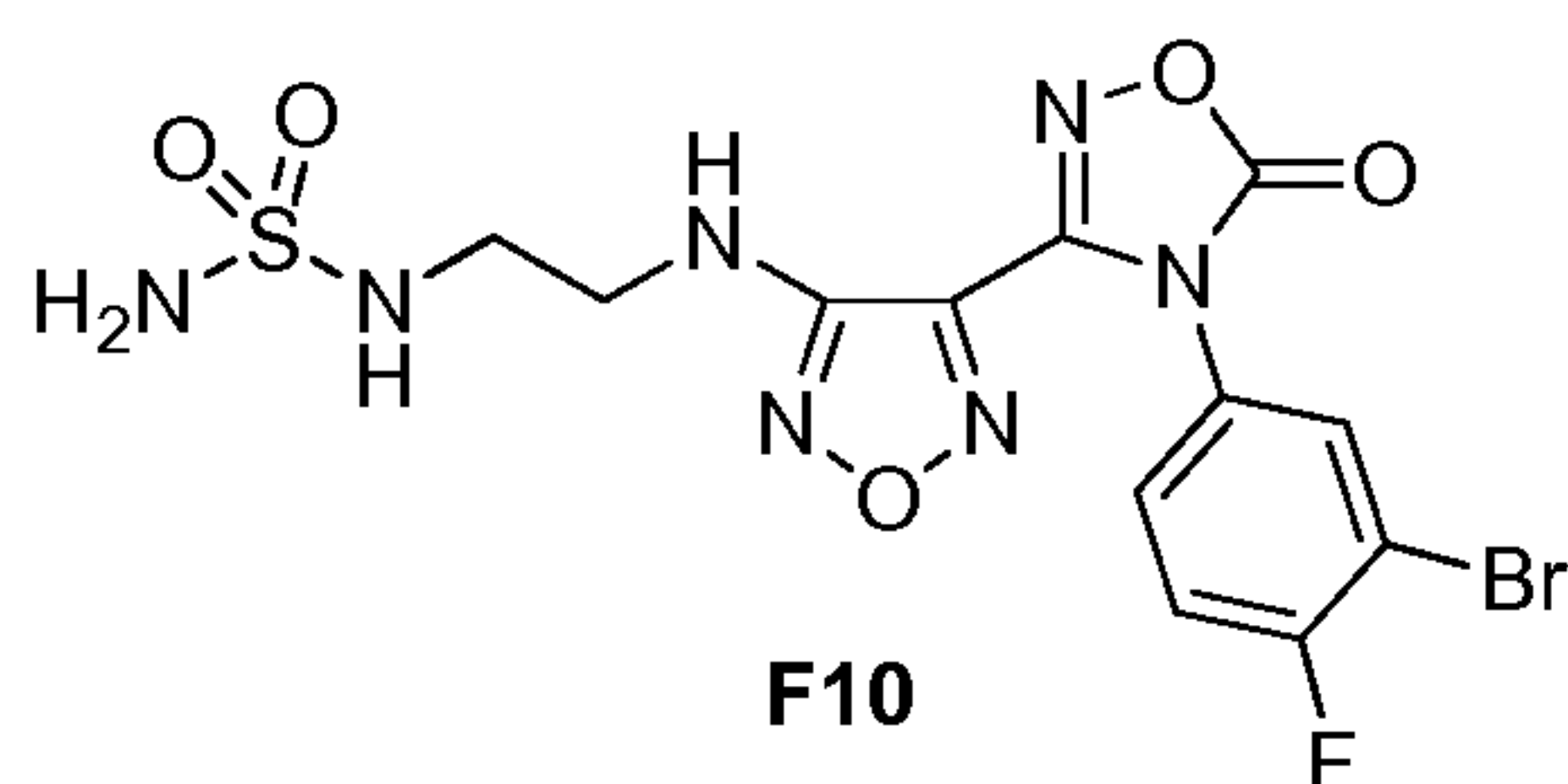
In some embodiments, said halogenated hydrocarbon solvent is dichloromethane.

In some embodiments, said organic base comprises a tri(C₁₋₆)alkylamine.

In some embodiments, said organic base is triethylamine.

In some embodiments, X is chloro.

In some embodiments, the invention further comprises deprotecting said compound of Formula F9 to afford a compound of Formula F10:



In some embodiments, suitable deprotecting agents may include those described above for deprotecting the compound of Formula F7.

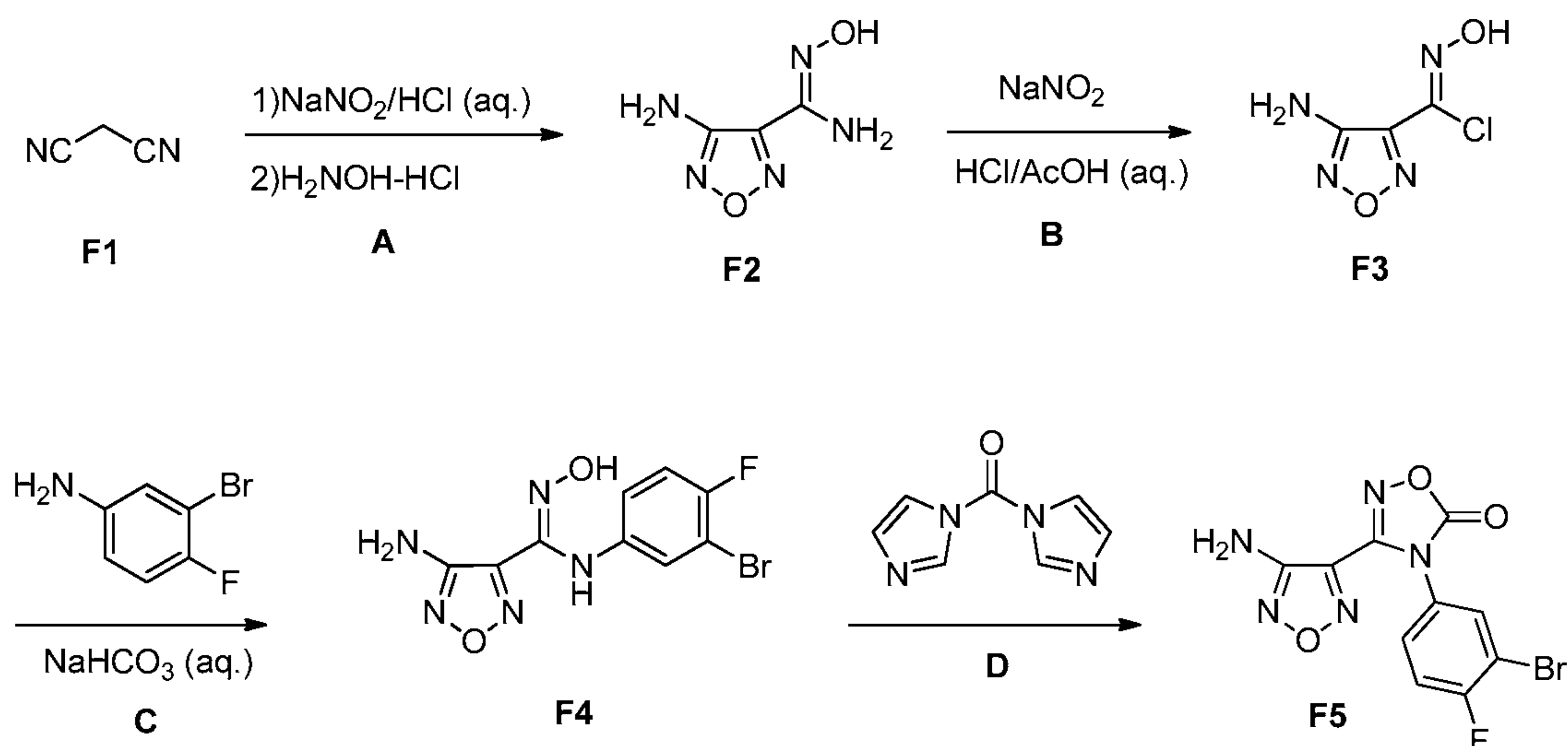
In some embodiments, said deprotecting comprises reacting a compound of Formula F9 with hydrochloric acid. In some embodiments, said deprotecting comprises reacting a compound of Formula F9 with hydrochloric acid in a solvent component comprising an alcohol. In some embodiments, said alcohol is ethanol. In some embodiments, said deprotecting comprises reacting a compound of Formula F9 with hydrochloric acid in a solvent component comprising ethyl acetate.

In some embodiments, the invention further comprises reacting said compound of Formula F10 with a base to afford a compound of Formula I:

I.

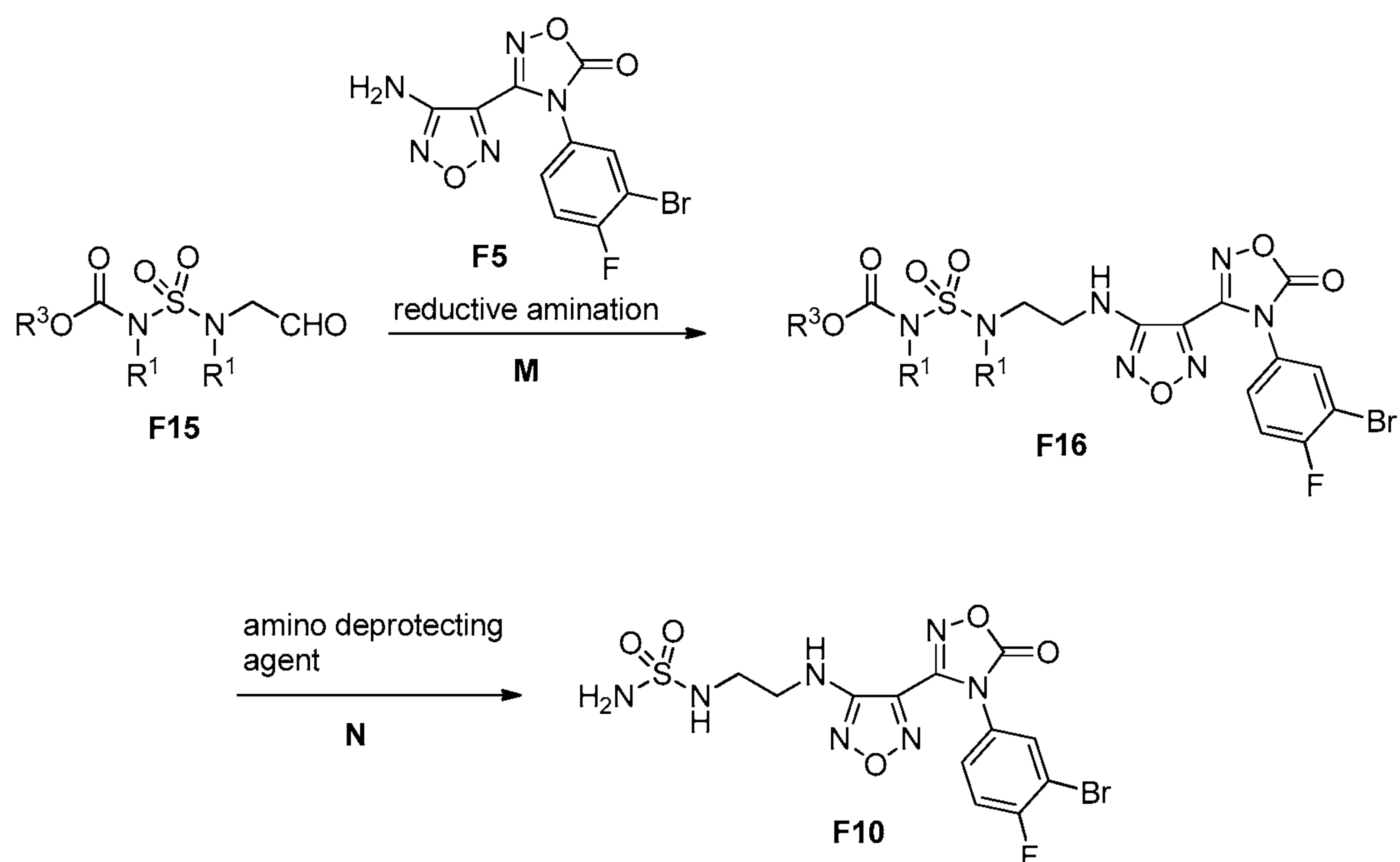
by addition of sodium bicarbonate, to provide an amidoxime of Formula F4. The amidoxime functionality of the compound of F4 can then be converted to an oxadiazolone or Formula F5 using N,N-carbonyldiimidazole (DCI) in a solvent (such as ethyl acetate, dioxane, THF and the like), at elevated temperatures such as about 50 °C, about 60 °C, about 70 °C, about 80 °C, about 90 °C, or about 100 °C.

Scheme 2

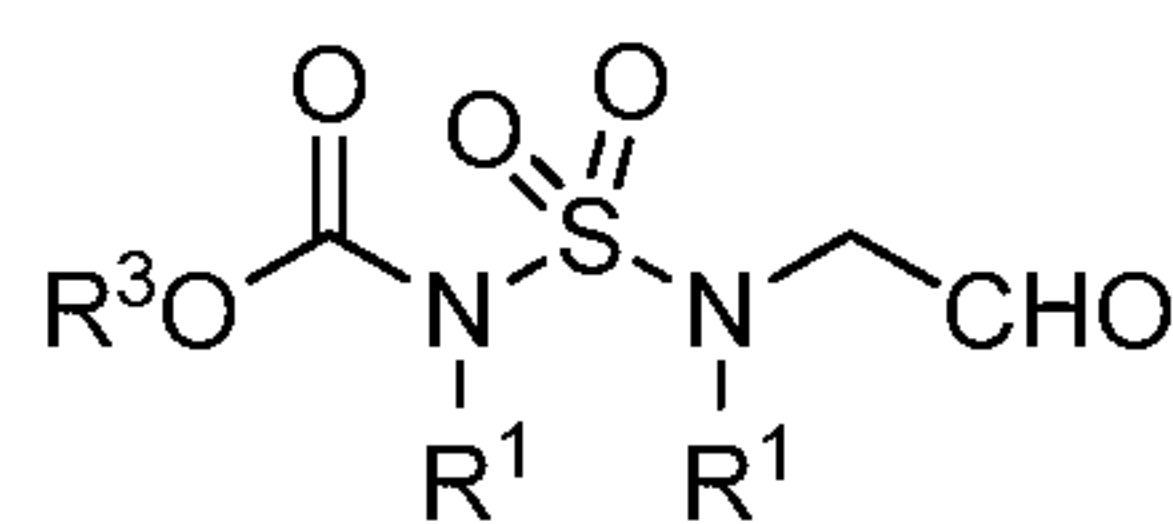


Alternately, the compound of Formula F10 can be obtained through a sequence of steps depicted in Scheme 3.

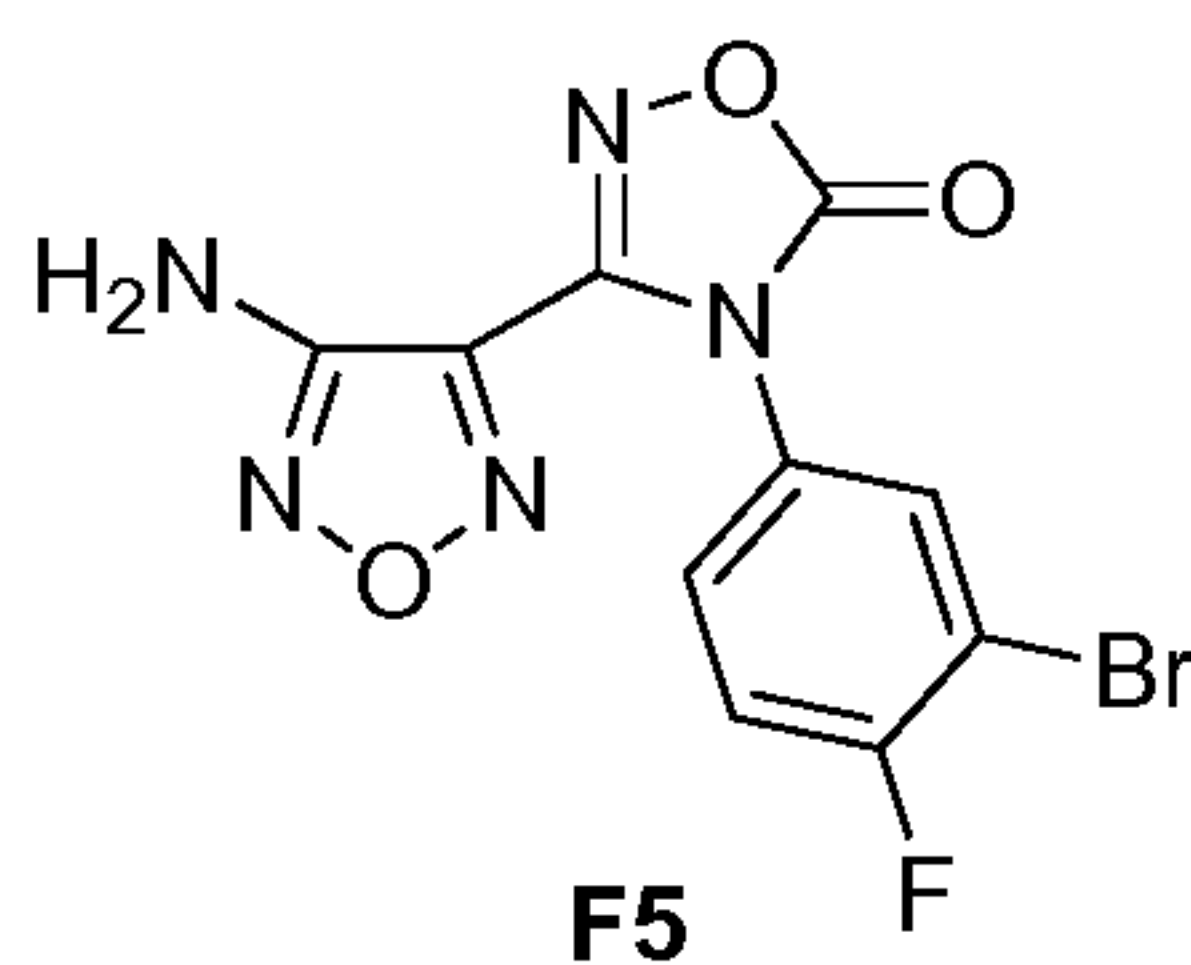
Scheme 3



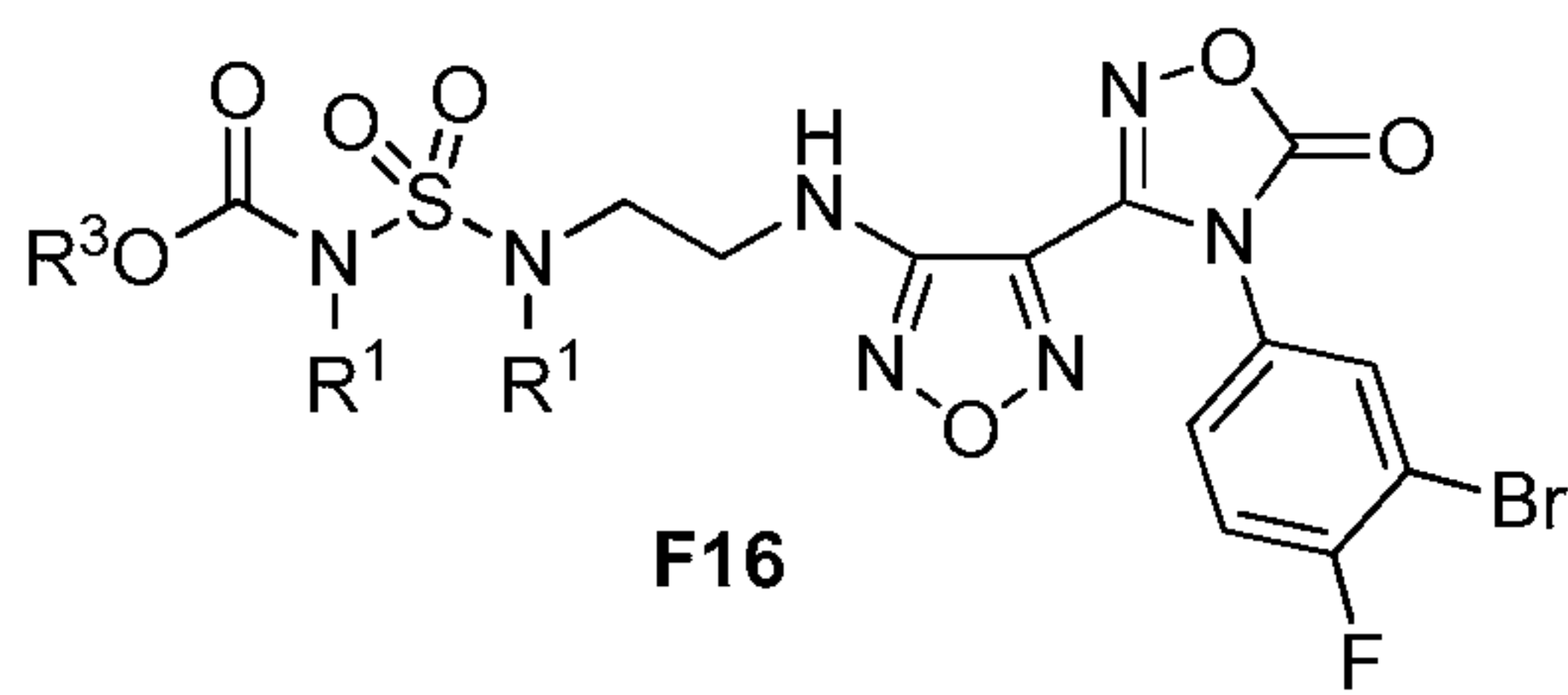
In some embodiments, the present application provides a process, comprising reacting a compound of Formula F15:

**F15**

with a compound of Formula F5:

**F5**

to afford a compound of Formula F16:

**F16**

wherein:

each R¹ is independently an amino protecting group; and

R³ is C₁-₆ alkyl or benzyl.

In some embodiments, R¹ is C₂-₄ alkenyl-C₁-₃ alkyl or phenyl-C₁-₃ alkyl, wherein said phenyl-C₁-₃ alkyl is optionally substituted by 1, 2, or 3 independently selected C₁-₄ alkoxy groups.

In some embodiments, R¹ is C₂-₄ alkenyl-C₁-₃ alkyl or phenyl-C₁-₃ alkyl, wherein said phenyl-C₁-₃ alkyl is optionally substituted by 1, 2, or 3 methoxy groups.

In some embodiments, R¹ is allyl.

In some embodiments, R¹ is 4-methoxybenzyl.

In some embodiments, R³ is C₁-₆ alkyl.

In some embodiments, R³ is *tert*-butyl.

In some embodiments, R³ is C₁-₄ alkyl.

In some embodiments, R³ is butyl.

Preferably, the reacting is performed in the presence of a reducing agent. The reducing agent can be any compound capable of reducing an organic compound to a lower

oxidation state. In some embodiments, the reducing agent can be hydrogen gas in the presence of a catalyst or a hydride reagent (such as $\text{NaB}(\text{OAc})_3\text{H}$, NaBH_4 , LiAlH_4 and the like); using triphenylphosphine; or using a combination of sodium iodide, chlorotrimethylsilane, and methanol. In some embodiments, this step can be performed in the presence of an acid such as trifluoroacetic acid. Suitable solvents for this step include isopropyl alcohol, THF, dioxane, or the like. In some embodiments, this step can be performed at a temperature from about -15°C to about 30°C , e.g., from about -15°C to about 0°C , from about -5°C to about 5°C , from about -5°C to about 0°C , from about 0 to 5°C , or from about 0°C to about 45°C .

In some embodiments, said reacting is performed in a solvent component comprising tetrahydrofuran.

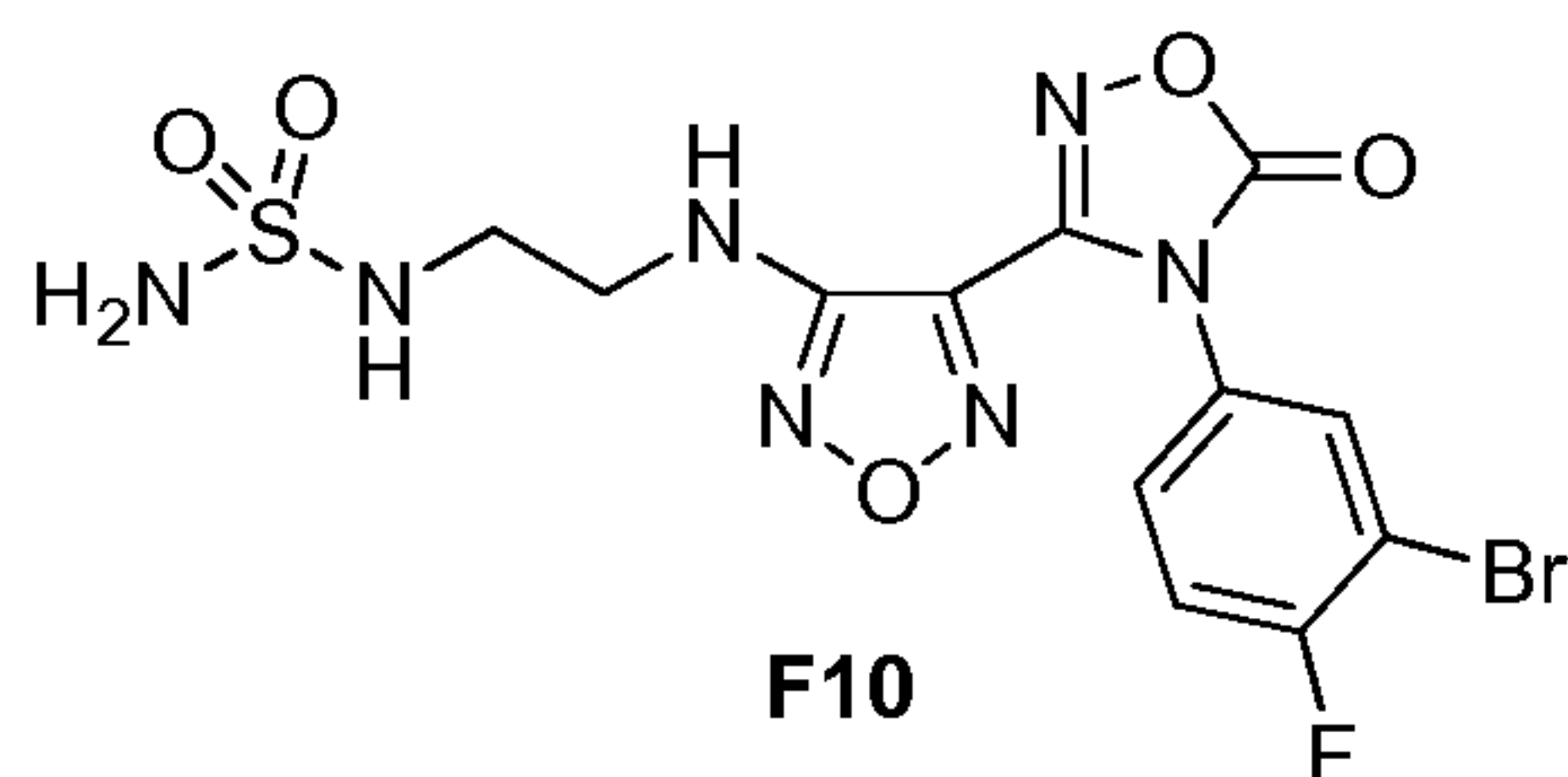
In some embodiments, said reacting is performed in the presence of a reducing agent.

In some embodiments, said reducing agent is a borohydride reducing agent.

In some embodiments, said borohydride reducing agent is sodium triacetoxymethylborohydride.

In some embodiments, said reacting is performed in the presence of trifluoroacetic acid.

In some embodiments, the invention further comprises deprotecting said compound of Formula F16 to afford a compound of Formula F10:



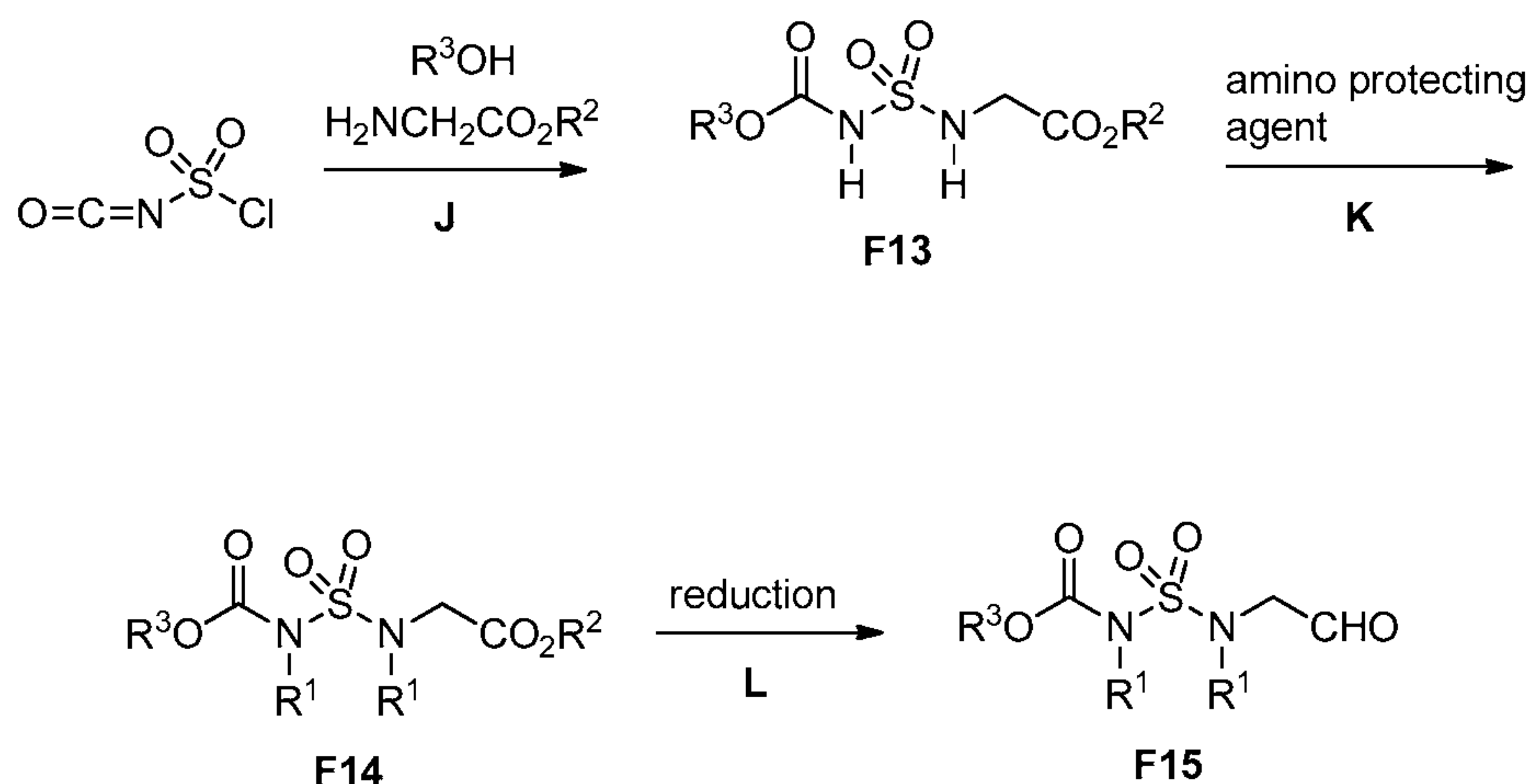
Treatment of a compound F16 to replace R^1N with NH_2 can be accomplished by methods for the deprotection of particular amine protecting groups known to one skilled in the art, such as those in Wuts and Greene, *Greene's Protective Groups in Organic Synthesis*, 4th Ed., pp 696-926, John Wiley & Sons: New York, 2006. In some embodiments, when R^1 is allyl, the deprotecting agent can be a palladium catalyst (e.g., $\text{Pd}(\text{Ph}_3\text{P})_4$, Pd/C , or $\text{Pd}(\text{dba})\text{DPPB}$). In some embodiments, when the R^1 is 4-methoxybenzyl, the deprotecting agent can include an organic acid (such as trifluoroacetic acid or methanesulfonic acid, and the like); an inorganic acid (such as hydrochloric acid); hydrogen and palladium; or sodium in liquid ammonia. The deprotecting can be performed at a temperature from about 30°C to about 90°C , e.g., from about 50°C to about 100°C , or from about 60°C to about 80°C .

In some embodiments, said deprotecting comprises reacting a compound of Formula F16 with trifluoroacetic acid.

In some embodiments, said deprotecting comprises reacting a compound of Formula F16 with is hydrochloric acid.

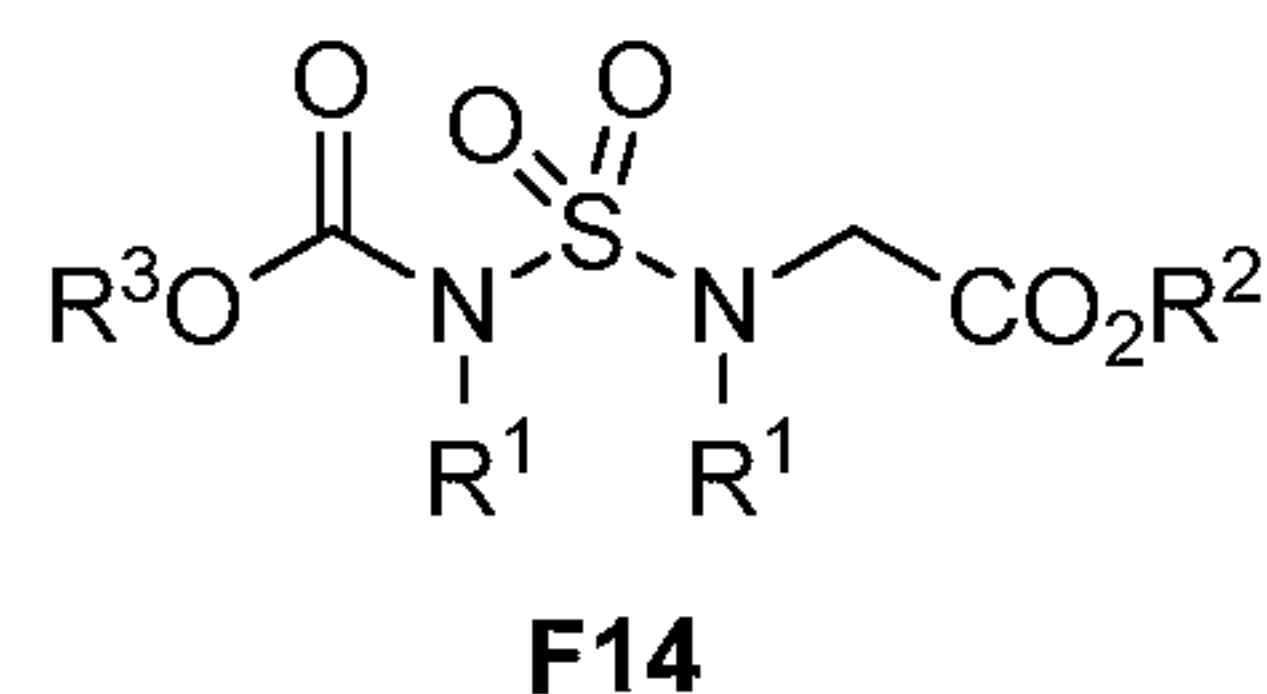
- 5 Compound F15 can be made by a three step process (Steps J, K and L) from chlorosulfonylisocyanate, as shown in Scheme 4.

Scheme 4



10

Accordingly, the present application further provides a process wherein said compound of Formula F15 is obtained by a process comprising treating a compound of Formula F14:



- 15 with a reducing agent to afford said compound of Formula F15; wherein R^2 is C_{1-4} alkyl; and R^3 is defined *supra*.

In some embodiments, R^2 is methyl.

In some embodiments, R^2 is ethyl.

- 20 In some embodiments, the reducing can be carried out with diisobutylaluminum hydride (DIBAL-H). Suitable solvents include halogenated hydrocarbon solvents such as dichloromethane, chloroform, dichloroethane, tetrachloroethane, and the like. In some embodiments, the reduction can be performed at about room temperature e.g., from about -80

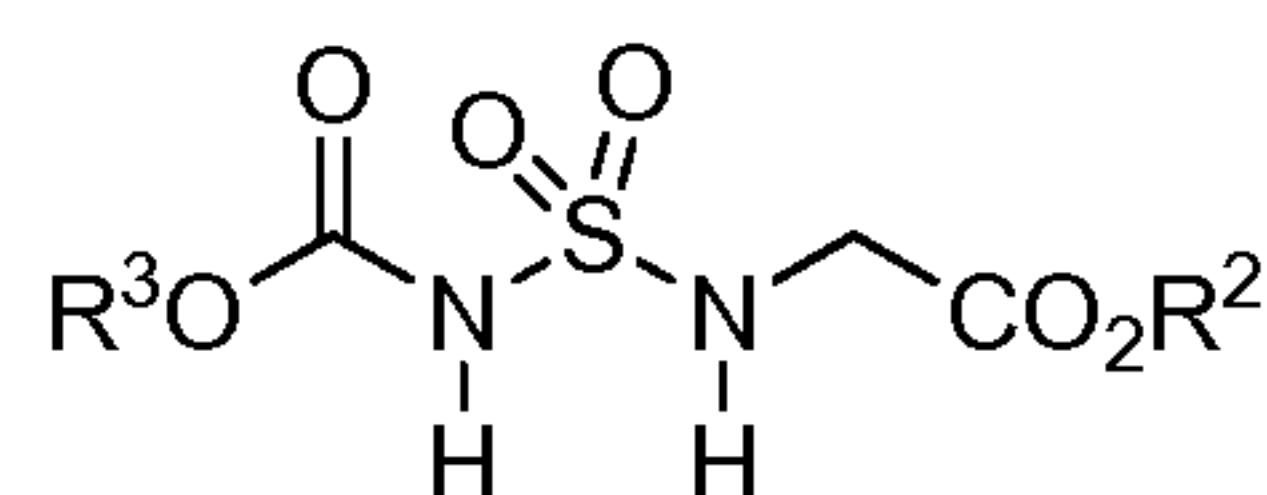
°C to about 30 °C, from about -78 °C to about 0 °C, from about 0 °C to about 30 °C, or from about 25 °C to about 30 °C.

In some embodiments, said treating is performed in a halogenated hydrocarbon solvent.

5 In some embodiments, said halogenated hydrocarbon solvent is dichloromethane.

In some embodiments, said reducing agent is diisobutylaluminum hydride.

In some embodiments, said compound of Formula F14 is obtained by a process comprising protecting a compound of Formula F13:



F13

10 with one or more independently selected amino protecting agents to afford a compound of Formula F14.

Protecting group R¹ on F14 can be selected from the various amino protecting groups known in the art (*supra*). In some embodiments, the amino protecting agent is allyl bromide or 4-methoxybenzyl chloride.

15 In some embodiments, said one or more amino protecting agents is selected from allyl bromide and 4-methoxybenzyl chloride.

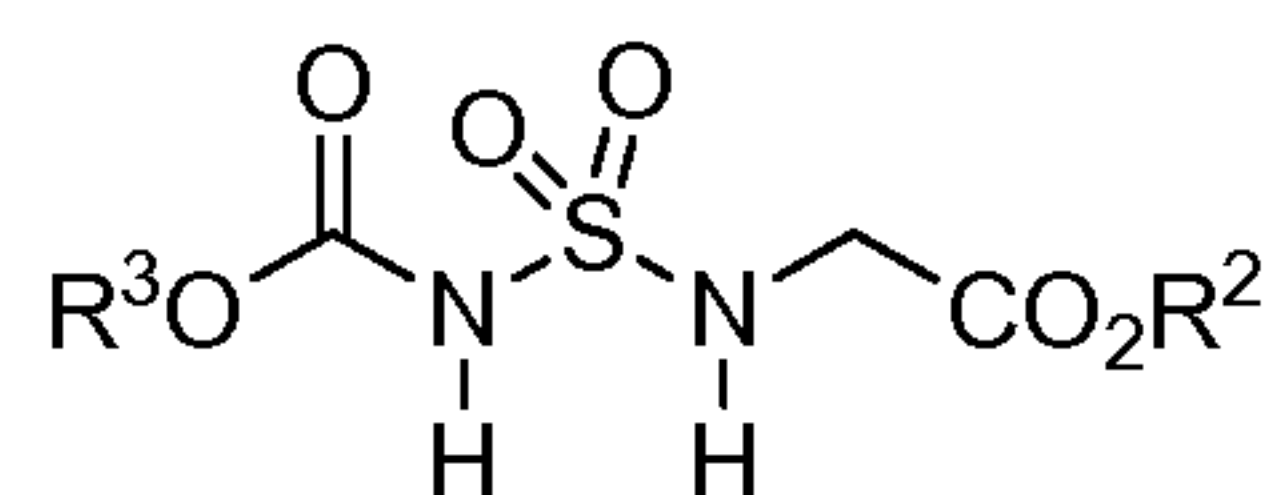
In some embodiments, said protecting is performed in the presence of a base.

In some embodiments, said base is potassium carbonate.

20 In some embodiments, said protecting is performed in a solvent component comprising acetonitrile.

In some embodiments, the preparation of the compound of F13 can be obtained by treating chlorosulfonylisocyanate with an alcohol R³OH (where R³ is defined above) and a glycine ester H₂NCH₂CO₂R², wherein R² is C₁₋₄ alkyl. In some embodiments, this Step J is carried out in the presence of an organic acid (such as acetic acid, benzoic acid, trifluoroacetic acid). Suitable solvents for this step include dichloromethane, chloroform, dichloroethane, tetrachloroethane, and the like.

25 In some embodiments, the present application provides a compound of Formula F13:

**F13**

wherein:

R² is C₁₋₄ alkyl; and

R³ is C₁₋₆ alkyl or benzyl.

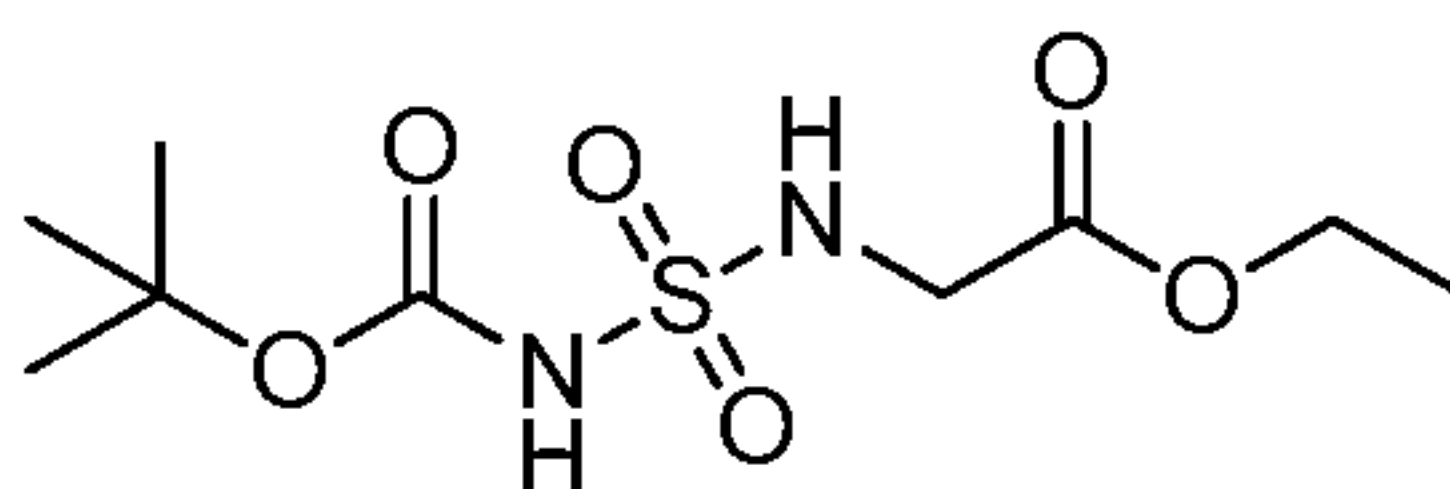
5 In some embodiments, R² is methyl.

In some embodiments, R² is ethyl.

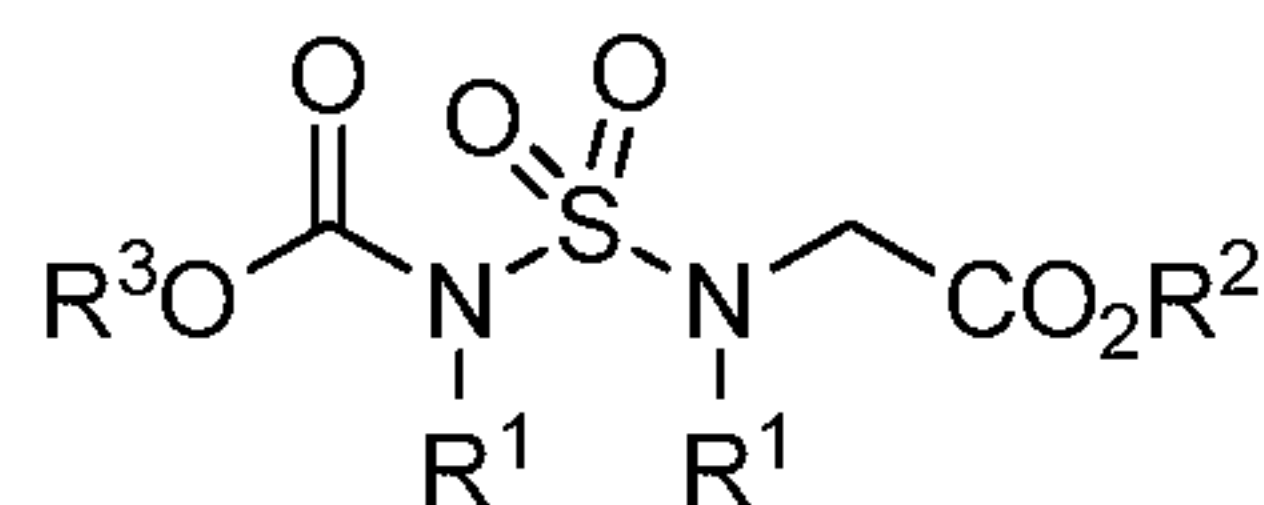
In some embodiments, R³ is C₁₋₆ alkyl.

In some embodiments, R³ is *tert*-butyl.

10 In some embodiments, the compound of Formula F13 is ethyl-2-((N-(*tert*-butoxycarbonyl)sulfamoyl)amino)acetate:



In some embodiments, the invention further provides a compound of Formula F14:

**F14**

wherein:

15 each R¹ is independently an amino protecting group;

R² is C₁₋₄ alkyl; and

R³ is C₁₋₆ alkyl or benzyl.

20 In some embodiments, R¹ is C₂₋₄ alkenyl-C₁₋₃ alkyl or phenyl-C₁₋₃ alkyl, wherein said phenyl-C₁₋₃ alkyl is optionally substituted by 1, 2, or 3 independently selected C₁₋₄ alkoxy groups.

In some embodiments, R¹ is allyl.

In some embodiments, R¹ is 4-methoxybenzyl.

In some embodiments, R² is methyl.

In some embodiments, R² is ethyl.

25 In some embodiments, R³ is C₁₋₆ alkyl.

In some embodiments, R³ is *tert*-butyl.

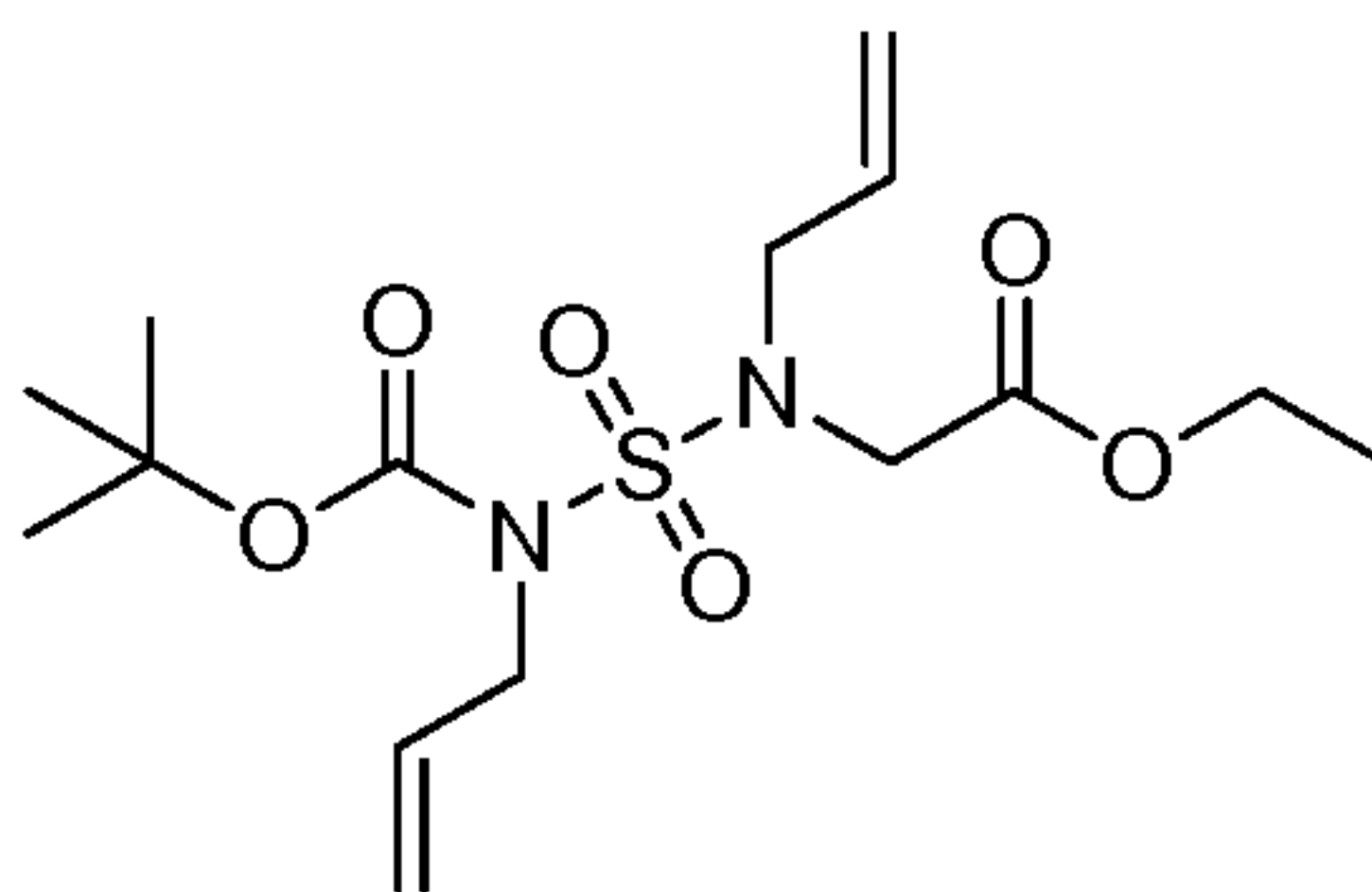
In some embodiments, R^3 is C_{1-4} alkyl.

In some embodiments, R^3 is butyl.

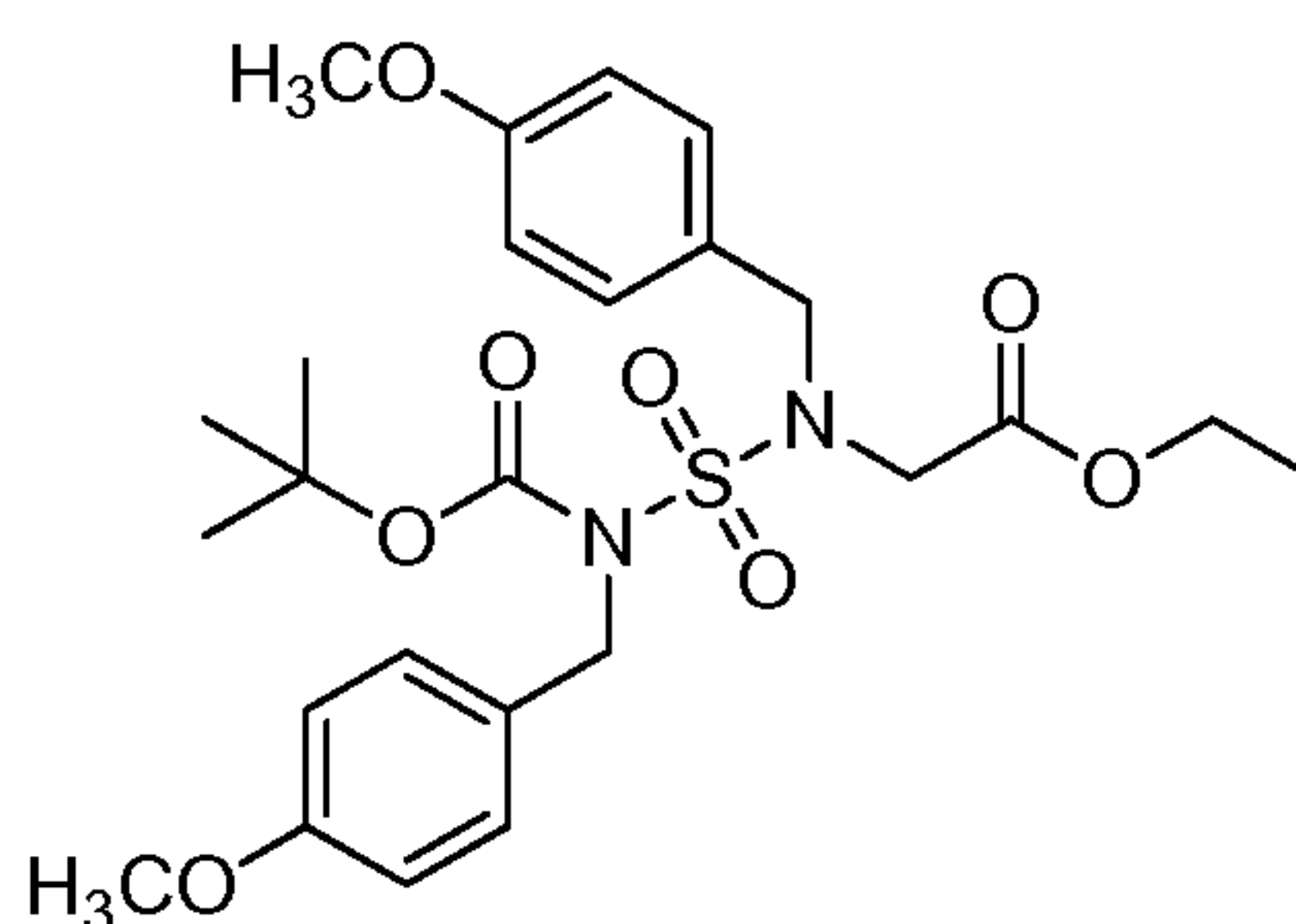
In some embodiments, R^3 is C_{1-4} alkyl.

In some embodiments, R^3 is butyl.

5 In some embodiments, the compound of Formula F14 is ethyl-2-(allyl(N-allyl-N-(*tert*-butoxycarbonyl)sulfamoyl)amino) acetate:

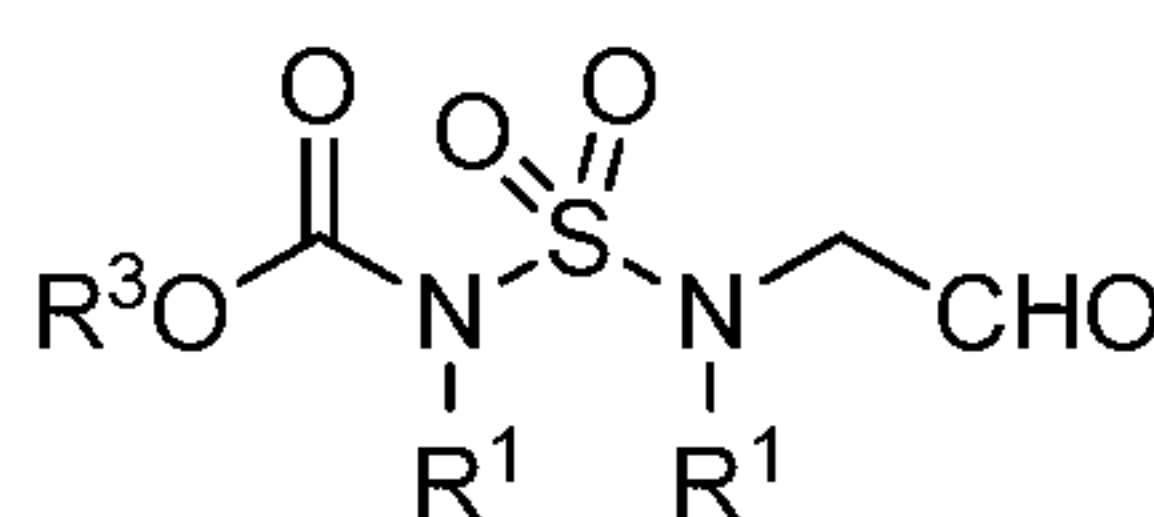


In some embodiments, the compound of Formula F14 is ethyl-2-(4-methoxybenzyl(N-4-methoxybenzyl-N-(*tert*-butoxycarbonyl) sulfamoyl)amino)acetate:



10

In some embodiments, the present application provides a compound of Formula F15:



F15

wherein:

R^3 is C_{1-6} alkyl or benzyl; and

15 each R^1 is independently an amino protecting group.

In some embodiments, R^1 is C_{2-4} alkenyl- C_{1-3} alkyl or phenyl- C_{1-3} alkyl, wherein said phenyl- C_{1-3} alkyl is optionally substituted by 1, 2, or 3 independently selected C_{1-4} alkoxy groups.

In some embodiments, R^1 is allyl.

20 In some embodiments, R^1 is 4-methoxybenzyl.

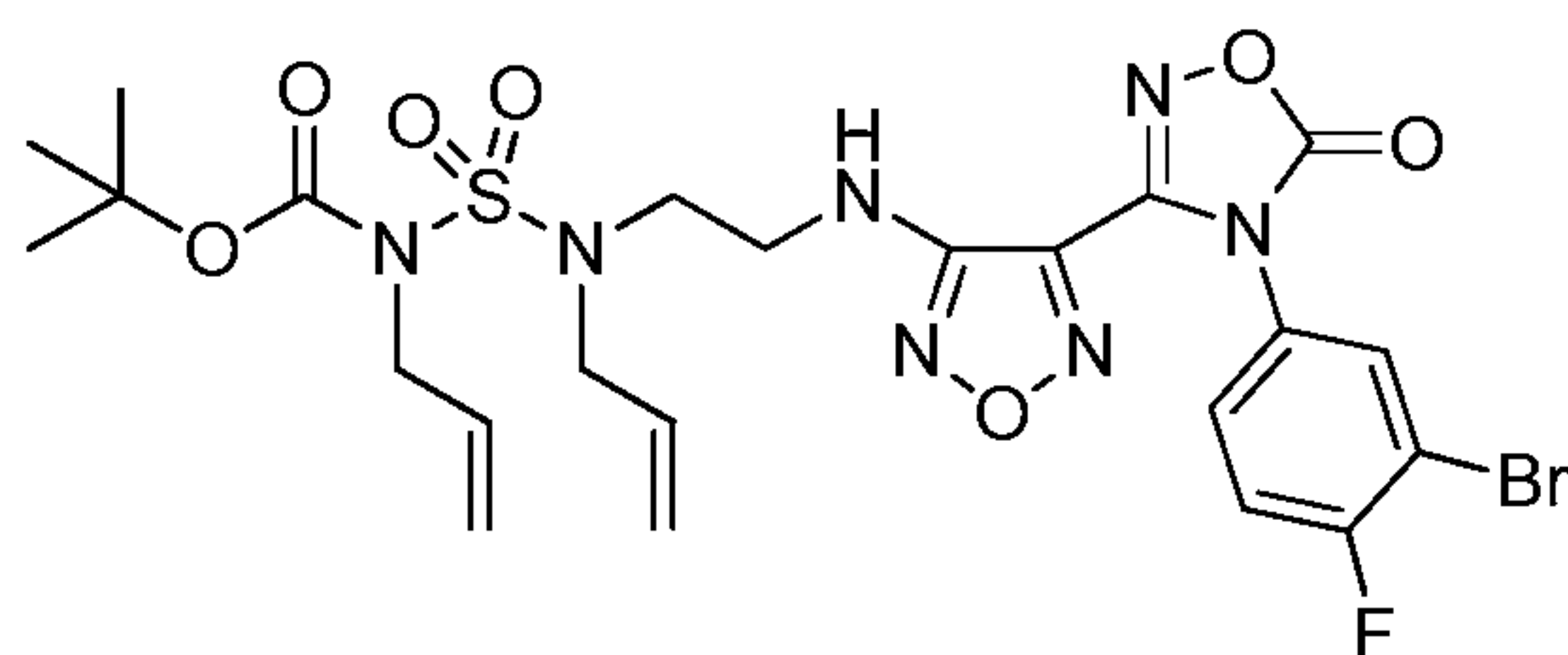
In some embodiments, R^3 is C_{1-6} alkyl.

In some embodiments, R^3 is *tert*-butyl.

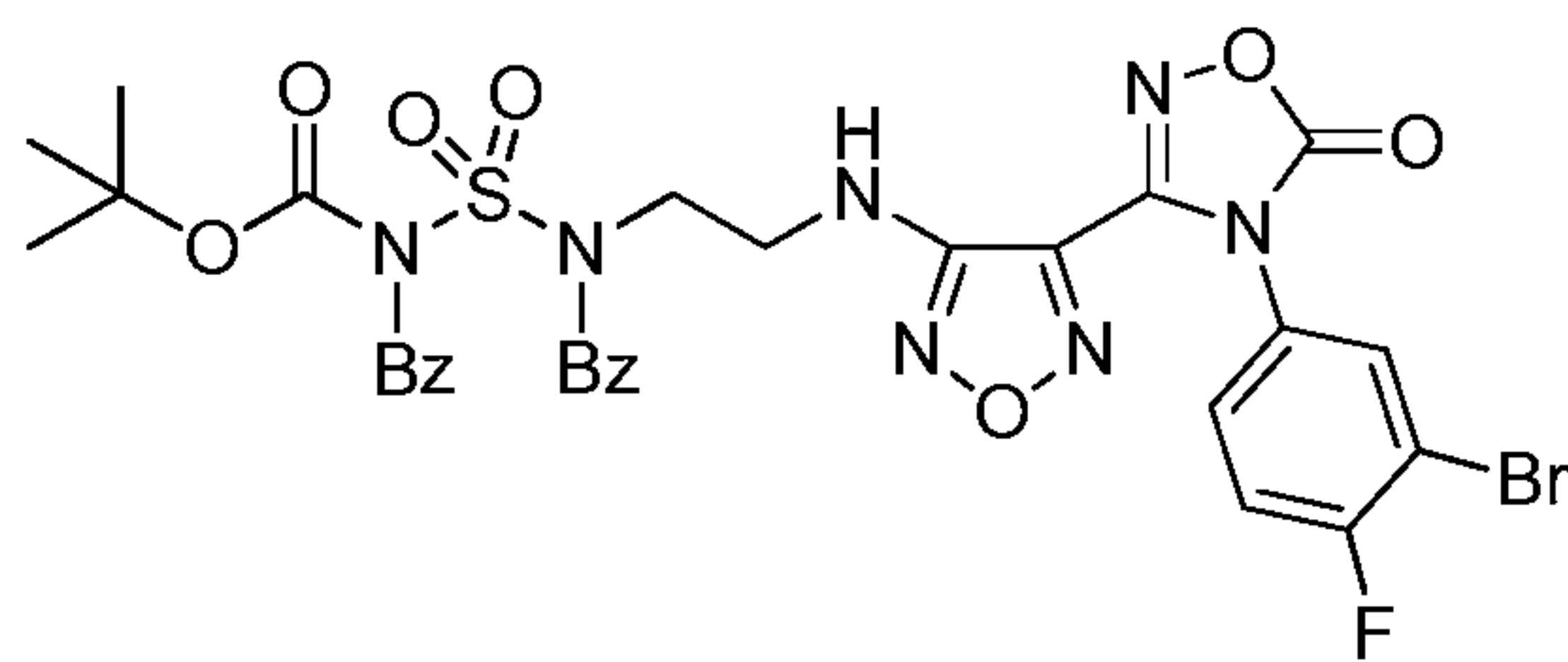
CC(C)(C)OC(=O)N(S(=O)(=O)NCC=O)CC=CCOc1ccc(cc1)CN(C(=O)OC(C)(C)C)S(=O)(=O)NCC=OBrC1=CC=C(C=C1)N2C(=O)N(C2)c3nc(NCCN(R1)S(=O)(=O)N(R1)C(=O)OR3)nc4ocn34

F16

21



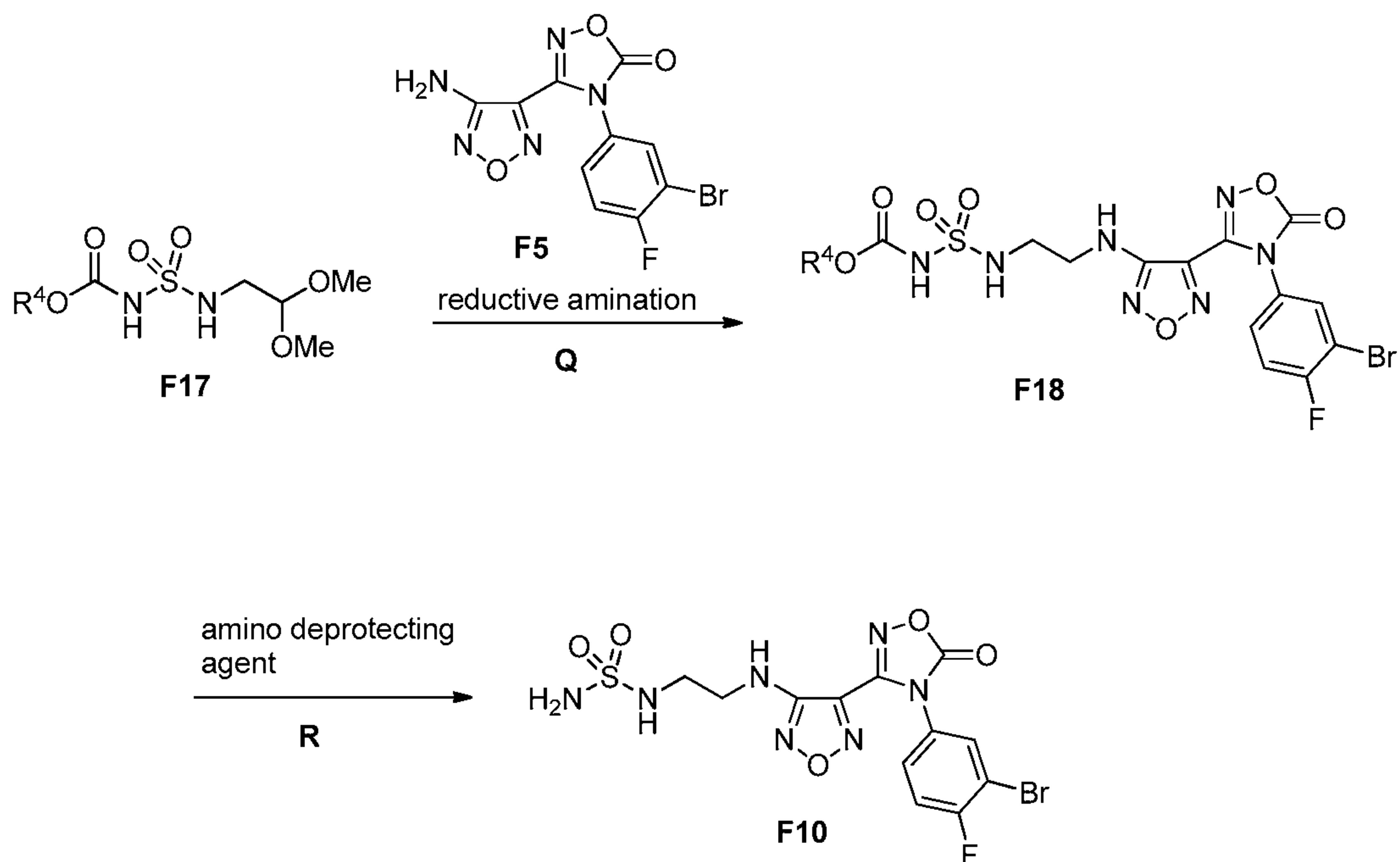
In some embodiments, the compound of Formula F16 is *tert*-butyl (4-methoxybenzyl)-(N-(4-methoxybenzyl)-N-(2-(4-(4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl)-1,2,5-oxadiazol-3-ylamino)ethyl)sulfamoyl)carbamate:



5

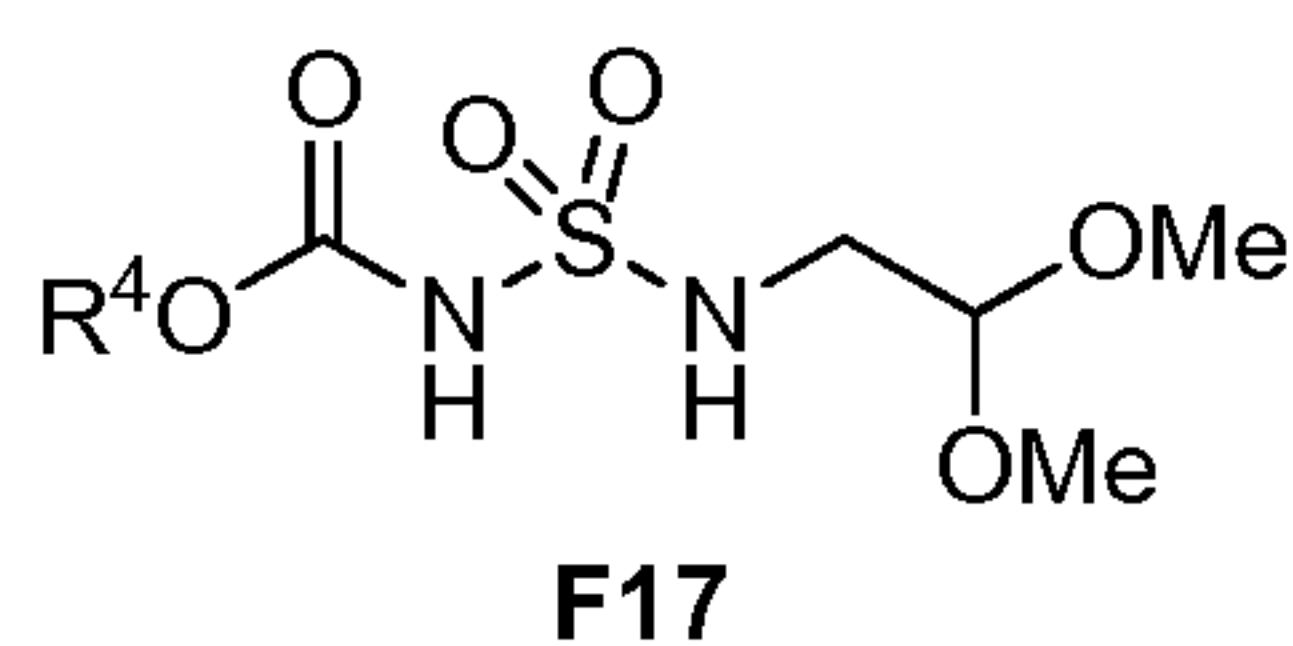
Scheme 5 delineates an alternative route for the preparation of the compound of Formula F10.

Scheme 5

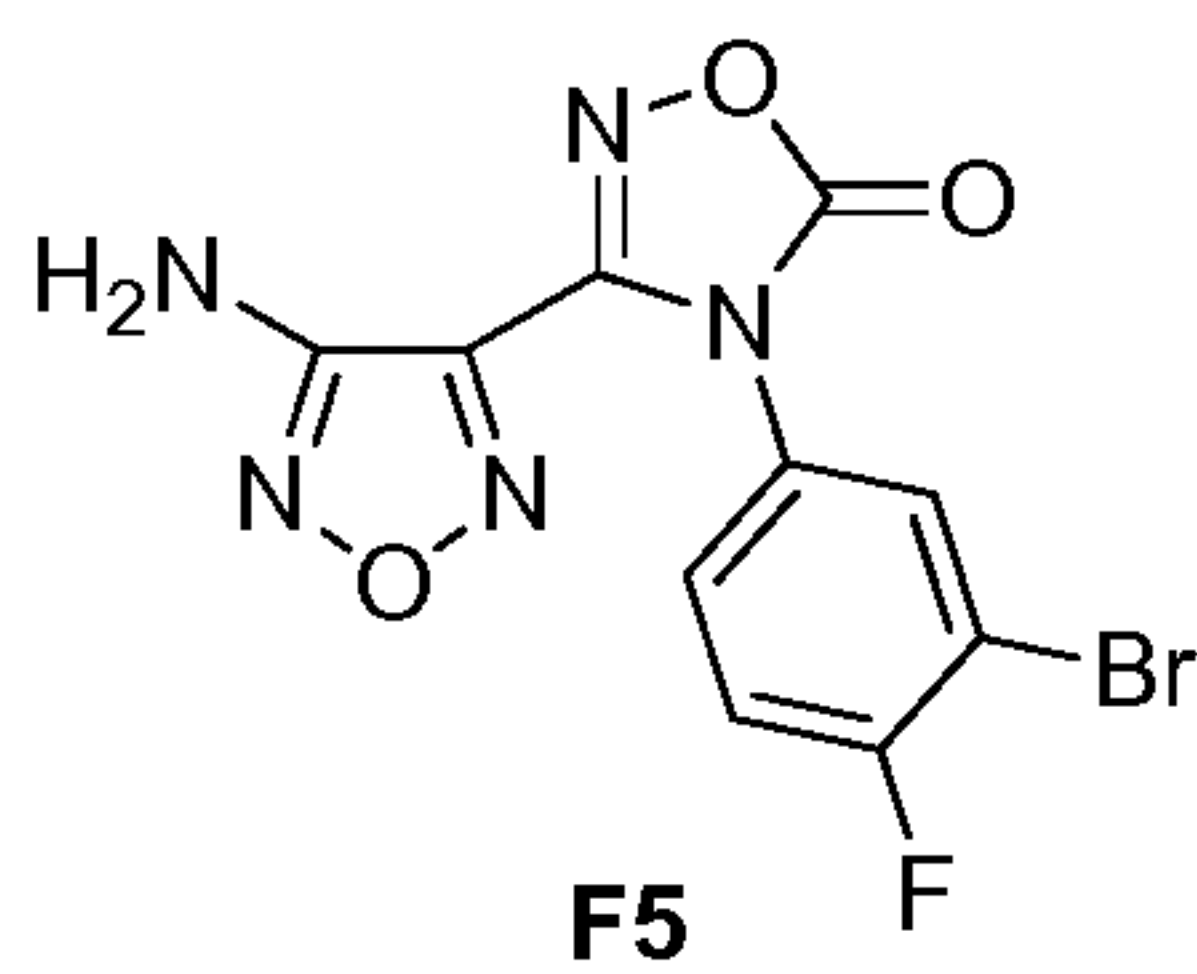


10

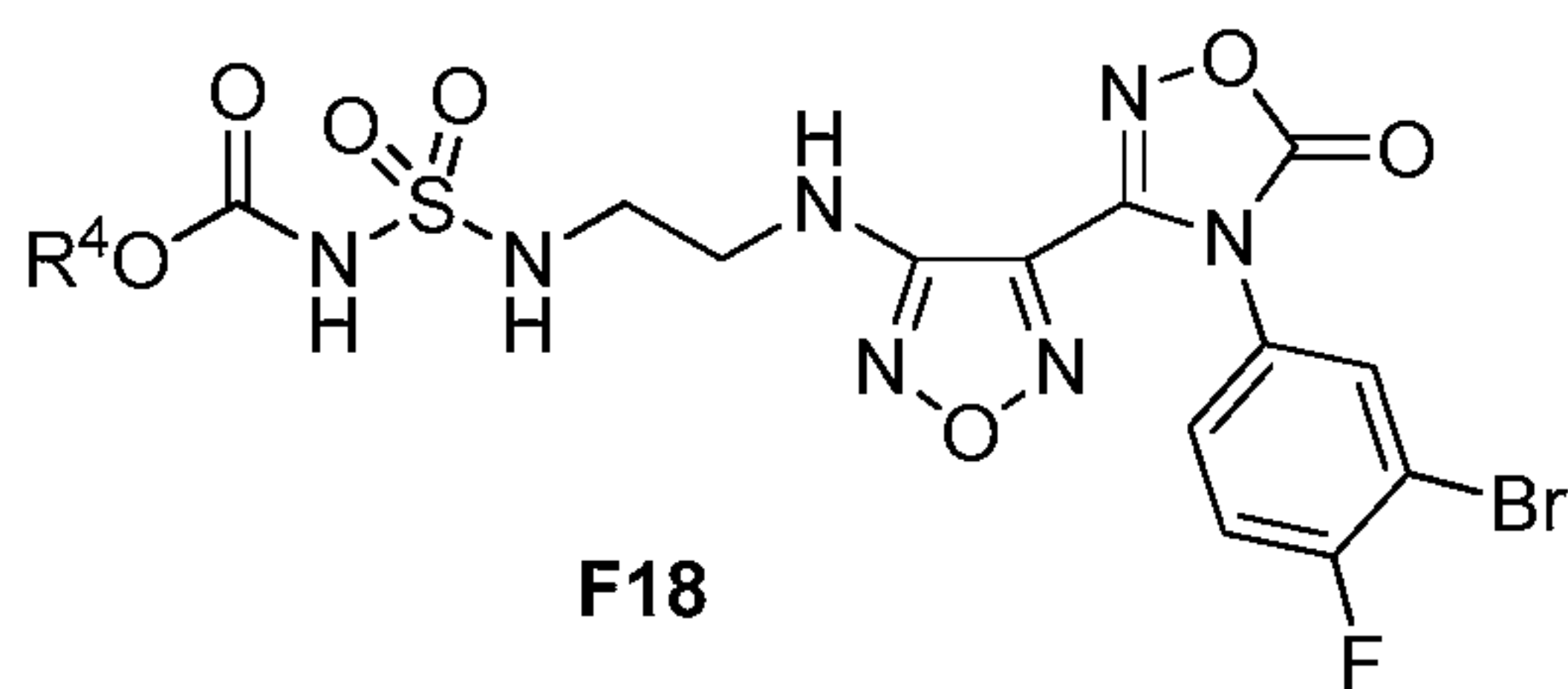
The present application also provides a process comprising reacting a compound of Formula F17:



wherein R⁴ is C₁₋₆ alkyl, C₁₋₆ haloalkyl, benzyl, or 9*H*-fluoren-9-ylmethyl with a compound of Formula F5:



5 to afford a compound of Formula F18:



In some embodiments, R⁴ is *tert*-butyl.

In some embodiments, R⁴ is benzyl.

In some embodiments, R⁴ is ethyl.

10 In some embodiments, R⁴ is C₁₋₃ haloalkyl.

In some embodiments, R⁴ is 2,2,2-trichloroethyl.

In some embodiments, R⁴ is 9*H*-fluoren-9-ylmethyl.

In this Step Q, compounds F18 can be prepared, in some embodiments, by reacting F17 with the amine compound of Formula F5 in the presence of a reducing agent.

15 In some embodiments, said reacting is carried out in the presence of a reducing agent.

The reducing agent can be any compound capable of reducing an organic compound to a lower oxidization state, for example by use of an organosilane such as tri(C₁₋₃ alkyl)silane (e.g., triethylsilane); elemental hydrogen or using a hydride reagent (such as NaB(OAc)₃H, NaBH₄, LiAlH₄ and the like); using triphenylphosphine; or using a
20 combination of sodium iodide, chlorotrimethylsilane, and methanol. In some embodiments, this step can be performed in the presence of an acid such as trifluoroacetic acid. Suitable solvents include, but are not limited halogenated hydrocarbon solvents (e.g., dichloromethane, chloroform, dichloroethane or tetrachloroethane). In some embodiments, the halogenated hydrocarbon solvent is 1,2-dichloroethane.

In some embodiments, said reducing agent is an organosilane.

In some embodiments, said reducing agent is tri(C₁₋₃ alkyl)silane.

In some embodiments, said reducing agent is triethylsilane.

In some embodiments, said reacting is carried out in the presence of an organic acid.

5 In some embodiments, said organic acid is trifluoroacetic acid.

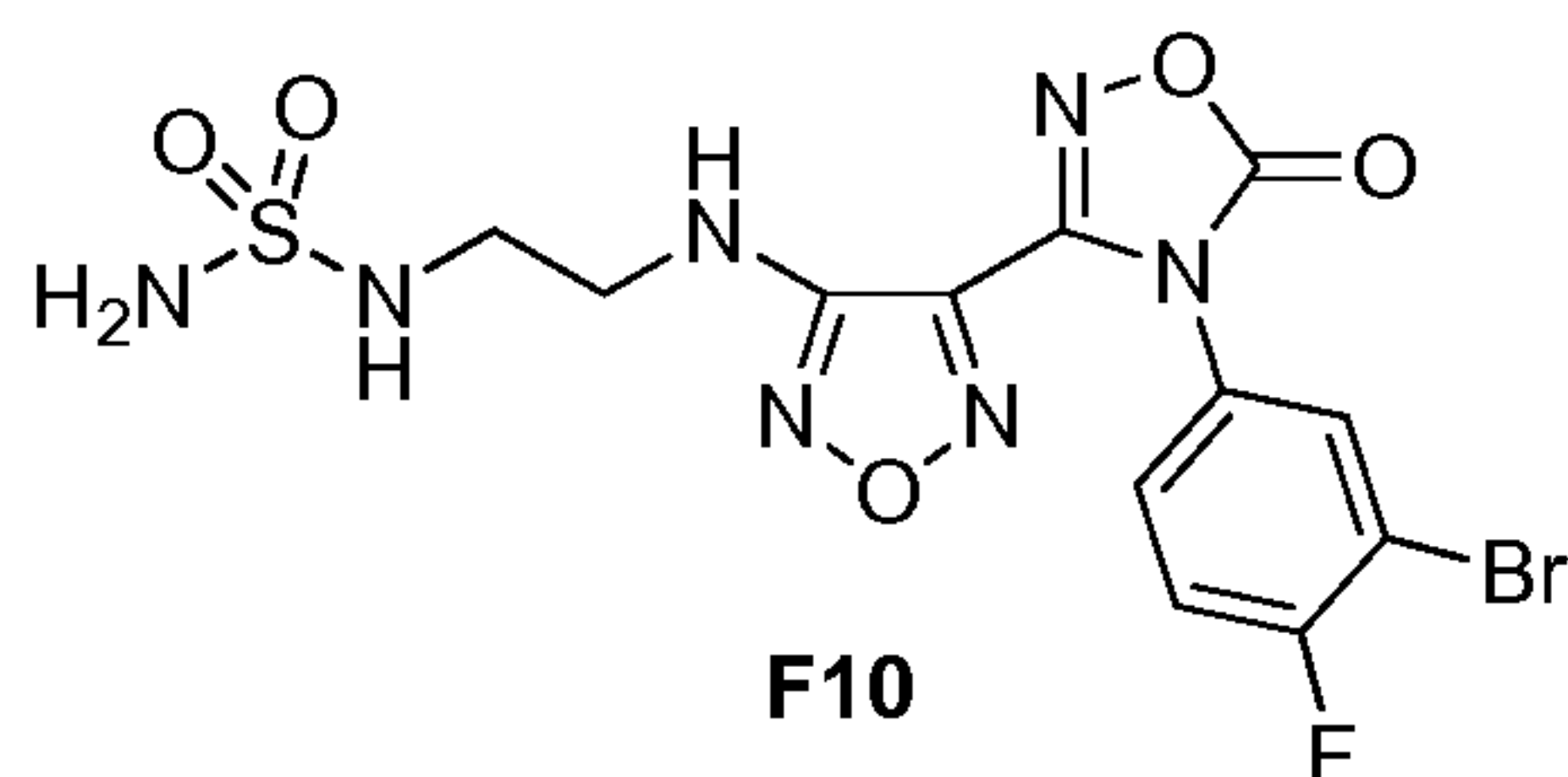
In some embodiments, said organic acid is methanesulfonic acid.

In some embodiments, said reacting is performed in a solvent component comprising a halogenated hydrocarbon solvent.

In some embodiments, said halogenated hydrocarbon solvent is dichloromethane.

10 In some embodiments, said halogenated hydrocarbon solvent is 1,2-dichloroethane.

In some embodiments, the process further comprises deprotecting said compound of Formula F18 to afford a compound of Formula F10:



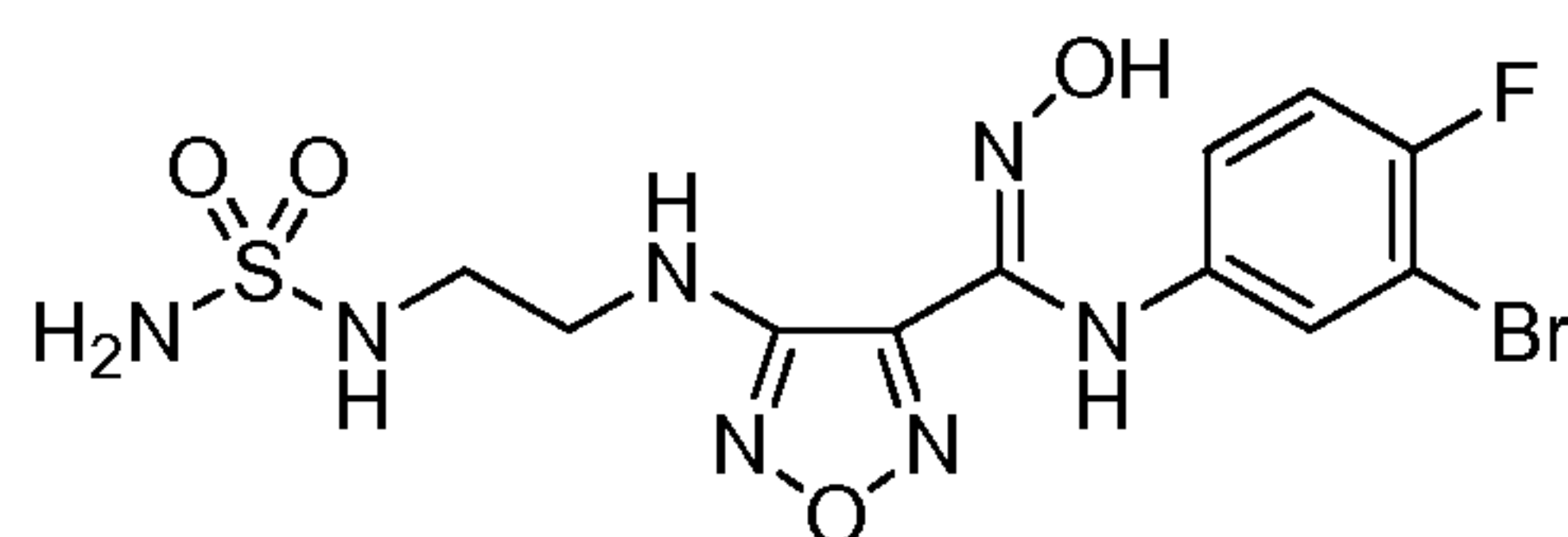
In some embodiments, methods for the deprotection of particular amine protecting groups (such as carbamates) are known to one skilled in the art, such as those in Wuts and Greene, *Greene's Protective Groups in Organic Synthesis*, 4th Ed., pp 696-926, John Wiley & Sons: New York, 2006. For example, the *tert*-butoxycarbonyl group (e.g., when R⁴ is *tert*-butyl) can be removed (e.g., hydrolyzed) from the nitrogen atom, for example, by treatment with an acid (such as hydrochloric acid, trifluoroacetic acid, toluenesulfonic acid, and the like); a combination of reagents (e.g., mixture of acetyl chloride and methanol) known to generate an acid; or a Lewis acid (e.g., BF₃·Et₂O). The benzyloxycarbonyl group (e.g., when R⁴ is benzyl) can be removed (e.g., hydrogenolyzed) from the nitrogen atom, for example, by treatment with hydrogen and a catalyst (such as palladium on carbon). The methoxycarbonyl and ethoxycarbonyl groups (i.e., when R⁴ is methyl or ethyl) can be removed by treatment with an inorganic base (such as KOH or K₂CO₃); a combination of reagents (e.g., mixture of acetyl chloride, sodium iodide and acetonitrile); or by treatment with an acid (e.g., HBr, AcOH). The 2,2,2-trichloroethoxycarbonyl group can be removed, for example by treatment with a catalyst (e.g., Zn/AcOH or Cd/AcOH). Suitable solvents for this step include, but are not limited to, methanol or tetrahydrofuran (THF), acetonitrile and the like. In some

embodiments, the treating is performed at a temperature from about 30 °C to about 90 °C, e.g., from about 50 °C to about 100 °C, or from about 60 °C to about 80 °C.

In some embodiments, said deprotecting comprises reacting the compound of Formula F18 with zinc in the presence of acetic acid.

5 In some embodiments, said deprotecting is performed in a solvent component comprising tetrahydrofuran.

In some embodiments, the process further comprises reacting said compound of Formula F10 with a base to afford a compound of Formula I:



10

I.

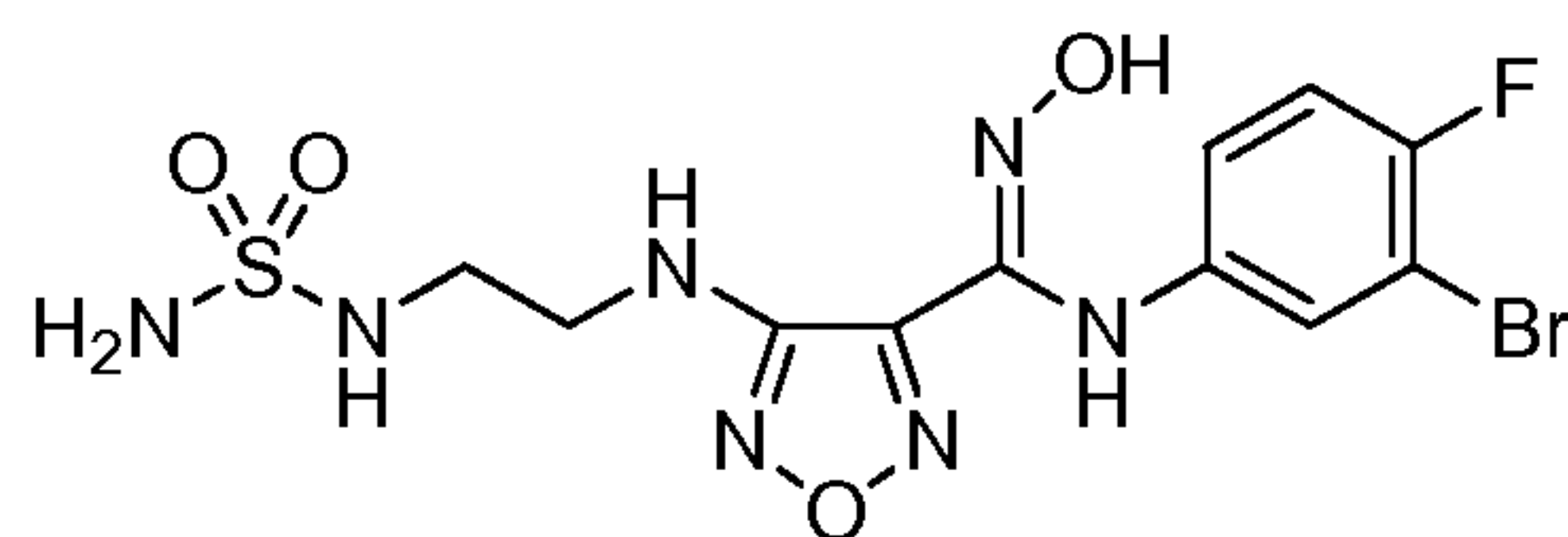
In some embodiments, said base comprises an alkali metal hydroxide.

In some embodiments, said alkali metal hydroxide is sodium hydroxide.

In some embodiments, said reacting is performed in a solvent component comprising tetrahydrofuran, water and ethanol.

15

In some embodiments, the process further comprises converting said compound of Formula F18 to a compound of Formula I:



I,

20 wherein said converting comprises combining the compound of Formula F18 with a base to form a first mixture. In some embodiments, the base is N,N-bis(2-aminoethyl)ethane-1,2-diamine.

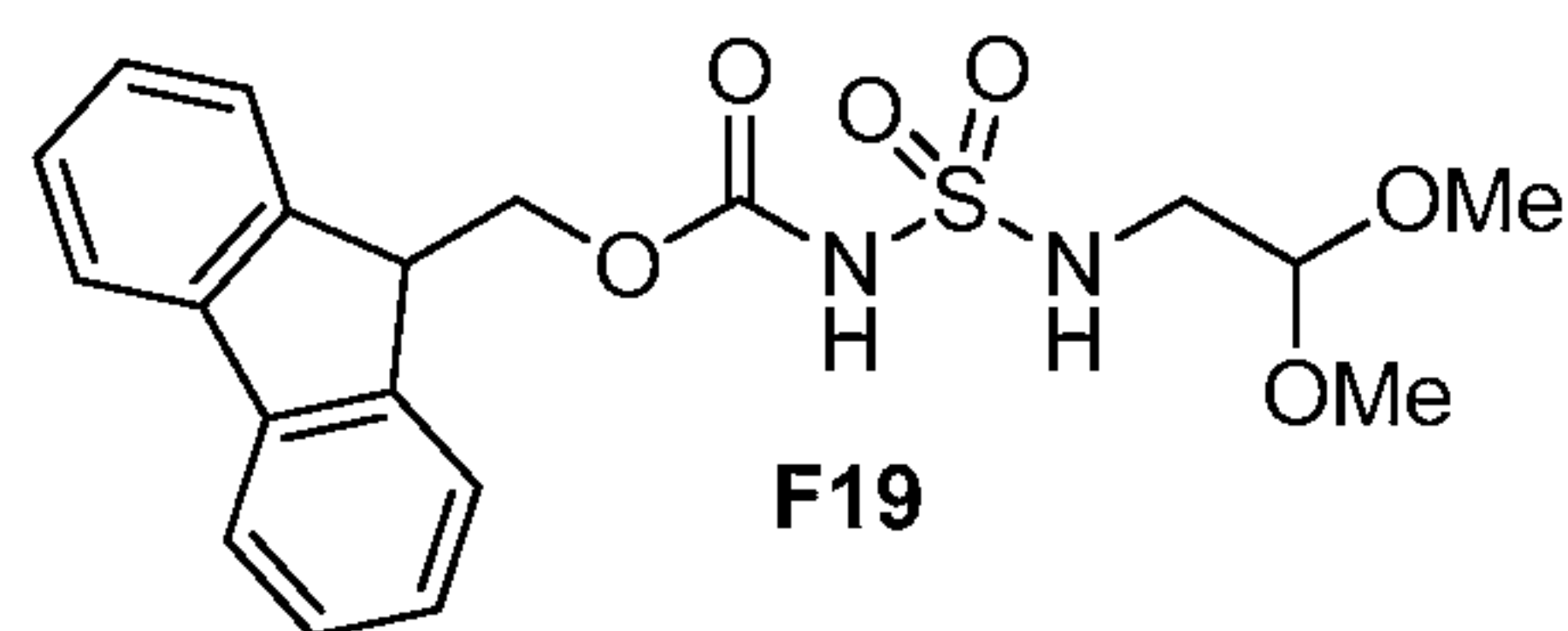
In some embodiments, the converting further comprises adding an acid to the first mixture. In some embodiments, said acid is an aqueous strong acid. In some embodiments, said aqueous strong acid is aqueous hydrochloric acid.

25

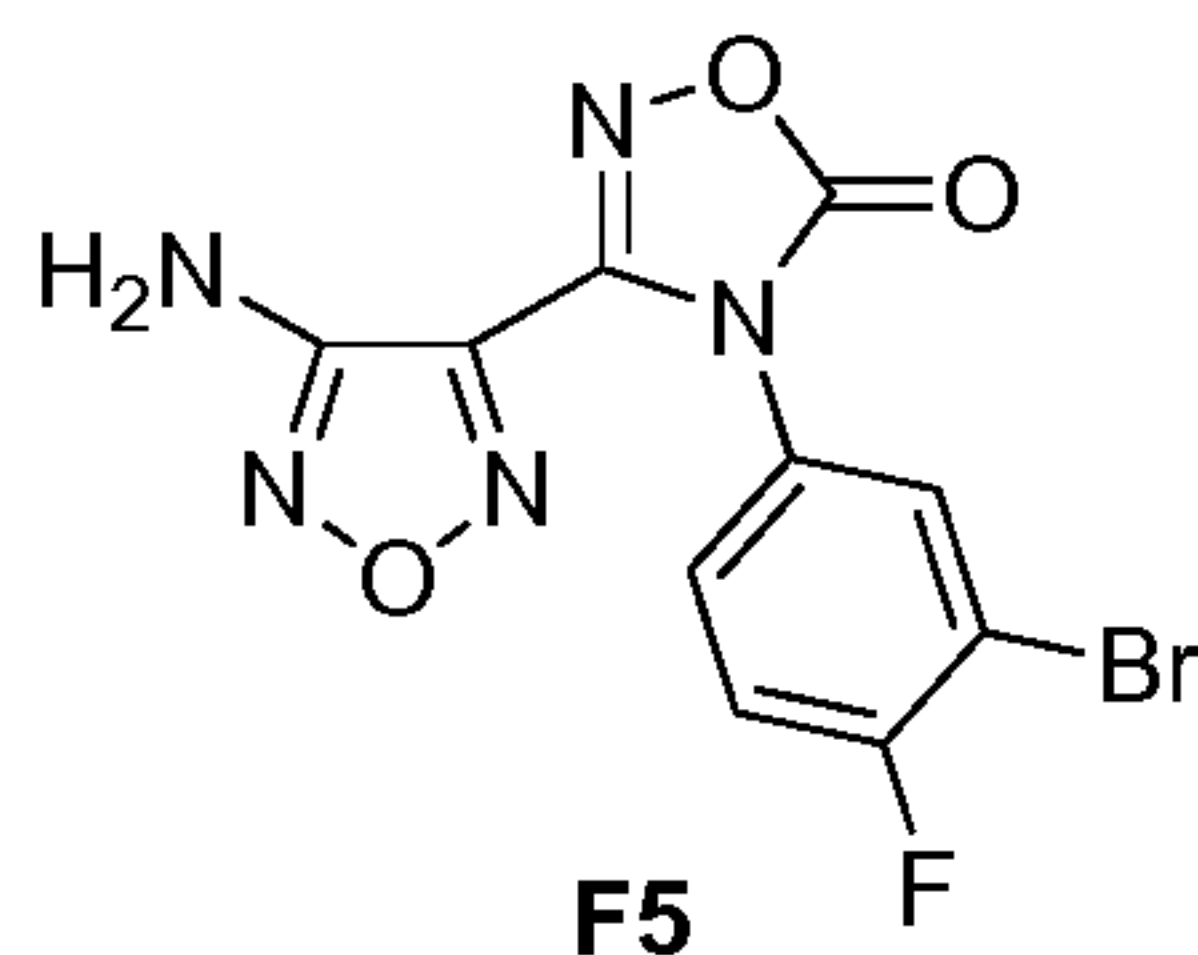
In some embodiments, said converting is performed in a solvent component comprising tetrahydrofuran and ethyl acetate.

The present application also provides a process comprising:

i) reacting a compound of Formula F19:

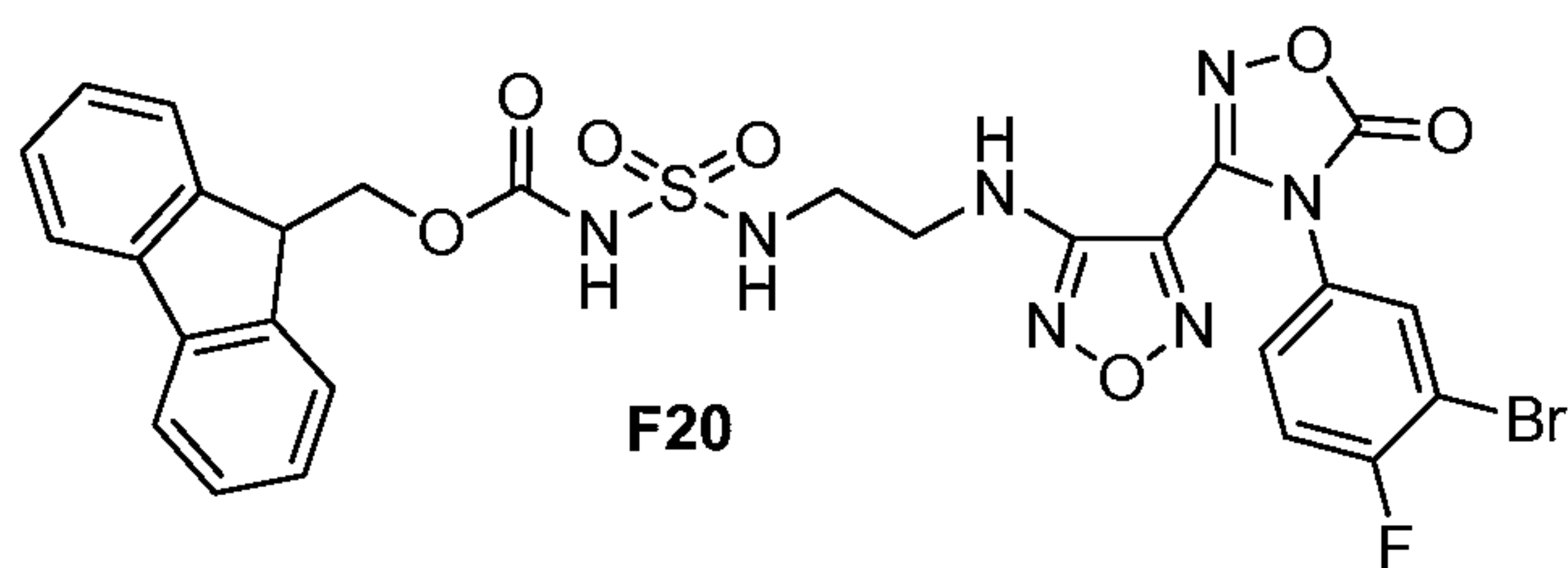


with a compound of Formula F5:



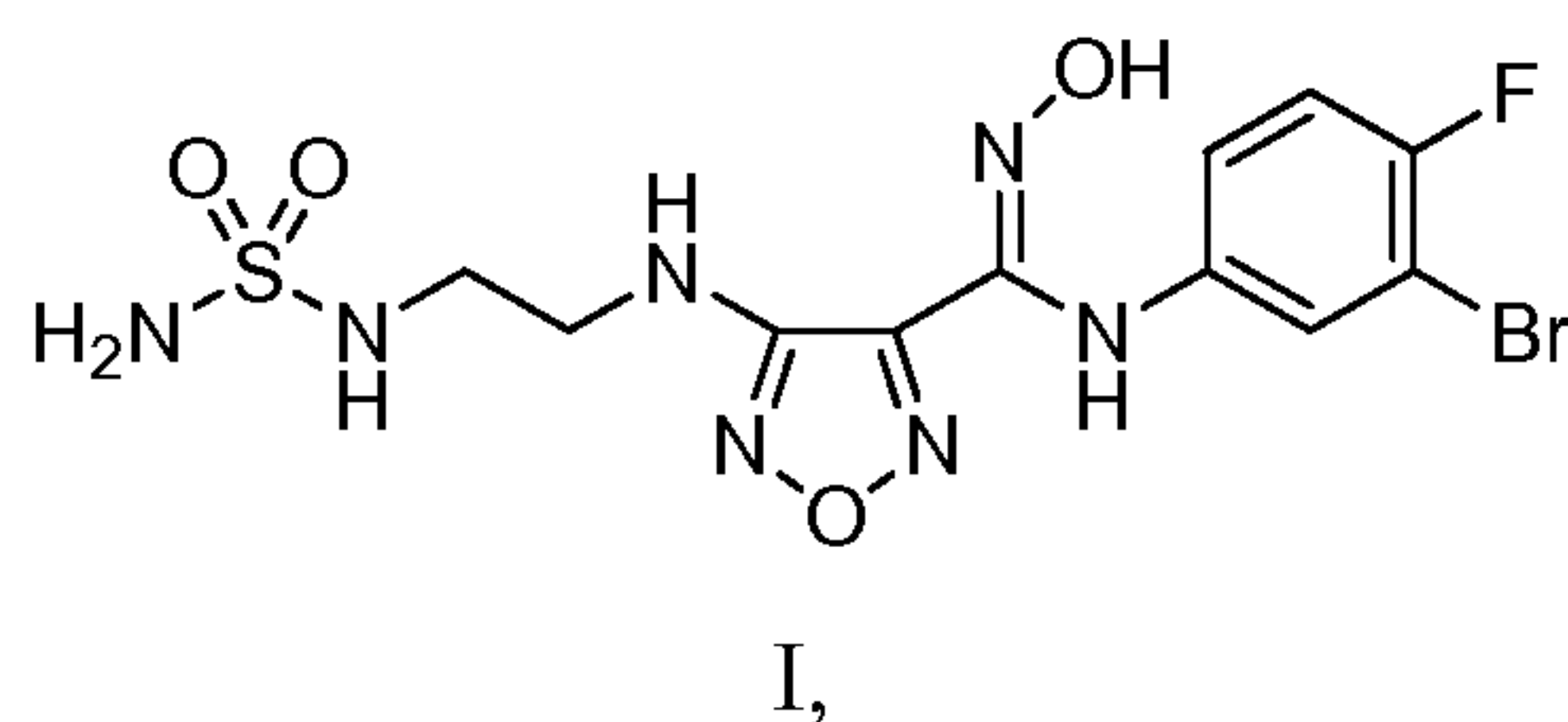
in the presence of triethylsilane and methanesulfonic acid to afford a compound of Formula

5 F20:



and

ii) converting said compound of Formula F20 to a compound of Formula I:

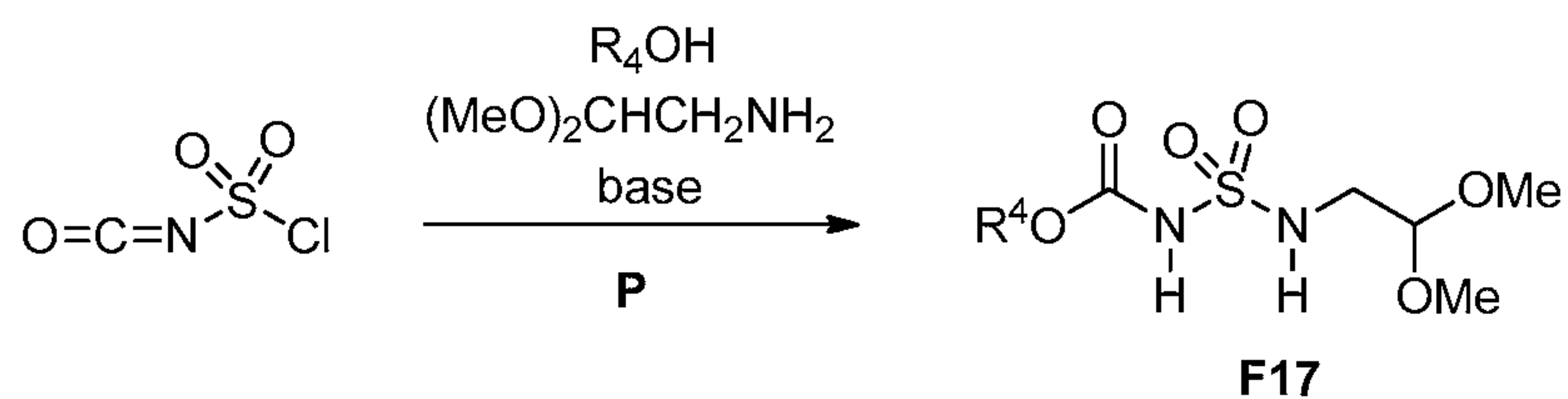


10

wherein said converting comprises combining said compound of Formula F20 with *N,N*-bis(2-aminoethyl)ethane-1,2-diamine. In some embodiments, said converting further comprises adding aqueous hydrochloric acid after said combining.

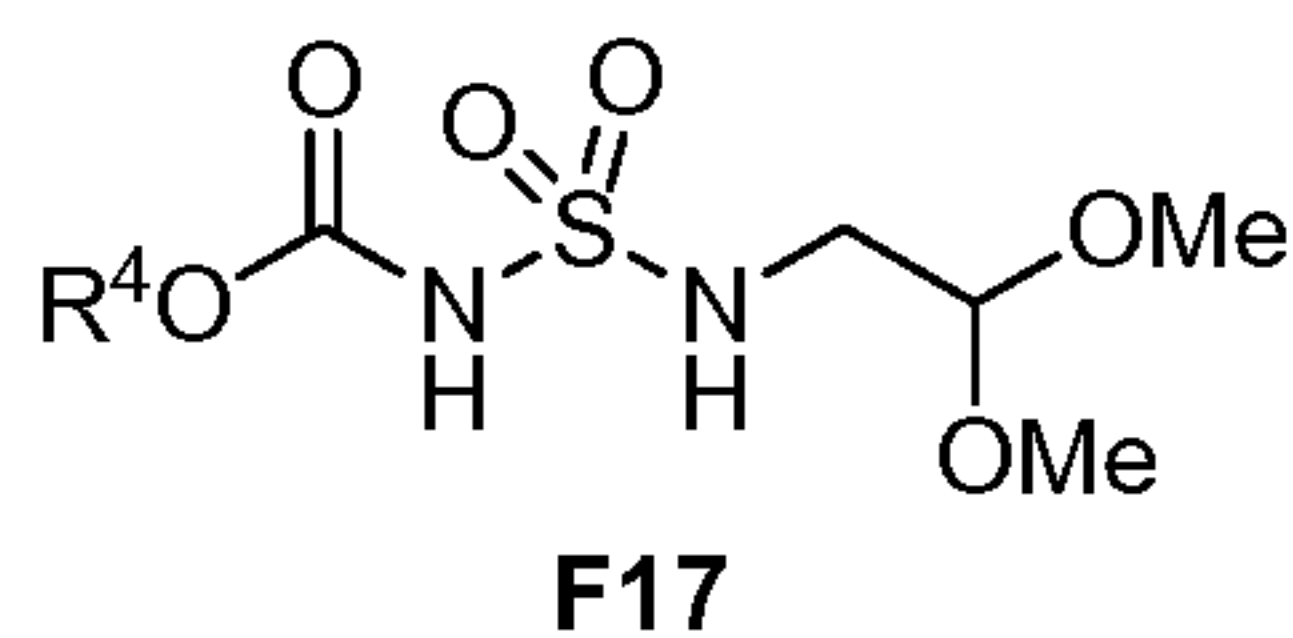
Compound F17 can be made by a one step process (Step P) from
15 chlorosulfonylisocyanate, as shown in Scheme 6.

Scheme 6



In some embodiments, the preparation of the compound of Formula F17 can be obtained by treating chlorosulfonylisocyanate with 2,2-dimethoxyethanamine and alcohol R⁴OH (R⁴ is defined as above), optionally in a solvent (e.g., a halogenated hydrocarbon solvent such as dichloromethane, chloroform, dichloroethane, tetrachloroethane). In some
 5 embodiments, this step carried out in the presence of a base. The base can be either an organic base such as an acyclic amine (e.g., triethylamine, diisopropylethylamine (DIPEA), etc.) or a cyclic amine (e.g., pyrrolidine, piperidine, etc.); or an inorganic base such as alkali (e.g., NaOH, LiOH, KOH, Mg(OH)₂, etc.). In some embodiments, the reaction is carried out in a solvent, for example, a halogenated hydrocarbon solvent such as dichloromethane,
 10 chloroform, dichloroethane, or tetrachloroethane.

In some embodiments, the present application further provides a compound of Formula F17:



wherein R⁴ is C₁₋₆ alkyl, C₁₋₆ haloalkyl, or benzyl.

15 In some embodiments, R⁴ is *tert*-butyl.

In some embodiments, R⁴ is benzyl.

In some embodiments, R⁴ is ethyl.

In some embodiments, R⁴ is C₁₋₃ haloalkyl.

In some embodiments, R⁴ is 2,2,2-trichloroethyl.

20 In some embodiments, R⁴ is 9*H*-fluoren-9-ylmethyl.

In some embodiments, the compound of Formula F17 is *tert*-butyl N-(2,2-dimethoxyethyl)sulfamoylcarbamate.

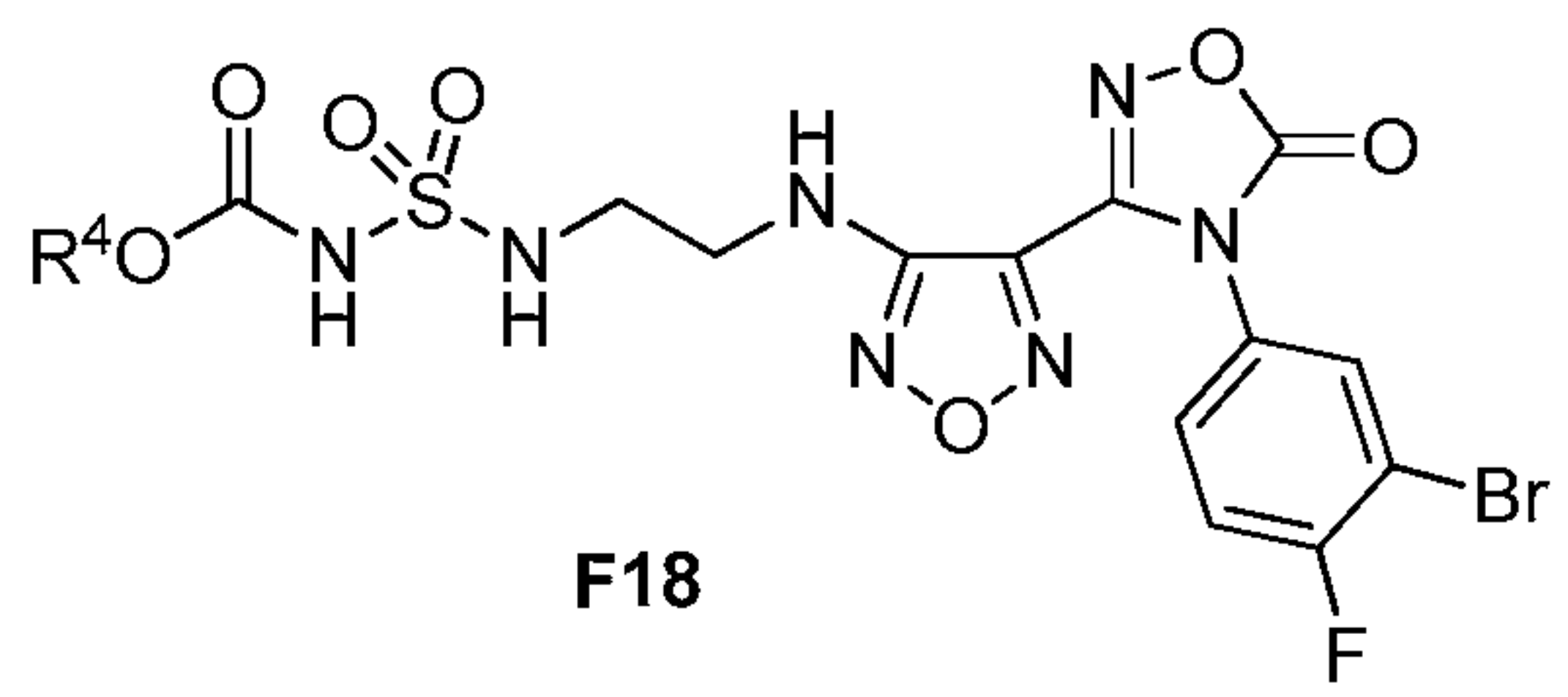
In some embodiments, the compound of Formula F17 is benzyl N-(2,2-dimethoxyethyl)sulfamoylcarbamate.

25 In some embodiments, the compound of Formula F17 is ethyl N-(2,2-dimethoxyethyl)sulfamoylcarbamate.

In some embodiments, the compound of Formula F17 is 2,2,2-trichloroethyl N-(2,2-dimethoxyethyl)sulfamoylcarbamate.

In some embodiments, the compound of Formula F17 is (9*H*-fluoren-9-yl)methyl N-(2,2-dimethoxyethyl)sulfamoylcarbamate.

In some embodiments, the present application further provides a compound of Formula F18:



wherein R^4 is C_{1-6} alkyl, C_{1-6} haloalkyl, or benzyl.

5 In some embodiments, R^4 is *tert*-butyl.

In some embodiments, R^4 is benzyl.

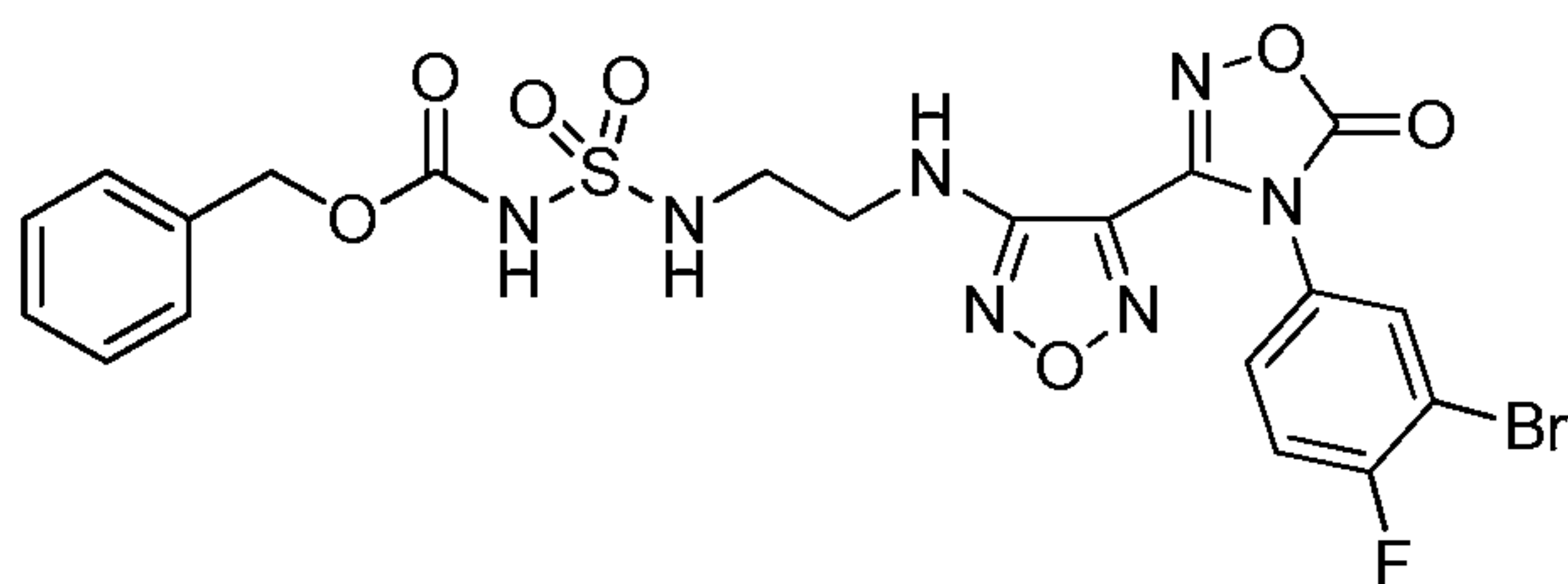
In some embodiments, R^4 is ethyl.

In some embodiments, R^4 is C_{1-3} haloalkyl.

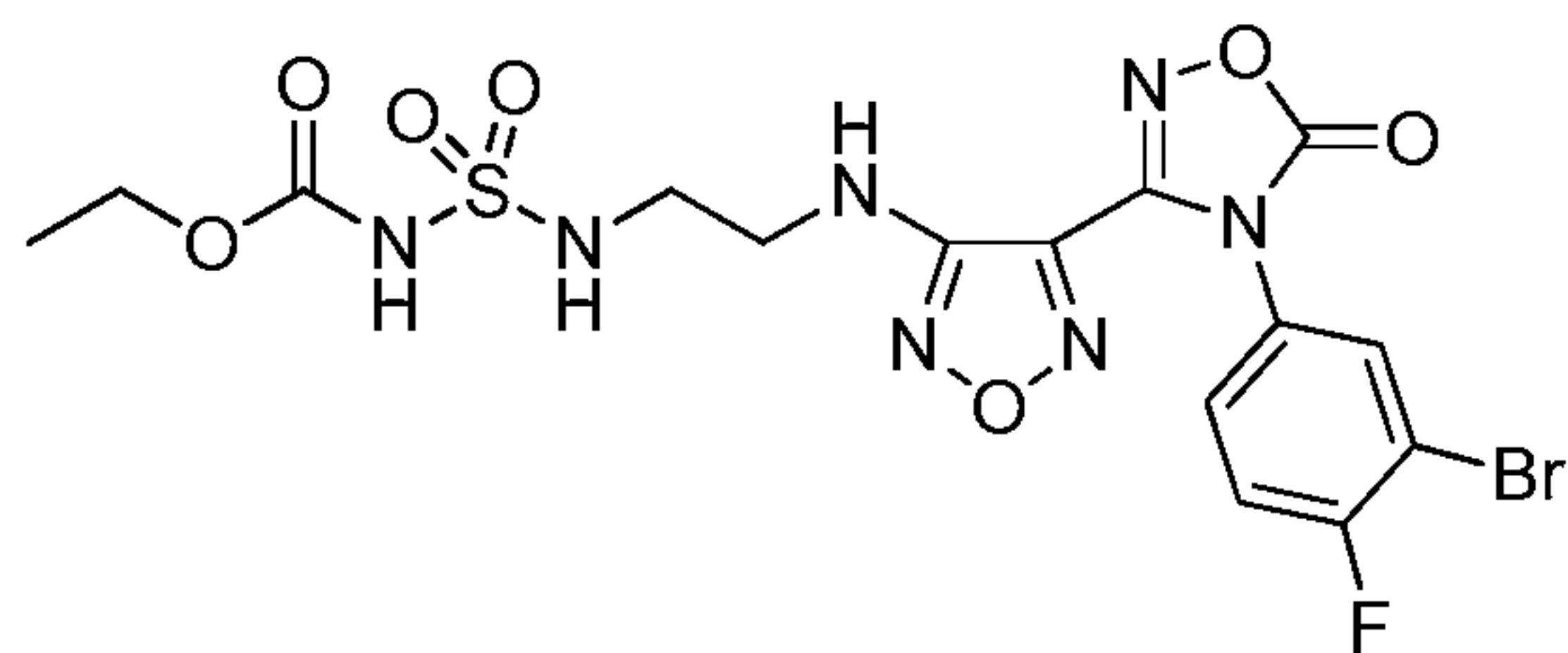
In some embodiments, R^4 is 2,2,2-trichloroethyl.

10 In some embodiments, R^4 is 9*H*-fluoren-9-ylmethyl.

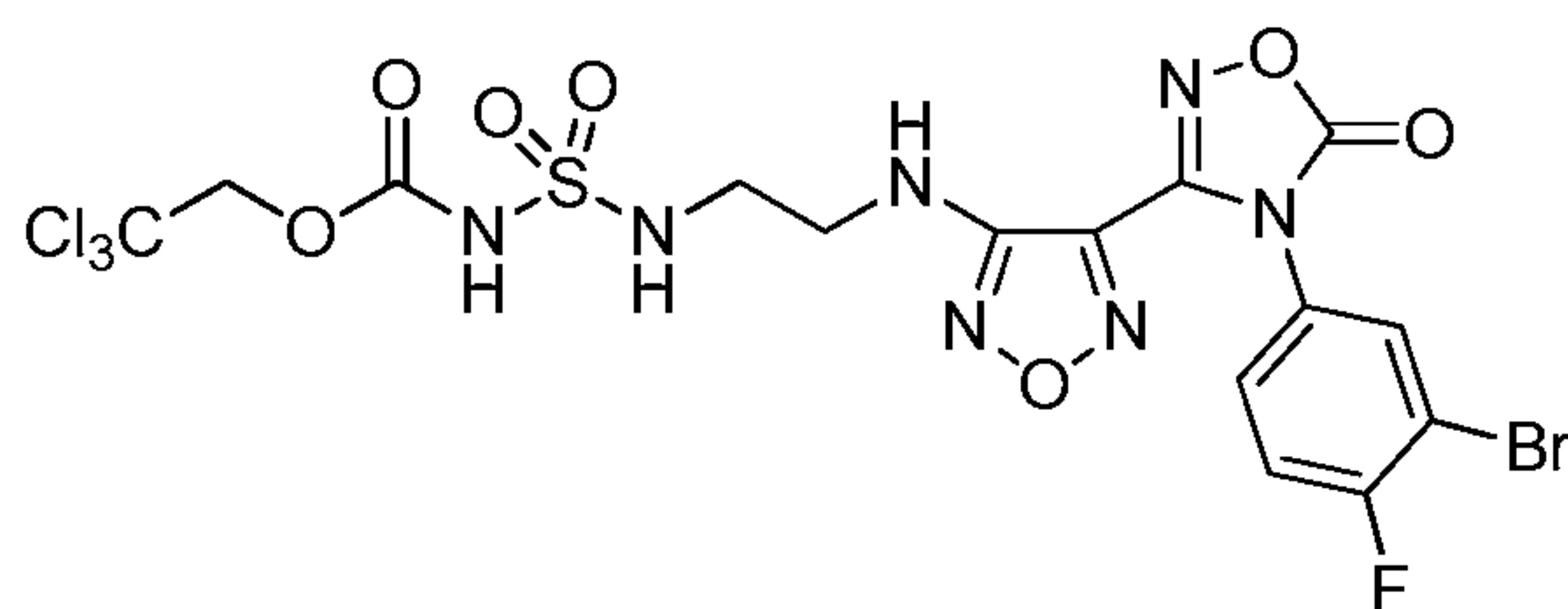
In some embodiments, the compound of Formula F18 is benzyl ({[2-({[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]amino}sulfonyl)carbamate:



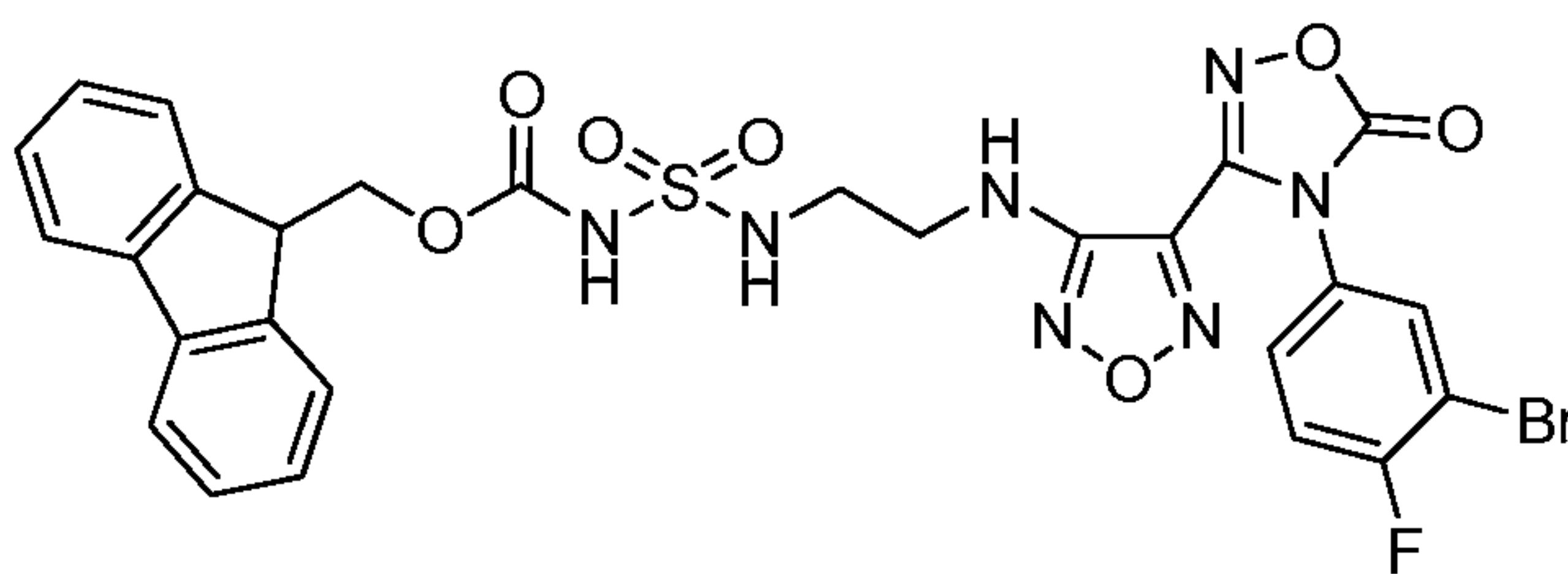
15 In some embodiments, the compound of Formula F18 is ethyl ({[2-({[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]amino}sulfonyl)carbamate:



20 In some embodiments, the compound of Formula F18 is 2,2,2-trichloroethyl ({[2-({[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]amino}sulfonyl)carbamate:



In some embodiments, the compound of Formula F18 is (9*H*-fluoren-9-yl)methyl N-(2-((4-(4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl)-1,2,5-oxadiazol-3-yl)amino)ethyl)sulfamoylcarbamate:



5

As used herein, the term “alkyl,” when used alone or together with additional moiety terms, refers to a straight-chained or branched, saturated hydrocarbon group having from 1 to 6 carbon atoms, 1 to 4 carbon atoms, or 1 to 3 carbon atoms. Example alkyl groups include methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, and the like.

10 As used herein, “alkenyl” refers to an alkyl group having one or more double carbon-carbon bonds. In some embodiments, said alkyl group has from 2 to 6 carbon atoms, 2 to 4 carbon atoms, or 2 to 3 carbon atoms. Example alkenyl groups include ethenyl (vinyl), propenyl, and the like.

As used herein, “alkenylalkyl” refers to a group of formula –alkyl-alkenyl. In some 15 embodiments, the alkenylalkyl group is allyl.

As used herein, the term “haloalkyl,” when used alone or together with an additional moiety, refers to an alkyl group substituted by one or more halogen atoms independently selected from F, Cl, Br, and I. Example haloalkyl groups include CF₃, CHF₂, CH₂CF₃, and the like.

20 As used herein, the term “alkoxy” refers to an –O-alkyl group. In some embodiments, the alkyl group has 1 to 6 carbon atoms, 1 to 4 carbon atoms, or 1 to 3 carbon atoms. Example alkoxy groups include methoxy, ethoxy, propoxy (*e.g.*, n-propoxy and isopropoxy), t-butoxy, and the like.

As used herein, “trialkylamine” refers to a nitrogen atom substituted by three 25 independently selected alkyl groups. In some embodiments, each alkyl group has from 2 to 6

carbon atoms, 2 to 4 carbon atoms, or 2 to 3 carbon atoms. Example trialkylamine groups include trimethylamine, triethylamine, and the like.

As used herein, the term “alkoxycarbonyl” refers to a group of formula $-C(O)-O-$ alkyl. In some embodiments, the alkyl group has from 2 to 6 carbon atoms, 2 to 4 carbon atoms, or 2 to 3 carbon atoms. Example alkoxycarbonyl groups include ethoxycarbonyl, *tert*-butoxycarbonyl (Boc), and the like.

Halogenated hydrocarbon solvents refer to halogenated alkanes, such as dichloromethane, chloroform, dichloroethane or tetrachloroethane, wherein the alkane can be branched or straight-chained having 1 to 12 carbon atoms, 1 to 6 carbon atoms, or 1 to 4 carbon atoms with one or more halo atoms. In some embodiments, the halogenated hydrocarbon solvent is a chlorinated alkane of 1 to 12 carbon atoms, 1 to 6 carbon atoms, or 1 to 4 carbon atoms.

At various places in the present specification, substituents of compounds of the invention may be disclosed in groups or in ranges. It is specifically intended that the invention include each and every individual subcombination of the members of such groups and ranges.

It is intended that the compounds of the invention are stable. As used herein “stable” refers to a compound that is sufficiently robust to survive isolation to a useful degree of purity from a reaction mixture, and preferably capable of formulation into an efficacious therapeutic agent.

It is further appreciated that certain features of the invention, which are, for clarity, described in the context of separate embodiments, can also be provided in combination in a single embodiment. Conversely, various features of the invention which are, for brevity, described in the context of a single embodiment, can also be provided separately or in any suitable subcombination.

The compounds of the invention are further intended to include all possible geometric isomers. Cis and trans geometric isomers of the compounds are described and may be isolated as a mixture of isomers or as separated isomeric forms.

Compounds of the invention also include tautomeric forms. Tautomeric forms result from the swapping of a single bond with an adjacent double bond together with the concomitant migration of a proton.

Compounds of the invention can also include all isotopes of atoms occurring in the intermediates or final compounds. Isotopes include those atoms having the same atomic

number but different mass numbers. For example, isotopes of hydrogen include tritium and deuterium.

In some embodiments, the compounds of the invention, and salts thereof, are substantially isolated. By “substantially isolated” is meant that the compound is at least partially or substantially separated from the environment in which it was formed or detected. Partial separation can include, for example, a composition enriched in the compound of the invention. Substantial separation can include compositions containing at least about 50%, at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least about 95%, at least about 97%, or at least about 99% by weight of the compound of the invention, or salt thereof. Methods for isolating compounds and their salts are routine in the art.

The present application also includes salts of the compounds described herein. As used herein, “salts” refers to derivatives of the disclosed compounds wherein the parent compound is modified by converting an existing acid or base moiety to its salt form. Examples of salts include, but are not limited to, mineral acid (such as HCl, HBr, H₂SO₄) or organic acid (such as acetic acid, benzoic acid, trifluoroacetic acid) salts of basic residues such as amines; alkali (such as Li, Na, K, Mg, Ca) or organic (such as trialkylammonium) salts of acidic residues such as carboxylic acids; and the like. The salts of the present application can be synthesized from the parent compound which contains a basic or acidic moiety by conventional chemical methods. Generally, such salts can be prepared by reacting the free acid or base forms of these compounds with a stoichiometric amount of the appropriate base or acid in water or in an organic solvent, or in a mixture of the two; generally, nonaqueous media like ether, ethyl acetate, ethanol, isopropanol, or acetonitrile (ACN) are preferred.

The present application also includes pharmaceutically acceptable salts of the compounds described herein. The “pharmaceutically acceptable salts” include a subset of the “salts” described above which are, conventional non-toxic salts of the parent compound formed, for example, from non-toxic inorganic or organic acids. Lists of suitable salts are found in *Remington's Pharmaceutical Sciences*, 17th ed., Mack Publishing Company, Easton, Pa., 1985, p. 1418 and *Journal of Pharmaceutical Science*, 66, 2 (1977), each of which is incorporated herein by reference in its entirety. The phrase “pharmaceutically acceptable” is employed herein to refer to those compounds, materials, compositions, and/or dosage forms which are, within the scope of sound medical judgment, suitable for use in contact with the tissues of human beings and animals without excessive toxicity, irritation, allergic response, or other problem or complication, commensurate with a reasonable benefit/risk ratio.

The processes described herein can be monitored according to any suitable method known in the art. For example, product formation can be monitored by spectroscopic means, such as nuclear magnetic resonance spectroscopy (e.g., ^1H or ^{13}C), infrared spectroscopy, spectrophotometry (e.g., UV-visible), or mass spectrometry; or by chromatography such as high performance liquid chromatography (HPLC) or thin layer chromatography. The compounds obtained by the reactions can be purified by any suitable method known in the art. For example, chromatography (medium pressure) on a suitable adsorbent (e.g., silica gel, alumina and the like), HPLC, or preparative thin layer chromatography; distillation; sublimation, trituration, or recrystallization. The purity of the compounds, in general, are determined by physical methods such as measuring the melting point (in case of a solid), obtaining a NMR spectrum, or performing a HPLC separation. If the melting point decreases, if unwanted signals in the NMR spectrum are decreased, or if extraneous peaks in an HPLC trace are removed, the compound can be said to have been purified. In some embodiments, the compounds are substantially purified.

Preparation of compounds can involve the protection and deprotection of various chemical groups. The need for protection and deprotection, and the selection of appropriate protecting groups can be readily determined by one skilled in the art. The chemistry of protecting groups can be found, for example, in Wuts and Greene, *Greene's Protective Groups in Organic Synthesis*, 4th Ed., John Wiley & Sons: New York, 2006, which is incorporated herein by reference in its entirety.

The reactions of the processes described herein can be carried out in suitable solvents which can be readily selected by one of skill in the art of organic synthesis. Suitable solvents can be substantially non-reactive with the starting materials (reactants), the intermediates, or products at the temperatures at which the reactions are carried out, i.e., temperatures which can range from the solvent's freezing temperature to the solvent's boiling temperature. A given reaction can be carried out in one solvent or a mixture of more than one solvent. Depending on the reaction step, suitable solvent(s) for that particular reaction step can be selected. Appropriate solvents include water, alkanes (such as pentanes, hexanes, heptanes, cyclohexane, etc., or a mixture thereof), aromatic solvents (such as benzene, toluene, xylene, etc.), alcohols (such as methanol, ethanol, isopropanol, etc.), ethers (such as dialkylethers, methyl *tert*-butyl ether (MTBE), tetrahydrofuran (THF), dioxane, etc.), esters (such as ethyl acetate, butyl acetate, etc.), halogenated hydrocarbon solvents (such as dichloromethane (DCM), chloroform, dichloroethane, tetrachloroethane), dimethylformamide (DMF), dimethylsulfoxide (DMSO), acetone, acetonitrile (ACN), hexamethylphosphoramide

(HMPA) and N-methyl pyrrolidone (NMP). Such solvents can be used in either their wet or anhydrous forms.

Resolution of racemic mixtures of compounds can be carried out by any of numerous methods known in the art. An example method includes fractional recrystallization using a
5 “chiral resolving acid” which is an optically active, salt-forming organic acid. Suitable resolving agents for fractional recrystallization methods are, for example, optically active acids, such as the D and L forms of tartaric acid, diacetyltartaric acid, dibenzoyltartaric acid, mandelic acid, malic acid, lactic acid or the various optically active camphorsulfonic acids. Resolution of racemic mixtures can also be carried out by elution on a column packed with an
10 optically active resolving agent (e.g., dinitrobenzoylphenylglycine). Suitable elution solvent composition can be determined by one skilled in the art.

Methods of Use

The compound of Formula I can inhibit activity of the enzyme indoleamine-2,3-
15 dioxygenase (IDO). For example, the compound of Formula I can be used to inhibit activity of IDO in cell or in an individual in need of modulation of the enzyme by administering an inhibiting amount of the compound of Formula I.

The compounds of Formula I can be used in methods of inhibiting the degradation of tryptophan in a system containing cells expressing IDO such as a tissue, living organism, or
20 cell culture. In some embodiments, the present application provides methods of altering (e.g., increasing) extracellular tryptophan levels in a mammal by administering an effective amount of the compound of Formula I. Methods of measuring tryptophan levels and tryptophan degradation are routine in the art.

The compounds of Formula I can be used in methods of inhibiting
25 immunosuppression such as IDO-mediated immunosuppression in a patient by administering to the patient an effective amount of the compound of Formula I. IDO-mediated immunosuppression has been associated with, for example, cancers, tumor growth, metastasis, viral infection, viral replication, etc.

The compounds of Formula I can also be used in methods of treating diseases
30 associated with activity or expression, including abnormal activity and/or overexpression, of IDO in an individual (e.g., patient) by administering to the individual in need of such treatment a therapeutically effective amount or dose of the compound of Formula I or a pharmaceutical composition thereof. Example diseases can include any disease, disorder or condition that is directly or indirectly linked to expression or activity of the IDO enzyme,

such as over expression or abnormal activity. An IDO-associated disease can also include any disease, disorder or condition that can be prevented, ameliorated, or cured by modulating enzyme activity. Examples of IDO-associated diseases include cancer, viral infection such as HIV infection, HCV infection, depression, neurodegenerative disorders such as Alzheimer's disease and Huntington's disease, trauma, age-related cataracts, organ transplantation (*e.g.*, organ transplant rejection), and autoimmune diseases including asthma, rheumatoid arthritis, multiple sclerosis, allergic inflammation, inflammatory bowel disease, psoriasis and systemic lupus erythematosus. Example cancers treatable by the methods herein include cancer of the colon, pancreas, breast, prostate, lung, brain, ovary, cervix, testes, renal, head and neck, lymphoma, leukemia, melanoma, and the like. The compound of Formula I can also be useful in the treatment of obesity and ischemia.

As used herein, the term "cell" is meant to refer to a cell that is *in vitro*, *ex vivo* or *in vivo*. In some embodiments, an *ex vivo* cell can be part of a tissue sample excised from an organism such as a mammal. In some embodiments, an *in vitro* cell can be a cell in a cell culture. In some embodiments, an *in vivo* cell is a cell living in an organism such as a mammal.

As used herein, the term "contacting" refers to the bringing together of indicated moieties in an *in vitro* system or an *in vivo* system. For example, "contacting" the IDO enzyme with the compound of Formula I includes the administration of the compound of Formula I to an individual or patient, such as a human, having IDO, as well as, for example, introducing the compound of Formula I into a sample containing a cellular or purified preparation containing the IDO enzyme.

As used herein, the term "individual" or "patient," used interchangeably, refers to any animal, including mammals, preferably mice, rats, other rodents, rabbits, dogs, cats, swine, cattle, sheep, horses, or primates, and most preferably humans.

As used herein, the phrase "therapeutically effective amount" refers to the amount of active compound or pharmaceutical agent that elicits the biological or medicinal response in a tissue, system, animal, individual or human that is being sought by a researcher, veterinarian, medical doctor or other clinician.

As used herein the term "treating" or "treatment" refers to 1) preventing the disease; for example, preventing a disease, condition or disorder in an individual who may be predisposed to the disease, condition or disorder but does not yet experience or display the pathology or symptomatology of the disease; 2) inhibiting the disease; for example, inhibiting a disease, condition or disorder in an individual who is experiencing or displaying the

pathology or symptomatology of the disease, condition or disorder (*i.e.*, arresting further development of the pathology and/or symptomatology), or 3) ameliorating the disease; for example, ameliorating a disease, condition or disorder in an individual who is experiencing or displaying the pathology or symptomatology of the disease, condition or disorder (*i.e.*,
 5 reversing the pathology and/or symptomatology).

Combination Therapy

One or more additional pharmaceutical agents or treatment methods such as, for example, anti-viral agents, chemotherapeutics or other anti-cancer agents, immune enhancers,
 10 immunosuppressants, radiation, anti-tumor and anti-viral vaccines, cytokine therapy (*e.g.*, IL2, GM-CSF, *etc.*), and/or tyrosine kinase inhibitors can be used in combination the compound of Formula I for treatment of IDO-associated diseases, disorders or conditions. The agents can be combined with the compound of Formula I in a single dosage form, or the agents can be administered simultaneously or sequentially as separate dosage forms.

15 Suitable antiviral agents contemplated for use in combination the compound of Formula I can comprise nucleoside and nucleotide reverse transcriptase inhibitors (NRTIs), non-nucleoside reverse transcriptase inhibitors (NNRTIs), protease inhibitors and other antiviral drugs.

Example suitable NRTIs include zidovudine (AZT); didanosine (ddl); zalcitabine
 20 (ddC); stavudine (d4T); lamivudine (3TC); abacavir (1592U89); adefovir dipivoxil [bis(POM)-PMEA]; lobucavir (BMS-180194); BCH-10652; emitricitabine [(-)-FTC]; beta-L-FD4 (also called beta-L-D4C and named beta-L-2', 3'-dicleoxy-5-fluoro-cytidene); DAPD, ((-)-beta-D-2,6,-diamino-purine dioxolane); and lodenosine (FddA). Typical suitable NNRTIs include nevirapine (BI-RG-587); delaviradine (BHAP, U-90152); efavirenz (DMP-266);
 25 PNU-142721; AG-1549; MKC-442 (1-(ethoxy-methyl)-5-(1-methylethyl)-6-(phenylmethyl)-(2,4(1H,3H)-pyrimidinedione); and (+)-calanolide A (NSC-675451) and B. Typical suitable protease inhibitors include saquinavir (Ro 31-8959); ritonavir (ABT-538); indinavir (MK-639); nelfinavir (AG-1343); amprenavir (141W94); lasinavir (BMS-234475); DMP-450; BMS-2322623; ABT-378; and AG-1 549. Other antiviral agents include hydroxyurea,
 30 ribavirin, IL-2, IL-12, pentafuside and Yisum Project No.11607.

Suitable chemotherapeutic or other anti-cancer agents include, for example, alkylating agents (including, without limitation, nitrogen mustards, ethylenimine derivatives, alkyl sulfonates, nitrosoureas and triazenes) such as uracil mustard, chlormethine, cyclophosphamide (CytoxanTM), ifosfamide, melphalan, chlorambucil, pipobroman,

triethylene-melamine, triethylenethiophosphoramine, busulfan, carmustine, lomustine, streptozocin, dacarbazine, and temozolomide.

In the treatment of melanoma, suitable agents for use in combination with the compound of Formula I include: dacarbazine (DTIC), optionally, along with other chemotherapy drugs such as carmustine (BCNU) and cisplatin; the “Dartmouth regimen,” which consists of DTIC, BCNU, cisplatin and tamoxifen; a combination of cisplatin, vinblastine, and DTIC; or temozolomide. Compounds according to the invention may also be combined with immunotherapy drugs, including cytokines such as interferon alpha, interleukin 2, and tumor necrosis factor (TNF) in the treatment of melanoma.

The compound of Formula I may also be used in combination with vaccine therapy in the treatment of melanoma. Antimelanoma vaccines are, in some ways, similar to the anti-virus vaccines which are used to prevent diseases caused by viruses such as polio, measles, and mumps. Weakened melanoma cells or parts of melanoma cells called antigens may be injected into a patient to stimulate the body's immune system to destroy melanoma cells.

Melanomas that are confined to the arms or legs may also be treated with a combination of agents including the compound of Formula I, using a hyperthermic isolated limb perfusion technique. This treatment protocol temporarily separates the circulation of the involved limb from the rest of the body and injects high doses of chemotherapy into the artery feeding the limb, thus providing high doses to the area of the tumor without exposing internal organs to these doses that might otherwise cause severe side effects. Usually the fluid is warmed to 102° to 104° F. Melphalan is the drug most often used in this chemotherapy procedure. This can be given with another agent called tumor necrosis factor (TNF) (see section on cytokines).

Suitable chemotherapeutic or other anti-cancer agents include, for example, antimetabolites (including, without limitation, folic acid antagonists, pyrimidine analogs, purine analogs and adenosine deaminase inhibitors) such as methotrexate, 5-fluorouracil, floxuridine, cytarabine, 6-mercaptopurine, 6-thioguanine, fludarabine phosphate, pentostatine, and gemcitabine.

Suitable chemotherapeutic or other anti-cancer agents further include, for example, certain natural products and their derivatives (for example, vinca alkaloids, antitumor antibiotics, enzymes, lymphokines and epipodophyllotoxins) such as vinblastine, vincristine, vindesine, bleomycin, dactinomycin, daunorubicin, doxorubicin, epirubicin, idarubicin, ara-C, paclitaxel (TAXOLTM), mithramycin, deoxycoformycin, mitomycin-C, L-asparaginase, interferons (especially IFN- α), etoposide, and teniposide.

Other cytotoxic agents include navelbene, CPT-11, anastrozole, letrozole, capecitabine, reloxafine, cyclophosphamide, ifosamide, and droloxafine.

Also suitable are cytotoxic agents such as epidophyllotoxin; an antineoplastic enzyme; a topoisomerase inhibitor; procarbazine; mitoxantrone; platinum coordination
5 complexes such as cis-platin and carboplatin; biological response modifiers; growth inhibitors; antihormonal therapeutic agents; leucovorin; tegafur; and haematopoietic growth factors.

Other anti-cancer agent(s) include antibody therapeutics such as trastuzumab (Herceptin), antibodies to costimulatory molecules such as CTLA-4, 4-1BB and PD-1, or
10 antibodies to cytokines (IL-10, TGF- β , etc.).

Other anti-cancer agents also include those that block immune cell migration such as antagonists to chemokine receptors, including CCR2 and CCR4.

Other anti-cancer agents also include those that augment the immune system such as adjuvants or adoptive T cell transfer.

15 Anti-cancer vaccines include dendritic cells, synthetic peptides, DNA vaccines and recombinant viruses.

Methods for the safe and effective administration of most of these chemotherapeutic agents are known to those skilled in the art. In addition, their administration is described in the standard literature. For example, the administration of many of the chemotherapeutic
20 agents is described in the "Physicians' Desk Reference" (PDR, e.g., 1996 edition, Medical Economics Company, Montvale, NJ), the disclosure of which is incorporated herein by reference as if set forth in its entirety.

Pharmaceutical Formulations and Dosage Forms

25 When employed as pharmaceuticals, the compound of Formula I can be administered in the form of pharmaceutical compositions which is a combination of the compound of Formula I and a pharmaceutically acceptable carrier. These compositions can be prepared in a manner well known in the pharmaceutical art, and can be administered by a variety of routes, depending upon whether local or systemic treatment is desired and upon the area to be
30 treated. Administration may be topical (including ophthalmic and to mucous membranes including intranasal, vaginal and rectal delivery), pulmonary (e.g., by inhalation or insufflation of powders or aerosols, including by nebulizer; intratracheal, intranasal, epidermal and transdermal), ocular, oral or parenteral. Methods for ocular delivery can

include topical administration (eye drops), subconjunctival, periocular or intravitreal injection or introduction by balloon catheter or ophthalmic inserts surgically placed in the conjunctival sac. Parenteral administration includes intravenous, intraarterial, subcutaneous, intraperitoneal, or intramuscular injection or infusion; or intracranial, e.g., intrathecal or intraventricular, administration. Parenteral administration can be in the form of a single bolus dose, or may be, for example, by a continuous perfusion pump. Pharmaceutical compositions and formulations for topical administration may include transdermal patches, ointments, lotions, creams, gels, drops, suppositories, sprays, liquids and powders. Conventional pharmaceutical carriers, aqueous, powder or oily bases, thickeners and the like may be necessary or desirable.

Pharmaceutical compositions containing the compound of Formula I can be prepared in combination with one or more pharmaceutically acceptable carriers. In making the compositions of the invention, the active ingredient is typically mixed with an excipient, diluted by an excipient or enclosed within such a carrier in the form of, for example, a capsule, sachet, paper, or other container. When the excipient serves as a diluent, it can be a solid, semi-solid, or liquid material, which acts as a vehicle, carrier or medium for the active ingredient. Thus, the compositions can be in the form of tablets, pills, powders, lozenges, sachets, cachets, elixirs, suspensions, emulsions, solutions, syrups, aerosols (as a solid or in a liquid medium), ointments containing, for example, up to 10 % by weight of the active compound, soft and hard gelatin capsules, suppositories, sterile injectable solutions, and sterile packaged powders.

In preparing a formulation, the active compound can be milled to provide the appropriate particle size prior to combining with the other ingredients. If the active compound is substantially insoluble, it can be milled to a particle size of less than 200 mesh. If the active compound is substantially water soluble, the particle size can be adjusted by milling to provide a substantially uniform distribution in the formulation, e.g. about 40 mesh.

Some examples of suitable excipients include lactose, dextrose, sucrose, sorbitol, mannitol, starches, gum acacia, calcium phosphate, alginates, tragacanth, gelatin, calcium silicate, microcrystalline cellulose, polyvinylpyrrolidone, cellulose, water, syrup, and methyl cellulose. The formulations can additionally include: lubricating agents such as talc, magnesium stearate, and mineral oil; wetting agents; emulsifying and suspending agents; preserving agents such as methyl- and propylhydroxy-benzoates; sweetening agents; and flavoring agents. The compositions of the invention can be formulated so as to provide quick,

sustained or delayed release of the active ingredient after administration to the patient by employing procedures known in the art.

The compositions can be formulated in a unit dosage form, each dosage containing from about 5 to about 100 mg, more usually about 10 to about 30 mg, of the active
5 ingredient. The term "unit dosage forms" refers to physically discrete units suitable as unitary dosages for human subjects and other mammals, each unit containing a predetermined quantity of active material calculated to produce the desired therapeutic effect, in association with a suitable pharmaceutical excipient.

The active compound can be effective over a wide dosage range and is generally
10 administered in a pharmaceutically effective amount. It will be understood, however, that the amount of the compound actually administered will usually be determined by a physician, according to the relevant circumstances, including the condition to be treated, the chosen route of administration, the actual compound administered, the age, weight, and response of the individual patient, the severity of the patient's symptoms, and the like.

15 For preparing solid compositions such as tablets, the principal active ingredient is mixed with a pharmaceutical excipient to form a solid pre-formulation composition containing a homogeneous mixture of the compound of Formula I. When referring to these pre-formulation compositions as homogeneous, the active ingredient is typically dispersed evenly throughout the composition so that the composition can be readily subdivided into
20 equally effective unit dosage forms such as tablets, pills and capsules. This solid pre-formulation is then subdivided into unit dosage forms of the type described above containing from, for example, 0.1 to about 500 mg of the active ingredient of the present application.

The tablets or pills containing the compound of Formula I can be coated or otherwise compounded to provide a dosage form affording the advantage of prolonged action. For
25 example, the tablet or pill can comprise an inner dosage and an outer dosage component, the latter being in the form of an envelope over the former. The two components can be separated by an enteric layer which serves to resist disintegration in the stomach and permit the inner component to pass intact into the duodenum or to be delayed in release. A variety of materials can be used for such enteric layers or coatings, such materials including a number
30 of polymeric acids and mixtures of polymeric acids with such materials as shellac, cetyl alcohol, and cellulose acetate.

The liquid forms in which the compounds and compositions of the present application can be incorporated for administration orally or by injection include aqueous solutions, suitably flavored syrups, aqueous or oil suspensions, and flavored emulsions with edible oils

such as cottonseed oil, sesame oil, coconut oil, or peanut oil, as well as elixirs and similar pharmaceutical vehicles.

Compositions for inhalation or insufflation include solutions and suspensions in pharmaceutically acceptable, aqueous or organic solvents, or mixtures thereof, and powders.

5 The liquid or solid compositions may contain suitable pharmaceutically acceptable excipients as described supra. In some embodiments, the compositions are administered by the oral or nasal respiratory route for local or systemic effect. Compositions in can be nebulized by use of inert gases. Nebulized solutions may be breathed directly from the nebulizing device or the nebulizing device can be attached to a face masks tent, or intermittent positive pressure
10 breathing machine. Solution, suspension, or powder compositions can be administered orally or nasally from devices which deliver the formulation in an appropriate manner.

The amount of compound or composition administered to a patient will vary depending upon what is being administered, the purpose of the administration, such as prophylaxis or therapy, the state of the patient, the manner of administration, and the like. In
15 therapeutic applications, compositions can be administered to a patient already suffering from a disease in an amount sufficient to cure or at least partially arrest the symptoms of the disease and its complications. Effective doses will depend on the disease condition being treated as well as by the judgment of the attending clinician depending upon factors such as the severity of the disease, the age, weight and general condition of the patient, and the like.

20 The compositions administered to a patient can be in the form of pharmaceutical compositions described above. These compositions can be sterilized by conventional sterilization techniques, or may be sterile filtered. Aqueous solutions can be packaged for use as is, or lyophilized, the lyophilized preparation being combined with a sterile aqueous carrier prior to administration. The pH of the compound preparations typically will be between 3 and
25 11, more preferably from 5 to 9 and most preferably from 7 to 8. It will be understood that use of certain of the foregoing excipients, carriers, or stabilizers will result in the formation of pharmaceutical salts.

The therapeutic dosage of the the compound of Formula I can vary according to, for example, the particular use for which the treatment is made, the manner of administration of
30 the compound, the health and condition of the patient, and the judgment of the prescribing physician. The proportion or concentration of the compound of Formula I in a pharmaceutical composition can vary depending upon a number of factors including dosage, chemical characteristics (e.g., hydrophobicity), and the route of administration. For example, the compound of Formula I can be provided in an aqueous physiological buffer solution

containing about 0.1 to about 10% w/v of the compound for parenteral administration. Some typical dose ranges are from about 1 µg/kg to about 1 g/kg of body weight per day. In some embodiments, the dose range is from about 0.01 mg/kg to about 100 mg/kg of body weight per day. The dosage is likely to depend on such variables as the type and extent of progression of the disease or disorder, the overall health status of the particular patient, the relative biological efficacy of the compound selected, formulation of the excipient, and its route of administration. Effective doses can be extrapolated from dose-response curves derived from *in vitro* or animal model test systems.

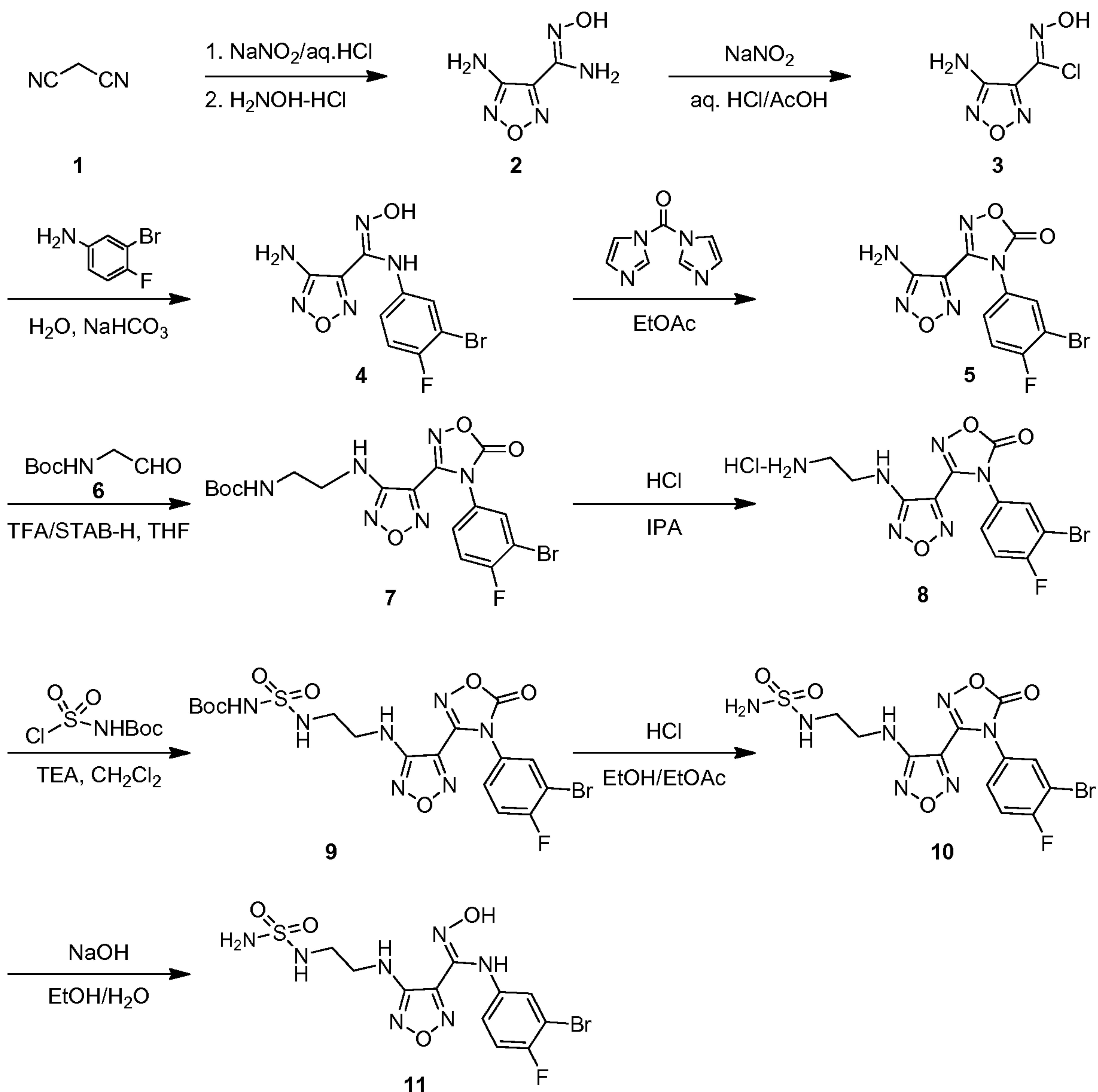
The compound of Formula I can also be formulated in combination with one or more additional active ingredients which can include any pharmaceutical agent such as anti-viral agents, vaccines, antibodies, immune enhancers, immune suppressants, anti-inflammatory agents and the like.

The present application also includes pharmaceutical kits useful, for example, in the treatment or prevention of IDO-associated diseases or disorders, obesity, diabetes and other diseases referred to herein which include one or more containers containing a pharmaceutical composition comprising a therapeutically effective amount of the compound of Formula I. Such kits can further include, if desired, one or more of various conventional pharmaceutical kit components, such as, for example, containers with one or more pharmaceutically acceptable carriers, additional containers, etc., as will be readily apparent to those skilled in the art. Instructions, either as inserts or as labels, indicating quantities of the components to be administered, guidelines for administration, and/or guidelines for mixing the components, can also be included in the kit.

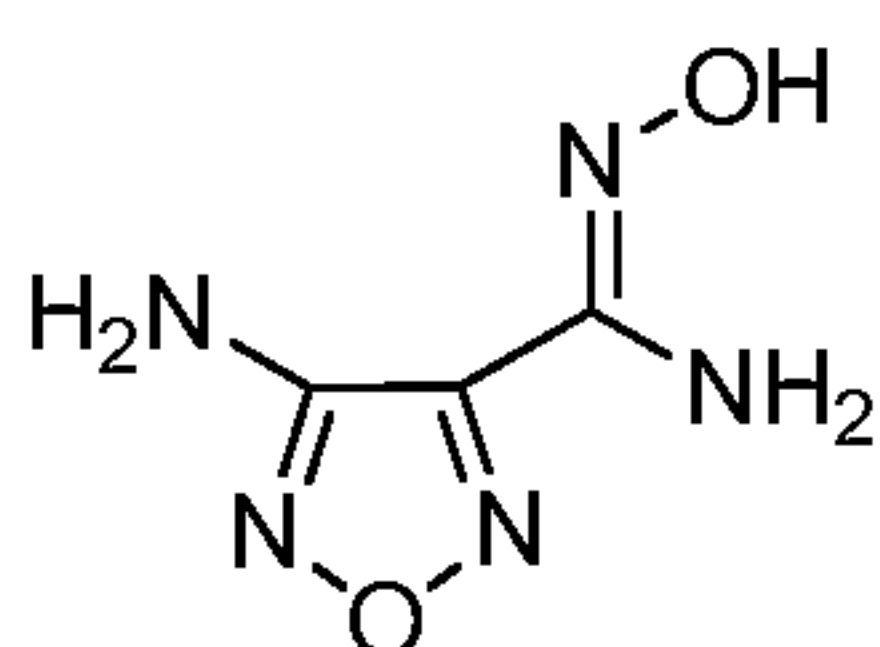
EXAMPLES

The invention will be described in greater detail by way of specific examples. The following examples are offered for illustrative purposes, and are not intended to limit the invention in any manner. Those of skill in the art will readily recognize a variety of non-critical parameters which can be changed or modified to yield essentially the same results.

Example 1. Synthesis of 4-({2-[(Aminosulfonyl)amino]ethyl}amino)-N-(3-bromo-4-fluorophenyl)-N'-hydroxy-1,2,5-oxadiazole-3-carboximidamide



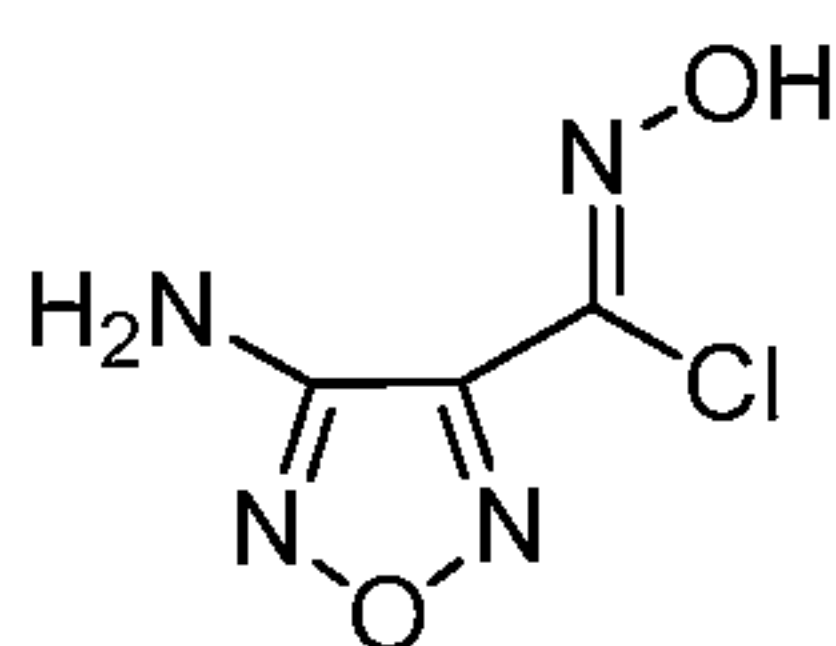
Step A: 4-Amino-*N*'-hydroxy-1,2,5-oxadiazole-3-carboximidamide (2)



- Malononitrile [Aldrich, product # M1407] (320.5 g, 5 mol) was added to water (7 L), preheated to 45 °C and stirred at 45 °C for 5 min. The resulting solution was cooled in an ice bath and sodium nitrite (380 g, 5.5 mol, 1.1 equiv.) was added. When the temperature reached 10 °C, 6 N hydrochloric acid (55 mL) was added. A mild exothermic reaction ensued with the temperature reaching 16 °C. After 15 min the cold bath was removed and the

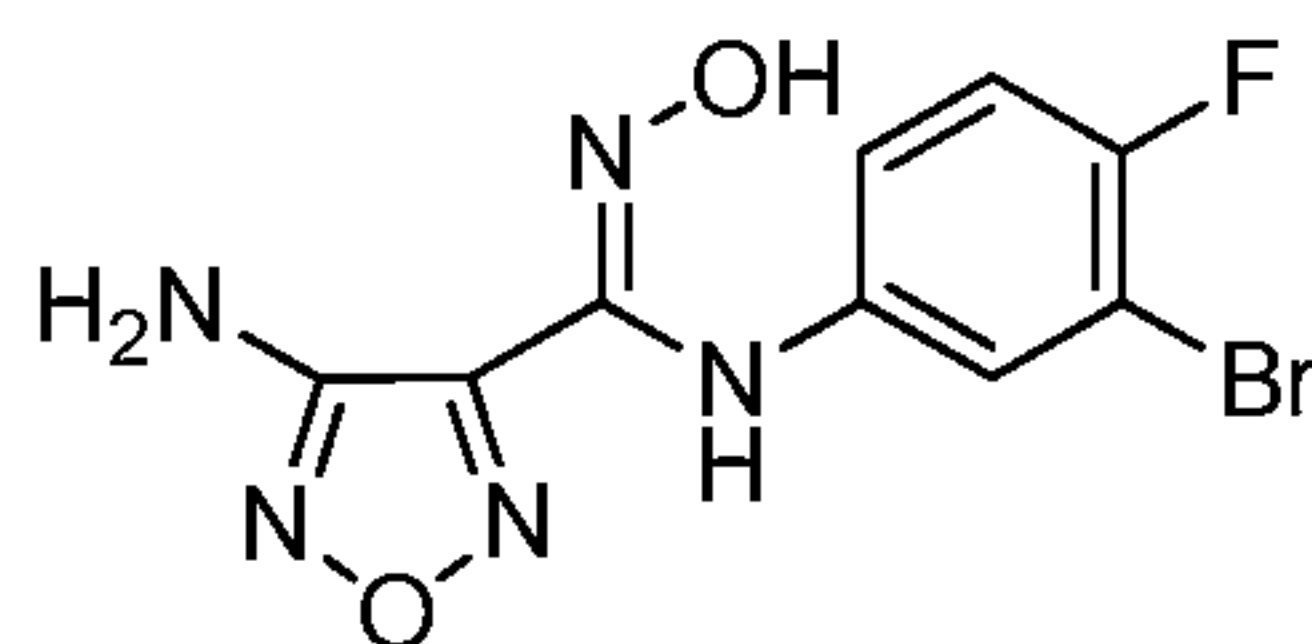
reaction mixture was stirred for 1.5 h at 16-18 °C. The reaction mixture was cooled to 13 °C and 50% aqueous hydroxylamine hydrochloride (990 g, 15 mol, 3.0 equiv.) was added all at once. The temperature rose to 26 °C. When the exothermic reaction subsided the cold bath was removed and stirring was continued for 1 h at 26 - 27 °C, then it was slowly brought to reflux. Reflux was maintained for 2 h and then the reaction mixture was allowed to gradually cool overnight. The reaction mixture was stirred in an ice bath and 6 N hydrochloric acid (800 mL) was added in portions over 40 min to adjust to pH 7.0. Stirring was continued in the ice bath at 5 °C. The precipitate was collected by filtration, washed well with water and dried in a vacuum oven (50 °C) to give the desired product (644 g, 90%) as an off-white solid. ¹³C NMR (75 MHz, CD₃OD): δ 156.0, 145.9, 141.3; C₃H₅N₅O₂ (MW 143.10), LCMS (EI) *m/e* 144.0 (M⁺ + H).

Step B: 4-Amino-N'-hydroxy-1,2,5-oxadiazole-3-carboximidoyl chloride (3)



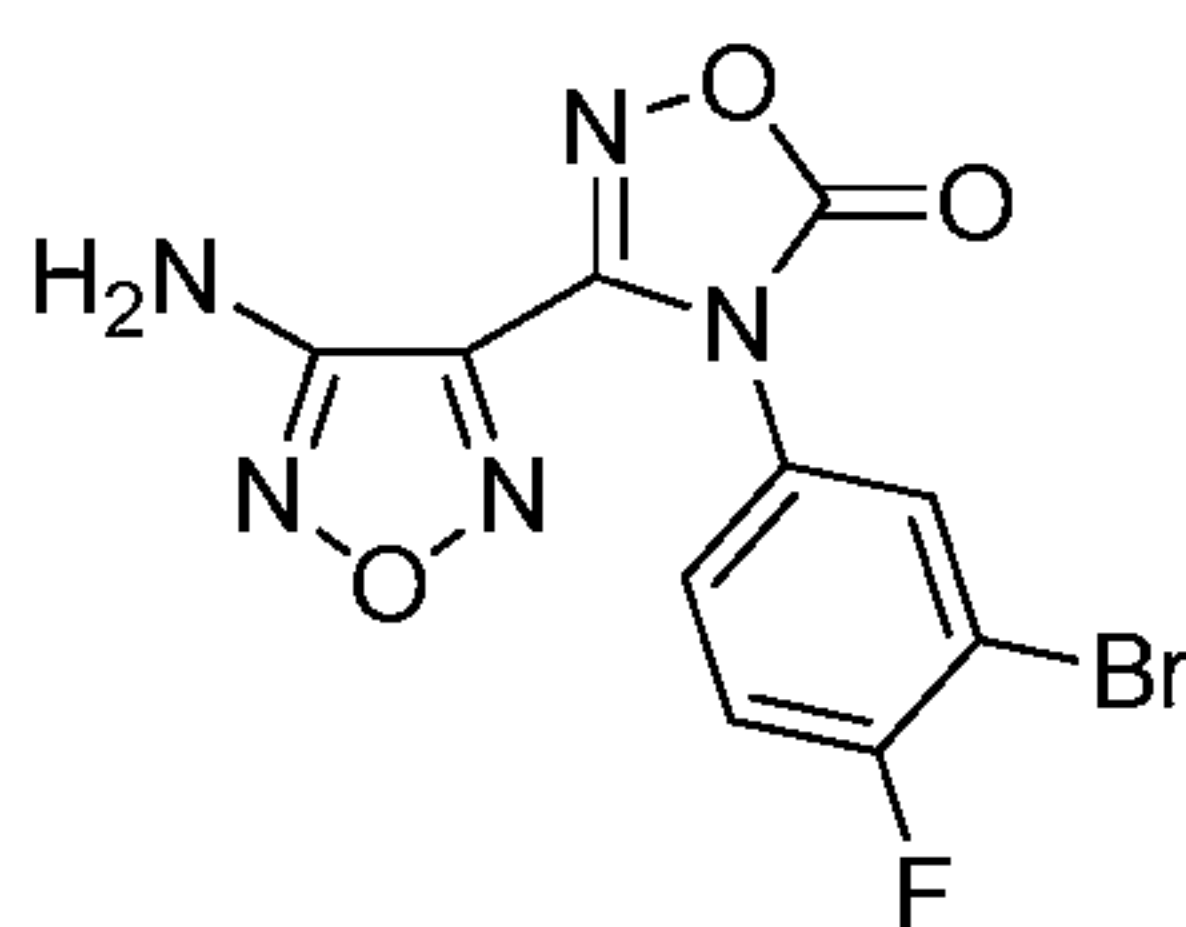
4-Amino-*N'*-hydroxy-1,2,5-oxadiazole-3-carboximidamide (422 g, 2.95 mol) was added to a mixture of water (5.9 L), acetic acid (3 L) and 6 N hydrochloric acid (1.475 L, 3.0 equiv.) and the suspension was stirred at 42 - 45 °C until a clear solution was achieved. Sodium chloride (518 g, 3.0 equiv.) was added and this solution was stirred in an ice/water/methanol bath. A solution of sodium nitrite (199.5 g, 0.98 equiv.) in water (700 mL) was added over 3.5 h while maintaining the temperature below 0 °C. After complete addition, stirring was continued in the ice bath for 1.5 h and then the reaction mixture was allowed to warm to 15 °C. The precipitate was collected by filtration, washed well with water, taken in ethyl acetate (3.4 L), treated with anhydrous sodium sulfate (500 g) and stirred at room temperature for 1 h. This suspension was filtered through sodium sulfate (200 g) and the filtrate was concentrated on a rotary evaporator. The residue was dissolved in methyl *tert*-butyl ether (5.5 L), treated with the activated charcoal (40 g), stirred at room temperature for 40 min and filtered through Celite. The solvent was removed in a rotary evaporator and the resulting product was dried in a vacuum oven (45 °C) to give the desired product (256 g, 53.4%) as an off-white solid. ¹³C NMR (100 MHz, CD₃OD) δ 155.8, 143.4, 129.7; C₃H₃ClN₄O₂ (MW 162.53), LCMS (EI) *m/e* 163/165 (M⁺ + H).

Step C: 4-Amino-N-(3-bromo-4-fluorophenyl)-N'-hydroxy-1,2,5-oxadiazole-3-carboximidamide (4)



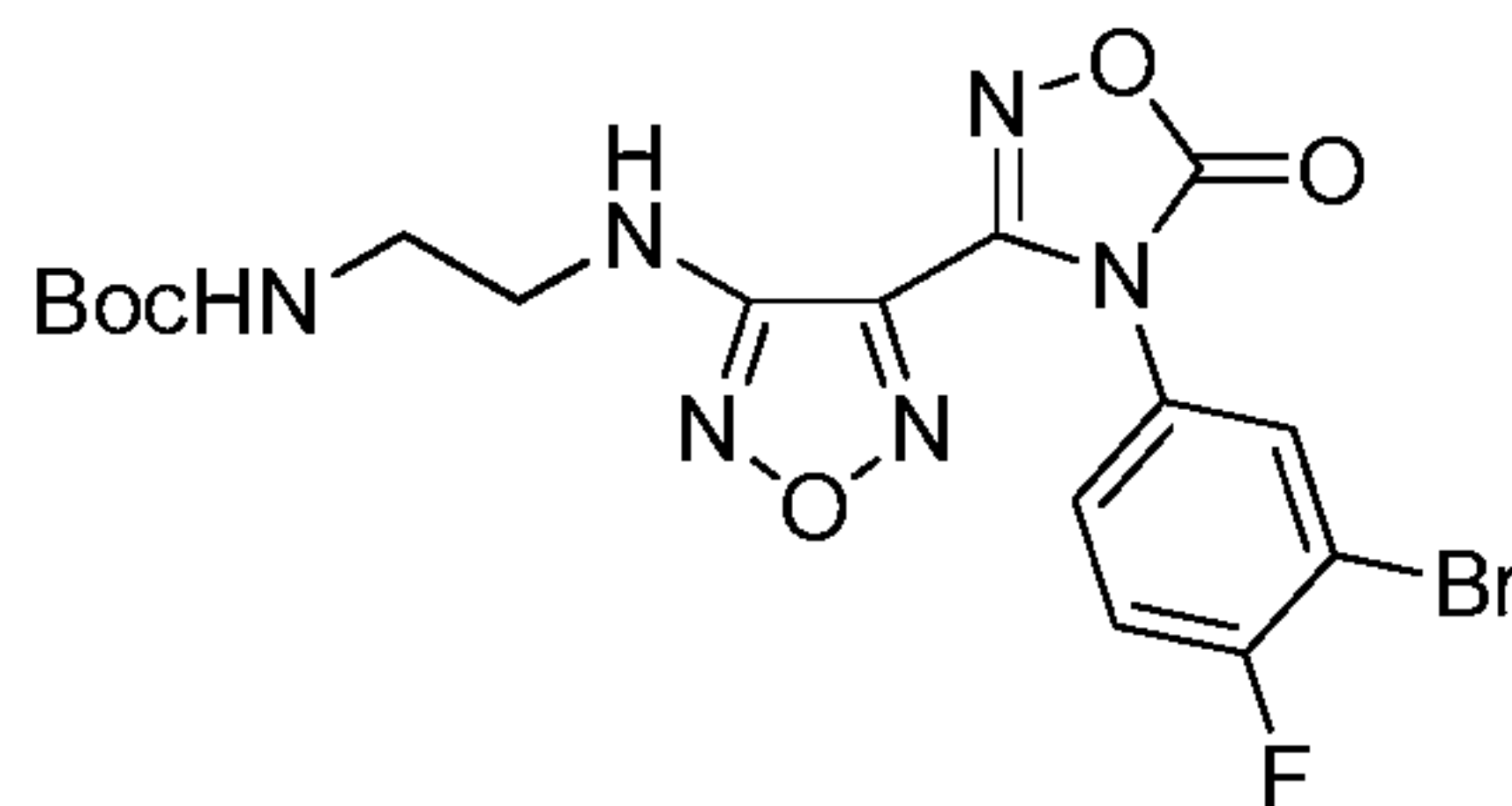
4-Amino-N-hydroxy-1,2,5-oxadiazole-3-carboximidoyl chloride (33.8 g, 208 mmol) was mixed with water (300 mL). At 60 °C, 3-bromo-4-fluoroaniline (Sigma-Aldrich) (43.6 g, 229 mmol, 1.1 equiv.) was added to the suspension with stirring for 10 min. A solution of sodium bicarbonate (26.3 g, 313 mmol, 1.5 equiv.) in water (300 mL) was added over 15 min with stirring at 60 °C. After stirring 20 min, LCMS indicated reaction completion. The reaction mixture was then cooled to room temperature and extracted with ethyl acetate (2 x 300 mL). The combined ethyl acetate solution was dried over anhydrous sodium sulfate and concentrated to give the desired product (65 g, 99%) as an off-white solid, which was used in the subsequent reaction without further purification. C₉H₇BrFN₅O₂ (MW 316.09), LCMS (EI) *m/e* 316/318 (M⁺ + H).

Step D: 3-(4-Amino-1,2,5-oxadiazol-3-yl)-4-(3-bromo-4-fluorophenyl)-1,2,4-oxadiazol-5(4H)-one (5)



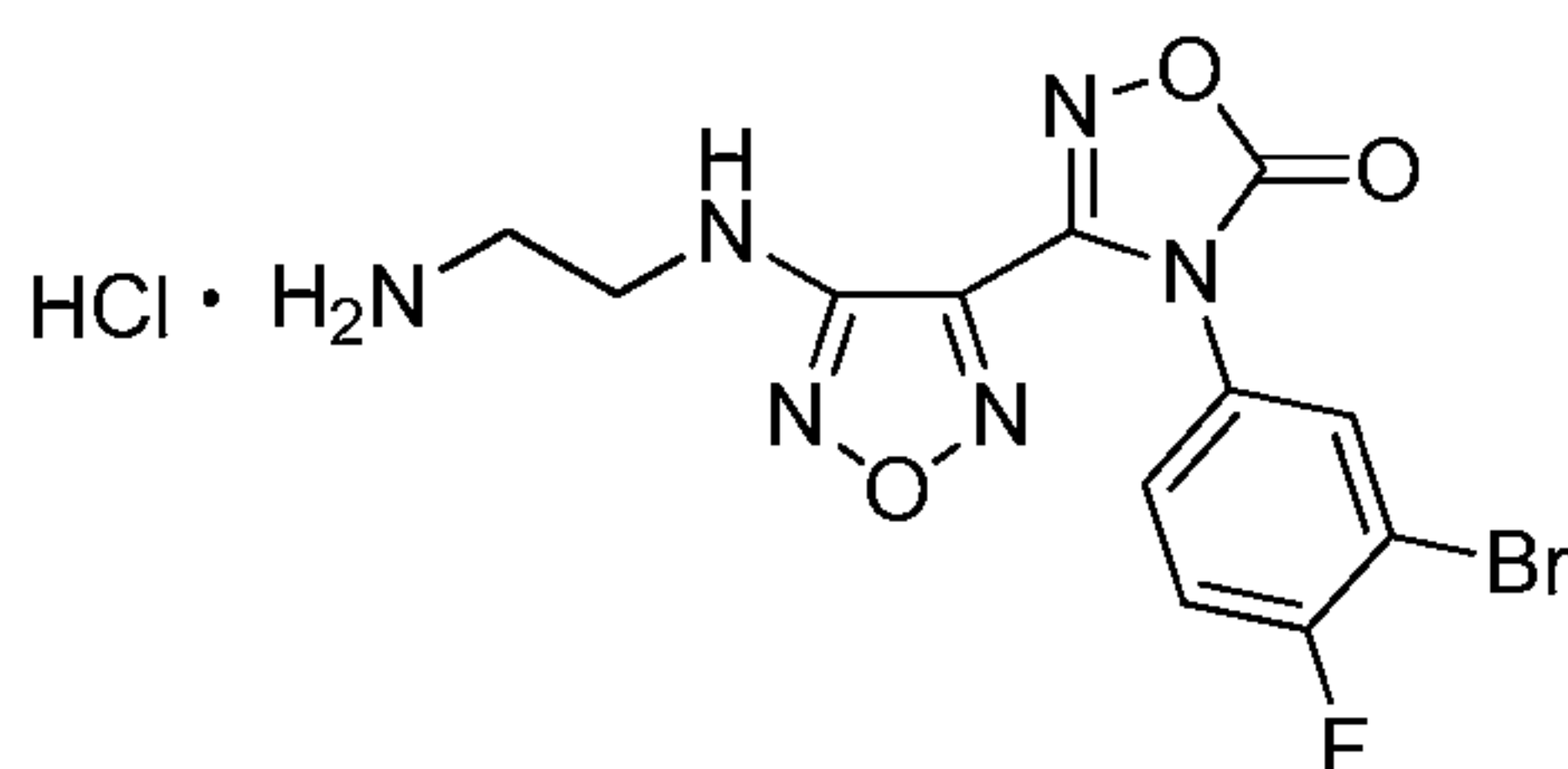
A mixture of 4-amino-N-(3-bromo-4-fluorophenyl)-N'-hydroxy-1,2,5-oxadiazole-3-carboximidamide (65.7 g, 208 mmol), *N,N*-carbonyldiimidazole (Sigma-Aldrich) (50.6 g, 312 mmol, 1.5 equiv.), and ethyl acetate (750 mL) was heated to 60 °C and stirred for 20 min. LCMS indicated reaction completed. The reaction was cooled to room temperature, washed with 1 N hydrochloric acid (2 x 750 mL), dried over sodium sulfate, and concentrated. The crude product was triturated with a mixture of dichloromethane, ethyl acetate, and diethyl ether to give the desired product (60.2 g, 85%) as an off-white solid. ¹H NMR (300 MHz, DMSO-*d*₆) δ 8.05 (m, 1H), 7.69 (m, 1H), 7.57 (t, 1H, *J* = 8.7 Hz), 6.58 (s, 2H); C₁₀H₅BrFN₅O₃ (MW 342.08), LCMS (EI) *m/e* 342/344 (M⁺ + H).

Step E: *tert*-Butyl [2-({4-[2-(3-bromo-4-fluorophenyl)-5-oxo-2,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]carbamate (7)



To a solution of trifluoroacetic acid (20.0 mL) and tetrahydrofuran (10.0 mL) was
 5 added sodium triacetoxyborohydride (10.59 g, 49.97 mmol, 10.0 equiv.) in portions with
 stirring under nitrogen. This mixture was stirred for 10 min at room temperature and then
 cooled to - 5 °C. A solution of 3-(4-amino-1,2,5-oxadiazol-3-yl)-4-(3-bromo-4-
 fluorophenyl)-1,2,4-oxadiazol-5(4*H*)-one (1.71 g, 5.0 mmol) and *tert*-butyl (2-
 oxoethyl)carbamate (Sigma-Aldrich) (1.99 g, 12.5 mmol, 2.5 equiv.) in THF (15.0 mL) was
 10 added drop-wise over 30 min with stirring while maintaining the temperature below 0 °C.
 The reaction was stirred at - 5 to 0 °C and additional portions of *tert*-butyl (2-
 oxoethyl)carbamate (0.20 g, 1.2 mmol, 0.24 equiv.) in THF (1.0 mL) were added drop-wise
 at 20 min, 40 min at 4 h intervals. HPLC indicated reaction completed after 5.25 h. The
 reaction mixture was poured into the ice-cold sodium bicarbonate (500 mL) and this solution
 15 was stirred at room temperature overnight. The precipitate was collected by filtration and
 washed with brine. The resulting residue was mixed with heptane (40 mL) and diethyl ether
 (40 mL) and stirred at room temperature for 5 h. The precipitate was collected by filtration,
 washed with diethyl ether and dried in a vacuum oven to give the desired product (1953 mg,
 80.5%) as an off-white solid. ¹H NMR (300 MHz, DMSO-*d*₆) δ 8.08 (m, 1H), 7.71 (m, 1H),
 20 7.59 (t, 1H, *J* = 8.7 Hz), 6.94 (m, 1H), 6.52 (m, 1H), 3.32 (m, 2H), 3.15 (m, 2H), 1.36 (s, 9H);
 C₁₇H₁₈BrFN₆O₅ (MW 485.26); LCMS (EI) *m/e* 507/509 (M⁺ + Na).

Step F: 3-{4-[(2-Aminoethyl)amino]-1,2,5-oxadiazol-3-yl}-4-(3-bromo-4-fluorophenyl)-
 1,2,4-oxadiazol-5(4*H*)-one hydrochloride (8)



Method A (prepared from tert-Butyl [2-(4-[2-(3-bromo-4-fluorophenyl)-5-oxo-2,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]carbamate; Step E):

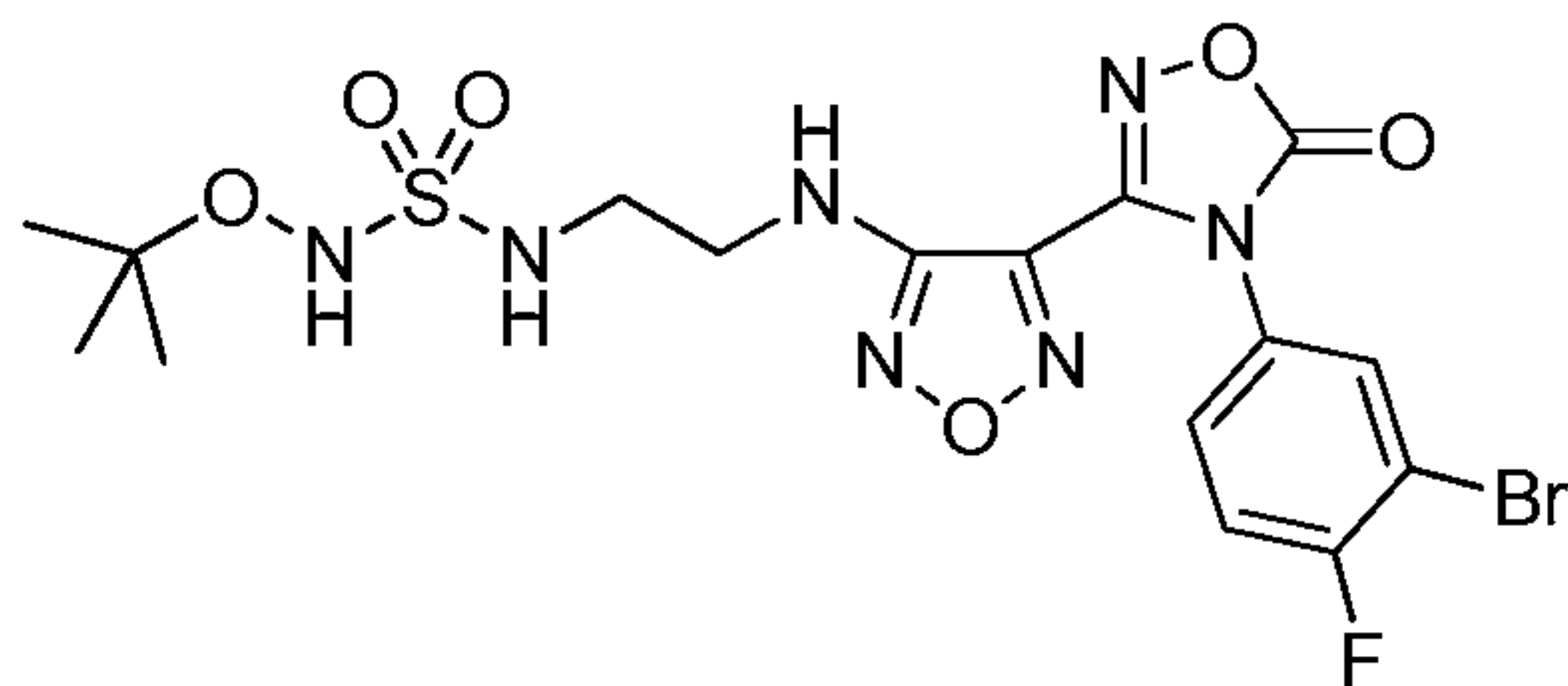
To a 500-mL flask was charged *tert*-butyl [2-(4-[2-(3-bromo-4-fluorophenyl)-5-oxo-2,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]carbamate (20 g, 41.2 mmol) and isopropanol (255 mL). The slurry was stirred at room temperature. Hydrogen chloride gas (7.55g, 207 mmol, 5.0 equiv.) was added to the slurry with a subsurface glass tube over 16 min. Ethyl acetate (111 mL) was then added to the batch and the reaction was heated to 43 °C and stirred for 7.5 h. The batch was cooled to 19 °C and ethyl acetate (44 mL) was added. The slurry was filtered and the resulting residue was washed with ethyl acetate (2 x 55 mL). The isolated solid was dried under reduced pressure at 45 °C for 15 h to afford the desired product (16.61 g, 95.5% yield) as an off-white to white solid. ¹H NMR (300 MHz, DMSO-*d*₆) δ 8.11 (bs, 3H), 7.78 (m, 1H), 7.73 (m, 1H), 7.59 (t, 1H, *J* = 8.7 Hz), 6.74 (t, 1H, *J* = 6.1 Hz), 3.50 (m, 2H), 3.02 (m, 2H); C₁₂H₁₁BrClFN₆O₃, (MW 421.61; C₁₂H₁₀BrFN₆O₃ for free base, MW 385.15), LCMS (EI) *m/e* 385/387 (*M*⁺ + H).

Method B (prepared directly from 3-(4-Amino-1,2,5-oxadiazol-3-yl)-4-(3-bromo-4-fluorophenyl)-1,2,4-oxadiazol-5(4H)-one; Step D):

Sodium triacetoxyborohydride (2.33 g, 11.0 mmol, 11.0 equiv.) was mixed with trifluoroacetic acid (12.0 mL, 155.8 mmol, 155.8 equiv.). The resulting solution was mixed at room temperature for 30 min. A solution of 3-(4-amino-1,2,5-oxadiazol-3-yl)-4-(3-bromo-4-fluorophenyl)-1,2,4-oxadiazol-5(4H)-one (**5**, 0.342 g, 1.0 mmol) and *tert*-butyl (2-oxoethyl)carbamate (Sigma-Aldrich) (1.04 g, 6.51 mmol, 6.5 equiv.) in dichloromethane (10.0 mL) and acetonitrile (6.0 mL) was stirred under N₂. This solution was cooled to -5 °C and the solution of sodium triacetoxyborohydride and trifluoroacetic acid was added drop-wise over 5 min. The reaction was stirred at room temperature for 4 h. HPLC and LC-MS (*M*⁺ - Boc + H: 385/387, bromide pattern) indicated a ratio of the desired product and the starting material was 4 to 1. The mixture was concentrated and diluted with dichloromethane (10 mL). The solution was cooled to 0 °C and 4 N sodium hydroxide was slowly added while maintaining the temperature at 0 – 5 °C to adjust the pH to 8 - 9. The aqueous layer was extracted with dichloromethane (3 x 10 mL). The combined dichloromethane solution was washed with sodium bicarbonate and brine, dried over sodium sulfate and concentrated. The crude residue was then dissolved in dichloromethane (6.0 mL) and the resulting solution was cooled to 0 °C. 4 N hydrochloric acid in dioxane (3.0 mL) was added drop-wise at 0 – 5 °C. The mixture was stirred at room temperature for 20 min. The precipitate was collected by

filtration, washed with diethyl ether, and dried in vacuum to afford the desired product (289 mg, 54%) as an off-white solid. ^1H NMR (300 MHz, DMSO- d_6) δ 8.11 (bs, 3H), 7.78 (m, 1H), 7.73 (m, 1H), 7.59 (t, 1H, $J = 8.7$ Hz), 6.74 (t, 1H, $J = 6.1$ Hz), 3.50 (m, 2H), 3.02 (m, 2H); $\text{C}_{12}\text{H}_{11}\text{BrClFN}_6\text{O}_3$, (MW 421.61; $\text{C}_{12}\text{H}_{10}\text{BrFN}_6\text{O}_3$ for free base, MW 385.15), LCMS (EI) m/e 385/387 ($\text{M}^+ + \text{H}$).

Step G: tert-Butyl ({[2-({4-[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]amino}sulfonyl)carbamate (9)



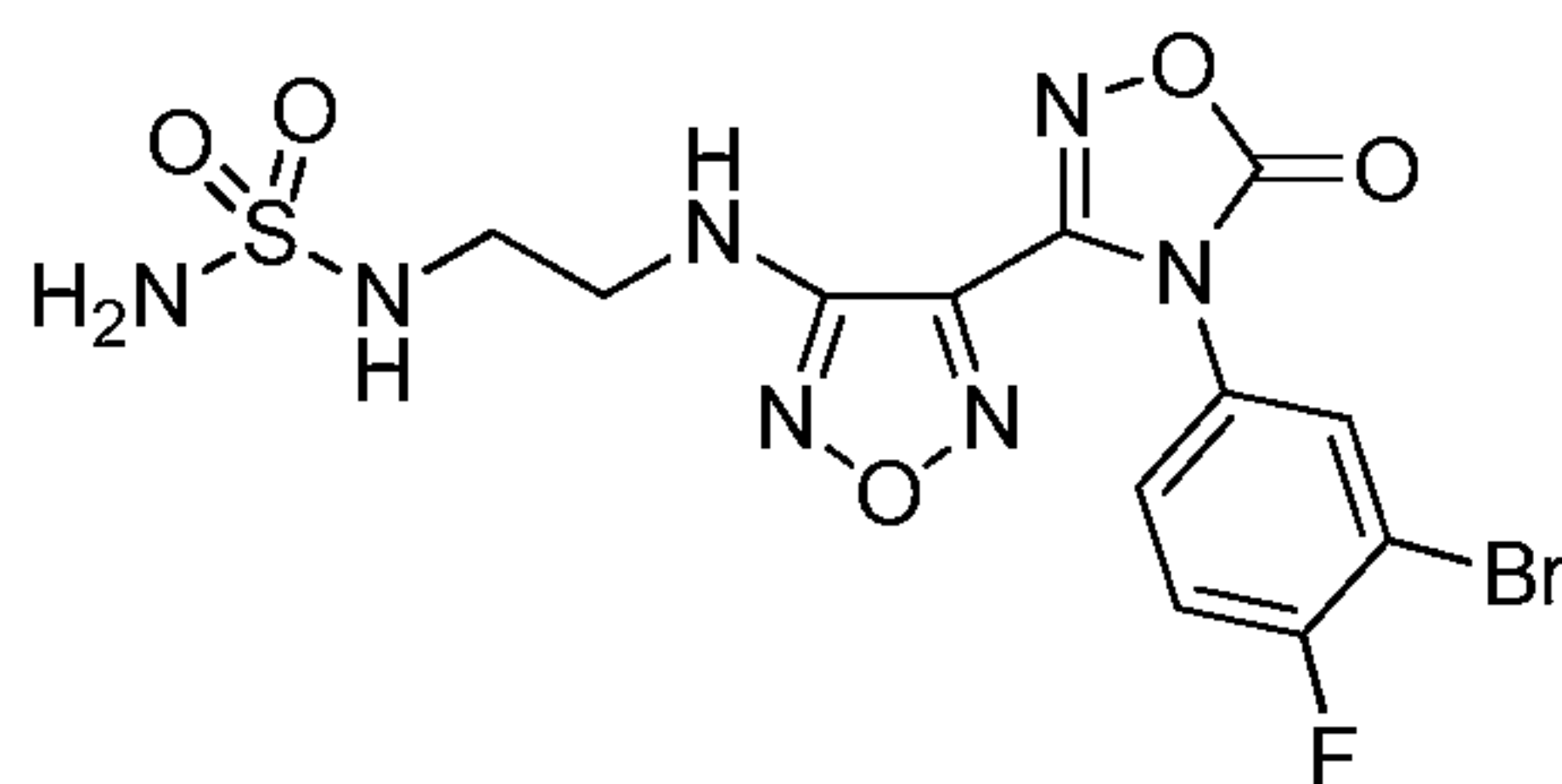
A 20-L glass reactor was charged with 3-{4-[(2-aminoethyl)amino]-1,2,5-oxadiazol-3-yl}-4-(3-bromo-4-fluorophenyl)-1,2,4-oxadiazol-5(4H)-one hydrochloride (1200 g, 2.846 mol) and dichloromethane (6.5 L) at room temperature. Triethylamine (950 g, 9.39 mol, 3.3 equiv.) was added to the batch over 7 minutes. The batch was then cooled to -14.6 $^{\circ}\text{C}$.

A 5-L round bottom flask was charged with *tert*-butanol (253 g, 3.41 mol, 1.2 equiv.) and dichloromethane (2.6 L). The solution was cooled to 0.9 $^{\circ}\text{C}$. To this solution was added chlorosulfonyl isocyanate (463 g, 3.27 mol, 1.15 equiv.) over 43 minutes while maintaining the batch temperature below 10 $^{\circ}\text{C}$. The resulting *tert*-butyl (chlorosulfonyl)carbamate solution was held at $3 - 5$ $^{\circ}\text{C}$ for 1 h.

The solution of *tert*-butyl (chlorosulfonyl)carbamate was added to the reactor over 73 min while maintaining the batch temperature below 0 $^{\circ}\text{C}$. The batch was then warmed to 10 $^{\circ}\text{C}$ over 1 h and was then stirred at $10 - 14$ $^{\circ}\text{C}$ for 1 h. Water (4.8 L) was added to the batch and the quenched reaction mixture was stirred at room temperature for 14.5 h. The batch was allowed to settle and phases separated. The dichloromethane solution was isolated kept in the reactor and was charged with acetic acid (513 g) over 25 min to precipitate the product. The resulting slurry was stirred at 20 $^{\circ}\text{C}$ for 2.5 h. The product was isolated by filtration and washed with dichloromethane (1.8 L). The product was dried under reduced pressure (-30 inHg) at 45 $^{\circ}\text{C}$ for 16 h to afford the desired product (1342 g, 83.5% yield) as a white solid. ^1H NMR (400 MHz, DMSO- d_6): δ 10.90 (s, 1 H), 8.08 (dd, $J = 6.2, 2.5$ Hz, 1 H), 7.72 (m, 1 H), 7.59 (t, $J = 8.6$ Hz, 1 H), 6.58 (t, $J = 5.7$ Hz, 1 H), 3.38 (dd, $J = 12.7, 6.2$ Hz, 2 H), 3.10

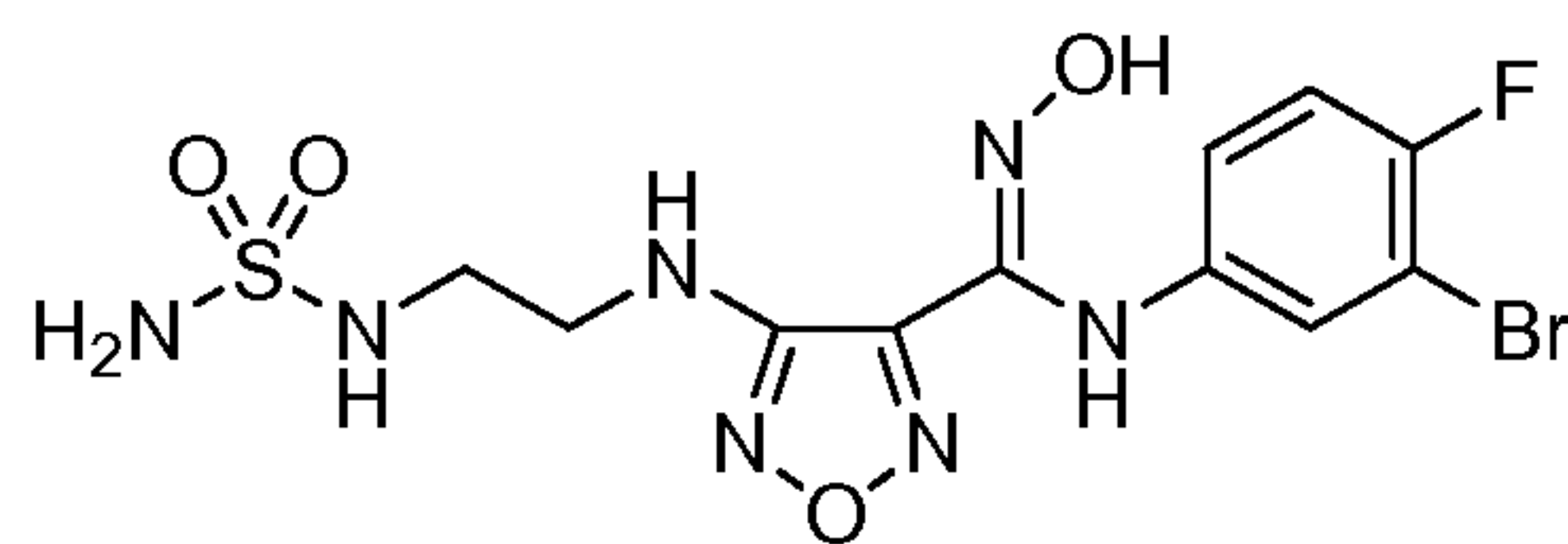
(dd, $J = 12.1, 5.9$ Hz, 2 H), 1.41 (s, 9 H). $C_{17}H_{19}BrFN_7O_7S$ (MW 564.34), LCMS (EI) m/e 585.9/587.9 ($M^+ + Na$).

Step H: *N*-[2-({4-[4-(3-Bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]sulfamide (10)



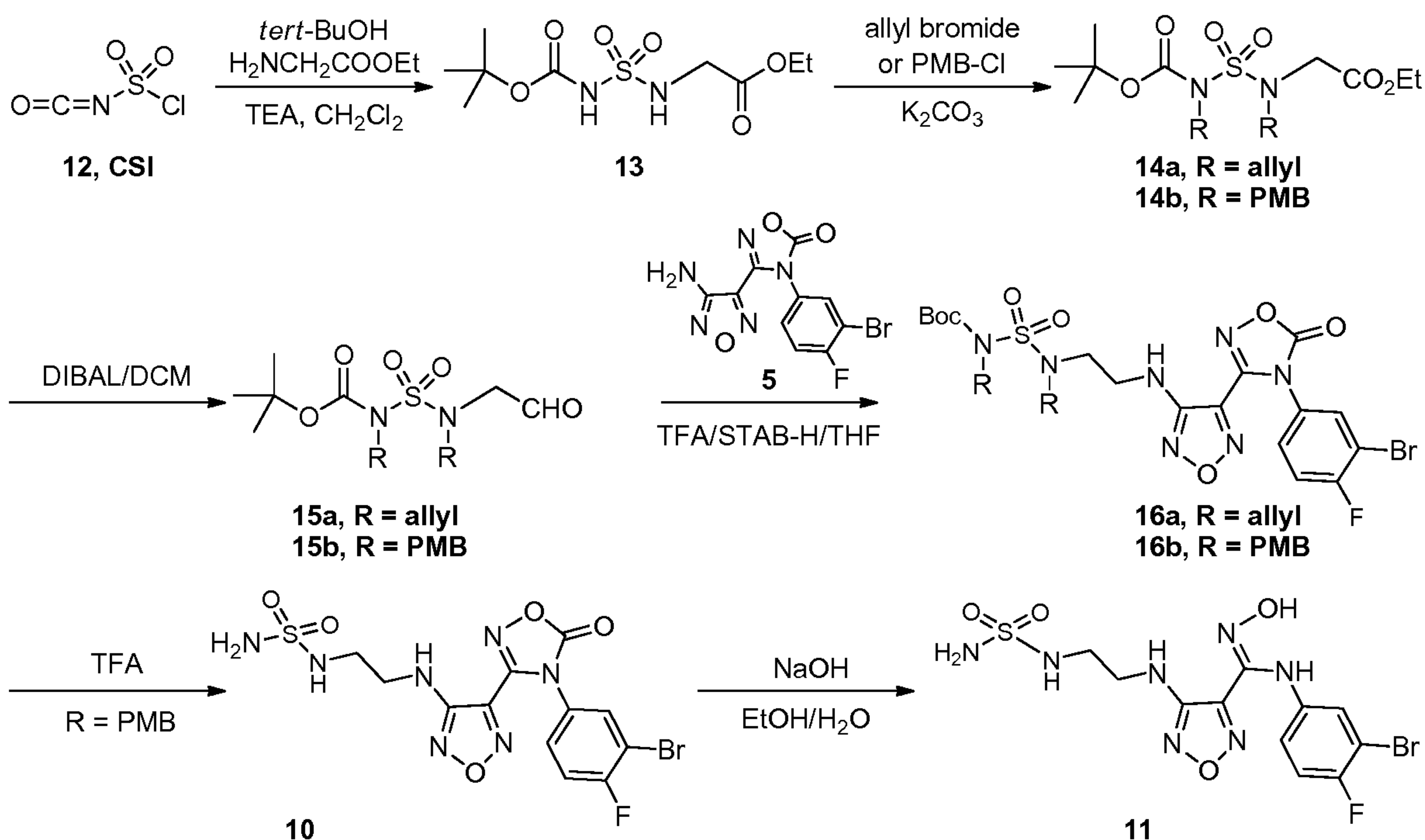
To a 20-L flask containing *tert*-butyl ({[2-({4-[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]amino}sulfonyl)carbamate (1200 g, 2.126 mol) was added ethanol (12 L) at 20 °C. The resulting mixture was stirred at room temperature and charged with hydrogen chloride gas (472 g, 12.9 mol, 6.07 equiv.). The batch was heated to 50 °C and the temperature was maintained for 3 h until reaction completion. Solvent was removed by vacuum distillation at 33 - 39 °C and 6 Kg of distillate was collected. Ethyl acetate (6.8 L, 6.1 Kg) was added to the batch and distilled to collect 5.1 Kg of distillate. Ethyl acetate (7.2 L, 6.48 Kg) was added to the batch and distilled to collect 5.1 Kg of distillate. Ethyl acetate (2.4 L, 2.14 Kg) was added to the batch to adjust the solvents ratio. ^1H NMR indicated the mole ratio of ethyl acetate to ethanol was 1.0: 0.1. The solution was heated to 65 °C. *n*-Heptane (4.1 kg) was added to the solution at 60 - 65 °C over 43 min. The resulting slurry was stirred at 65 °C for 1 h. The slurry was cooled to 20 °C over 2.5 h and stirred at that temperature for 15 h. The product was collected by filtration and washed with *n*-heptane (2.42 L). The product was dried under reduced pressure at 45 °C for 65 h to afford the desired product (906 g, 91.8% yield) as an off-white solid. ^1H NMR (400 MHz, DMSO- d_6) δ 8.08 (dd, $J = 6.2$) 7.72 (m, 1 H), 7.59 (t, $J = 8.7$ Hz, 1 H), 6.67 (t, $J = 5.9$ Hz, 1H), 6.55 (s, 2H) 6.52 (t, $J = 6.0$ Hz, 1 H), 3.38 (dd, $J = 12.7, 6.3$ Hz, 2 H), 3.11 (dd, $J = 12.3, 6.3$ Hz, 2H); $C_{12}H_{11}BrFN_7O_5S$ (MW 464.23), LCMS (EI) m/e 485.8/487.8 ($M^+ - C_5H_8O_2 + Na$).

Step I: 4-({2-[(Aminosulfonyl)amino]ethyl}amino)-*N*-(3-bromo-4-fluorophenyl)-*N'*-hydroxy-1,2,5-oxadiazole-3-carboximidamide (11)

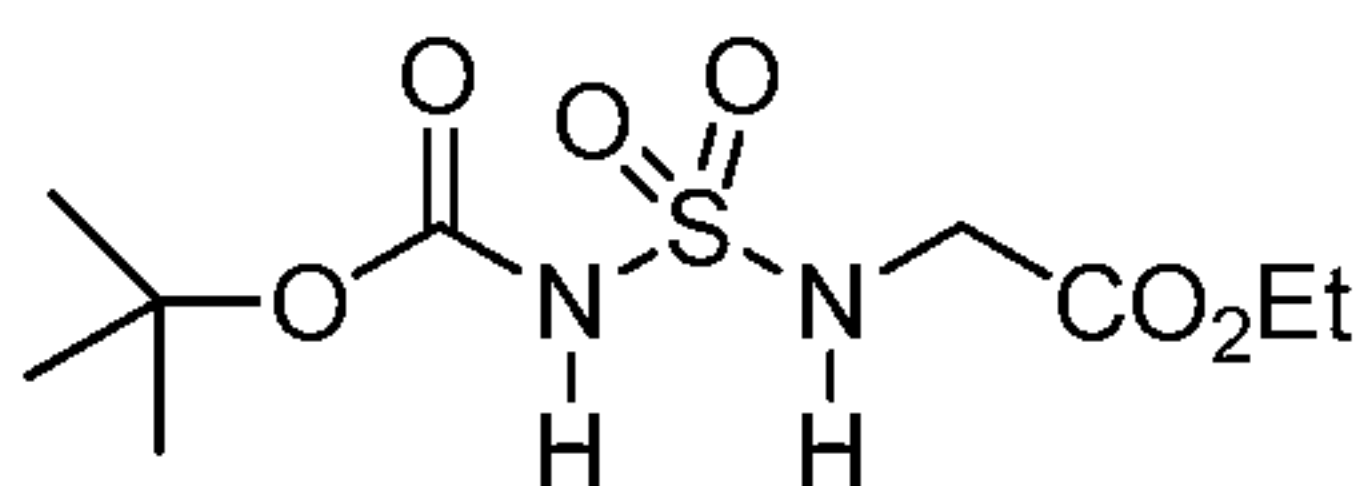


To a 20-L glass reactor was added *N*-[2-({4-[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]sulfamide (799.4 g, 1.72 mol) and THF (3.2 L). The resulting solution was stirred at 20 °C for 7 min. and then charged
 5 with water (1.6 L). The batch was cooled to 2 °C and was charged with 30 wt % sodium hydroxide solution (475 mL, 666.4 g, 4.99 mol, 2.9 equiv.) over 8 minutes. The batch was warmed to 20 °C and the temperature was maintained for 16 h. The batch was then charged with methyl *tert*-butyl ether (8.0 L) over 23 minutes. Water (2.7 L) was added and the batch was cooled to about 0 °C. The batch was then charged with 85 wt % phosphoric acid (370.7
 10 g, 3.22 mol, 1.9 equiv.) over 9 minutes. The batch was warmed to 20 °C and stirred for 1 h. The batch was allowed to settle and phases were separated. The organic layer was retained in the reactor and charged with water (2.9 L) and 85 wt% phosphoric acid (370.7g, 3.22 mol) and stirred at 20 °C for 1 h. The batch was allowed to settle and phases were separated. The organic layer was retained in the reactor and charged with water (3.2 L) and stirred at 20 °C
 15 for 1 h. The batch was allowed to settle and phases were separated. The organic solution was retained in the reactor and distilled under reduced pressure at 20 °C to remove 3.4 Kg of distillate. Ethanol (4.8 L) was charged to the batch and the batch was distilled to a volume of 3.2 L. This distillation process was repeated one more time. Ethanol (0.6 L) was added to the batch to adjust the batch volume to 4 L. The batch was stirred at 20 °C for 16 h and then
 20 charged with water (6.39 L). The resultant slurry was stirred at 20 °C for 5 h. The product was collected by filtration and was washed twice with a mixture of ethanol (529 mL) and water (1059 mL). The product was dried under reduced pressure at 45 °C for 65 h to afford the desired product (719.6 g, 95.4%) as a white solid. ¹H NMR (400 MHz, DMSO-*d*₆): δ 11.51 (s, 1 H), 8.90 (s, 1 H), 7.17 (t, *J* = 8.8 Hz, 1 H), 7.11 (dd, *J* = 6.1, 2.7 Hz, 1 H), 6.76 (m, 1 H), 6.71 (t, *J* = 6.0 Hz, 1 H), 6.59 (s, 2 H), 6.23 (t, *J* = 6.1 Hz, 1 H), 3.35 (dd, *J* = 10.9, 7.0 Hz, 2 H), 3.10 (dd, *J* = 12.1, 6.2 Hz, 2 H); C₁₁H₁₃BrFN₇O₄S (MW 438.23), LCMS (EI) *m/e* 437.9/439.9 (M⁺ + H).

Example 2. Alternate preparation of *N*-[2-({4-[4-(3-Bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]sulfamide

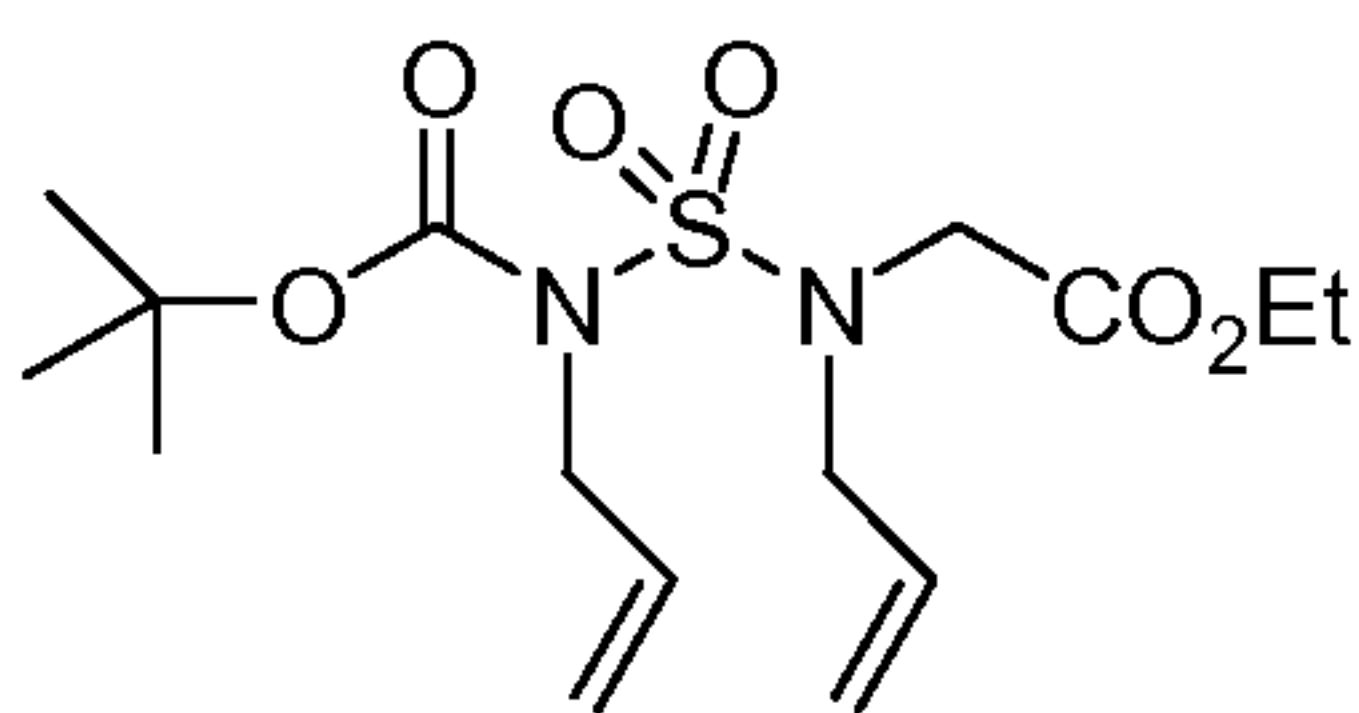


Step 1: Ethyl {[*tert*-butoxycarbonyl]-amino}sulfonyl}aminoacetate (13)



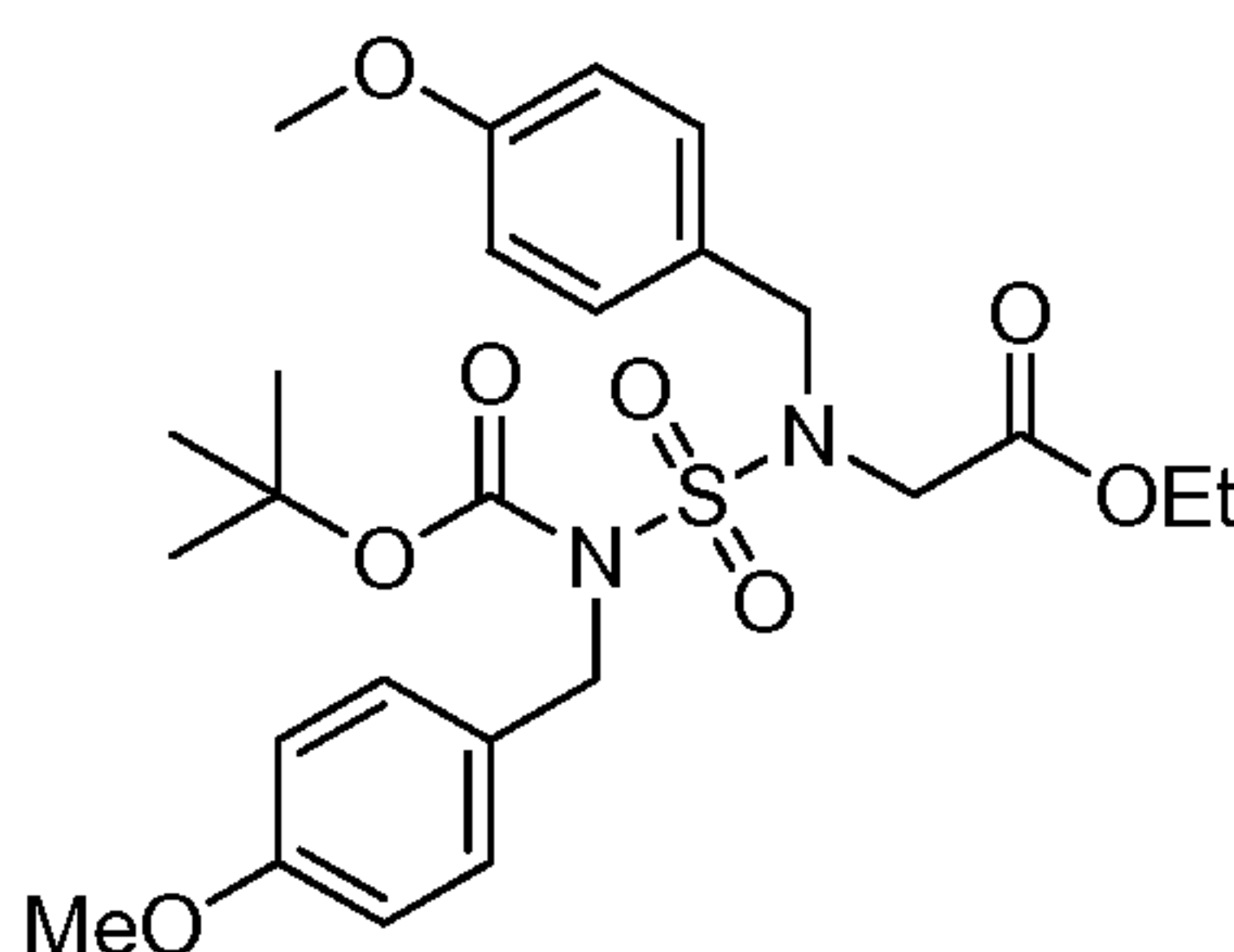
- 5 A solution of chlorosulfonylisocyanate (Sigma-Aldrich) (5.0 mL, 57.4 mmol) in dichloromethane (100 mL) was cooled to 0 °C. *Tert*-butyl alcohol (4.26 g, 57.4 mmol, 1.0 equiv.) was added dichloromethane (100 mL) via an addition funnel. This solution was stirred at 0 °C for 30 min. Glycine ethyl ester hydrochloride (8.82 g, 63.2 mmol, 1.1 equiv.) was added followed by drop-wise addition of triethylamine (20.0 mL, 144 mmol, 2.5 equiv.)
- 10 at 0 °C. This reaction mixture was stirred at room temperature for 4 h. The reaction was diluted with dichloromethane (100 mL) and washed with 0.1 N hydrochloric acid and brine. The organic layer was dried over sodium sulfate and concentrated to give the desired product (13.2 g, 81.4 %) as a crude off-white solid, which was used in the subsequent reaction without further purification. ¹H NMR (300 MHz, DMSO-*d*₆) δ 10.88 (s, 1H), 8.07 (t, 1H, *J* = 6.1 Hz), 4.08 (q, 2H, *J* = 7.1 Hz), 3.78 (d, 2H, *J* = 6.1 Hz), 1.40 (s, 9H), 1.18 (t, 3H, *J* = 7.1 Hz).
- 15

Step 2a. Ethyl (allyl{[allyl(*tert*-butoxycarbonyl)amino}sulfonyl}amino)acetate (14a)



Ethyl ({[(*tert*-butoxycarbonyl)amino]sulfonyl}aminoacetate (1.0 g, 3.54 mmol) was mixed with potassium carbonate (2.45 g, 17.7 mmol, 5.0 equiv.) and acetonitrile (23.0 mL) under N₂ at room temperature. Allyl bromide (1.84 mL, 21.2 mmol, 6.0 equiv.) was added drop-wise. This reaction mixture was heated to 70 °C and stirred at that temperature for 14 h. HPLC and LCMS indicated reaction completion. The reaction was filtered and the filtrate was concentrated. The residue was dissolved in dichloromethane and washed with sodium bicarbonate and brine. The organic layer was dried over sodium sulfate and concentrated to give the desired product (1.11 g, 87 %) as crude off-white solid, which was used in the subsequent reaction without further purification. ¹H NMR (300 MHz, DMSO-*d*₆) δ 5.75 (m, 2H), 5.20 (m, 4H), 4.12 (m, 6H), 3.89 (m, 2H), 1.43 (s, 9H), 1.18 (t, 3H, J = 8.7 Hz).

*Step 2b. Ethyl {[[(*tert*-butoxycarbonyl)(4-methoxybenzyl)amino]sulfonyl}(4-methoxybenzyl)amino]acetate (14b)*

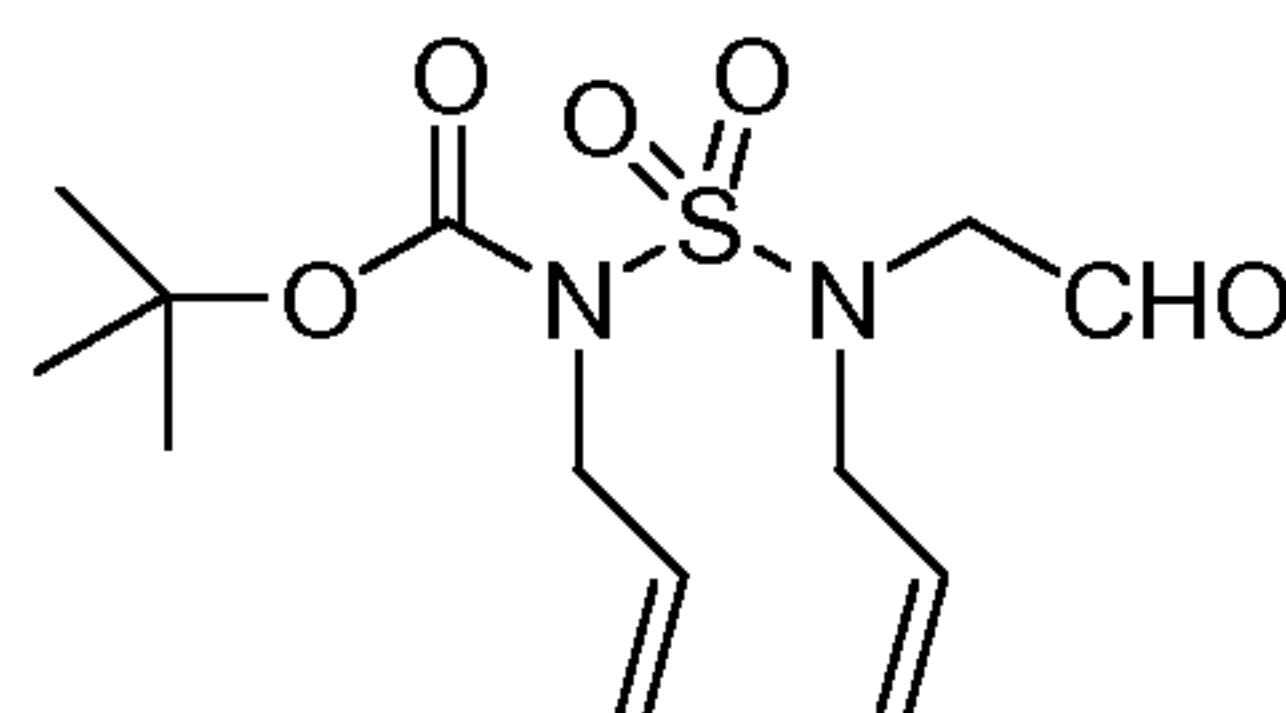


Ethyl ({[(*tert*-butoxycarbonyl) amino]sulfonyl}amino)acetate (1.00 g, 4.0 mmol) was mixed with *N,N*-dimethylformamide (DMF, 6.0 mL) and stirred at room temperature. Sodium iodide (0.01 g, 0.1 mmol, 0.025 equiv.), potassium carbonate (2.40 g, 20 mmol, 5.0 equiv.) and *para*-methoxybenzyl chloride (2.64 mL, 19.5 mmol, 4.875 equiv.) were added to the mixture. This reaction was warmed to 80 °C and stirred at 80 °C for 2 h. LCMS indicated reaction completion. The reaction was cooled to room temperature and filtered through Celite. The Celite bed was washed with dichloromethane and the combined organic filtrates were concentrated. The concentrated residue was dissolved in dichloromethane (20 mL) and washed with sodium bicarbonate (5 x 12 mL) and brine. The organic layer was dried over sodium sulfate and concentrated. The residue was purified on silica gel (0 – 40% ethyl

acetate/hexane gradient elution) to give the desired product (1.39 g, 80%) as an off-white solid. ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 7.22 (m, 2H), 7.14 (m, 2H), 6.88 (m, 4H), 4.64 (s, 2H), 4.33 (s, 2H), 4.03 (q, 2H, $J = 7.1$ Hz), 3.92 (s, 2H), 3.72 (s, 3H), 3.71 (s, 3H), 1.39 (s, 9H), 1.14 (t, 3H, $J = 7.1$ Hz).

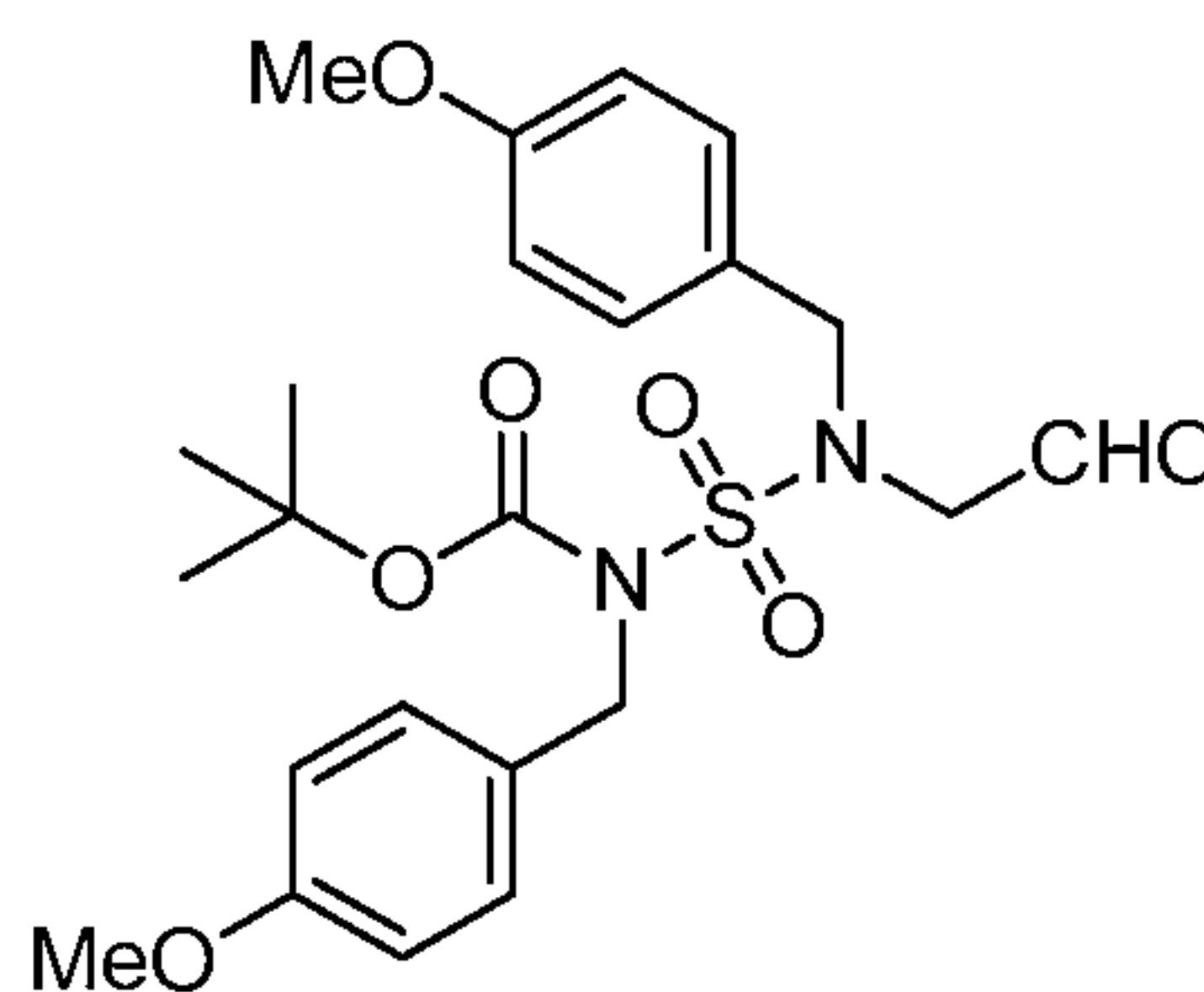
5

Step 3a. tert-Butyl allyl{[allyl(2-oxoethyl)amino]sulfonyl}carbamate (15a)



A solution of ethyl (allyl{[allyl(*tert*-butoxycarbonyl)amino]sulfonyl}amino)acetate (1.11 g, 3.05 mmol) in dichloromethane (15 mL) at -78°C under N_2 was treated with 1.0 M diisobutylaluminum hydride in dichloromethane (3.66 mL, 3.66 mmol, 1.2 equiv.). The reaction mixture was stirred at -78°C for 1 h and then quenched with methanol (1.5 mL) and treated with a saturated solution of sodium potassium tartrate (65 mL). This solution was stirred at room temperature overnight. The aqueous layer was then extracted with dichloromethane (3 x 20 mL). The combined dichloromethane solution was washed with brine, dried over sodium sulfate, filtered, and concentrated to give the desired product (0.62 g, 64 %) as a crude thick colorless oil, which was used in subsequent reaction without further purification. ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 9.45 (s, 1H), 5.76 (m, 2H), 5.18 (m, 4H), 4.15 (m, 4H), 3.72 (m, 2H), 1.43 (s, 9H).

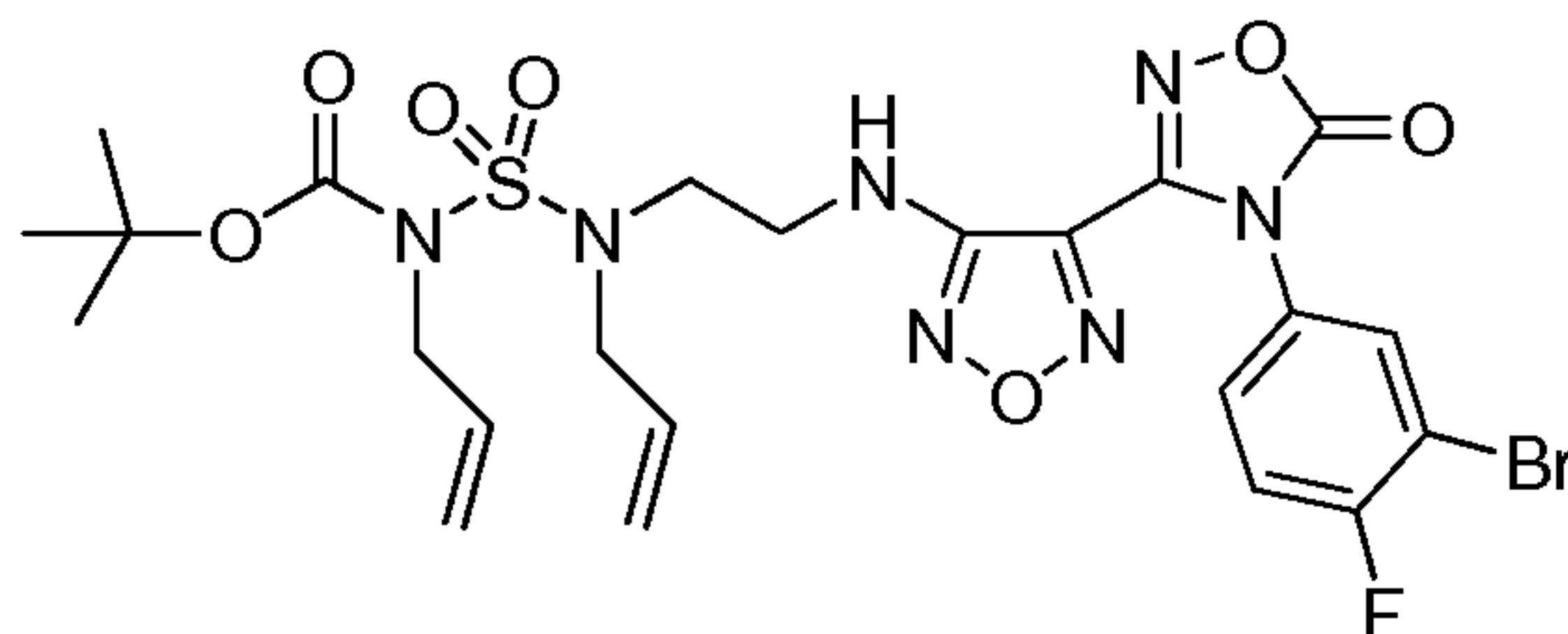
Step 3b. tert-Butyl (4-methoxybenzyl){[(4-methoxybenzyl)(2-oxoethyl)amino]sulfonyl}carbamate (15b)



A solution of ethyl [{[(*tert*-butoxycarbonyl)(4-methoxybenzyl)amino]sulfonyl} (4-methoxybenzyl)amino]acetate (5.30 g, 10 mmol) in dichloromethane (20.0 mL) at -78°C under N_2 was treated with 1.0 M diisobutylaluminum hydride in dichloromethane (12.2 mL,

12.2 mmol, 1.22 equiv.). The reaction mixture was stirred at -78 °C for 3 h. The reaction was then quenched with methanol (3 mL) and treated with dichloromethane (100 mL) and a saturated solution of sodium potassium tartrate (150 mL). This solution was stirred at room temperature overnight. The aqueous layer was then extracted with dichloromethane (3 x 20 mL). The combined dichloromethane solution was washed with brine, dried over sodium sulfate and concentrated. The residue was then purified on silica gel (0 – 30% ethyl acetate/hexane gradient elution) to give the desired product (3.45 g, 71%) as an off-white solid. ¹H NMR (300 MHz, DMSO-*d*₆) δ 9.24 (s, 1H), 7.23 (m, 4H), 6.88 (m, 4H), 4.68 (s, 2H), 4.31 (s, 2H), 4.07 (s, 2H), 3.72 (s, 3H), 3.71 (s, 3H), 1.40 (s, 9H).

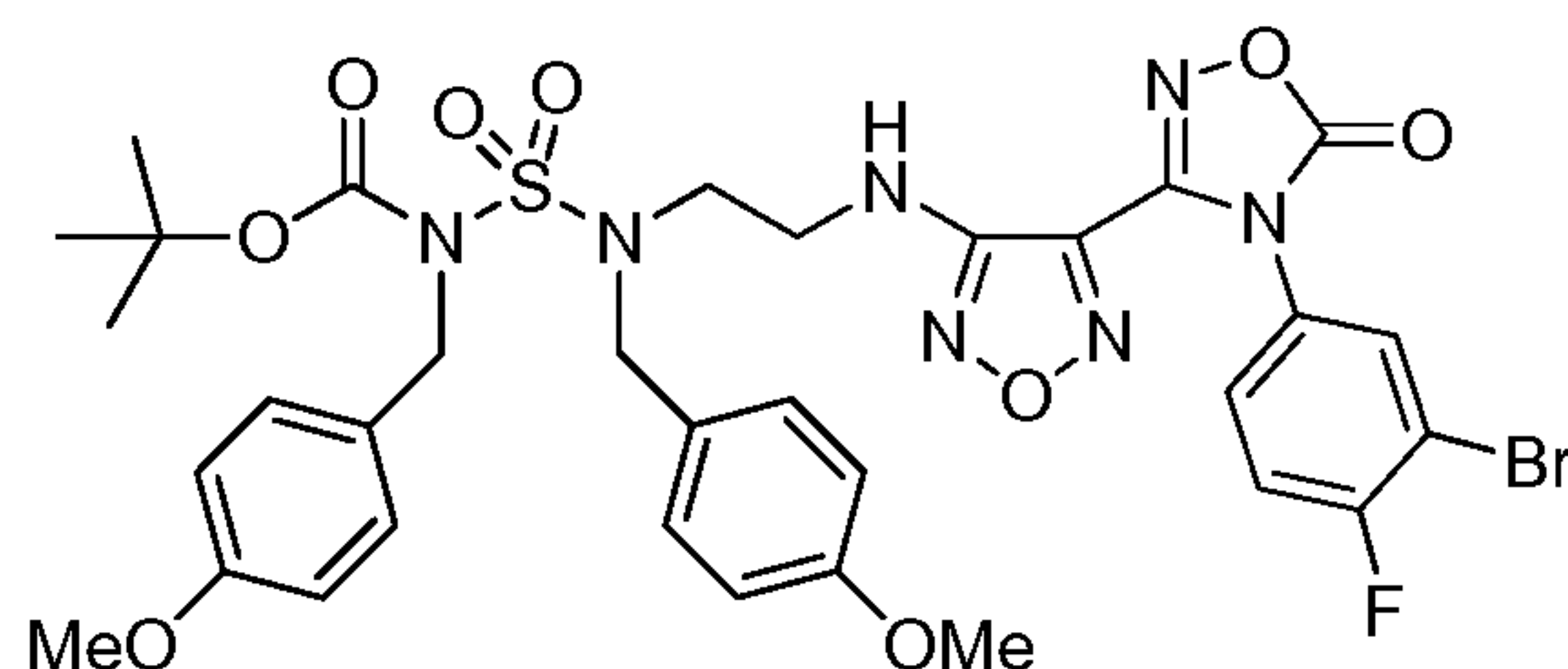
Step 4a. tert-Butyl allyl(N-allyl-N-(2-(4-(4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl)-1,2,5-oxadiazol-3-ylamino)ethyl)sulfamoyl)carbamate (16a)



To a 50-mL flask was added sodium triacetoxyborohydride (1.06 g, 5.0 mmol, 1.0 equiv.), trifluoroacetic acid (TFA, 2.0 mL, 26 mmol) and tetrahydrofuran (THF, 1.0 mL) at ambient temperature. This mixture was cooled to - 5 °C under N₂ and stirred at 0 - 5 °C for 10 min. To this solution was added 3-(4-amino-1,2,5-oxadiazol-3-yl)-4-(3-bromo-4-fluorophenyl)-1,2,4-oxadiazol-5(4*H*)-one (0.171 g, 5.0 mmol; Step D) and *tert*-butyl allyl{[allyl(2-oxoethyl)amino]sulfonyl}carbamate (0.398 g, 2.5 mmol, 0.5 equiv.) in THF (1.5 mL) drop-wise at 0 - 5 °C over 5 min. The resulting reaction mixture was stirred under N₂ at 0 - 5 °C. At 20 min, 40 min, and 2.5 h time points, a solution of *tert*-butyl allyl{[allyl(2-oxoethyl)amino]sulfonyl}carbamate (0.040 g, 0.125 mmol, 0.25 equiv.) in THF (0.20 mL) was added drop-wise at 0 - 5 °C. At 2.5 h, a solution of sodium triacetoxyborohydride (0.211 g, 1.0 mmol, 0.2 equiv.) in trifluoroacetic acid (TFA, 1.5 mL, 9.5 mmol) was added at 0 - 5 °C. The reaction was warmed to room temperature and stirred overnight. The reaction mixture was then poured into an ice-cold saturated solution of sodium carbonate (50 mL) and extracted with dichloromethane (3 x 20 mL). The combined dichloromethane extracts were washed with brine, dried over sodium sulfate, and concentrated. The residue was then purified on silica gel (0 – 75% ethyl acetate/hexane gradient elution) to give the desired product (0.239 g, 74.2%) as an off-white solid. ¹H NMR

(300 MHz, DMSO-*d*₆) δ 8.07 (m, 1H), 7.71 (m, 1H), 7.58 (t, 1H, *J* = 8.7 Hz), 6.62 (m, 1H), 5.77 (m, 2H), 5.19 (m, 4H), 4.17 (m, 2H), 3.89 (m, 2H), 3.44 (m, 2H), 3.38 (m, 2H), 1.42 (s, 9H); C₂₃H₂₇BrFN₇O₇S (MW 644.47), LCMS (EI) *m/e* 544/546 (M⁺ - Boc + H).

- 5 *Step 4b. tert-Butyl N-(2-(4-(4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl)-1,2,5-oxadiazol-3-ylamino)ethyl)-N-(4-methoxybenzyl)sulfamoyl(4-methoxybenzyl)carbamate (16b)*

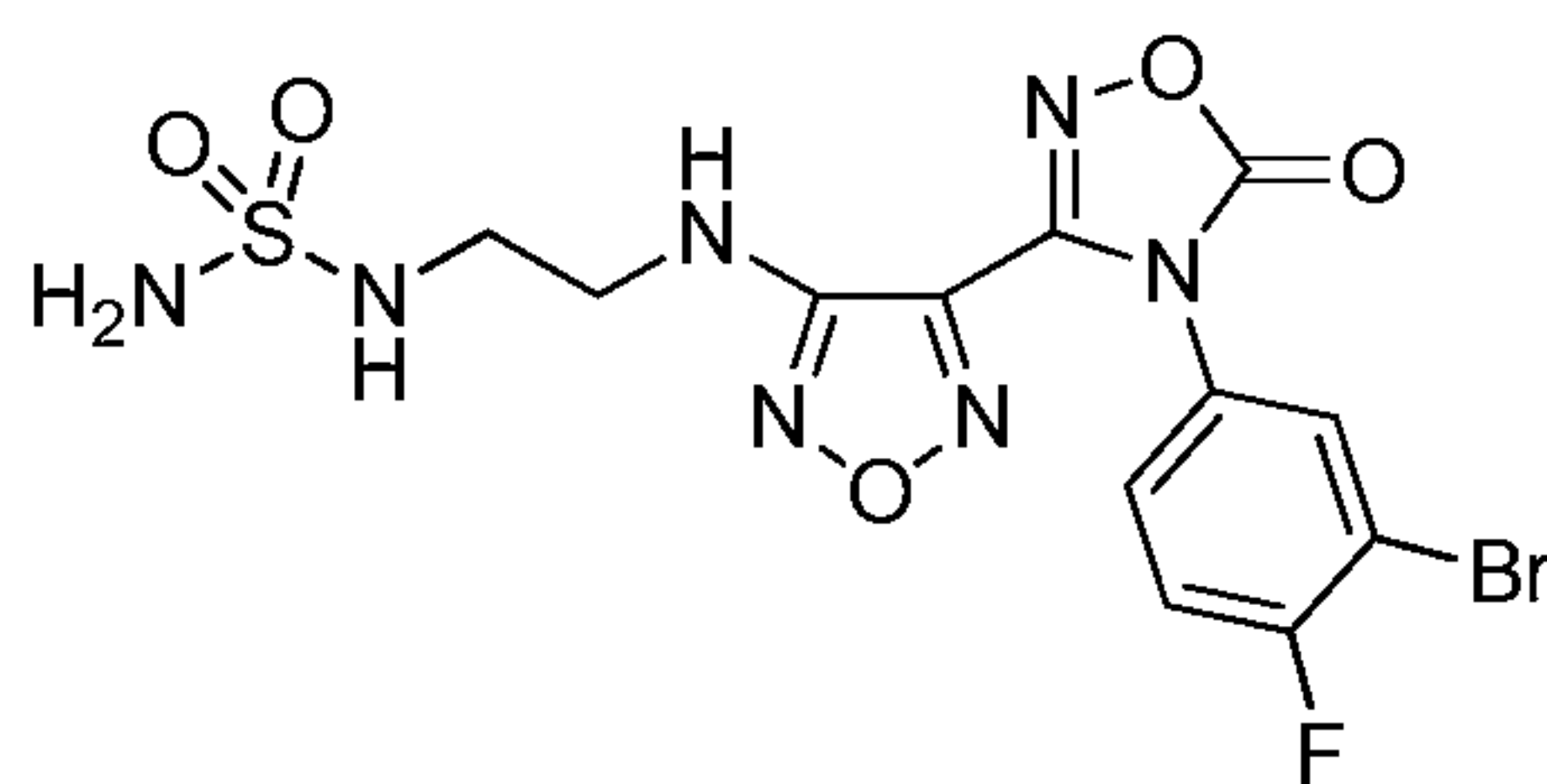


- To a 50-mL flask was added sodium triacetoxyborohydride (0.50 g, 2.37 mmol, 4.74 equiv.), trifluoroacetic acid (TFA, 1.0 mL, 13 mmol) and tetrahydrofuran at ambient temperature. This mixture was cooled to 0 - 5 °C under N₂ and stirred at 0 - 5 °C for 10 min. To this solution was added *tert*-butyl (4-methoxybenzyl){[(4-methoxybenzyl)(2-oxoethyl)amino]sulfonyl}carbamate (0.40 g, 0.84 mmol, 1.68 equiv) and 3-(4-amino-1,2,5-oxadiazol-3-yl)-4-(3-bromo-4-fluorophenyl)-1,2,4-oxadiazol-5(4*H*)-one (0.17 g, 0.50 mmol; Step D) in tetrahydrofuran (THF, 1.50 mL) at 0 - 5 °C. The reaction was stirred at 0 - 5 °C for 45 min and a solution of *tert*-butyl (4-methoxybenzyl){[(4-methoxybenzyl)(2-oxoethyl)amino]sulfonyl}carbamate (0.12 g, 0.20 mmol, 0.4 equiv.) in THF (0.50 mL) was then added at 0 - 5 °C. After stirring at 0 - 5 °C for 1 h, the reaction was gradually warmed to room temperature with stirring. At 2.5 h and 4.5 h time points, trifluoroacetic acid (0.25 mL) was added. At 5 h, a solution of *tert*-butyl (4-methoxybenzyl){[(4-methoxybenzyl)(2-oxoethyl)amino]sulfonyl}carbamate (0.060 g, 0.1 mmol, 0.2 equiv.) in THF (0.20 mL) was added. At 6.5 h, a solution of sodium triacetoxyborohydride (0.060 g, 0.24 mmol, 0.48 equiv.) in trifluoroacetic acid (0.25 mL) was added. HPLC indicated approximately 4% of the 3-(4-amino-1,2,5-oxadiazol-3-yl)-4-(3-bromo-4-fluorophenyl)-1,2,4-oxadiazol-5(4*H*)-one starting material (from Step D) still remaining. The reaction mixture was stirred at room temperature overnight. HPLC indicated reaction completion. The reaction mixture was poured into an ice-cold saturated solution of sodium carbonate (50 mL) and the mixture was extracted with dichloromethane (3 x 20 mL). The combined dichloromethane extracts were washed with brine, dried over sodium sulfate, and concentrated. The residue was then purified on silica gel (0 - 30% ethyl acetate/hexane gradient elution) to give the desired

product (0.33 g, 82.5%) as an off-white solid. ^1H NMR (300 MHz, DMSO- d_6) δ 8.06 (m, 1H), 7.69 (m, 1H), 7.57 (t, 1H, $J = 8.7$ Hz), 7.22 (m, 4H), 6.87 (m, 4H), 6.48 (m, 1H), 4.72 (s, 2H), 4.36 (s, 2H), 3.70 (s, 6H), 3.39 (m, 2H), 3.31 (m, 2H), 1.37 (s, 9H); $\text{C}_{33}\text{H}_{35}\text{BrFN}_7\text{O}_9\text{S}$ (MW 804.64), LCMS (EI) m/e 826/828 ($\text{M}^+ - \text{Boc} + \text{Na}$).

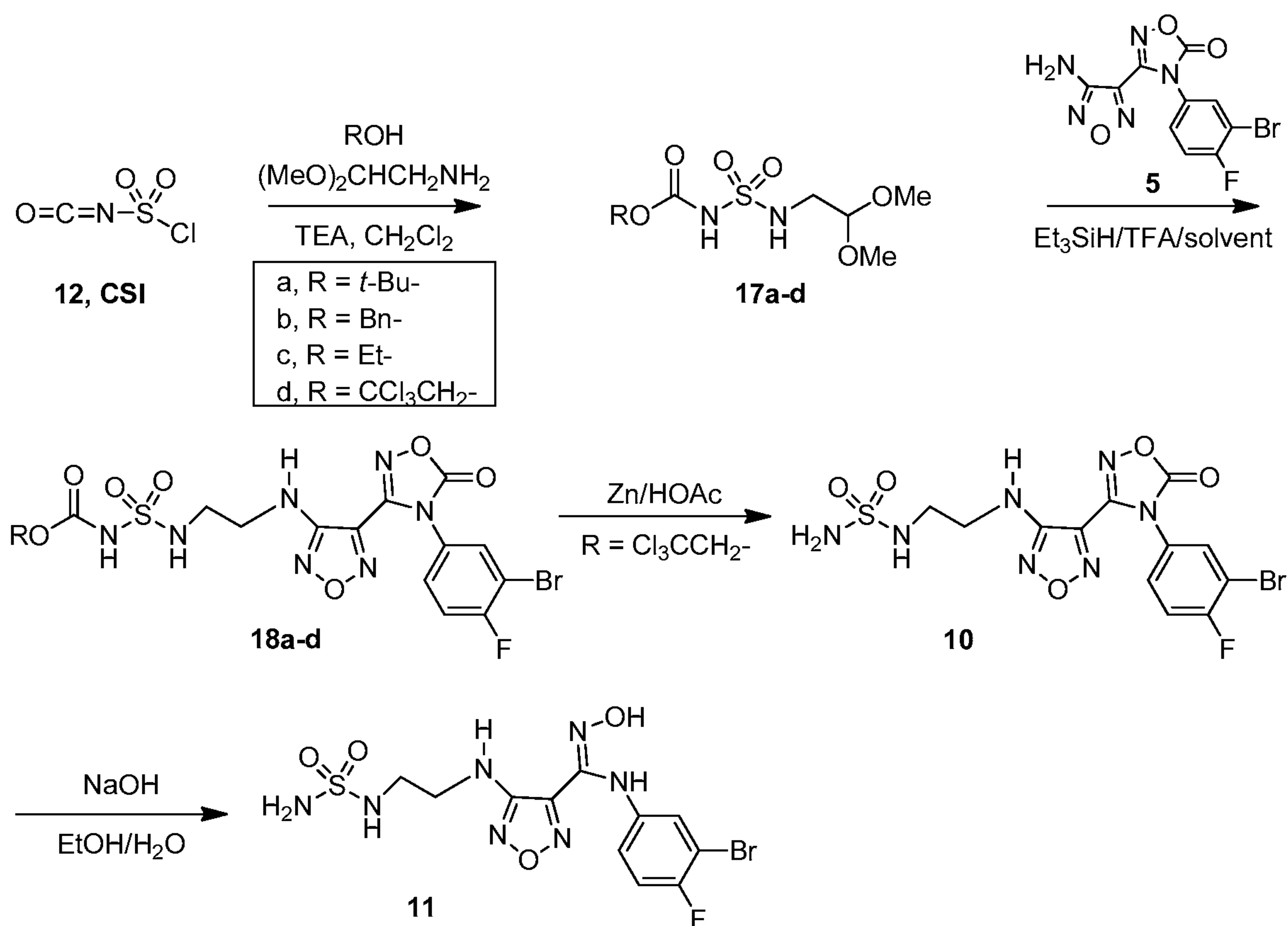
5

Step 5: *N*-[2-({4-[4-(3-Bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]sulfamide (10)

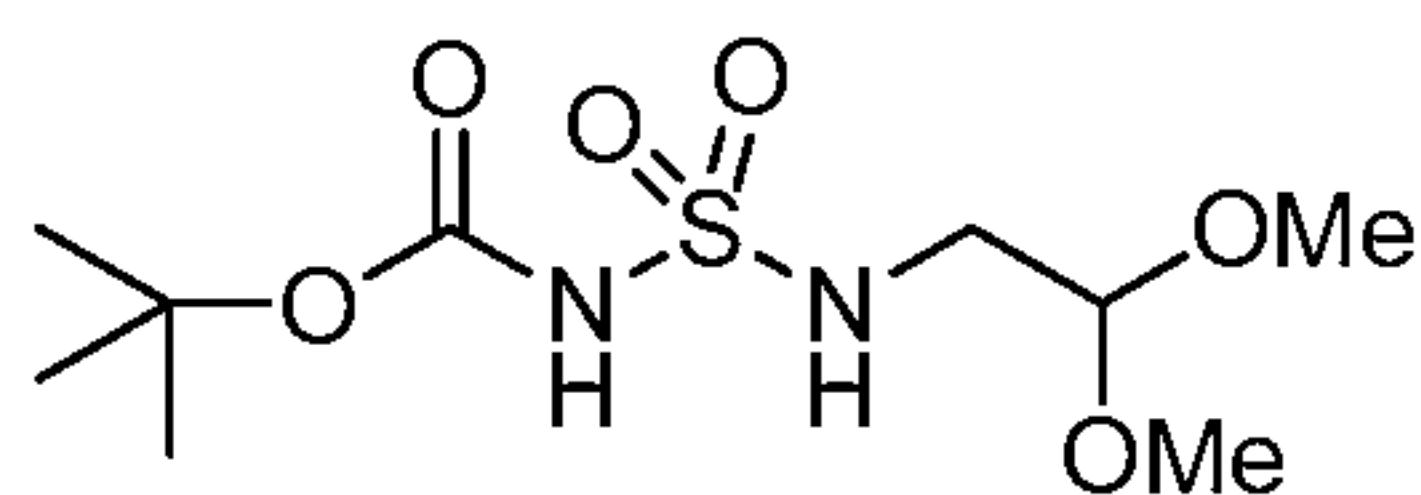


To a 25-mL flask was added *tert*-butyl {[2-({4-[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl](4-methoxybenzyl)amino]sulfonyl}(4-methoxybenzyl)carbamate (40.2 mg, 0.050 mmol) in trifluoroacetic acid (TFA, 0.50 mL, 6.5 mmol) at ambient temperature. This reaction mixture was heated to 70 °C under N_2 and stirred for 1 h. HPLC indicated reaction completed. The reaction mixture was cooled to room temperature and the TFA was evaporated. The residual TFA was removed by treatment with dichloromethane (3 x 10 mL) followed by evaporation in vacuum. The residue was then triturated with dichloromethane and methanol to give the desired product (20 mg, 87%) as a crude off-white solid. ^1H NMR (400 MHz, DMSO- d_6) δ 8.08 (dd, $J = 6.2, 2.5$ Hz, 1 H), 7.72 (m, 1 H), 7.59 (t, $J = 8.7$ Hz, 1 H), 6.67 (t, $J = 5.9$ Hz, 1H), 6.55 (s, 2H) 6.52 (t, $J = 6.0$ Hz, 1 H), 3.38 (dd, $J = 12.7, 6.3$ Hz, 2 H), 3.11 (dd, $J = 12.3, 6.3$ Hz, 2H); $\text{C}_{12}\text{H}_{11}\text{BrFN}_7\text{O}_5\text{S}$ (MW 464.23), LCMS (EI) m/e 487.8/489.8 ($\text{M}^+ + \text{Na}$).

Example 3. Alternate preparation of *N*-[2-({4-[4-(3-Bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]sulfamide

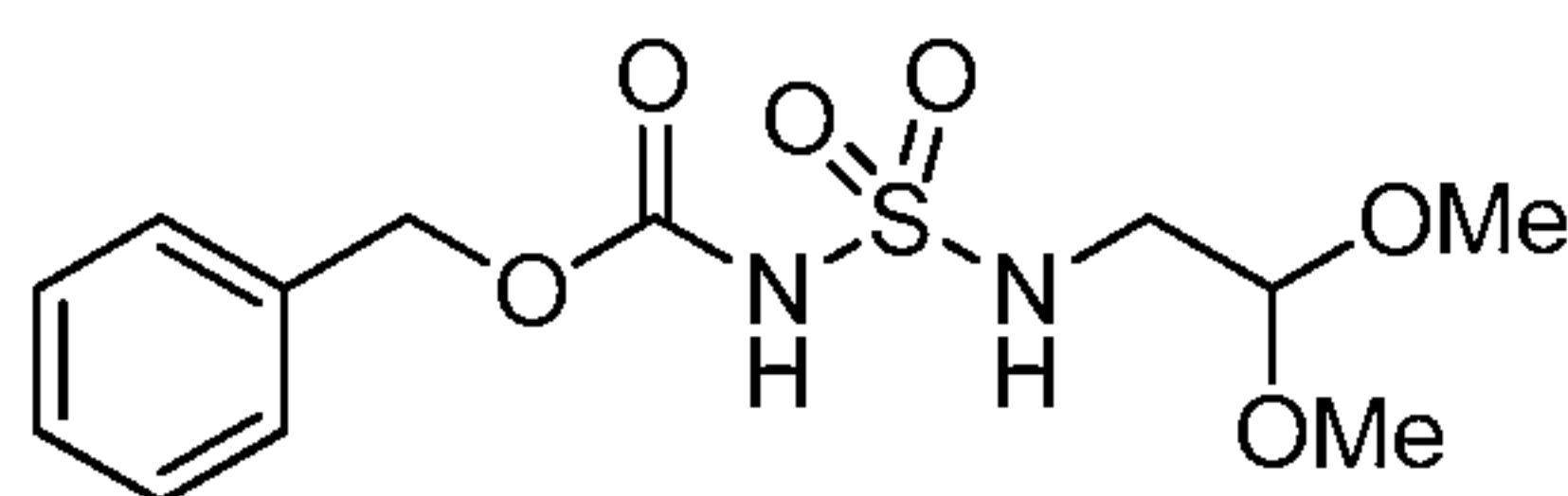


Step 1a. tert-Butyl N-(2,2-dimethoxyethyl)sulfamoylcarbamate (17a)



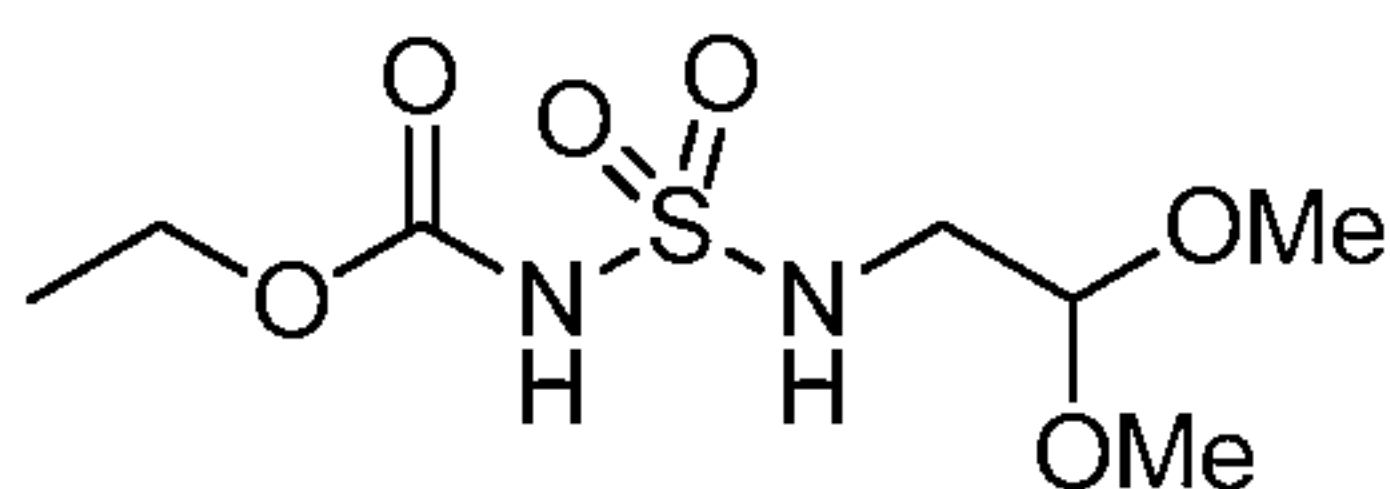
- 5 A solution of chlorosulfonylisocyanate (11.32 g, 80 mmol) in dichloromethane (120 mL) was cooled to 0 °C. *Tert*-butyl alcohol (7.65 mL, 80.0 mmol, 1.0 equiv.) was added via addition funnel. The mixture was stirred at 0 °C for 1.5 h. To this mixture, a solution of aminoacetaldehyde dimethyl acetal (8.76 mL, 80.0 mmol, 1.0 equiv.) and triethylamine (TEA, 33.4 mL, 240 mmol, 3.0 equiv.) in methylene chloride (DCM 120.0 mL) was added
- 10 drop-wise via addition funnel. The reaction was warmed to room temperature and stirred overnight. The reaction was treated with 0.1 N hydrochloric acid and the organic layer was washed with brine, dried over sodium sulfate and concentrated to give the desired product (15.6 g, 68.5 %) as a crude off-white solid, which was used for the subsequent reaction without further purification: ¹H NMR (300 MHz, DMSO-*d*₆) δ 10.84 (s, 1H), 7.62 (t, 1H, *J* = 6.0 Hz), 4.38 (t, 1H, *J* = 5.5 Hz), 3.24 (s, 6H), 2.96 (dd, 2H, *J* = 5.8 Hz), 1.41(s, 9H).
- 15

Step 1b. Benzyl N-(2,2-dimethoxyethyl)sulfamoylcarbamate (17b)



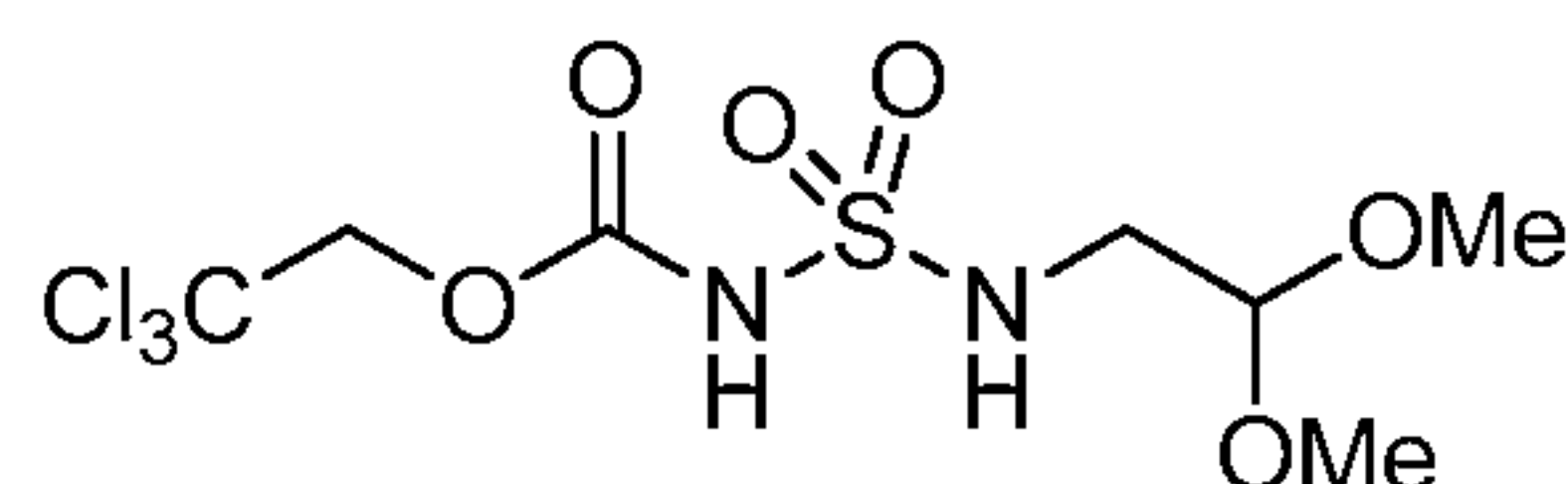
A solution of chlorosulfonylisocyanate (16.26 g, 114.9 mmol) in dichloromethane (100 mL) was cooled to 0 °C. Benzyl alcohol (12.44g, 115.0 mmol, 1.0 equiv.) was added via an addition funnel. The mixture was stirred at 0 °C for 0.5 h. To this mixture was added a mixture of aminoacetaldehyde dimethyl acetal (13.25 g, 126.0 mmol, 1.1 equiv.) and triethylamine (TEA, 17.4 g, 172 mmol, 1.5 equiv.) dropwise via an addition funnel at below 15 °C. The reaction was warmed to room temperature and stirred overnight. The reaction mixture was treated with 0.5 N hydrochloric acid (100 mL) and the collected organic phase was washed with brine, dried over sodium sulfate and concentrated in vacuum to give the desired product (23.5 g, 64.3 %) as a crude off-white solid. ¹H NMR (300 MHz, DMSO-*d*₆) δ 11.29 (s, 1H), 7.90 (t, 1H, *J* = 6.0 Hz), 7.37 (m, 5H), 5.12 (s, 2H), 4.35 (t, 1H, *J* = 5.5 Hz), 3.21 (s, 6H), 2.97 (dd, 2H, *J* = 5.8 Hz).

Step 1c. Ethyl N-(2,2-dimethoxyethyl)sulfamoylcarbamate (17c)



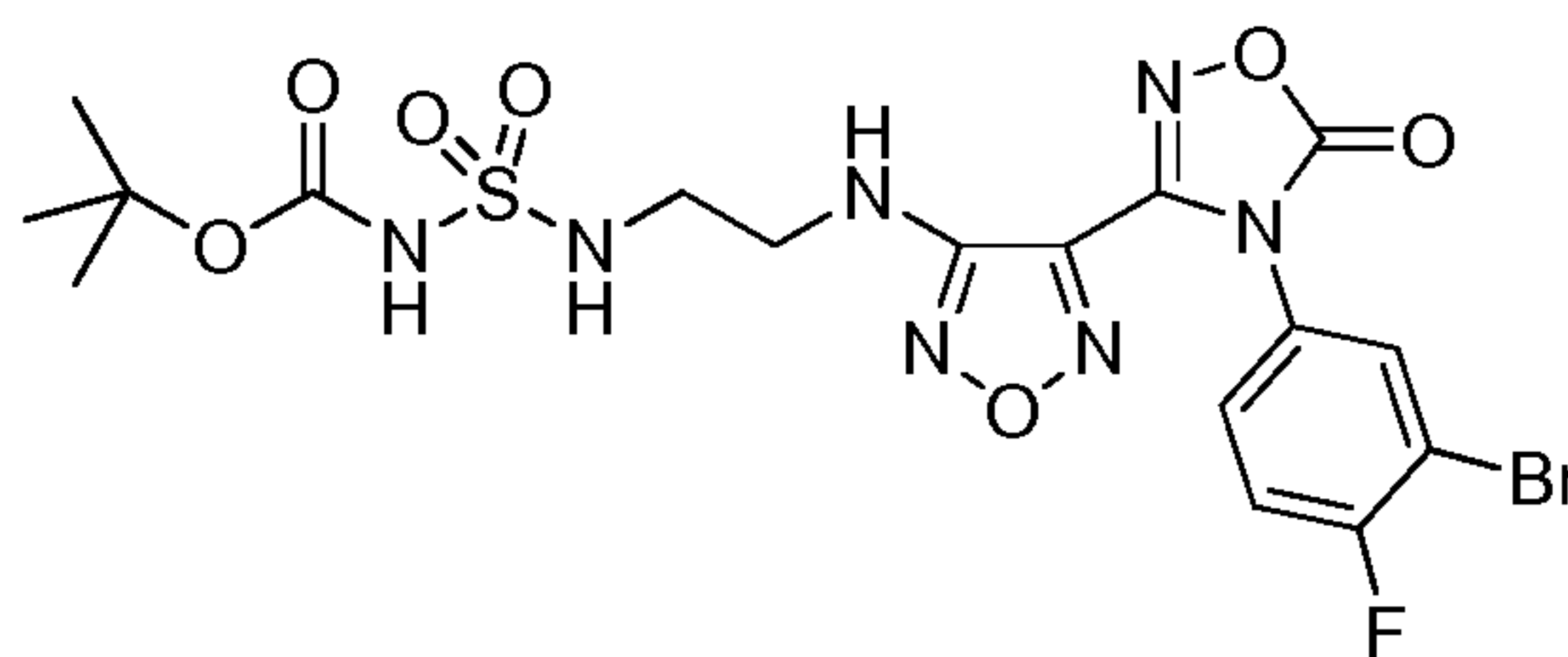
A solution of chlorosulfonylisocyanate (11.32 g, 80 mmol) in dichloromethane (120 mL) was cooled to 0 °C. Ethanol (4.67 mL, 80.0 mmol, 1.0 equiv.) was added via addition funnel. The mixture was stirred at 0 °C for 1.5 h. To this mixture was added a solution of aminoacetaldehyde dimethyl acetal (8.76 mL, 80.0 mmol, 1.0 equiv.), triethylamine (TEA, 33.4 mL, 240 mmol, 3.0 equiv.) in dichloromethane (DCM, 120.0 mL) drop-wise via addition funnel 0 °C. The reaction was warmed to room temperature and stirred overnight. The reaction was treated with 0.1 N hydrochloric acid and the collected organic phase was washed with brine, dried over sodium sulfate and concentrated in vacuum to give the desired product (11.2 g, 55 %) as a crude off-white solid. ¹H NMR (300 MHz, DMSO-*d*₆) δ 11.13 (s, 1H), 7.81 (t, 1H, *J* = 6.0 Hz), 4.37 (t, 1H, *J* = 5.5 Hz), 4.09 (q, 2H, *J* = 7.1 Hz), 3.23 (s, 6H), 2.97 (dd, 2H, *J* = 5.8 Hz), 1.19 (t, 3H, *J* = 7.1 Hz).

Step 1d. 2,2,2-Trichloroethyl N-(2,2-dimethoxyethyl)sulfamoylcarbamate (17d)



A solution of chlorosulfonylisocyanate (6.96 mL, 80 mmol) in dichloromethane (120 mL) was cooled to 0 °C. 2,2,2-trichloroethanol (7.67 mL, 80.0 mmol, 1.0 equiv.) was added via addition funnel at 0 °C. This mixture was stirred at 0 °C for 1.5 h. To this mixture was
 5 then added a solution of aminoacetaldehyde dimethyl acetal (8.76 mL, 80.0 mmol, 1.0 equiv.) and triethylamine (TEA, 33.4 mL, 240 mmol, 3.0 equiv.) in dichloromethane (DCM, 120.0 mL) added drop-wise via addition funnel at 0 °C. The reaction was warmed to room temperature and stirred at room temperature overnight. The reaction was treated with 0.1 N hydrochloric acid and the collected organic phase was washed with brine, dried over Na₂SO₄,
 10 and concentrated to give the desired product (28.01 g, 97 %) as a crude off-white solid, which was used in the subsequent reaction without further purification. ¹H NMR (300 MHz, DMSO-*d*₆) δ 11.79 (s, 1H), 8.08 (t, 1H, *J* = 5.9 Hz), 4.90 (s, 2H), 4.37 (t, 1H, *J* = 5.5 Hz), 3.23 (s, 6H), 3.00 (dd, 2H, *J* = 5.7 Hz).

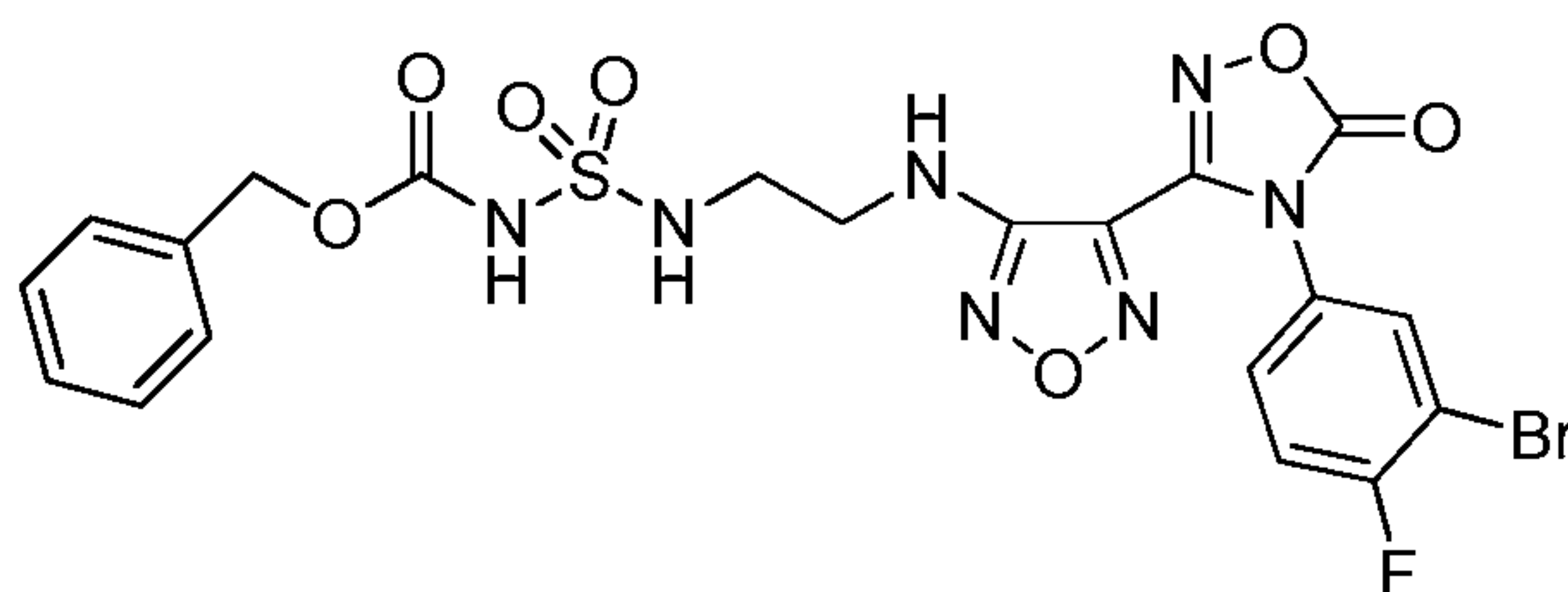
Step 2a. Tert-butyl ({[2-({[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]amino}sulfonyl)carbamate (18a)
 15



A mixture of 3-(4-amino-1,2,5-oxadiazol-3-yl)-4-(3-bromo-4-fluorophenyl)-1,2,4-oxadiazol-5(4*H*)-one (103 mg, 0.302 mmol, 1.5 equiv.; Step D) and *tert*-butyl *N*-(2,2-dimethoxyethyl)sulfamoylcarbamate (57.2 mg, 0.201 mmol) in dichloromethane (1.0
 20 mL) was stirred under N₂ at room temperature. To this mixture was added trifluoroacetic acid (0.50 mL, 6.5 mmol) and triethylsilane (80.2 μL, 0.502 mmol, 2.5 equiv.) drop-wise. This reaction mixture was stirred at room temperature for 2 h. HPLC indicated approximately 30% conversion. The reaction mixture was cooled to 0 °C and quenched with saturated sodium bicarbonate to pH ~ 8. The mixture was extracted in ethyl acetate (3 x 10
 25 mL). The combined organic extracts were washed with brine, dried over sodium sulfate and concentrated. The residue was purified by preparative TLC (50% ethyl acetate/hexane) to

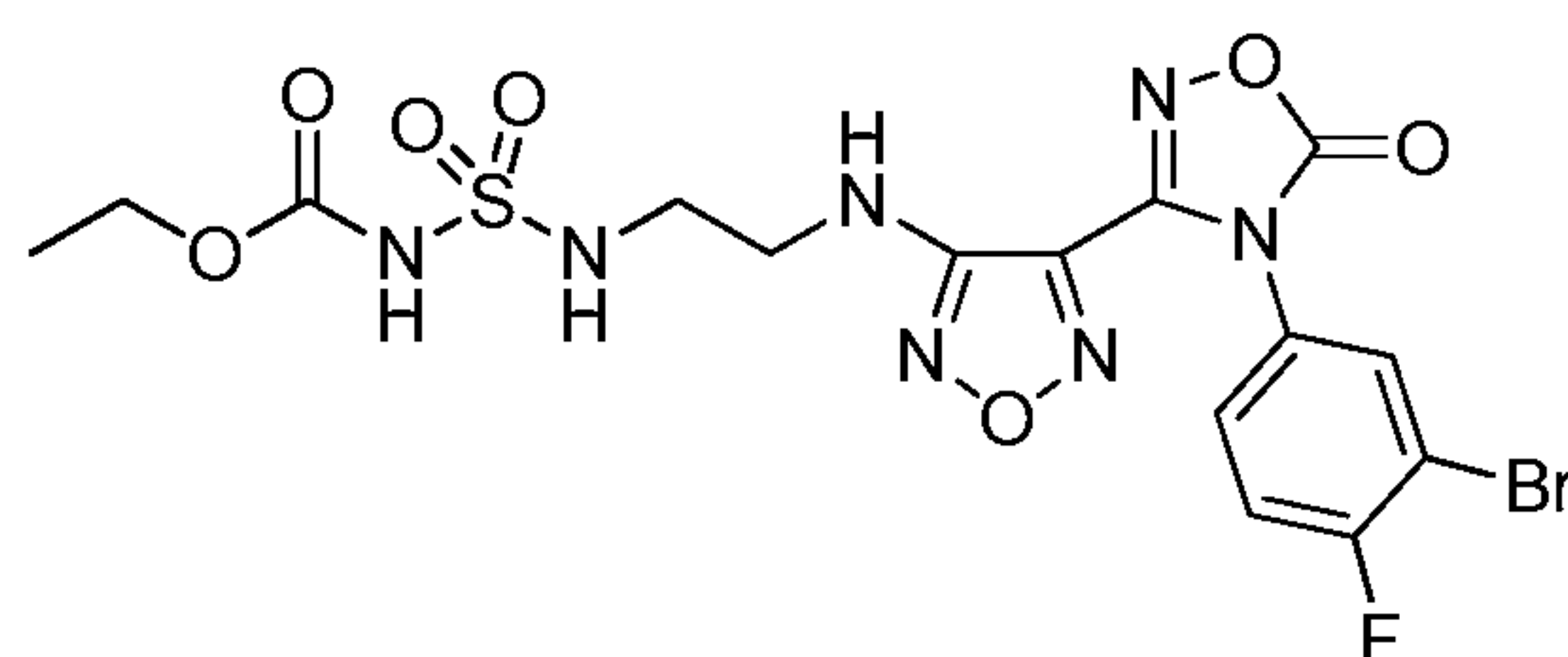
give the desired product (27.5 mg, 29.5%) as an off-white solid. ¹H NMR (DMSO-d₆, 400 MHz): δ 10.90 (s, 1H), 8.08 (dd, *J* = 6.2, 2.5 Hz, 1H), 7.72 (m, 1H), 7.59 (t, *J* = 8.6 Hz, 1H), 6.58 (t, *J* = 5.7 Hz, 1H), 3.38 (dd, *J* = 12.7, 6.2 Hz, 2H), 3.10 (dd, *J* = 12.1, 5.9 Hz, 2H), 1.41 (s, 9H). C₁₇H₁₉BrFN₇O₇S (MW 564.34), LCMS (EI) *m/e* 485.8/487.8 (M⁺ - C₅H₈O₂ + Na).

5 *Step 2b. Benzyl* ({[2-({[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]amino}sulfonyl)carbamate (18b)



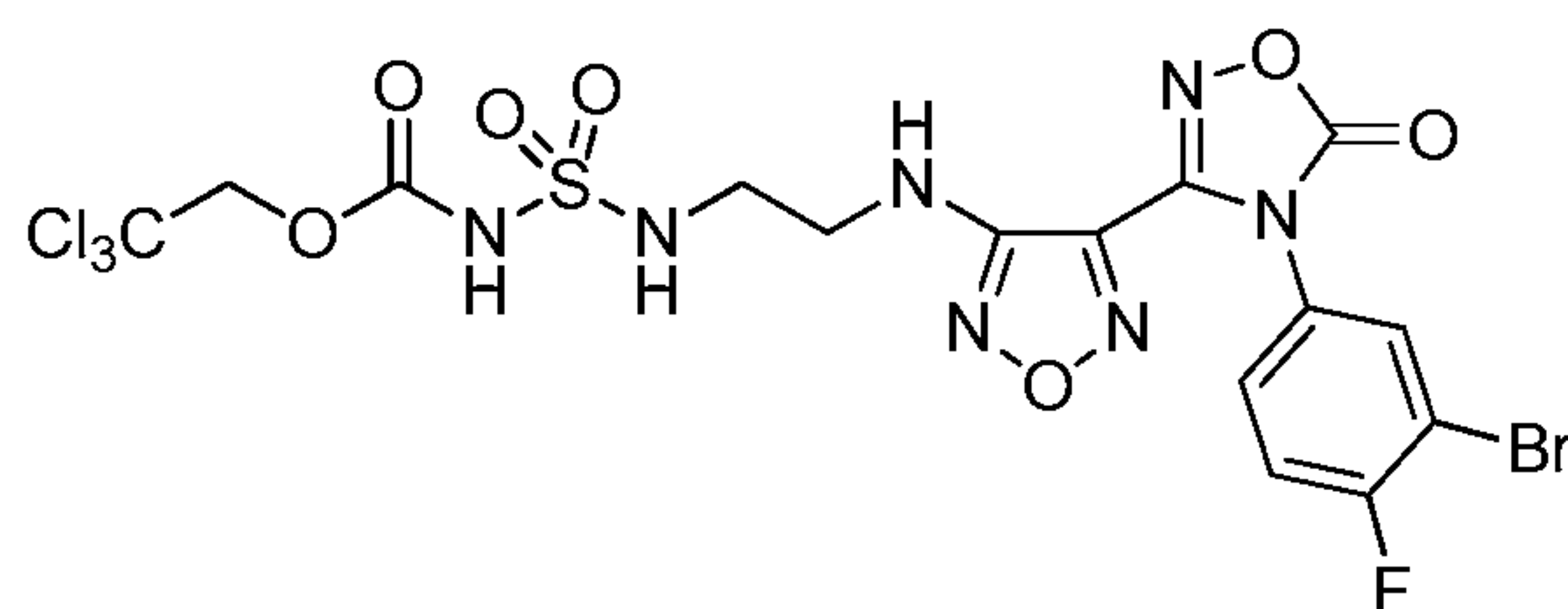
A mixture of 3-(4-amino-1,2,5-oxadiazol-3-yl)-4-(3-bromo-4-fluorophenyl)-1,2,4-oxadiazol-5(4*H*)-one (68 mg, 0.20 mmol; from Step D) and benzyl {[2,2-dimethoxyethyl)amino]sulfonyl}carbamate (191 mg, 0.60 mmol, 3.0 equiv.) in 1,2-dichloroethane (3.0 mL) was cooled to 0 °C. To this mixture was added trifluoroacetic acid (1.0 mL, 13.0 mmol) and triethylsilane (105 μL, 0.66 mmol, 3.3 equiv.) drop-wise. This reaction mixture was stirred at 0 °C for 2 h. HPLC indicated reaction completion. The reaction mixture was cooled to 0 °C and quenched with saturated sodium bicarbonate to pH ~ 8. and the quenched reaction mixture was extracted with EtOAc (3 x 10 mL). The combined organic extracts were washed with brine, dried over sodium sulfate and concentrated. The residue was then stirred in a mixture of heptane and diethyl ether overnight. The solids were collected by filtration, washed with heptane and dried in vacuum to give the desired product (125 mg, 99%) as a crude off-white solid. ¹H NMR (300 MHz, DMSO-*d*₆) δ 11.31 (s, 1H), 8.05 (m, 1H), 7.87 (m, 1H), 7.68 (m, 1H), 7.56 (m, 1H), 7.32 (m, 5H), 6.54 (m, 1H), 5.07 (s, 2H), 3.29 (m, 2H), 3.09 (m, 2H); C₂₀H₁₇BrFN₇O₇S (MW 598.36), LCMS *m/e* 598/600 (M⁺ + H).

25 *Step 2c. Ethyl* ({[2-({[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]amino}sulfonyl)carbamate (18c)



A mixture of 3-(4-amino-1,2,5-oxadiazol-3-yl)-4-(3-bromo-4-fluorophenyl)-1,2,4-oxadiazol-5(4H)-one (68 mg, 0.20 mmol; from Step D) and ethyl {[2,2-dimethoxyethyl)amino]sulfonyl} carbamate (154 mg, 0.600 mmol, 3.0 equiv.) in 1,2-dichloroethane (2.50 mL, 31.7 mmol) was stirred at 0 °C. To this mixture was added trifluoroacetic acid (1.00 mL, 13.0 mmol) and triethylsilane (105 µL, 0.66 mmol, 3.3 equiv.) = drop-wise. The reaction mixture was stirred at 0 °C for 3 h. HPLC indicated 97.5% conversion to the desired product. The reaction mixture was cooled to 0 °C and quenched with saturated sodium bicarbonate to pH ~ 8. The mixture was extracted in ethyl acetate (3 x 10 mL). The combined organic extracts were washed with brine, dried over sodium sulfate and concentrated. The residue was stirred in a mixture of heptane and diethyl ether overnight. The solids were collected by filtration, washed with heptane to give the desired product (95 mg, 88%) as a crude off-white solid. ¹H NMR (300 MHz, DMSO-*d*₆) δ 11.18 (s, 1H), 8.08 (m, 1H), 7.70 (m, 2H), 7.59 (t, 1H, *J* = 8.7 Hz), 6.56 (s, 1H), 4.04 (d, 2H, *J* = 7.2 Hz), 3.35 (m, 2H), 3.11 (m, 2H), 1.15 (t, 3H, *J* = 7.2 Hz); C₁₅H₁₅BrFN₇O₇S (MW 536.29), LCMS (EI) *m/e* 536/538 (M⁺ + H).

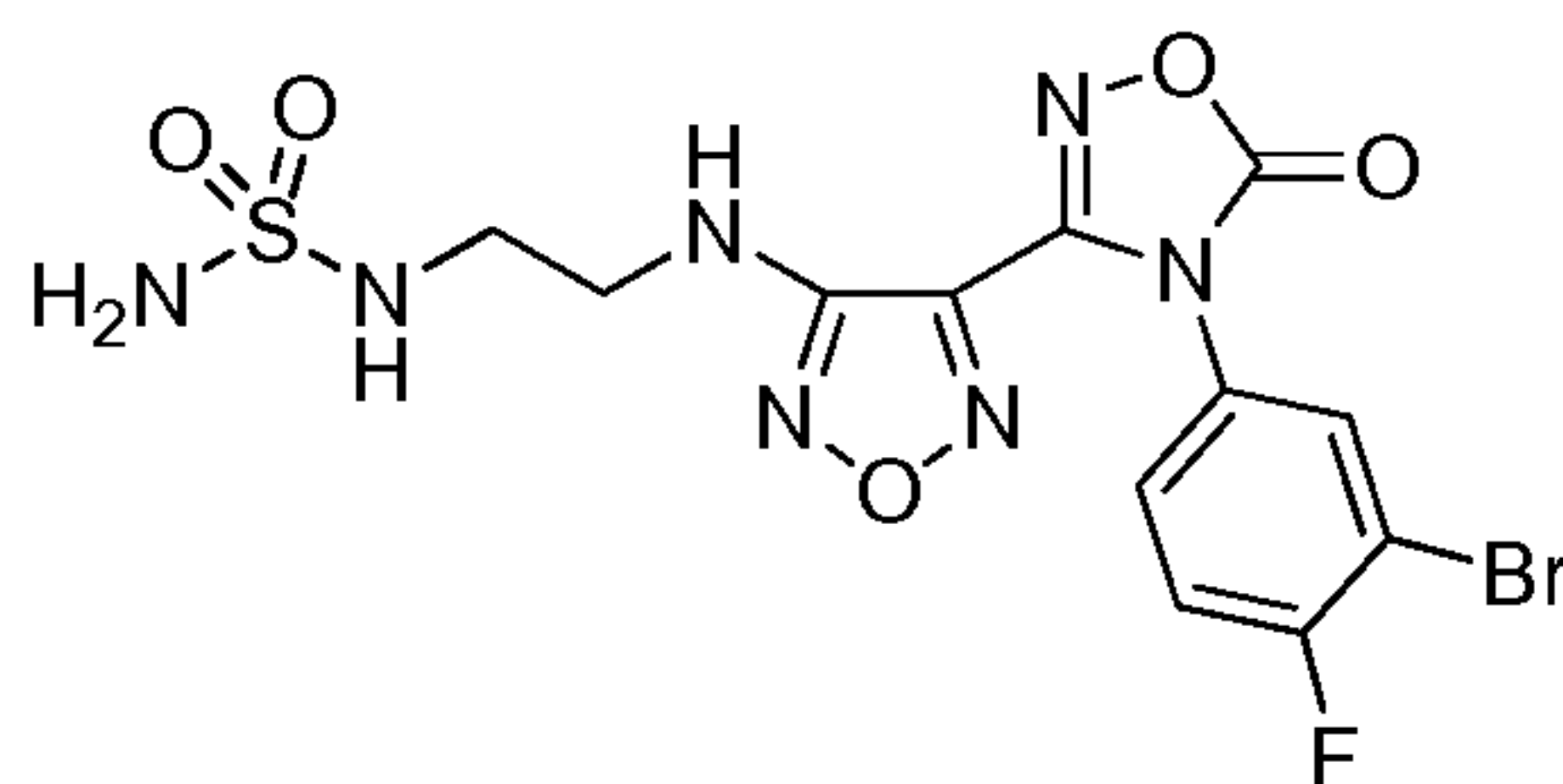
Step 2d. 2,2,2-Trichloroethyl ({[2-({4-[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]amino}sulfonyl)carbamate (18d)



A suspension of 3-(4-amino-1,2,5-oxadiazol-3-yl)-4-(3-bromo-4-fluorophenyl)-1,2,4-oxadiazol-5(4H)-one (**5**, 0.680 g, 1.99 mmol) and 2,2,2-trichloroethyl {[2,2-dimethoxyethyl)amino]sulfonyl} carbamate (**17d**, 2.22 g, 6.17 mmol, 3.1 equiv.) in dichloromethane (DCM, 6.0 mL) was stirred at room temperature. To this mixture was added triethylsilane (1.27 mL, 7.95 mmol, 4.0 equiv.) and a solution of trifluoroacetic acid (TFA, 3.0 mL, 39.0 mmol) in dichloromethane (DCM, 2.0 mL) while maintaining the

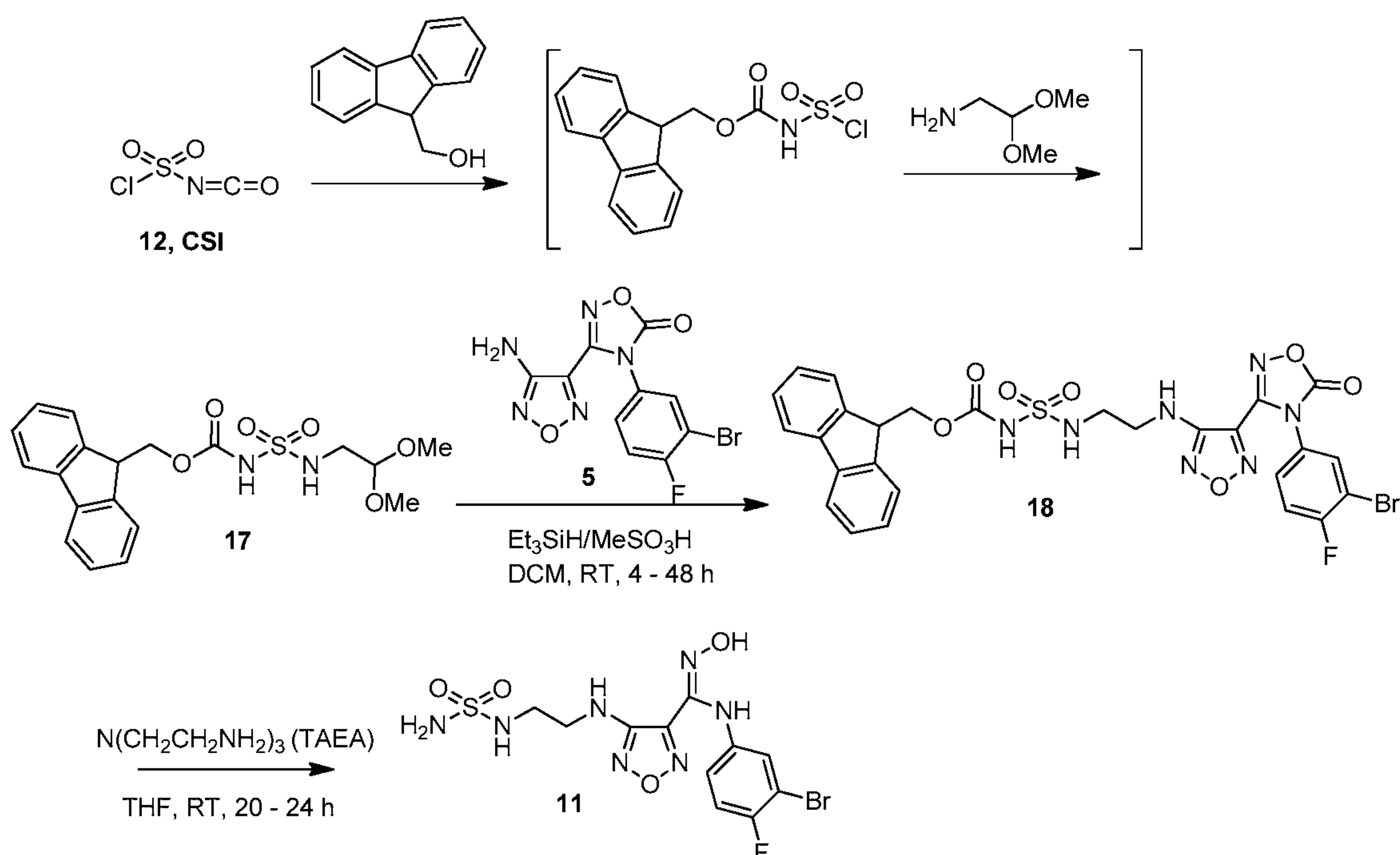
reaction temperature below 30 °C. The reaction mixture became homogenous after 5 min with agitation at room temperature and was stirred at room temperature for 1 h. HPLC indicated reaction completion. The reaction was filtered and the precipitate was suspended in a mixture of dichloromethane and heptane (ratio of dichloromethane to heptane was 1 to 9 by volume). The suspension was stirred at room temperature overnight. The precipitate was collected by filtration and washed with 10% dichloromethane in heptane and dried in vacuum to give the desired product (1.15 g, 90.4%) as an off-white solid, which was used in the subsequent reaction without further purification. ¹H NMR (300 MHz, DMSO-*d*₆) δ 11.85 (s, 1H), 8.07 (m, 2H), 7.70 (m, 1H), 7.57 (t, 1H, *J* = 8.7 Hz), 6.56 (m, 1H), 4.88 (m, 2H), 3.37 (m, 2H), 3.16 (m, 2H); C₁₅H₁₂BrCl₃FN₇O₇S (MW 639.62), LCMS (EI) *m/e* 638/640/642 (M⁺ + H).

Step 3. N-[2-({4-[4-(3-Bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]sulfamide (10)

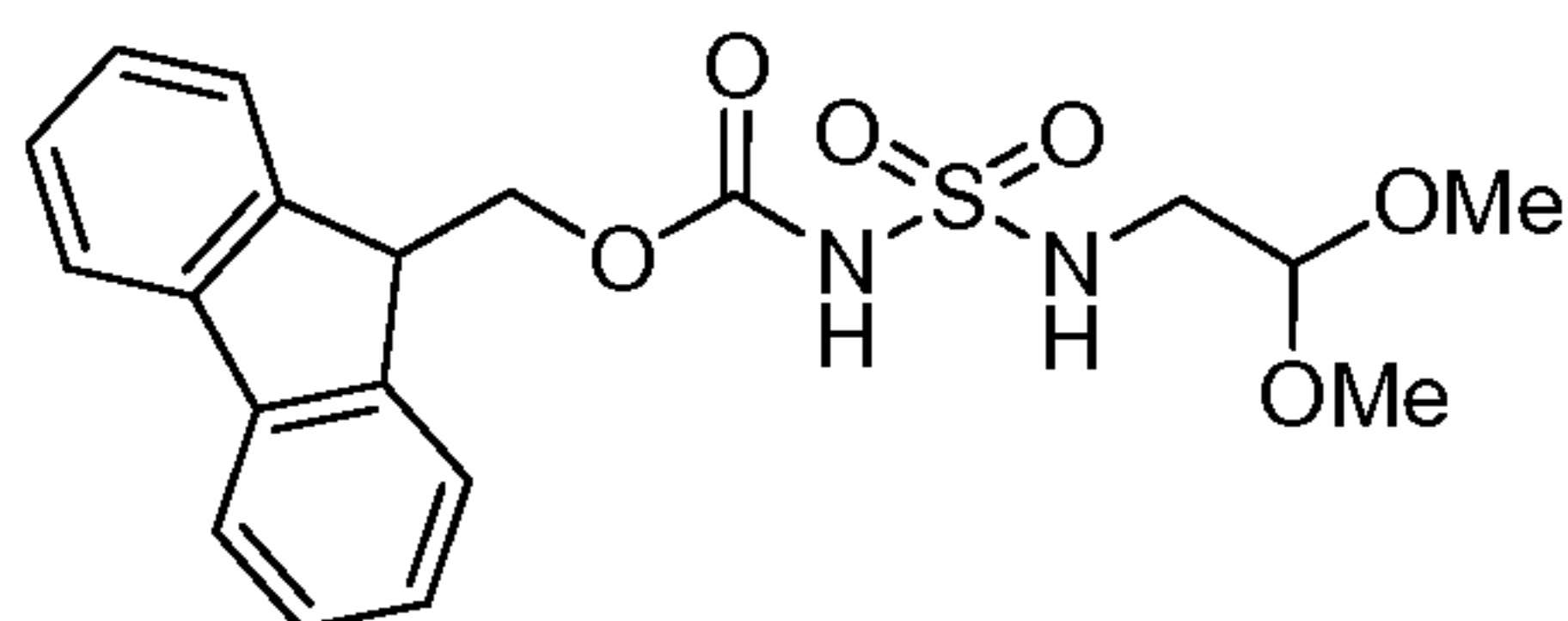


A solution of 2,2,2-trichloroethyl ({[2-({4-[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]amino}sulfonyl)carbamate (320 mg, 0.50 mmol; from Step Q, Method D) in tetrahydrofuran (THF, 4.0 mL) was stirred at room temperature. Acetic acid (0.30 mL, 5.3 mmol) and zinc flakes (160 mg, 2.5 mmol, 5.0 equiv.) were sequentially added. This reaction mixture was stirred at room temperature for 3 h. HPLC indicated reaction completion. The reaction mixture was filtered through Celite and the Celite was washed with THF. The combined filtrate was concentrated in vacuum and the resulting residue was dissolved in ethyl acetate (20 mL). The ethyl acetate solution was washed with saturated sodium carbonate and brine, dried over sodium sulfate and concentrated. The crude material was crystallized from ethyl acetate and diethyl ether to give the desired product (147 mg, 63%) as an off-white solid.

Example 4. Alternate Preparation of 4-({2-[(Aminosulfonyl)amino]ethyl}amino)-N-(3-bromo-4-fluorophenyl)-N'-hydroxy-1,2,5-oxadiazole-3-carboximidamide



Step 1. (9H-fluoren-9-yl)methyl N-(2,2-dimethoxyethyl)sulfamoylcarbamate



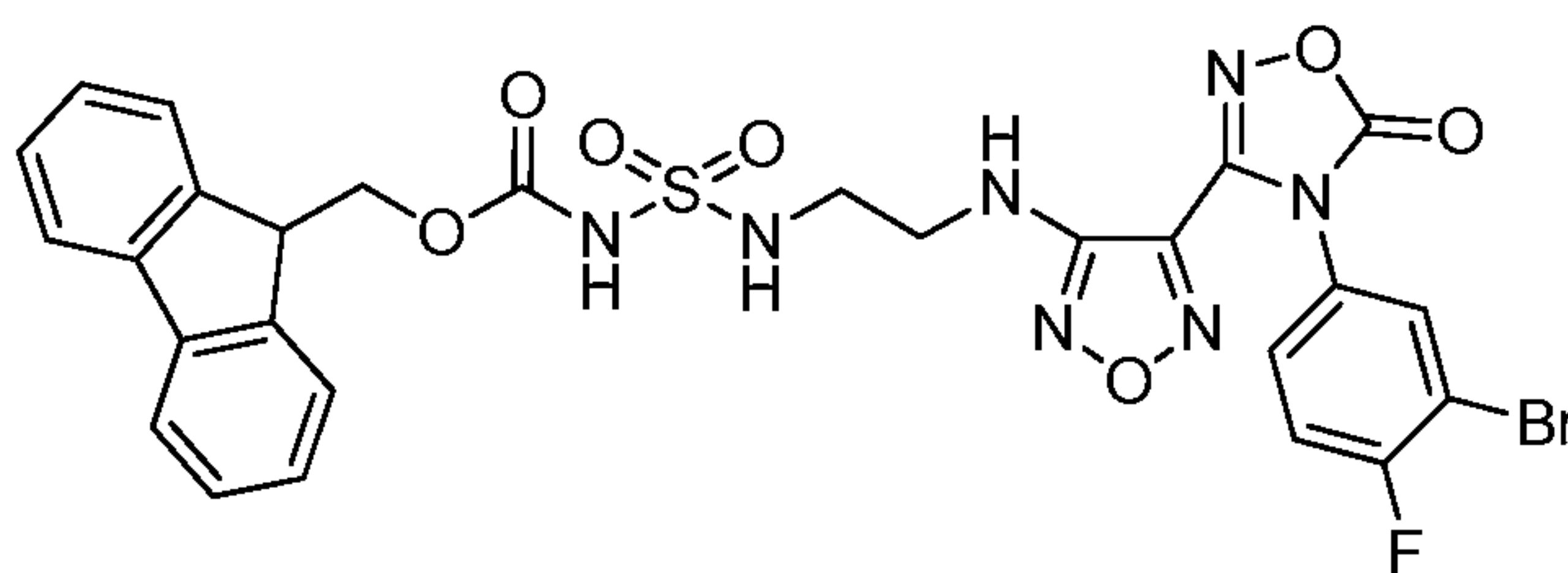
5

Into an oven dried 2 L 4-neck round bottom flask was charged 9-fluorenylmethanol (50.0 g, 255 mmol) and anhydrous DCM (382 mL) at room temperature. The resulting slurry was cooled in an ice-bath to about 0 - 5 °C. A solution of chlorosulfonyl isocyanate (CSI, 23.0 mL, 264 mmol) in anhydrous DCM (127 mL) was added dropwise to the slurry through an addition funnel over 22 minutes, maintaining the reaction mixture temperature at < 5°C. The resulting mixture was stirred at 0 - 5 °C for 1.75 h, producing a thick white slurry. A solution of aminoacetaldehyde dimethyl acetal (27.9 mL, 255 mmol) in anhydrous DCM (382 mL) and 4-methylmorpholine (84.0 mL, 764 mmol) were added to the mixture at about 0 - 5 °C over 71 minutes. The resulting reaction mixture was then stirred in the ice bath for 1.5 hours. When HPLC showed the the reaction was complete, the reaction mixture was acidified by the dropwise addition of a 1.0 M phosphoric acid (H_3PO_4 , aq., 640 mL) over 22 minutes to pH 1 - 2. Water (300 mL), EtOAc (2150 mL) and heptane (250 mL) were then added and the resulting mixture was stirred for 10 minutes. The two phases were separated and the organic phase was washed sequentially with water (500 mL), heptane (300 mL) and water (2 x 500 mL) and dried over MgSO_4 . The filtrate was concentrated under

20

vacuum to dryness. The resulting solids were redissolved in EtOAc (600 mL) at 65 °C and the warm solution was filtered into a clean 3 L round bottom flask. The filtrate was cooled to room temperature and stirred for 2.5 h before heptane (1200 mL) was then added *via* an addition funnel over 80 min. After stirring overnight at room temperature, the mixture was then cooled in an ice bath for 1 h. The resulting solids were collected by filtration, washed with 25% EtOAc/heptane (250 mL), and dried overnight at about 40 - 45°C under vacuum to afford 9*H*-fluoren-9-ylmethyl {[2-(4-[[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl]amino)ethyl]amino}sulfonyl}carbamate (91.3 g, 88% yield) as a white powder. ¹H NMR (300 MHz, DMSO-*d*₆) δ 11.43 (s, 1H), 7.98 – 7.85 (m, 3H), 7.76 (d, *J* = 7.5 Hz, 2H), 7.43 (t, *J* = 7.2 Hz, 2H), 7.33 (td, *J* = 7.4, 1.1 Hz, 2H), 4.44 – 4.33 (m, 3H), 4.33 – 4.22 (m, 1H), 3.23 (s, 6H), 2.99 (t, *J* = 5.8 Hz, 2H) ppm.

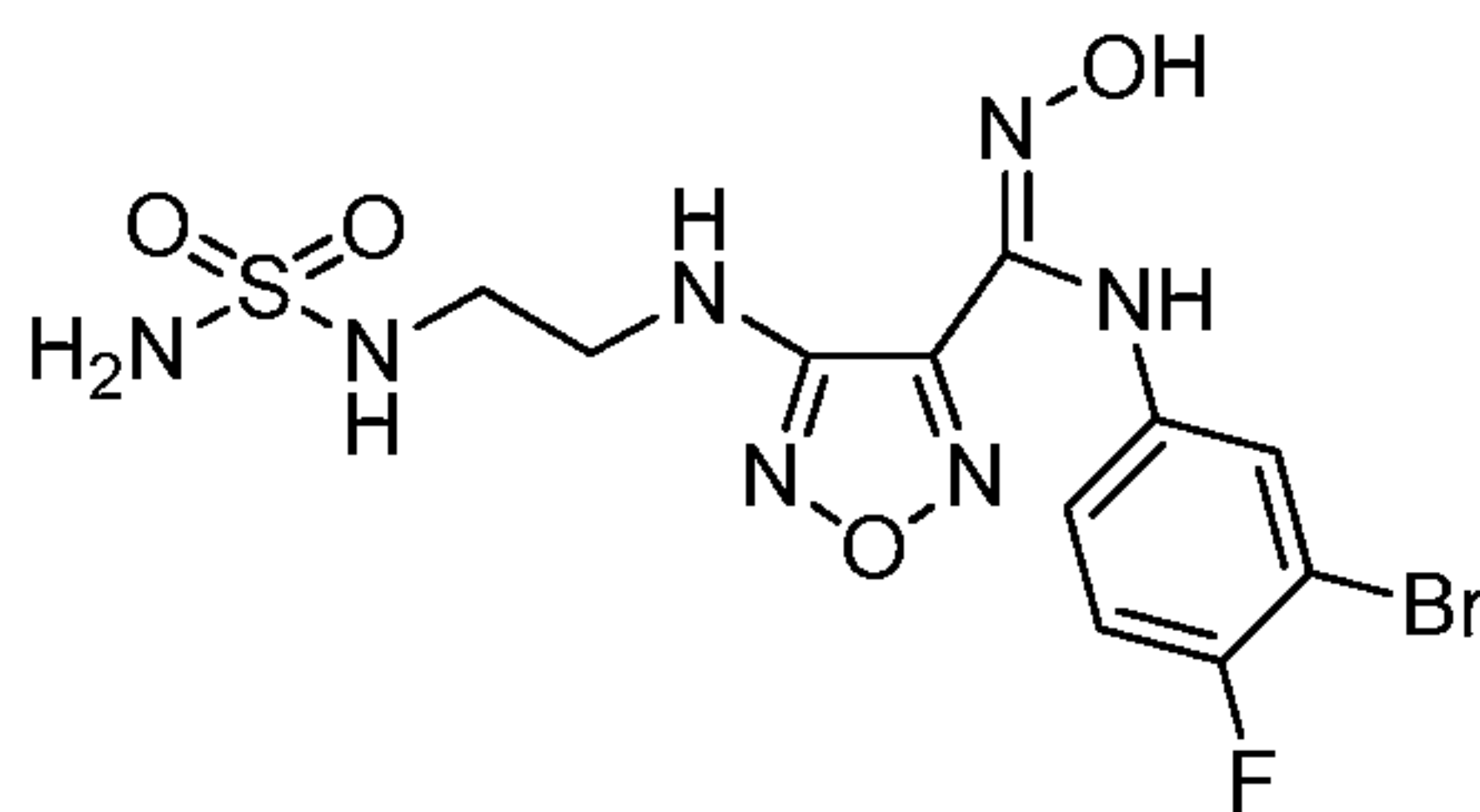
Step 2. 9H-Fluoren-9-ylmethyl ({[2-(4-[[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl]amino)ethyl]amino}sulfonyl}carbamate



To a stirred suspension of 3-(4-amino-1,2,5-oxadiazol-3-yl)-4-(3-bromo-4-fluorophenyl)-1,2,4-oxadiazol-5(4*H*)-one (10.00 g, 29.23 mmol) in DCM (160 mL) was added methanesulfonic acid (MeSO₃H, 8.46 g, 88.04 mmol) and triethylsilane (Et₃SiH, 8.37 g, 71.96 mmol) at ambient temperature over 10 minutes to give a slurry. Solid 9*H*-fluoren-9-ylmethyl {[2-(2,2-dimethoxyethyl)amino]sulfonyl}carbamate (12.25 g, 30.14 mmol) was added portionwise (1 g/3 - 4 min; over 1 h) while maintaining the internal temperature at less than about 20 °C using a water bath. After the addition, the resulting mixture was stirred at about 13 to 22 °C for 3 days. Additional triethylsilane (Et₃SiH, 0.1755 g, 1.51 mmol) and 9*H*-fluoren-9-ylmethyl {[2-(2,2-dimethoxyethyl)amino]sulfonyl}carbamate (0.3082 g, 0.76 mmol) were added and the resulting mixture was stirred at ambient temperature for an additional 23 h. Isopropyl alcohol (IPA, 15 mL) was added and the resulting mixture was stirred at ambient temperature for 1 h. Heptane (100 mL) was added and the mixture was stirred at ambient temperature for an additional 2 h. The solids were collected by filtration, washed with IPA/heptane (1/5; 2 x 30 mL) and heptane (2 x 30 mL), and dried under vacuum to afford 9*H*-fluoren-9-ylmethyl ({[2-(4-[[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl]amino)ethyl]amino}sulfonyl}carbamate as a white

solid (18.30 g, 91.1% yield). ^1H NMR (500 MHz, $\text{DMSO}-d_6$) δ 11.44 (s, 1H), 8.07 (dd, J = 6.2, 2.5 Hz, 1H), 7.90 (t, J = 5.6 Hz, 1H), 7.88 (d, J = 7.6 Hz, 2H), 7.72 (d, J = 7.0 Hz, 2H), 7.71 (ddd, J = 8.9, 4.3, 2.6 Hz, 1H), 7.57 (dd, J = 8.7, 8.7 Hz, 1H), 7.40 (t, J = 7.4 Hz, 2H), 7.31 (td, J = 7.4, 1.0 Hz, 2H), 6.55 (t, J = 6.0 Hz, 1H), 4.35 (d, J = 7.3 Hz, 2H), 4.25 (t, J = 7.2 Hz, 1H), 3.39 (q, J = 6.4 Hz, 2H), 3.15 (q, J = 6.3 Hz, 2H); ^{13}C NMR (126 MHz, $\text{DMSO}-d_6$) δ 159.03 (d, J = 248.7 Hz), 156.61 (s), 155.22 (s), 151.55 (s), 148.67 (s), 143.29 (s), 140.68 (s), 133.82 (s), 133.39 (s), 130.05 (d, J = 8.5 Hz), 128.54 (d, J = 3.2 Hz), 127.73 (s), 127.07 (s), 125.24 (s), 120.11 (s), 117.42 (d, J = 24.0 Hz), 108.19 (d, J = 22.5 Hz), 66.70 (s), 46.17 (s), 43.34 (s), 40.79 (s) ppm; ^{19}F NMR (376 MHz, $\text{DMSO}-d_6$) δ -103.99 – -107.39 (m) ppm.

Step 3. 4-({2-[(Aminosulfonyl)amino]ethyl}amino)-N-(3-bromo-4-fluorophenyl)-N'-hydroxy-1,2,5-oxadiazole-3-carboximidamide



Into a 1 L 4-neck round bottom flask was charged 9H-fluoren-9-ylmethyl({[2-({4-[4-(3-bromo-4-fluorophenyl)-5-oxo-4,5-dihydro-1,2,4-oxadiazol-3-yl]-1,2,5-oxadiazol-3-yl}amino)ethyl]amino}sulfonyl)carbamate (25.0 g, 36.4 mmol) and anhydrous THF (250 mL) at ambient temperature to produce a homogeneous solution. The solution was then cooled to 0 – 5 °C in an ice bath before *N,N*-bis(2-aminoethyl)ethane-1,2-diamine (114 mL, 728 mmol) was added dropwise over 35 minutes *via* an addition funnel. The addition funnel was rinsed with anhydrous THF (50 mL) and the rinse was added to the reaction mixture. The cold bath was removed and the reaction was gradually warmed to ambient temperature and stirred at ambient temperature for 2.5 h. EtOAc (400 mL) was added and the resulting mixture was transferred to a 2 L 4-neck round bottom flask and cooled to about 0 - 5 °C in an ice bath. A solution of 2.0 M aqueous HCl (400 mL, 800.0 mmol) was added dropwise *via* an addition funnel, while maintaining the internal temperature at below 10 °C. The two phases were separated and the aqueous phase was extracted with EtOAc (200 mL). The organic fractions were combined and cooled to about 6-7 °C. A solution of 2.0 M aqueous HCl (200.0 mL, 400.0 mmol) was added dropwise to the cold organic fraction, maintaining the internal temperature at below 10°C. The two phases were separated and the organic

phase was washed with water (2 x 400 mL), dried over MgSO₄, and concentrated under reduced pressure to a light yellow syrup. The syrup was dissolved in EtOAc (60.0 mL) to yield a homogeneous solution. To the solution was added a solution of DCM (250.0 mL) and *tert*-butyl methyl ether (TBME, 100.0 mL) dropwise. The resulting slurry was stirred
5 overnight at room temperature, then cooled in an ice bath for 1 h. The solids were collected by filtration, washed with an ice cold 250 mL solution of DCM (150 mL) and TBME (100 mL), and dried under vacuum to give 14.4 g of the crude desired product as white solids.

The crude product was dissolved in EtOAc (140.0 mL) at 60 °C and the warm solution was filtered. The filtrate was cooled to room temperature before heptane (100.0
10 mL) was added dropwise over 55 min. The resulting mixture was then stirred overnight at room temperature. The solids were collected by filtration, washed with a 2:1 mixture of heptane and EtOAc (75 mL), and dried under vacuum at 40 – 50 °C to constant weight to afford 4-({2-[(aminosulfonyl)amino]ethyl}amino)-*N*-(3-bromo-4-fluorophenyl)-*N'*-hydroxy-1,2,5-oxadiazole-3-carboximidamide (12.9 g, 81% yield) as a white solid.

Example A: Human indoleamine 2,3-dioxygenase (IDO) enzyme assay

Human indoleamine 2,3-dioxygenase (IDO) with an N-terminal His tag was expressed in *E.coli* and purified to homogeneity. IDO catalyzes the oxidative cleavage of the pyrrole ring of the indole nucleus of tryptophan to yield *N'*-formylkynurenine. The assays
20 were performed at room temperature as described in the literature using 95 nM IDO and 2 mM D-Trp in the presence of 20 mM ascorbate, 5 μM methylene blue and 0.2 mg/mL catalase in 50 mM potassium phosphate buffer (pH 6.5). The initial reaction rates were recorded by continuously following the absorbance increase at 321 nm due to the formation of *N'*-formylkynurenine (See: Sono, M., *et al.*, 1980, *J. Biol. Chem.* 255, 1339-1345). The
25 compound of Formula I was tested in the assay of Example A and found to have an IC₅₀ of < 200 nM.

Example B: Determination of inhibitor activity in HeLa cell-based indoleamine 2,3-dioxygenase (IDO)/Kynurenine assay

30 HeLa cells (#CCL-2) were obtained from the American Type Tissue Culture Collection (ATCC, Manassas, VA) and routinely maintained in minimum essential medium (eagle) with 2 mM L-glutamine and Earle's BSS adjusted to contain 1.5 g/L sodium bicarbonate, 0.1 mM non-essential amino acids, 1 mM sodium pyruvate and 10 % fetal

bovine serum (all from Invitrogen). Cells were kept at 37 °C in a humidified incubator supplied with 5 % CO₂. The assay was performed as follows: HeLa cells were seeded in a 96 well culture plate at a density of 5 x 10³ per well and grown overnight. On the next day, IFN-γ (50 ng/mL final concentration) and serial dilutions of compounds (in total volume of 200 μL culture medium) were added into cells. After 48 hours of incubation, 140 μL of the supernatant per well was transferred to a new 96 well plate. 10 μL of 6.1 N trichloroacetic acid (#T0699, Sigma) was mixed into each well and incubated at 50 °C for 30 min to hydrolyze *N*-formylkynurenine produced by indoleamine 2,3-dioxygenase to kynurenine. The reaction mixture was then centrifuged for 10 min at 2500 rpm to remove sediments. 100 μL of the supernatant per well was transferred to another 96 well plate and mixed with 100 μL of 2% (w/v) *p*-dimethylaminobenzaldehyde (#15647-7, Sigma-Aldrich) in acetic acid. The yellow color derived from Kynurenine was measured at 480 nm using a SPECTRAmax 250 microplate reader (Molecular Devices). L-kynurenine (#K8625, Sigma) was used as standard. The standards (240, 120, 60, 30, 15, 7.5, 3.75, 1.87 μM) were prepared in 100 μL culture media and mixed with equal volume of 2 % (w/v) *p*-dimethylaminobenzaldehyde. The percent inhibition at individual concentrations was determined and the average values of duplicates were obtained. The data was analyzed by using nonlinear regression to generate IC₅₀ values (Prism Graphpad). See: Takikawa O, *et al.*, 1988, *J. Biol. Chem.*, 263(4): 2041-8.

20 **Example C: Determination of effect of IDO inhibitors on T cell proliferation that is suppressed by IDO-expressing dendritic cells**

Monocytes were collected from human peripheral mononuclear cells by leukaphoresis. Monocytes were then seeded at a density of 1 x 10⁶ cells/well in a 96 well plate, using RPMI 1640 medium supplemented with 10 % fetal bovine serum and 2 mM L-glutamine (all from Invitrogen). Adherent cells were retained on the plate after overnight culture at 37 °C. Adherent monocytes were then stimulated for 5-7 days with 100 ng/ml GM-CSF (# 300-03, PeproTech) and 250 ng/ml IL-4 (#200-04, PeproTech), followed by activation with 5 μg/mL LPS from *Salmonella typhimurium* (#437650, Sigma) and 50 ng/mL IFN-γ (# 285-IF, R&D Systems) for additional 2 days to induce dendritic cell maturation.

30 After dendritic cell activation, the medium was replaced with completed RPMI 1640 supplemented with 100-200 U/mL IL-2 (#CYT-209, ProSpec-Tany TechnoGene) and 100 ng/mL anti-CD3 antibody (#555336, PharMingen), T cells (2-3 x 10⁵ cells/well), and serial dilutions of IDO compounds. After incubation for 2 more days, T cell proliferation was

measured by BrdU incorporation assay, using a colorimetric Cell Proliferation ELISA kit per manufacturer's instruction (#1647229, Roche Molecular Biochemicals). Cells were continuously cultured for 16-18 hrs in presence of 10 μ M BrdU labeling solution. Then, the labeling medium was removed, and 200 μ L FixDenat per well was added to the cells and incubated for 30 minutes at room temperature. The FixDenat solution was removed and 100 μ L/well anti-BrdU-POD antibody conjugate working solution was added. The reaction was carried out for 90 minutes at room temperature. The antibody conjugate was then removed, and cells were rinsed three times with 200 μ L/well washing solution. Finally, 100 μ L/well of substrate solution was added and the results were obtained using a microplate reader (Spectra Max PLUS, Molecular Devices) during color development. Multiple readings at various time points were obtained to ensure the data was within the linear range. The data was routinely obtained from replicated experiments, and appropriate controls were included. See: Terness P, *et al.* 2002, *J. Exp. Med.*, 196(4): 447-57; and Hwu, P, *et al.* 2000, *J. Immunol.*, 164(7): 3596-9.

Example D: *In vivo* testing of IDO inhibitors for antitumor activity

In vivo anti-tumor efficacy can be tested using modified tumor allograft/xenograft protocols. For instance, it has been described in the literature that IDO inhibition can synerize with cytotoxic chemotherapy in immune-competent mice (Muller, A.J., *et al.* 2005, *Nat. Med.* 11:312-319). This synergy was shown to be dependent on T-cells by comparison of the synergistic effects of an investigational IDO inhibitor in murine tumor xenograft models (e.g. B16 and related variants, CT-26, LLC) grown in immune competent syngenic mice to that observed in syngenic mice treated with neutralizing anti-CD4 antibodies, or the same tumors grown in immune-compromised mice (e.g. nu/nu).

The concept of differential anti-tumor effects in immune-competent versus immune-compromised mice may also permit testing of investigational IDO inhibitors as single agents. For instance, LLC tumors grow well in their syngenic host strain, C57Bl/6. However, if these mice are treated with the IDO inhibitor 1-MT (versus placebo) the formation of tumors is markedly delayed, implying that IDO inhibition was growth inhibitory (Friberg, M., *et al.* 2002, *Int. J. Cancer* 101:151-155). Following this logic, one can examine the efficacy of IDO inhibition in the LLC xenograft tumor model grown in C57Bl/6 immune competent mice and compare that to the effects of IDO inhibitors on LLC tumor growth in nude or SCID mice (or C57Bl/6 mice treated with antibodies that neutralize T-cell activity). As the

effects of relieving the tumor-mediated immune suppressive activity of IDO will likely differ depending on the immunogenic potential of different tumor models, genetic modifications can be made to the tumor cells to increase their immunogenic potential. For instance, expression of GM-CSF in B16.F10 cells increases their immunogenic potential (Dranoff, G.,
5 *et al.* 1993, *Proc. Natl. Acad. Sci., USA*, 90:3539-3543). As such, in some tumor models (e.g. B16.F10) one can generate [poly]clones that express immune stimulatory proteins such as GM-CSF and test the growth inhibitory effects of IDO inhibitors against tumors established from these tumor cells in both immune-competent and –compromised mice.

A third avenue for assessing the efficacy of IDO inhibitors *in vivo* employs ‘pre-
10 immunization’ murine tumor allograft/xenograft models. In these models, immune-competent mice are sensitized to a specific tumor antigen or antigens to mimic a therapeutic anti-tumor vaccination. This primes the mice for an anti-tumor response mediated by the immune system when mice are subsequently challenged with murine tumor cell lines (possessing similar tumor antigens to those used for immunization) in xenograft experiments.
15 Expression of IDO has been shown to blunt the anti-tumor response and allow xenografts to grow more rapidly. Importantly, the growth of tumors in this model is inhibited by the IDO inhibitor 1-MT (Uyttenhove, C., *et al.* 2003, *Nat. Med.* 9:1269-1274). This model is particularly attractive as IDO activity is permissive for P815 tumor growth and specific inhibition of IDO should therefore growth inhibitory.

20 Lastly, therapeutic immunization may be used to evaluate the impact of IDO inhibitors *in vivo*. For example, it has been demonstrated using B16-BL6 cells that one can challenge Blk/6 mice with an intravenous injection of tumor cells followed by treatment with a well characterized immunogenic peptide (e.g. TRP-2) expressed by the tumor cells (Ji, *et al.*, 2005, *J. Immunol*, 175: 1456-63). Importantly, immune system modifiers, such as anti-
25 CTL-4 antibody, can improve responses to such therapeutic immunizations. The impact of IDO inhibitors may be evaluated in a similar manner – tumor peptide immunization with or without IDO inhibitor. Efficacy is assessed by animal survival (time to morbidity) or by the measurement of tumor metastases to the lungs and/or other organs at defined timepoints.

In any/all of the above mentioned models, it may also be possible to directly and/or
30 indirectly measure the number and/or activity of tumor reactive immune cells. Methods for measuring the number and/or activity of tumor reactive immune cells are well established and can be performed using techniques familiar to those schooled in the art (Current Protocols in Immunology, Vol. 4, Coligan, J. E., *et al.*; *Immunotherapy of Cancer*, Human Press, 2006, Disis, M.L. and references therein). Conceptually, a reduction in the immune suppressive

effects of IDO may result in increased numbers or reactivity of tumor specific immune cells. Further, IDO inhibition may further increase the number or reactivity of tumor reactive immune cells when combined with other therapeutics, for example chemotherapeutics and/or immune modulators (e.g. anti-CTLA4 antibody).

5 All allograft/xenograft experiments can be performed using standard tumor techniques (reviewed by Corbett, *et al.*, In *Cancer Drug Discovery and Development: Anticancer Drug Development Guide: Preclinical Screening, Clinical Trials, and Approval*, 2nd Ed. Teicher, B.A. and Andrews, P.A., Humana Press Inc.: Totowa, NJ, 2004). The cloning and introduction of genes (e.g. IDO, GM-CSF) into tumor cell lines, can be
10 performed using techniques familiar to those schooled in the art (reviewed in Sambrook, J. and Russel, D., *Molecular Cloning: A laboratory Manual (3rd edition)*, Cold Spring Harbor Laboratory Press: Cold Spring Harbor, NY, 2001).

Example E: *In vivo* testing of IDO inhibitors in human immunodeficiency virus-1 (HIV-

15 1) encephalitis model

1. Cell isolation and viral infection

Monocytes and PBL can be obtained by countercurrent centrifugal elutriation of leukopheresis packs from HIV-1, 2 and hepatitis B seronegative donors. Monocytes are cultivated in suspension culture using Teflon flasks in Dulbecco's Modified Eagle's Medium
20 (DMEM, Sigma-Aldrich) supplemented with 10 % heat-inactivated pooled human serum, 1 % glutamine, 50 µg/mL gentamicin, 10 µg/mL ciprofloxacin (Sigma), and 1000 U/mL highly purified recombinant human macrophage colony stimulating factor. After seven days in culture, MDM are infected with HIV-1_{ADA} at multiplicity of infection of 0.01.

25 2. Hu-PBL-NOD/SCID HIVE mice

Four-wk old male NOD/C.B-17 SCID mice can be purchased (Jackson Laboratory). Animals are maintained in sterile microisolator cages under pathogen-free conditions. All animals are injected intraperitoneally with rat anti-CD122 (0.25 mg/mouse) three days before PBL transplantation and twice with rabbit asialo-GM1 antibodies (0.2 mg/mouse) (Wako)
30 one day before and three days after PBL injection (20×10^6 cells/mouse). HIV-1_{ADA}-infected MDM (3×10^5 cells in 10 µL) are injected intracranially (i.c.) eight days following PBL reconstitution generating hu-PBL-NOD/SCID HIVE mice. Immediately following i.c. injection of HIV-1 infected MDM the hu-PBL-NOD/SCID HIVE mice are subcutaneously

(s.c) implanted with control (vehicle) or compound pellets (14 or 28 day slow release, Innovative Research). Initial experiments are designed to confirm the induction of virus-specific CTL in the hu PBL-NOD/SCID HIVE animals treated with IDO compounds. This is confirmed by tetramer staining and neuropathologic analyses of MDM elimination from the brain tissue. Then, the experiment is designed to analyze human lymphocyte reconstitution, humoral immune responses, and neuropathological alterations. In these experiments, animals are bled on day 7 and sacrificed at 14 and 21 days after i.c. injection of human MDM. Blood collected in EDTA-containing tubes is used for flow cytometry and plasma is used for detection of HIV-1 p24 using ELISA (Beckman Coulter™). HIV-1-specific antibodies are detected by Western blot tests according to the manufacturer instructions (Cambridge Biotech HIV-1 Western blot kit, Calypte Biomedical). Similar amount of virus-specific antibodies are detected in control and compound-treated animals. A total of three independent experiments can be performed using three different human leukocyte donors.

3. *FACScan of peripheral blood and spleen in hu PBL-NOD/SCID HIVE mice*

Two-color FACS analysis can be performed on peripheral blood at wk 1-3 and splenocytes at wk 2 and 3 after i.c. injection of human MDM. Cells are incubated with fluorochrome-conjugated monoclonal Abs (mAbs) to human CD4, CD8, CD56, CD3, IFN- γ (eBioscience) for 30 min at 4 °C. To evaluate the cellular immune response, IFN- γ intracellular staining is performed in combination with anti-human CD8 and FITC-conjugated anti-mouse CD45 to exclude murine cells. To determine the Ag-specific CTL, allophycocyanin-conjugated tetramer staining for HIV-1^{gag} (p17 (aa77-85) SLYNTVATL, SL-9) and HIV-1^{pol} [(aa476-485) ILKEPVHGV, IL-9] is performed on phytohemagglutinin/interleukin-2 (PHA/IL-2)-stimulated splenocytes. Cells are stained following the recommendation of the NIH/National Institute of Allergy and Infectious Disease, National Tetramer Core Facilities. Data were analyzed with a FACS Calibur™ using CellQuest software (Becton Dickinson Immunocytometry System).

4. *Histopathology and image analyses*

Brain tissue is collected at days 14 and 21 after i.c. injection of MDM, fixed in 4 % phosphate-buffered paraformaldehyde and embedded in paraffin or frozen at -80 °C for later use. Coronal sections from the embedded blocks are cut in order to identify the injection site. For each mouse, 30-100 (5- μ m-thick) serial sections are cut from the human MDM injection

site and 3-7 slides (10 sections apart) are analyzed. Brain sections are deparaffinized with xylene and hydrated in gradient alcohols. Immunohistochemical staining follows a basic indirect protocol, using antigen retrieval by heating to 95 °C in 0.01 mol/L citrate buffer for 30 min for antigen retrieval. To identify human cells in mouse brains, mAb to vimentin (1:50, clone 3B4, Dako Corporation), which identifies all human leukocytes is used. Human MDM and CD8⁺ lymphocytes are detected with CD68 (1:50 dilution, clone KP 1) and CD8 (1:50 dilution, clone 144B) antibodies, respectively. Virus-infected cells are labeled with mAb to HIV-1 p24 (1:10, clone Kal-1, all from Dako). Reactive murine microglial cells are detected with Iba-1 antibody (1:500, Wako). Expression of human IDO (huIDO) is visualized with Abs obtained from the Department of Cell Pharmacology, Central Research Institute, Graduate School of Medicine, Hokkaido University, Sapporo, Japan. Primary antibodies are detected with the appropriate biotinylated secondary antibodies and visualized with avidin-biotin complexes (Vectastain Elite ABC kit, Vector Laboratories) and horseradish peroxidase (HRP) coupled dextran polymer (EnVision, Dako Corporation). Immunostained sections are counterstained with Mayer's hematoxylin. Sections from which primary antibody is deleted or irrelevant IgG isotype is incorporated served as controls. Two independent observers in a blinded fashion count the numbers of CD8⁺ lymphocytes, CD68⁺ MDM and HIV-1 p24⁺ cells in each section from each mouse. Light microscopic examination is performed with a Nikon Eclipse 800 microscope (Nikon Instruments Inc). Semi-quantitative analysis for Iba1 (percentage of area occupied by immunostaining) is carried out by computer-assisted image analysis (Image-Pro®Plus, Media Cybernetics) as previously described.

5. Statistical analysis

Data can be analyzed using Prism (Graph Pad) with Student *t*-test for comparisons and ANOVA. *P*-values < 0.05 were considered significant.

6. Reference

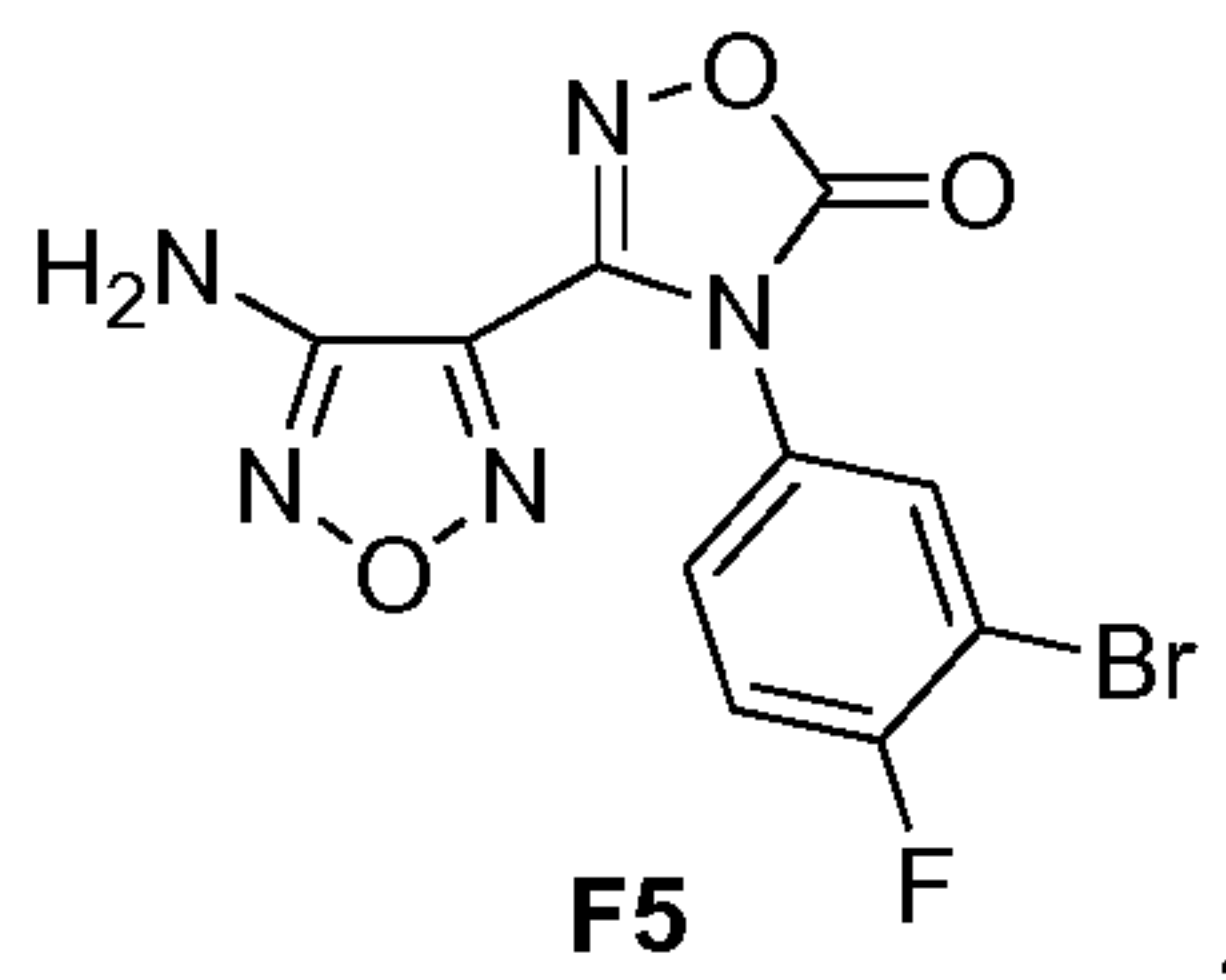
Poluektova LY, Munn DH, Persidsky Y, and Gendelman HE (2002). Generation of cytotoxic T cells against virus-infected human brain macrophages in a murine model of HIV-1 encephalitis. *J. Immunol.* 168(8):3941-9.

Various modifications of the invention, in addition to those described herein, will be apparent to those skilled in the art from the foregoing description. Such modifications are

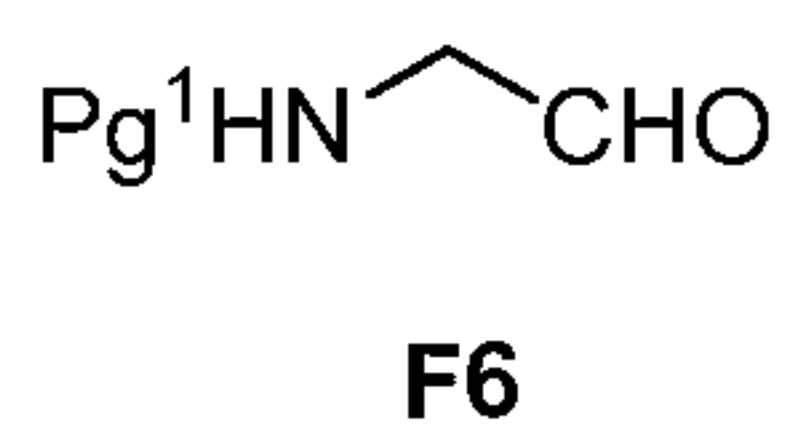
also intended to fall within the scope of the appended claims. Each reference, including all patent, patent applications, and publications, cited in the present application is incorporated herein by reference in their entirety.

What it claimed is:

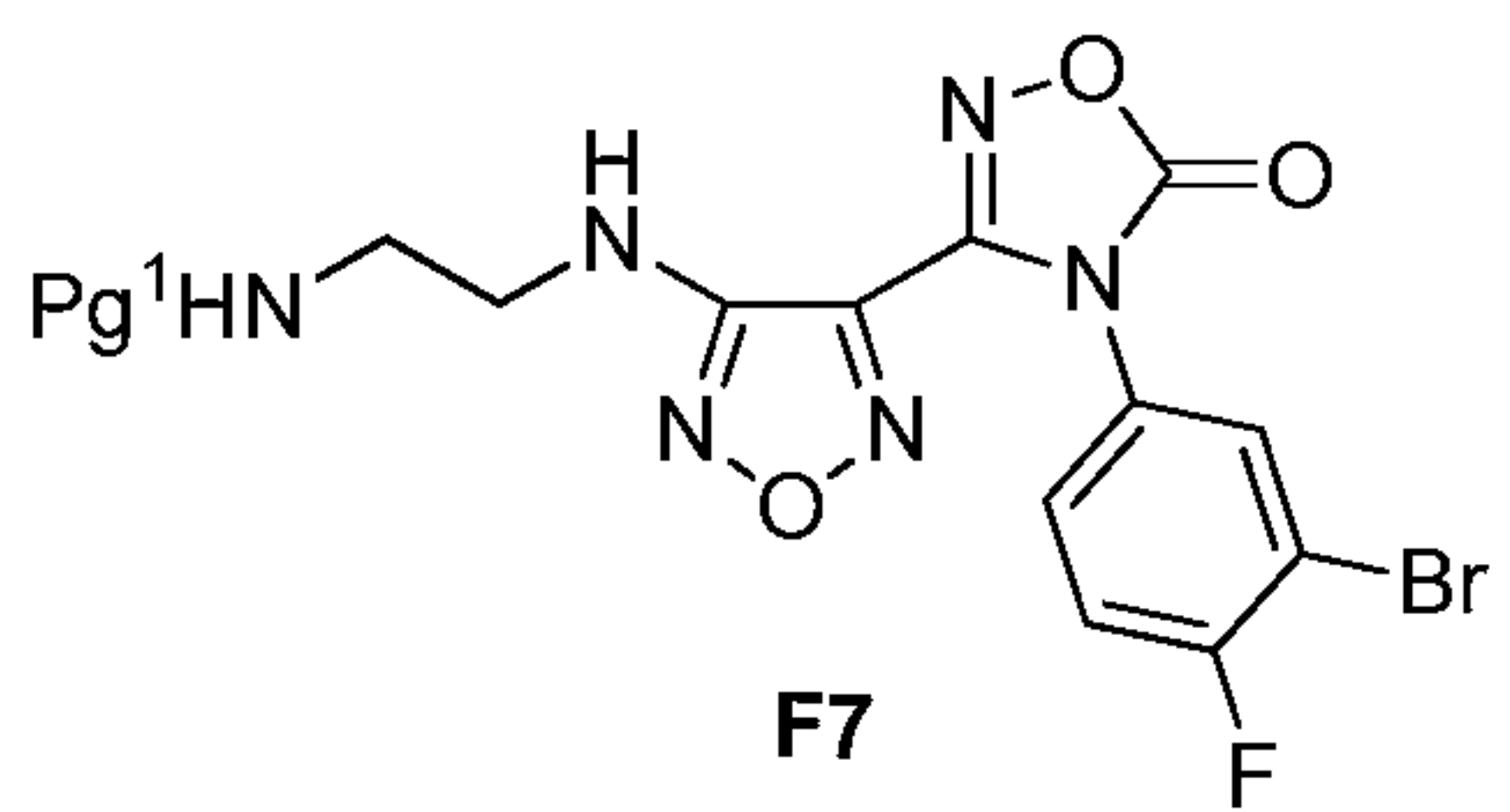
1. A process comprising reacting a compound of Formula F5:



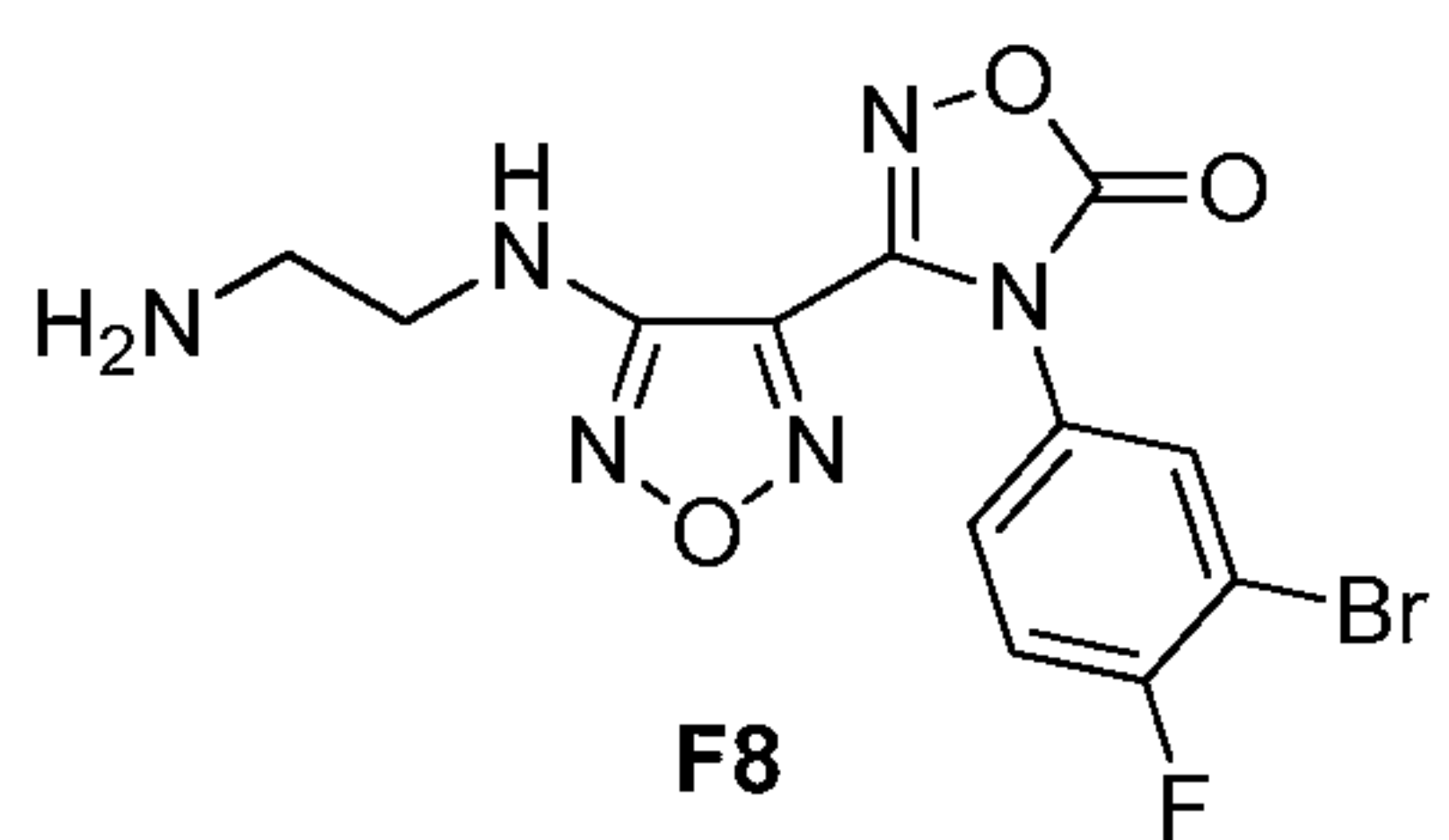
with an aldehyde of Formula F6:



wherein Pg¹ is an amino protecting group, to afford a compound of Formula F7:

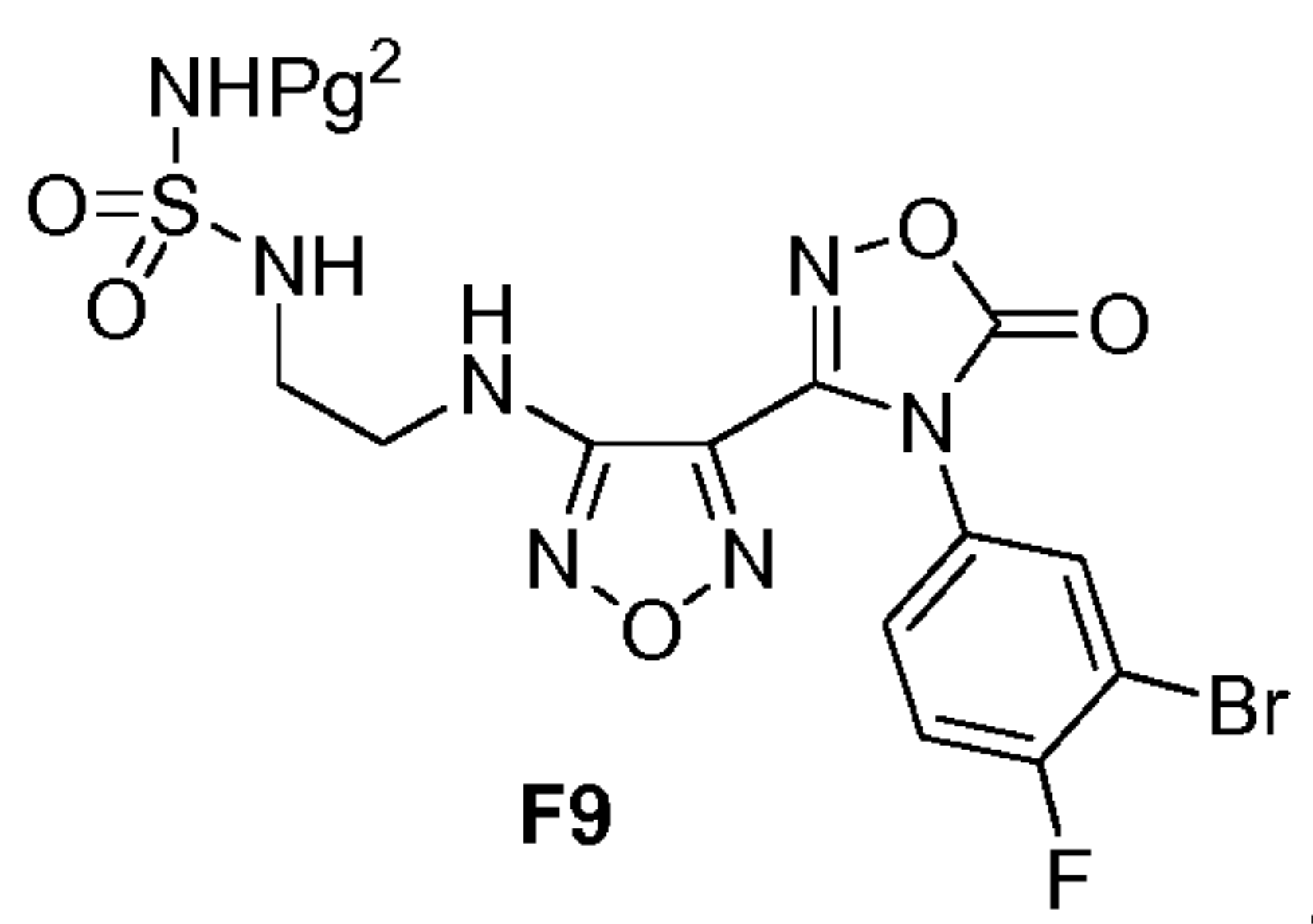


2. The process of claim 1, wherein Pg¹ is ethoxycarbonyl, *tert*-butoxycarbonyl, benzyloxycarbonyl, or 9-fluorenylmethyloxycarbonyl.
3. The process of claim 1, wherein Pg¹ is *tert*-butoxycarbonyl.
4. The process of any one of claims 1-3, wherein said reacting is performed in the presence of a reducing agent.
5. The process of claim 4, wherein said reducing agent is a borohydride reducing agent.
6. The process of claim 5, wherein said borohydride reducing agent is sodium triacetoxyborohydride.
7. The process of any one of claims 1-6, further comprising deprotecting said compound of Formula F7 to afford a compound of Formula F8:



8. The process of claim 7, wherein said deprotecting comprises reacting the compound of Formula F7 with hydrochloric acid.

9. The process of claim 7 or 8, further comprising reacting said compound of Formula F8, with $\text{Pg}^2\text{-NH-SO}_2\text{-X}$, in the presence of an organic base to afford a compound of Formula F9:



wherein:

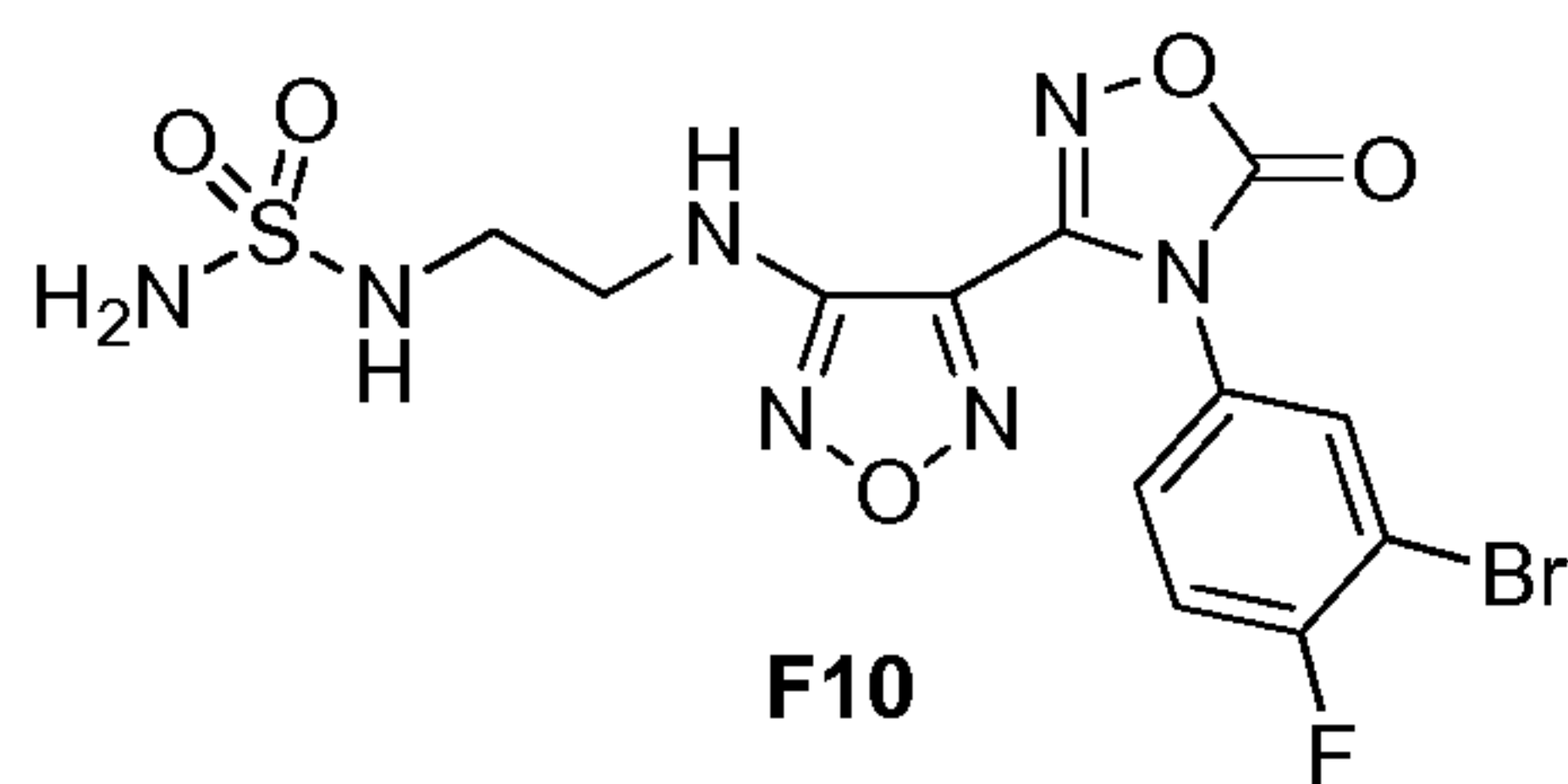
Pg^2 is an amino protecting group; and
X is halo.

10. The process of claim 9, wherein Pg^2 is ethoxycarbonyl, *tert*-butoxycarbonyl, benzyloxycarbonyl, or 9-fluorenylmethyloxycarbonyl.

11. The process of claim 9, wherein Pg^2 is *tert*-butoxycarbonyl.

12. The process of any one of claims 9-11, wherein X is chloro.

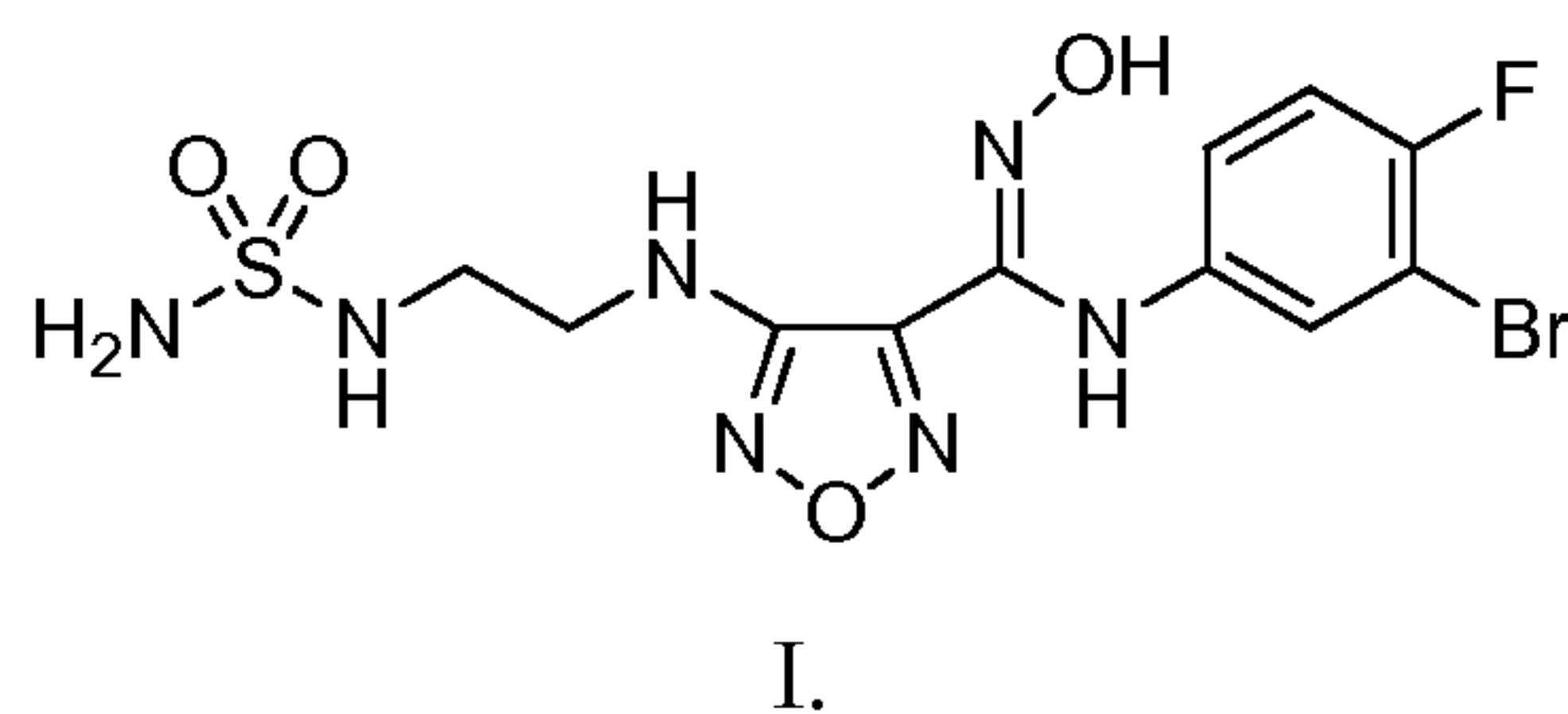
13. The process of any one of claims 9-12, further comprising deprotecting said compound of Formula F9 to afford a compound of Formula F10:



14. The process of claim 13, wherein said deprotecting comprises reacting a compound of Formula F9 with hydrochloric acid.

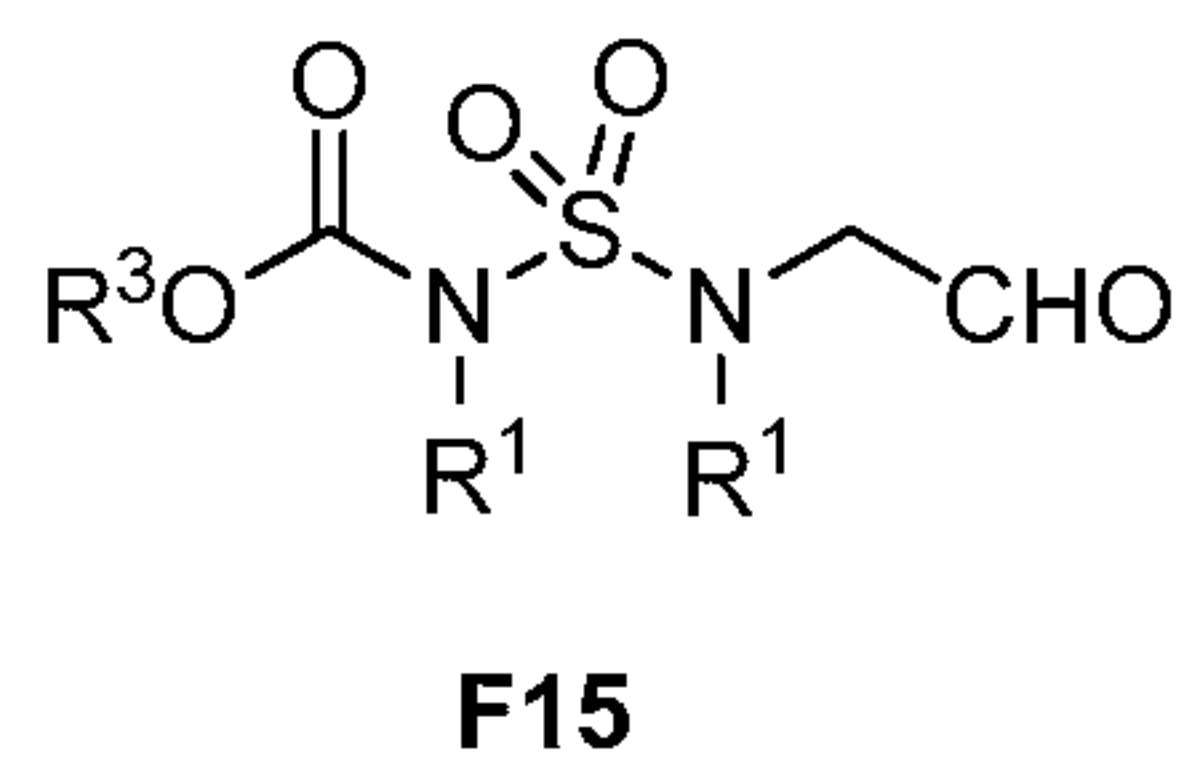
15. The process of claim 13, wherein said deprotecting comprises reacting a compound of Formula F9 with hydrochloric acid in a solvent component comprising ethyl acetate.

16. The process of any one of claims 13-15, further comprising reacting said compound of Formula F10 with a base to afford a compound of Formula I:

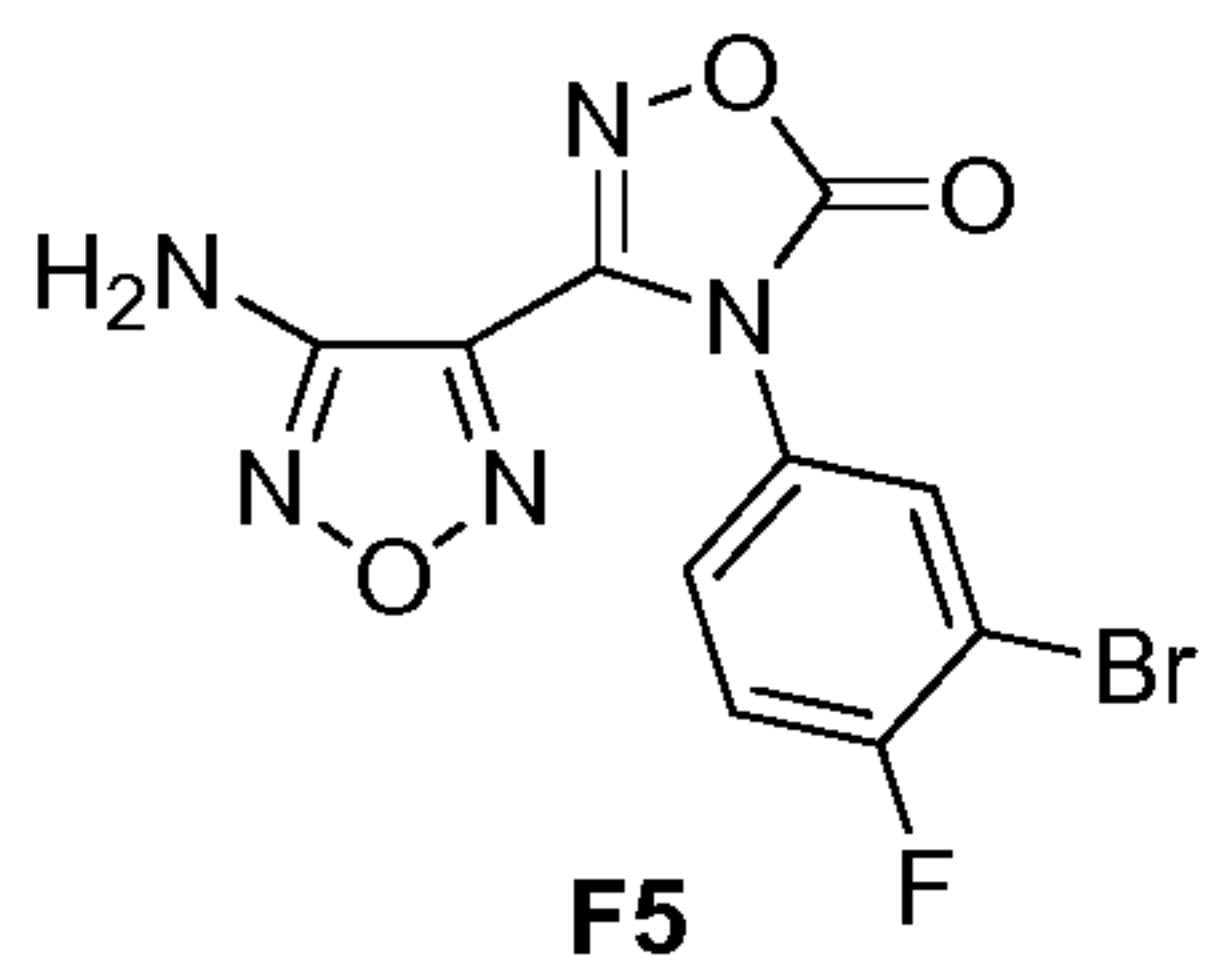


17. The process of claim 16, wherein said base is sodium hydroxide.

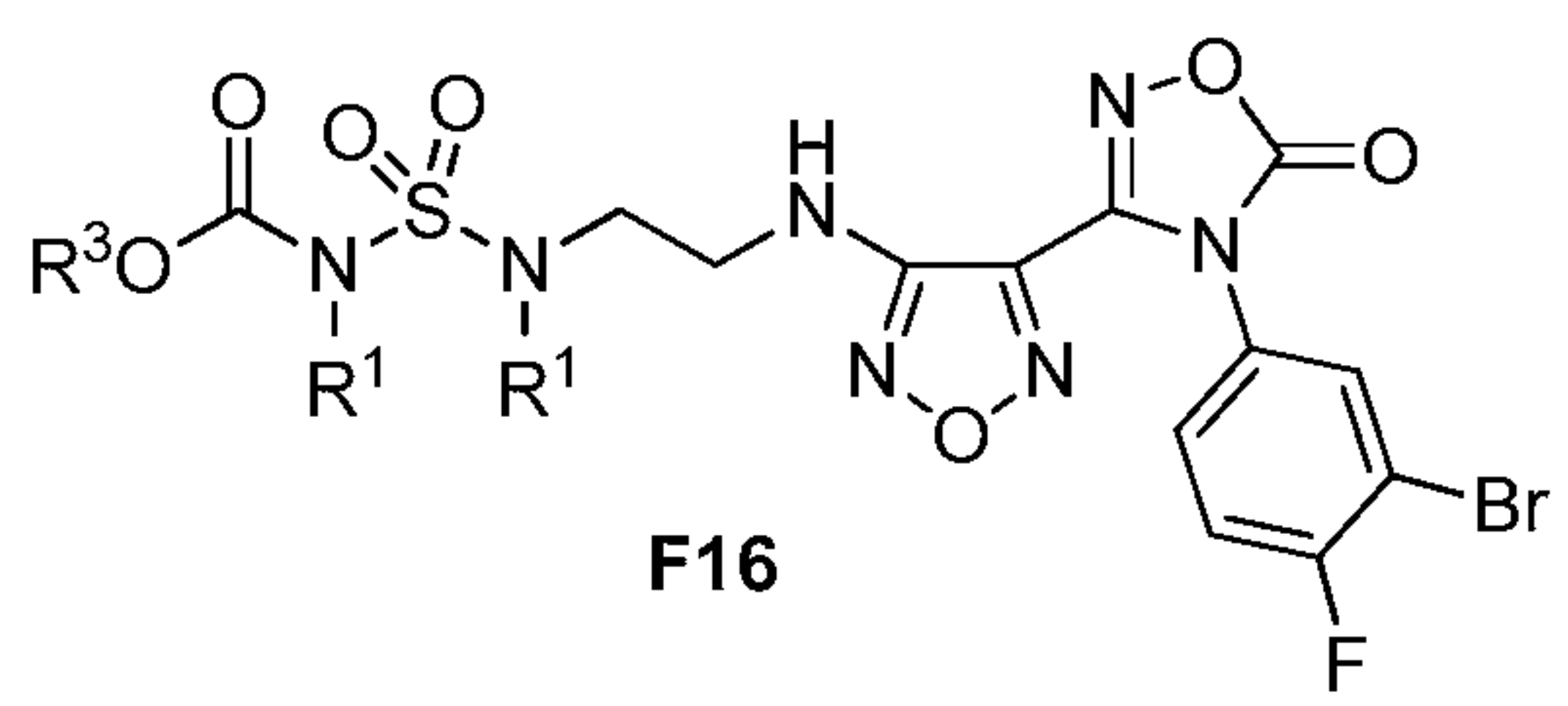
18. A process, comprising reacting a compound of Formula F15:



with a compound of Formula F5:



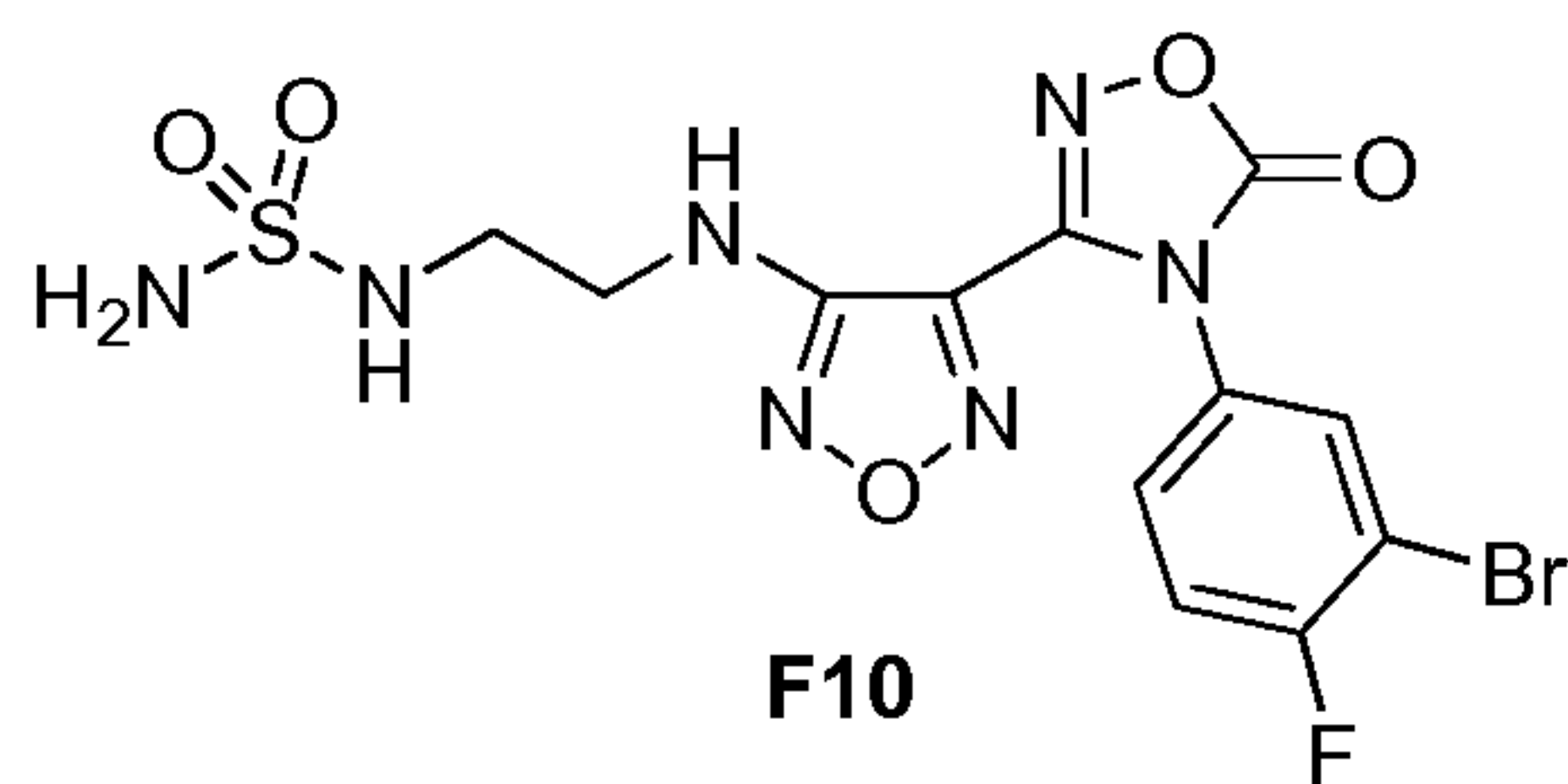
to afford a compound of Formula F16:



wherein:

each R^1 is independently an amino protecting group; and
 R^3 is C_{1-6} alkyl or benzyl.

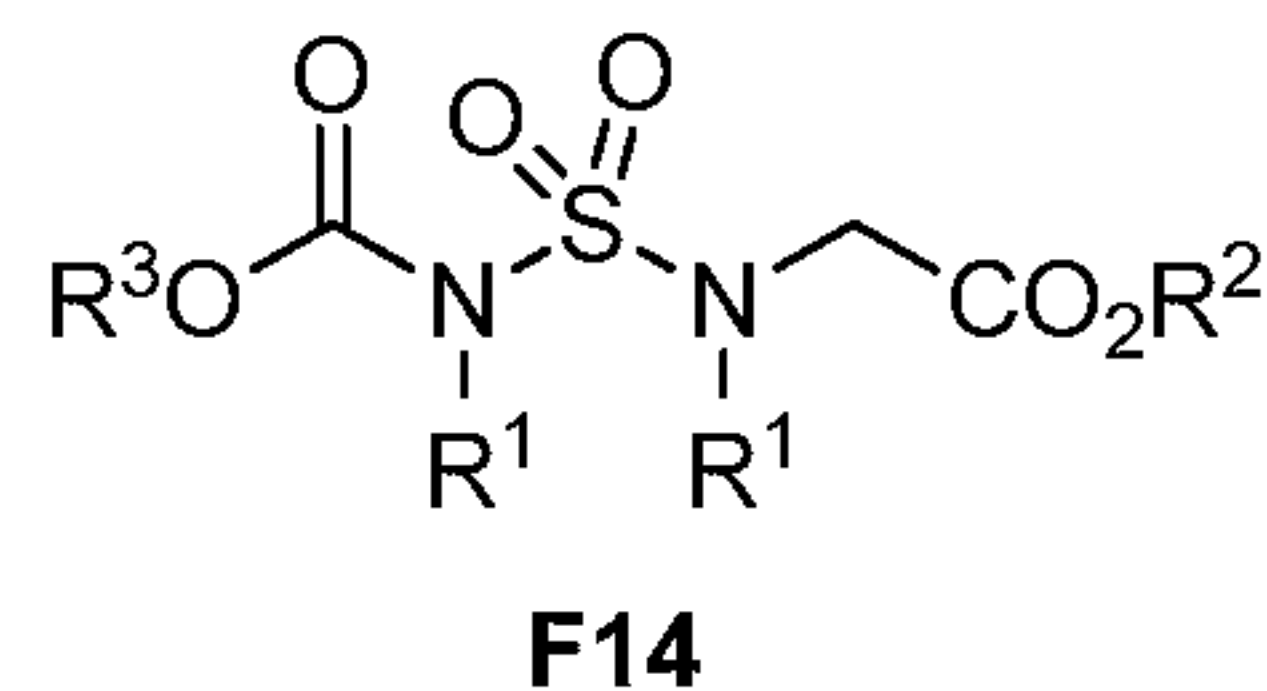
19. The process of claim 18, wherein each R^1 is C_{2-4} alkenyl- C_{1-3} alkyl or phenyl- C_{1-3} alkyl, wherein said phenyl- C_{1-3} alkyl is optionally substituted by 1, 2, or 3 independently selected C_{1-4} alkoxy groups.
20. The process of claim 18, wherein each R^1 is allyl.
21. The process of claim 18, wherein each R^1 is 4-methoxybenzyl.
22. The process of claim 18, wherein R^3 is C_{1-6} alkyl.
23. The process of claim 18, wherein R^3 is *tert*-butyl.
24. The process of any one of claims 18-23, wherein said reacting is performed in the presence of a reducing agent.
25. The process of claim 24, wherein said reducing agent is a borohydride reducing agent.
26. The process of claim 25, wherein said borohydride reducing agent is sodium triacetoxyborohydride.
27. The process of any one of claims 24-26, wherein said reacting is performed in the presence of trifluoroacetic acid.
28. The process of any one of claims 18-27, further comprising deprotecting said compound of Formula F16 to afford a compound of Formula F10:



29. The process of claim 28, wherein said deprotecting comprises reacting a compound of Formula F16 with trifluoroacetic acid.

30. The process of claim 28, wherein said deprotecting comprises reacting a compound of Formula F16 with is hydrochloric acid.

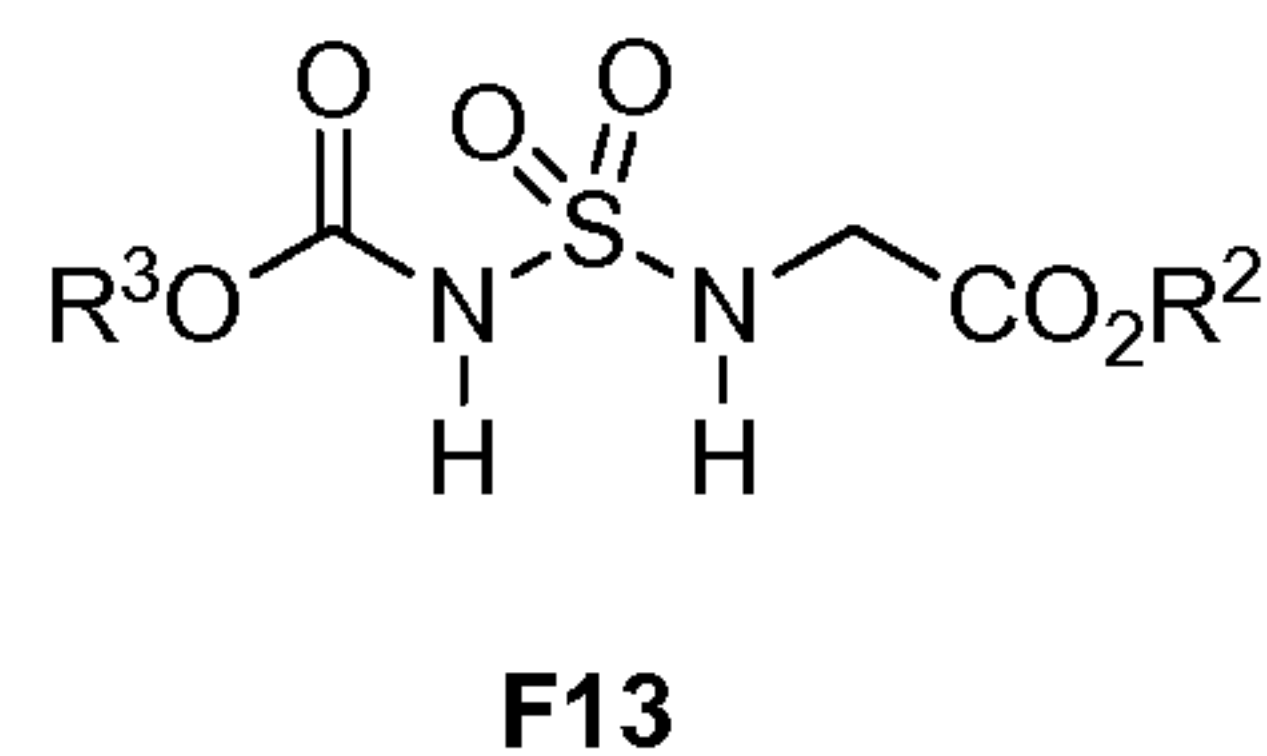
31. The process of any one of claims 18-30, wherein said compound of Formula F15 is obtained by a process comprising treating a compound of Formula F14:



with a reducing agent to afford said compound of Formula F15; wherein R² is C₁₋₄ alkyl.

32. The process of claim 31, wherein said reducing agent is diisobutylaluminum hydride.

33. The process of claim 31 or 32, wherein said compound of Formula F14 is obtained by a process comprising protecting a compound of Formula F13:

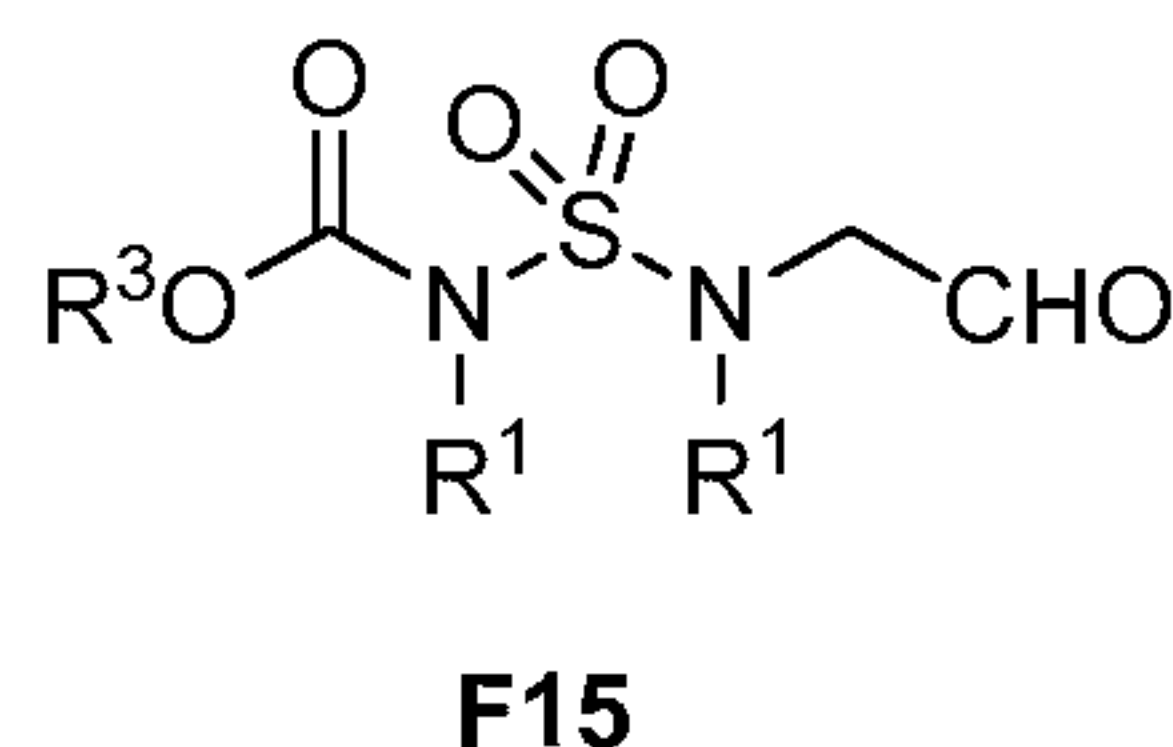


with one or more independently selected amino protecting agents to afford a compound of Formula F14.

34. The process of claim 33, wherein said one or more amino protecting agents is selected from allyl bromide and 4-methoxybenzyl chloride.

35. The process of claim 33 or 34, wherein said protecting is performed in the presence of a base.

36. A compound of Formula F15:



wherein:

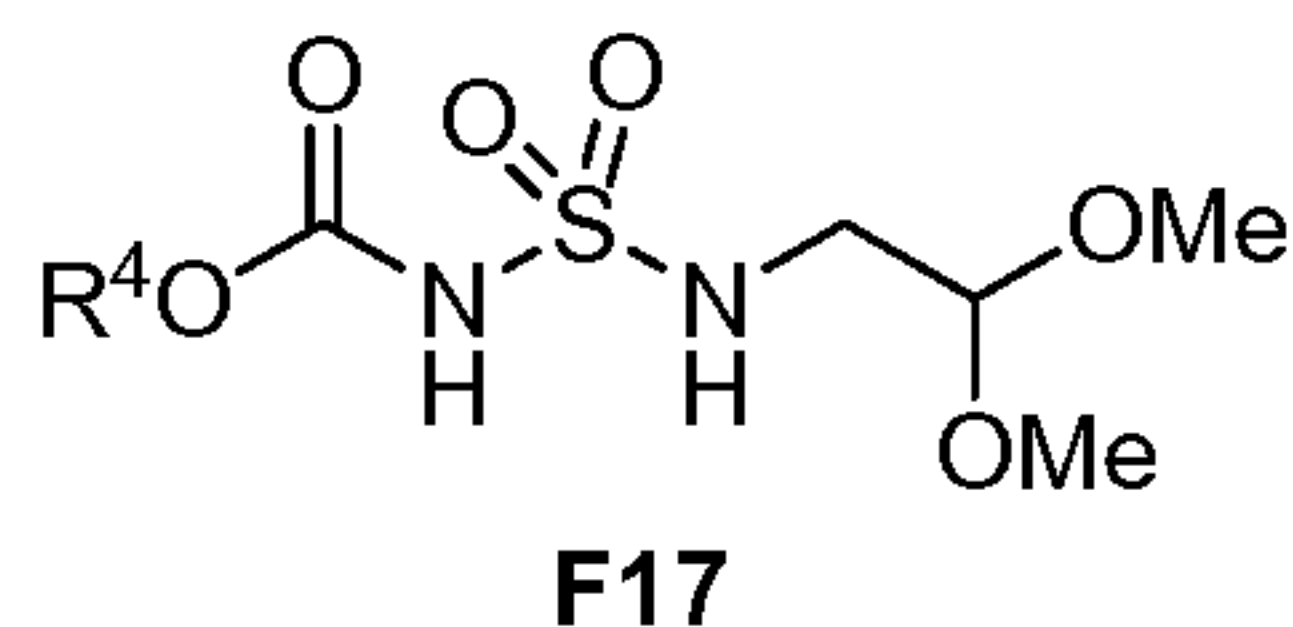
R^3 is C_{1-6} alkyl or benzyl; and

each R^1 is independently an amino protecting group.

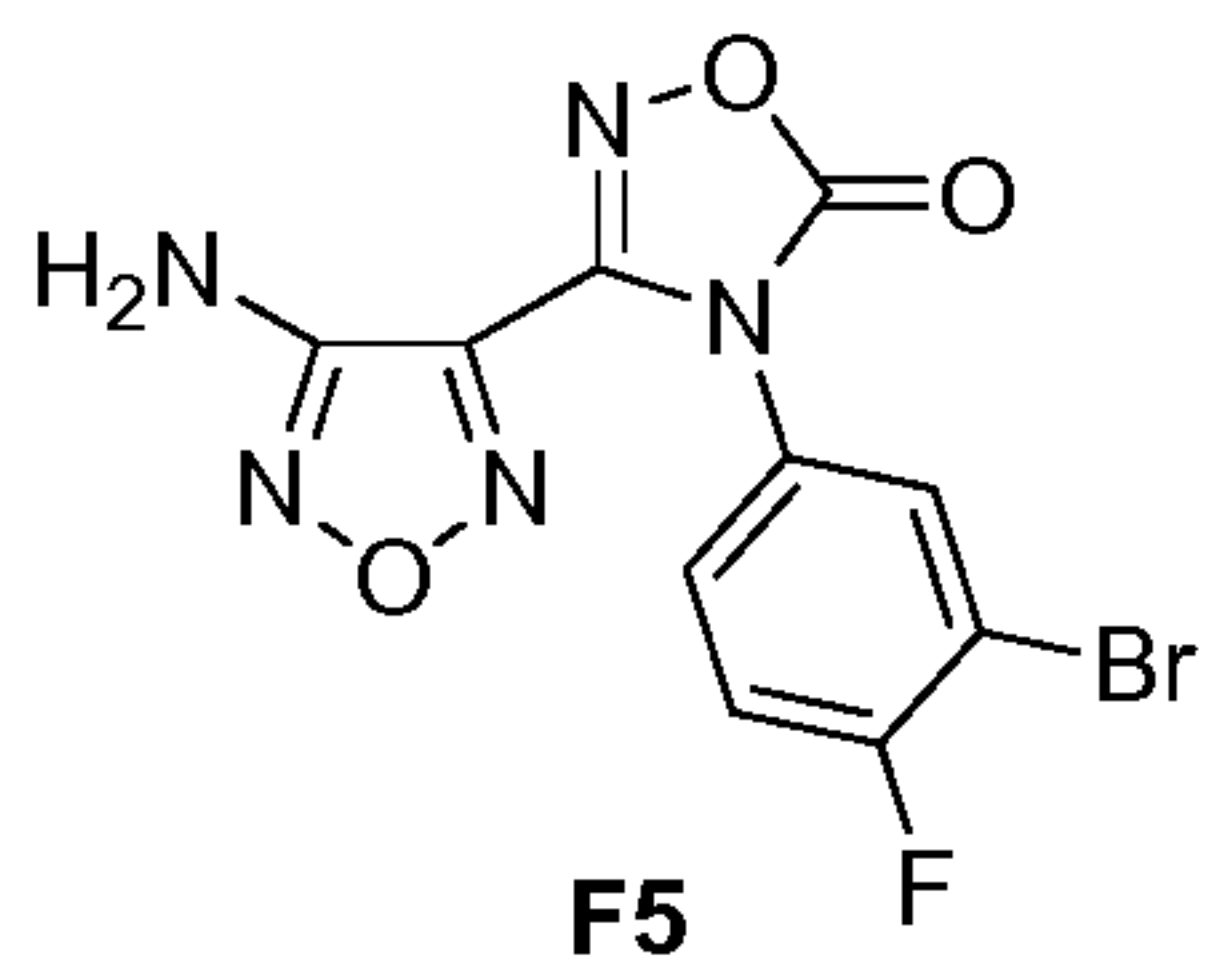
37. The compound of claim 36, which is *tert*-butyl allyl {[allyl(2-oxoethyl)amino]sulfonyl}carbamate.

38. The compound of claim 36, which is *tert*-butyl(4-methoxybenzyl){[(4-methoxybenzyl)(2-oxoethyl)amino]sulfonyl}carbamate.

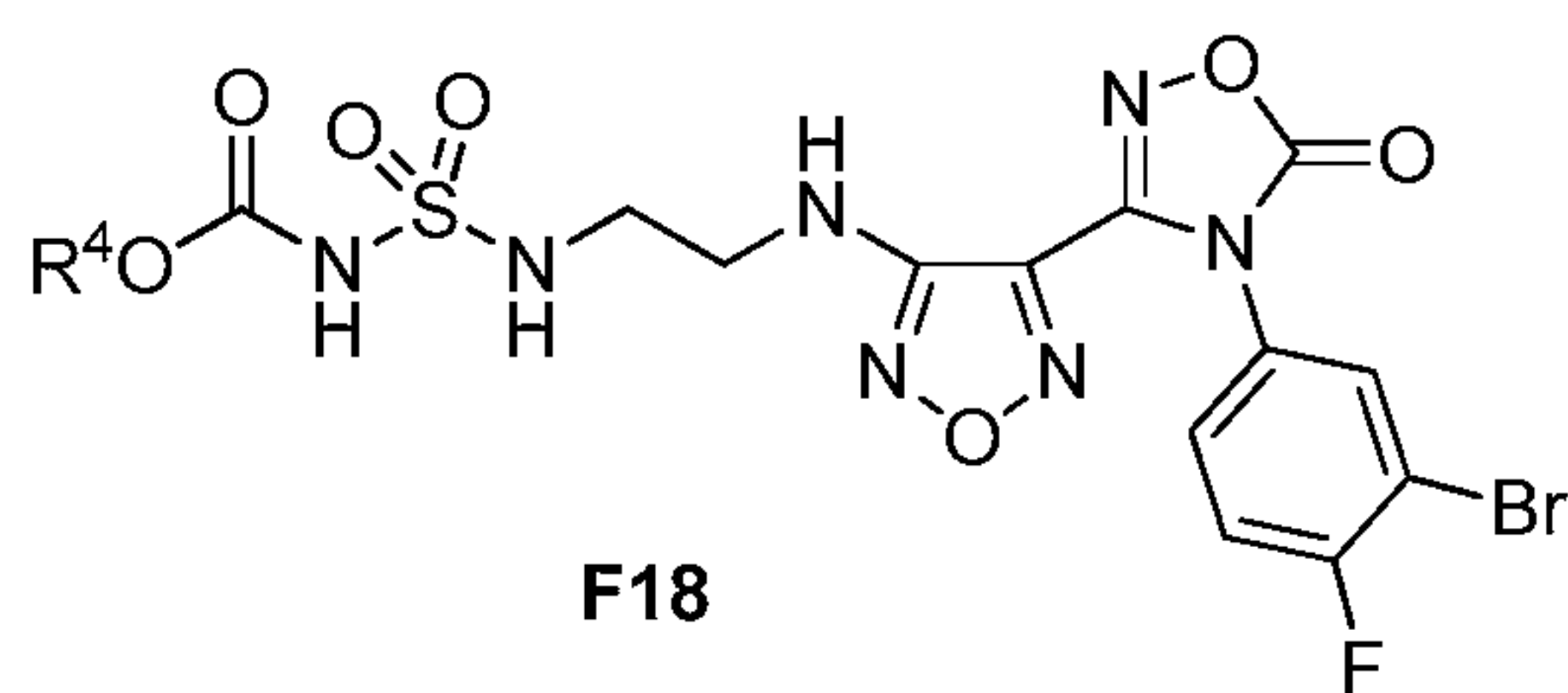
39. A process, comprising reacting a compound of Formula F17:



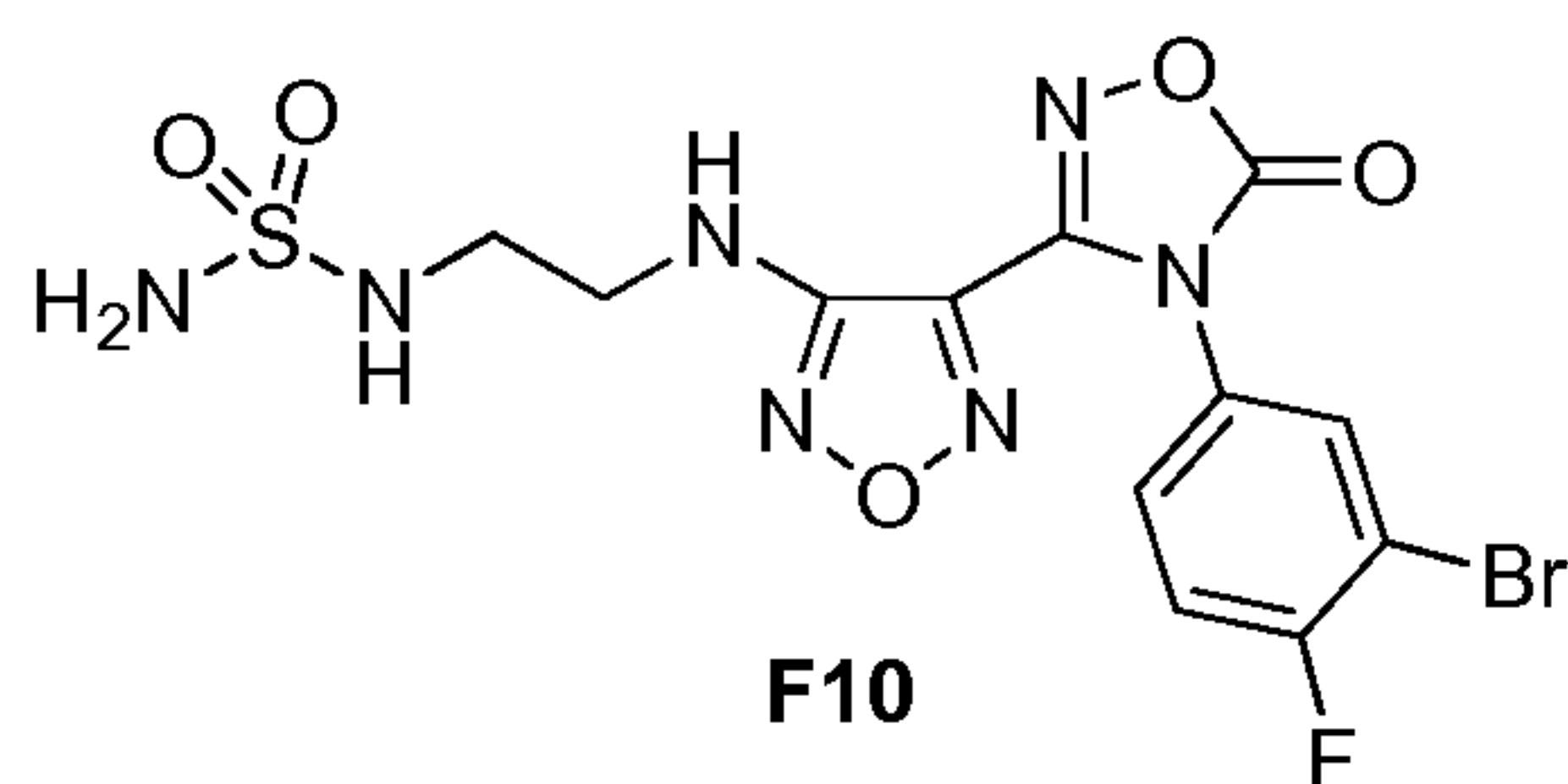
wherein R^4 is C_{1-6} alkyl, C_{1-6} haloalkyl, benzyl, or 9*H*-fluoren-9-ylmethyl with a compound of Formula F5:



to afford a compound of Formula F18:

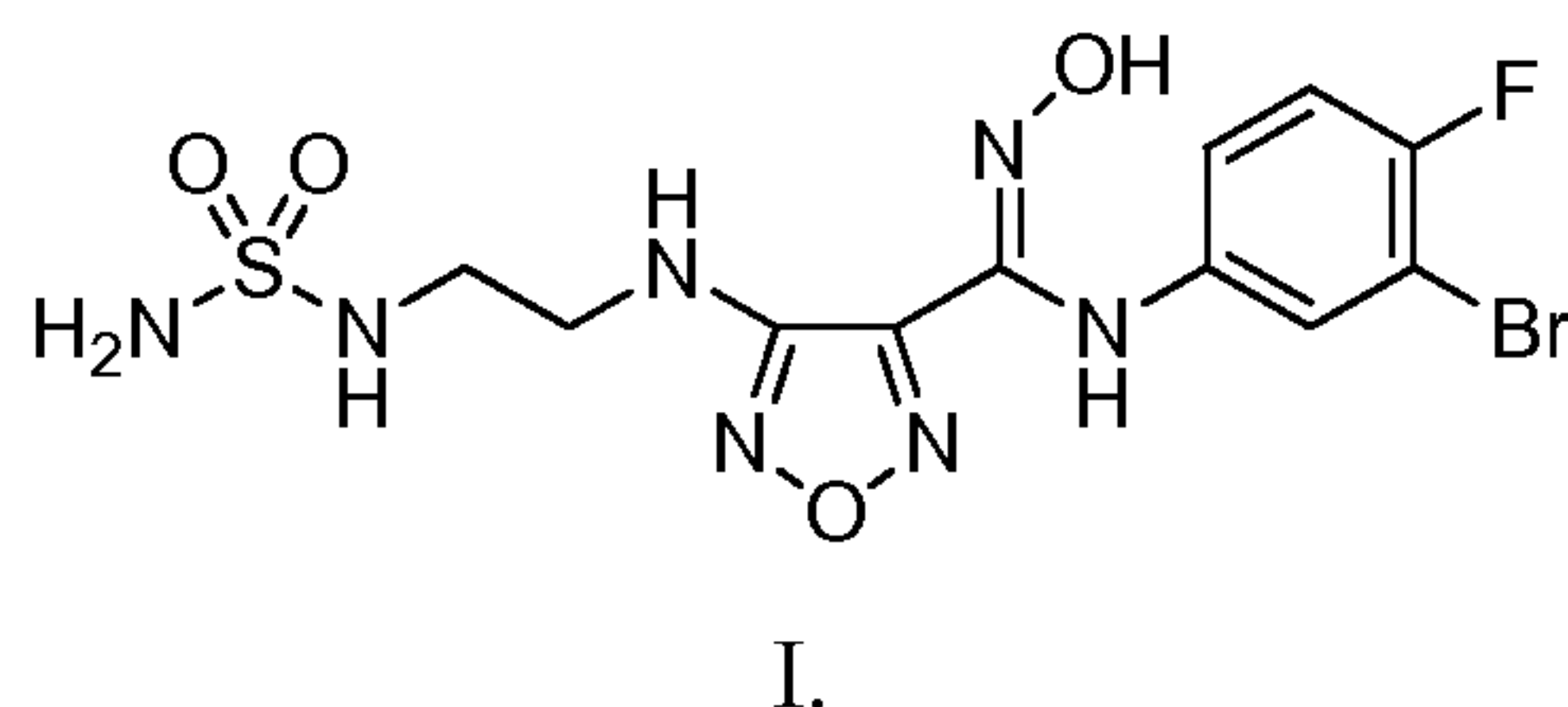


40. The process of claim 39, wherein R⁴ is *tert*-butyl.
41. The process of claim 39, wherein R⁴ is benzyl.
42. The process of claim 39, wherein R⁴ is ethyl.
43. The process of claim 39, wherein R⁴ is 2,2,2-trichloroethyl.
44. The process of any one of claims 40-43, wherein said reacting is carried out in the presence of a reducing agent.
45. The process of claim 44, wherein said reducing agent is triethylsilane.
46. The process of claim 44 or 45, wherein said reacting is carried out in the presence of an organic acid.
47. The process of claim 46, wherein said organic acid is trifluoroacetic acid.
48. The process of any one of claims 40-47, further comprising deprotecting said compound of Formula F18 to afford a compound of Formula F10:



49. The process of claim 48, wherein said deprotecting comprises reacting the compound of Formula F18 with zinc in the presence of acetic acid.

50. The process of claim 48 or 49, further comprising reacting said compound of Formula F10 with a base to afford a compound of Formula I:



51. The process of claim 50, wherein said base is sodium hydroxide.

52. The process of claim 39, wherein R^4 is 9H-fluoren-9-ylmethyl.

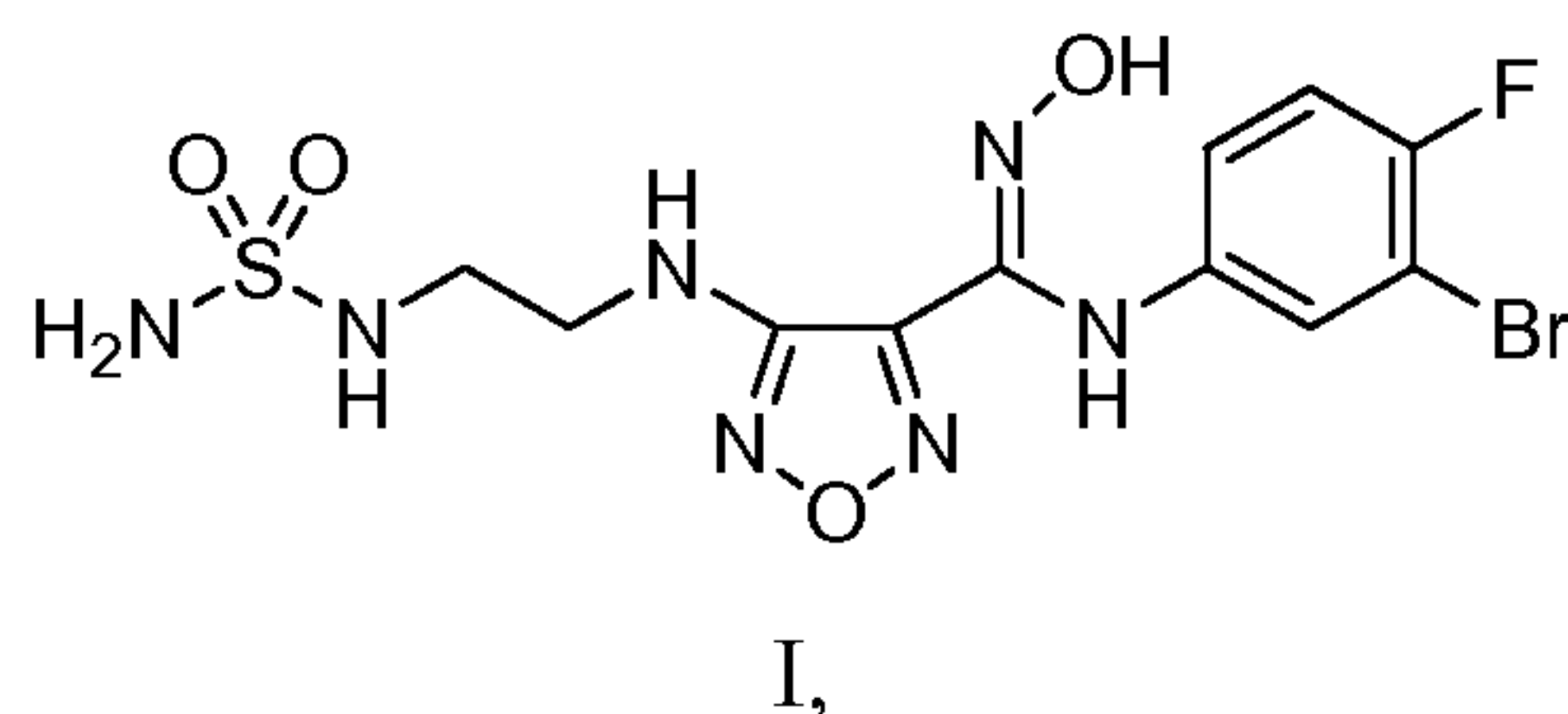
53. The process of claim 52, wherein said reacting is carried out in the presence of a reducing agent.

54. The process of claim 53, wherein said reducing agent is triethylsilane.

55. The process of claim 53 or 54, wherein said reacting is carried out in the presence of an organic acid.

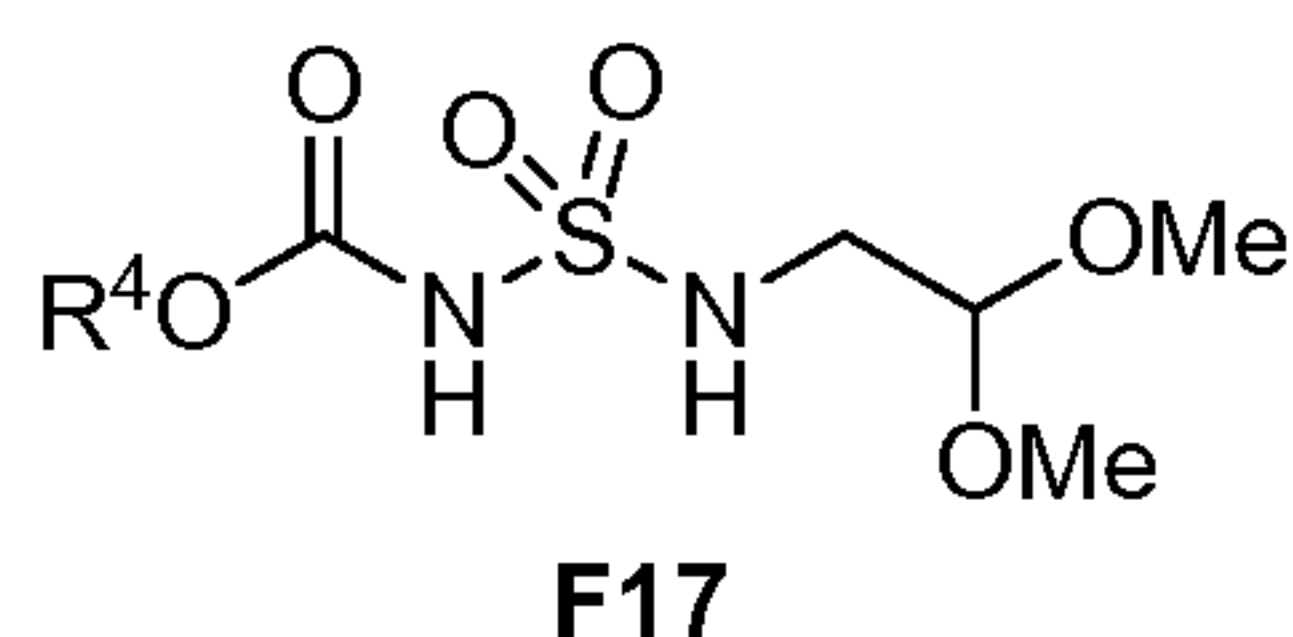
56. The process of claim 55, wherein said organic acid is methanesulfonic acid.

57. The process of any one of claims 52-56, further comprising converting said compound of Formula F18 to a compound of Formula I:



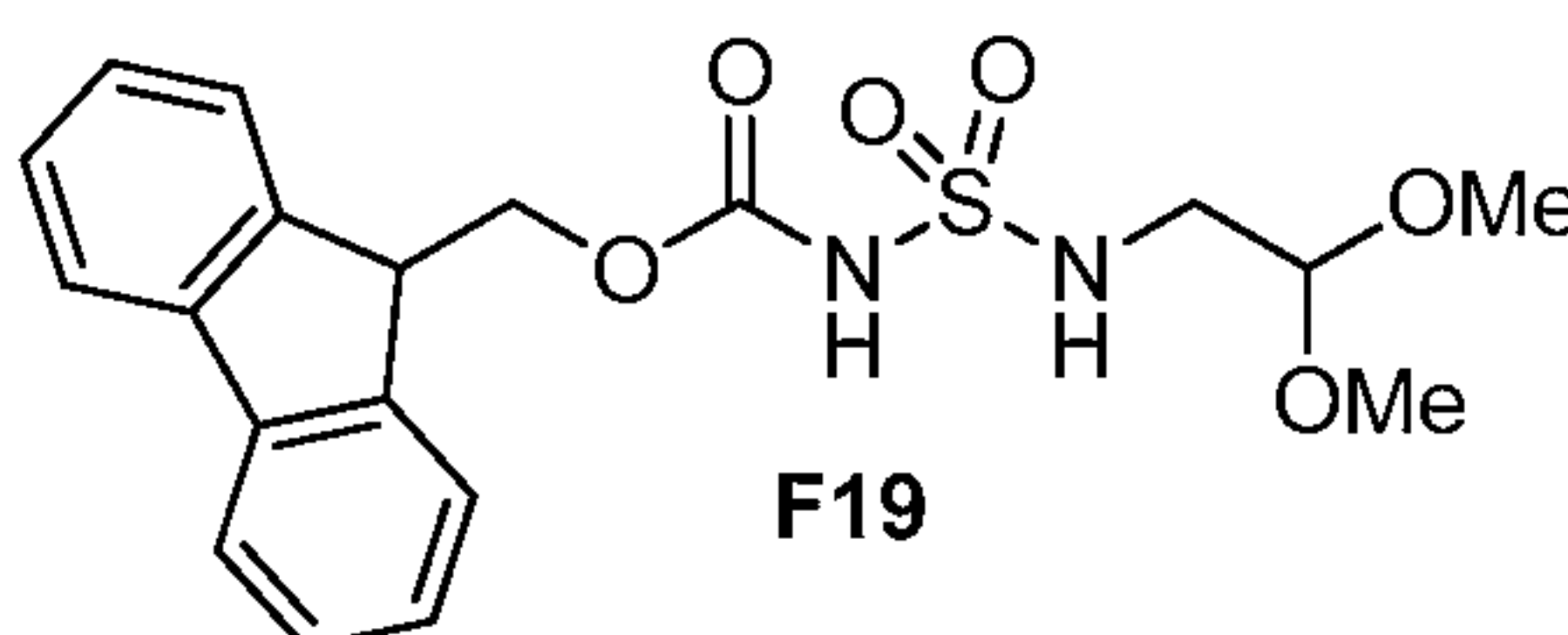
wherein said converting comprises combining the compound of Formula F18 with a base to form a first mixture.

58. The process of claim 57, wherein said base is *N,N*-bis(2-aminoethyl)ethane-1,2-diamine.
59. The process of any one of claims 57 to 58, wherein said converting further comprises adding aqueous hydrochloric acid to said first mixture.
60. A compound of Formula F17:

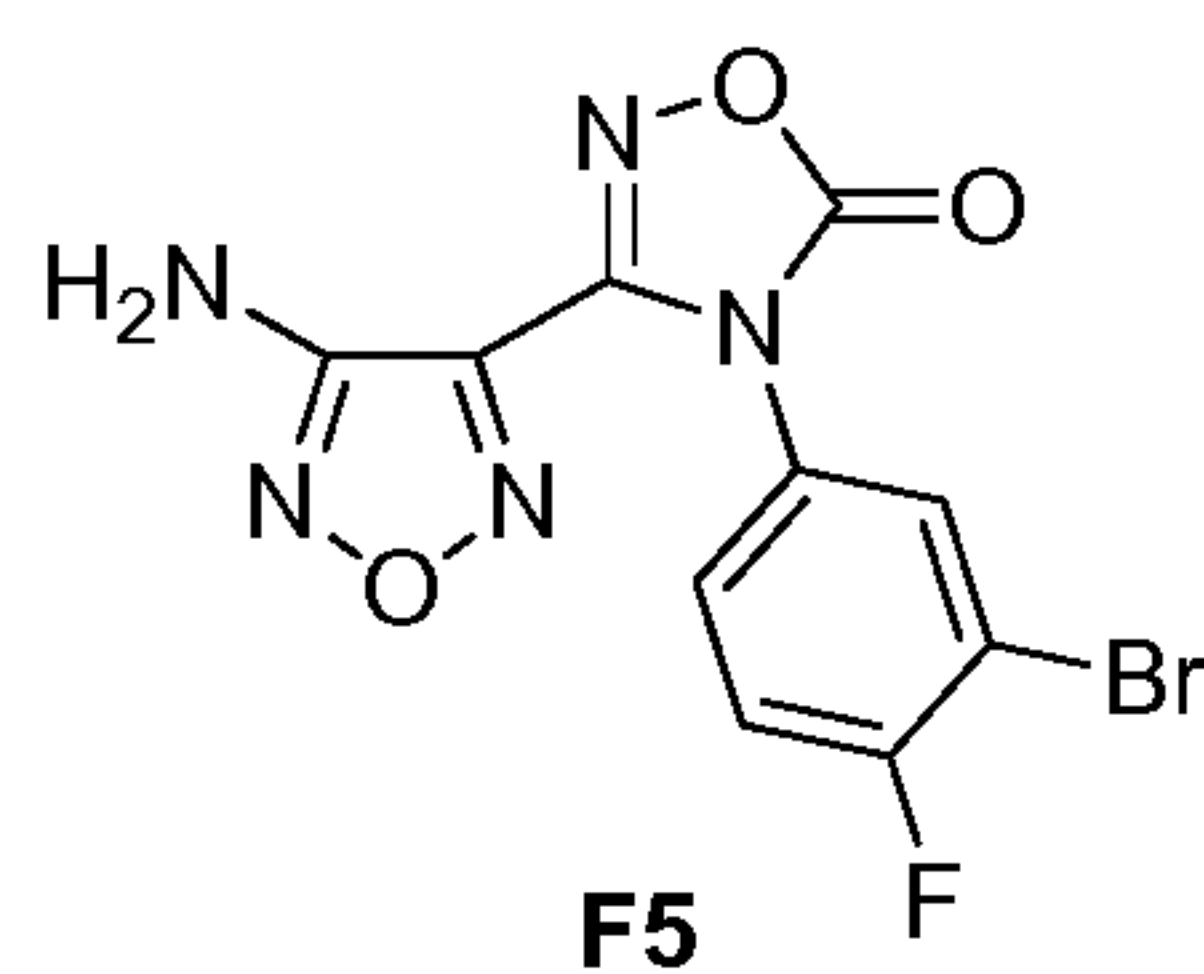


wherein R⁴ is C₁₋₆ alkyl, C₁₋₆ haloalkyl, benzyl, or 9*H*-fluoren-9-ylmethyl.

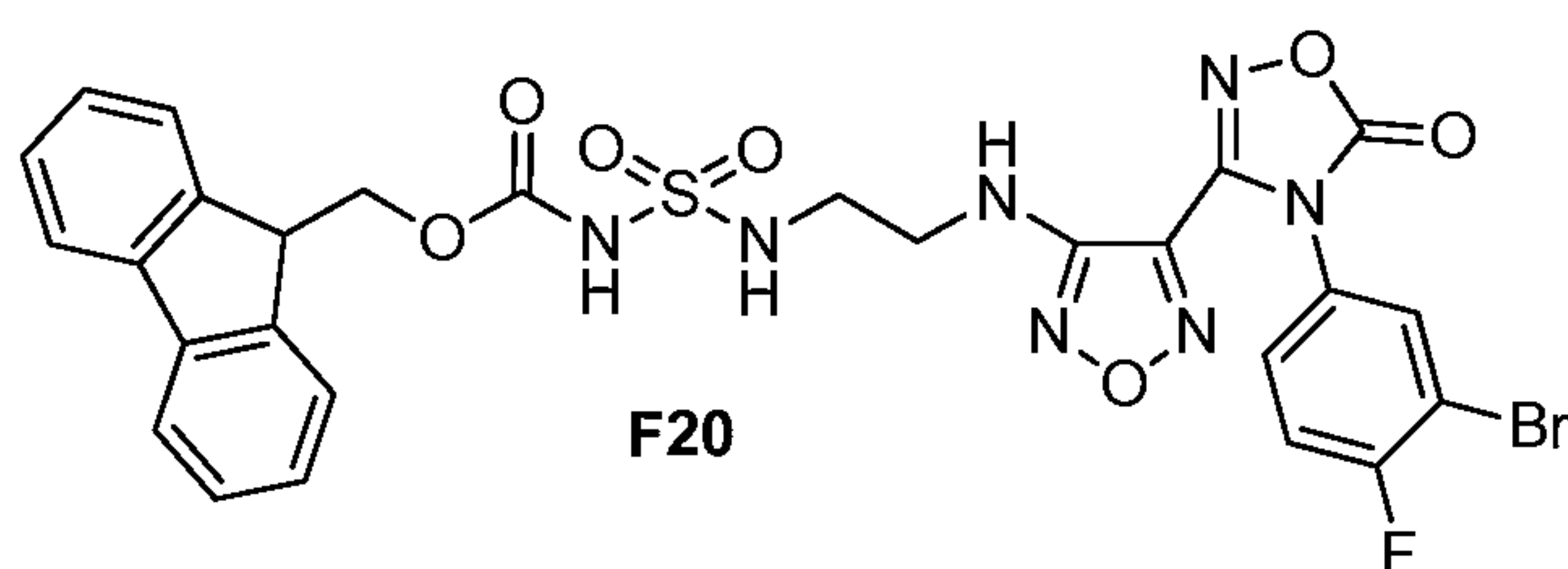
61. The compound of claim 60, which is tert-butyl *N*-(2,2-dimethoxyethyl)sulfamoylcarbamate.
62. The compound of claim 60, which is benzyl *N*-(2,2-dimethoxyethyl)sulfamoylcarbamate.
63. The compound of claim 60, which is ethyl *N*-(2,2-dimethoxyethyl)sulfamoylcarbamate.
64. The compound of claim 60, which is 2,2,2-trichloroethyl *N*-(2,2-dimethoxyethyl)sulfamoylcarbamate.
65. The compound of claim 60, which is (9*H*-fluoren-9-yl)methyl *N*-(2,2-dimethoxyethyl)sulfamoylcarbamate.
66. A process, comprising:
i) reacting a compound of Formula F19:



with a compound of Formula F5:

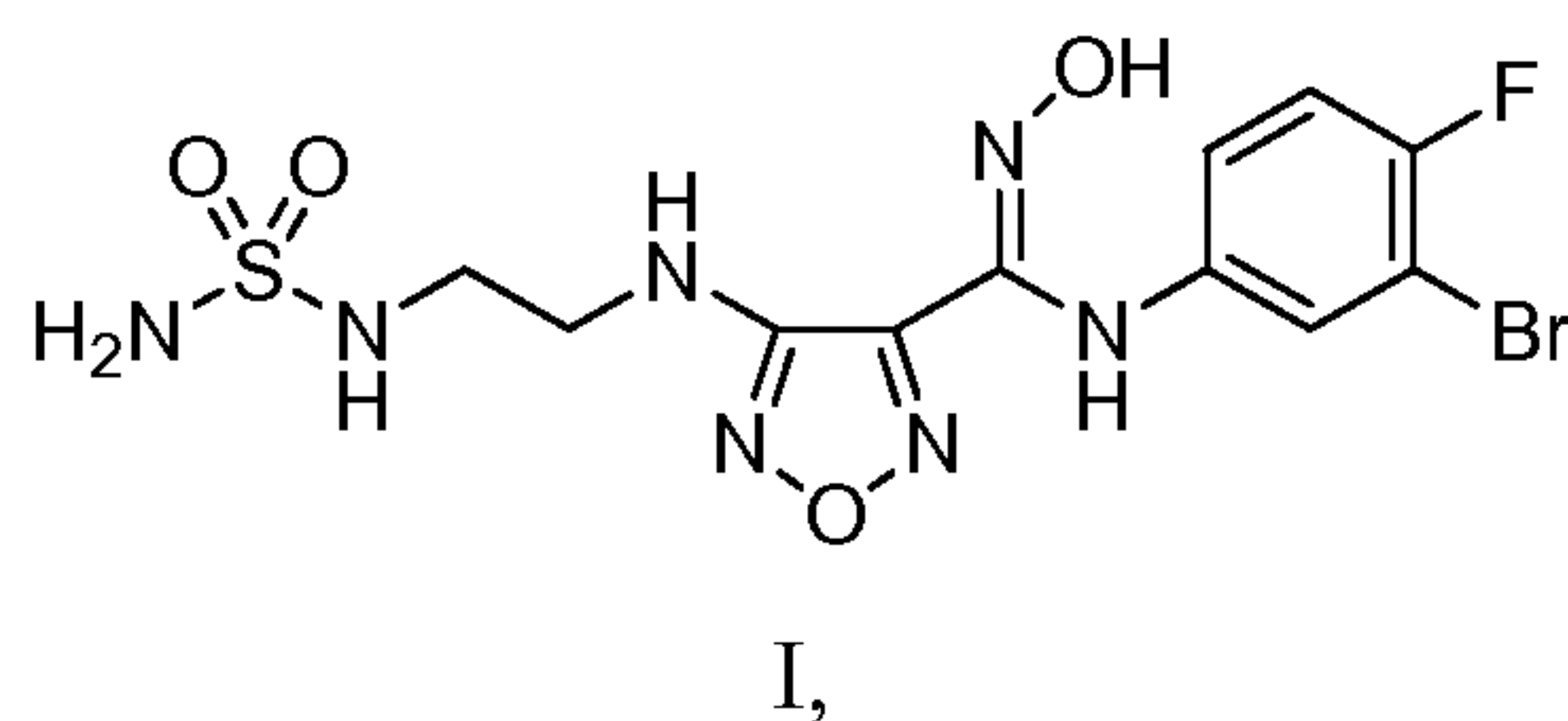


in the presence of triethylsilane and methanesulfonic acid to afford a compound of Formula F20:



and

ii) converting said compound of Formula F20 to a compound of Formula I:



wherein said converting comprises combining said compound of Formula F20 with N,N-bis(2-aminoethyl)ethane-1,2-diamine.