

TRIAZOLONE COMPOUNDS AS mPGES-1 INHIBITORS

Related Applications

This application claims the benefit of Indian Provisional Application Nos. 1733/MUM/2012 filed on 15 June 2012; 3319/MUM/2012 filed on 19 November 2012; and 387/MUM/2013 filed on 08 February 2013; and U.S. Provisional Application Nos. 61/668,146 filed on 05 July 2012; 61/735,679 filed on 11 December 2012; and 61/792,225 filed on 15 March 2013, each of which is hereby incorporated by reference in its entirety.

Technical Field

The present application relates to triazolone compounds as mPGES-1 inhibitors.

Background of the Invention

There are many diseases or disorders that are inflammatory in their nature. One of the major problems associated with existing treatments of inflammatory conditions is inadequate efficacy and/or the prevalence of side effects. Inflammatory diseases that affect the population include asthma, inflammatory bowel disease, rheumatoid arthritis, osteoarthritis, rhinitis, conjunctivitis and dermatitis. Inflammation is also a common cause of pain.

The enzyme cyclooxygenase (COX) converts arachidonic acid to an unstable intermediate, prostaglandin H_2 (PGH_2), which is further converted to other prostaglandins, including PGE_2 , $PGF_{2\alpha}$, PGD_2 , prostacyclin and thromboxane A_2 . These arachidonic acid metabolites are known to have pronounced physiological and pathophysiological activity, including pro-inflammatory effects. The COX enzyme exists in two forms, one that is constitutively expressed in many cells and tissues (COX-1), and another that in most cells and tissues is induced by pro-inflammatory stimuli, such as cytokines, during an inflammatory response (COX-2).

Among all prostaglandin metabolites, PGE_2 is particularly known to be a strong pro-inflammatory mediator, and is also known to induce fever and pain. Consequently, numerous drugs have been developed with a view to inhibiting the formation of PGE_2 ,

including “NSAIDs” (non-steroidal anti-inflammatory drugs) and “coxibs” (selective COX-2 inhibitors). These drugs act predominantly by inhibition of COX-1 and/or COX-2, thereby reducing the formation of PGE₂. However, the inhibition of COXs has the disadvantage of reducing the formation of all metabolites of PGH₂, thereby decreasing the beneficial properties of some of the metabolites. In view of this, drugs which act by inhibition of COXs are suspected to cause adverse biological effects. For example, the non-selective inhibition of COXs by NSAIDs may give rise to gastrointestinal side-effects and affect platelet and renal function. Even the selective inhibition of COX-2 by coxibs, whilst reducing such gastrointestinal side-effects, is believed to give rise to cardiovascular problems.

A combination of pharmacological, genetic and neutralizing antibody approaches demonstrates the importance of PGE₂ in inflammation. The conversion of PGH₂ to PGE₂ by prostaglandin E synthases (PGES) may, therefore, represent a pivotal step in the propagation of inflammatory stimuli. There are two microsomal prostaglandin E synthases (mPGES-1 and mPGES-2), and one cytosolic prostaglandin E synthase (cPGES). mPGES-1 is an inducible PGES after exposure to pro-inflammatory stimuli. mPGES-1 is induced in the periphery and CNS by inflammation, and represents therefore a target for acute and chronic inflammatory disorders. PGE₂ is a major prostanoid, produced from arachidonic acid liberated by phospholipases (PLAs), which drives the inflammatory processes. Arachidonic acid is transformed by the action of prostaglandin H synthase (PGH synthase, cyclooxygenase) into PGH₂ which is a substrate for mPGES-1, the terminal enzyme transforming PGH₂ to the pro-inflammatory PGE₂.

Agents that are capable of inhibiting the action of mPGES-1, and thus reducing the formation of the specific arachidonic acid metabolite PGE₂, are beneficial in the treatment of inflammation. Further, agents that are capable of inhibiting the action of the proteins involved in the synthesis of the leukotrienes are also beneficial in the treatment of asthma and COPD.

Blocking the formation of PGE₂ in animal models of inflammatory pain results in reduced inflammation, pain and fever response (Kojima et. al, *The Journal of Immunology* **2008**, 180, 8361-6; Xu et. al., *The Journal of Pharmacology and Experimental Therapeutics* **2008**, 326, 754-63). In abdominal aortic aneurism,

inflammation leads to connective tissue degradation and smooth muscle apoptosis ultimately leading to aortic dilation and rupture. In animals lacking mPGES-1 a slower disease progression and disease severity has been demonstrated (Wang et. al., *Circulation*, **2008**, 117, 1302-1309).

Several lines of evidence indicate that PGE₂ is involved in malignant growth. PGE₂ facilitates tumor progression by stimulation of cellular proliferation and angiogenesis and by modulation of immunosuppression. In support of a role for PGE₂ in cancers, genetic deletion of mPGES-1 in mice suppresses intestinal tumorigenesis (Nakanishi et. al., *Cancer Research* **2008**, 68(9), 3251-9). In human beings, mPGES-1 is also upregulated in cancers such as colorectal cancer (Schroder, *Journal of Lipid Research* **2006**, 47, 1071-80).

Myositis is a chronic muscle disorder characterized by muscle weakness and fatigue. Proinflammatory cytokines and prostanoids have been implicated in the development of myositis. In skeletal muscle tissue from patients suffering from myositis an increase in cyclooxygenases and mPGES-1 has been demonstrated, implicating mPGES-1 as a target for treating this condition. (Korotkova, *Annals of the Rheumatic Diseases* **2008**, 67, 1596- 1602).

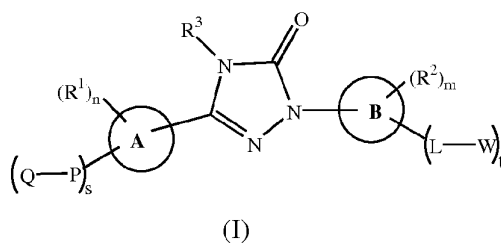
In atherosclerosis, inflammation of the vasculature leads to atheroma formation that eventually may progress into infarction. In patients with carotid atherosclerosis an increase in mPGES-1 in plaque regions has been reported (Gomez-Hernandez *Atherosclerosis* **2006**, 187, 139-49). In an animal model of atherosclerosis, mice lacking the mPGES-1 receptor were found to show a retarded atherogenesis and a concomitant reduction in macrophage-derived foam cells together with an increase in vascular smooth muscle cells (Wang, *Proceedings of National Academy of Sciences* **2006**, 103(39), 14507-12).

International Publication Nos. WO 2006/063466, WO 2007/059610, WO 2010/034796, WO 2010/100249, WO 2012/055995, WO 2012/110860 and WO 2013/038308 disclose numerous heterocyclic compounds which are stated to be inhibitors of the microsomal prostaglandin E synthase- 1 (mPGES-1) enzyme.

The present application is directed to compounds that act as inhibitors of the mPGES-1 enzyme and, therefore, are useful for the treatment of pain and inflammation in a variety of diseases or conditions.

Summary of the Invention

In one aspect, the present invention relates to a compound of formula (I)



or a pharmaceutically acceptable salt thereof,

wherein,

ring A is selected from C₆₋₁₄aryl, 5-14 membered heteroaryl, C₃₋₁₂cycloalkyl and 3-15 membered heterocyclyl;

ring B is selected from C₆₋₁₄aryl, 5-14 membered heteroaryl, C₃₋₁₂cycloalkyl and 3-15 membered heterocyclyl;

each occurrence of L is independently selected from -(CR^xR^y)_qNR^xC(O)-, and -(CR^xR^y)_qNR^xS(O)₂-;

each occurrence of P is independently selected from -(CR^xR^y)_qNR^xC(O)-, and -(CR^xR^y)_qNR^xS(O)₂-;

each occurrence of Q is independently selected from C₁₋₈alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₁₋₈alkoxy, C₁₋₈alkoxyC₁₋₈alkyl, haloC₁₋₈alkyl, hydroxyC₁₋₈alkyl, carboxylC₁₋₈alkyl, C₃₋₁₂cycloalkyl, C₆₋₁₄aryl, C₆₋₁₄arylC₁₋₈alkyl, 3-15 membered heterocyclyl, and 5-14 membered heteroaryl;

each occurrence of W is independently selected from C₁₋₈alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₁₋₈alkoxy, C₁₋₈alkoxyC₁₋₈alkyl, haloC₁₋₈alkyl, hydroxyC₁₋₈alkyl, carboxylC₁₋₈alkyl, C₃₋₁₂cycloalkyl, C₆₋₁₄aryl, C₆₋₁₄arylC₁₋₈alkyl, 3-15 membered heterocyclyl, and 5-14 membered heteroaryl;

each occurrence of R¹ is independently selected from halogen, cyano, C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxyC₁₋₈alkyl, haloC₁₋₈alkoxyC₁₋₈alkyl, haloC₁₋₈alkyl, haloC₁₋₈alkoxy,

hydroxyC₁₋₈alkyl, C₃₋₁₂cycloalkyl, C₃₋₈cycloalkylC₁₋₈alkyl, C₆₋₁₄aryl, C₆₋₁₄arylC₁₋₈alkyl, C₆₋₁₄aryloxy, 3-15 membered heterocyclyl, 3-15 membered heterocyclylC₁₋₈alkyl, 5-14 membered heteroaryl, 5-14 membered heteroarylC₁₋₈alkyl, -C(O)NHR, -S(O)₂NHR, -NHC(O)R, -CH₂S(O)NHR and -C≡CR;

each occurrence of R² is independently selected from halogen, cyano, C₁₋₈alkyl, C₁₋₈alkoxy, C₁₋₈alkoxyC₁₋₈alkyl, haloC₁₋₈alkyl, haloC₁₋₈alkoxy, hydroxyC₁₋₈alkyl, C₃₋₁₂cycloalkyl, C₃₋₈cycloalkylC₁₋₈alkyl, C₆₋₁₄aryl, C₆₋₁₄arylC₁₋₈alkyl, C₆₋₁₄aryloxy, 3-15 membered heterocyclyl, 3-15 membered heterocyclylC₁₋₈alkyl, 5-14 membered heteroaryl, 5-14 membered heteroarylC₁₋₈alkyl, -NHR, -C(O)NHR, -S(O)₂NHR, -NHC(O)R, -CH₂S(O)NHR and -C≡CR;

each occurrence of R is independently selected from C₁₋₈alkyl, C₃₋₁₂cycloalkyl, C₃₋₈cycloalkylC₁₋₈alkyl, C₆₋₁₄aryl, C₆₋₁₄arylC₁₋₈alkyl, 3-15 membered heterocyclyl and 5-14 membered heteroaryl;

R³ is independently selected from hydrogen, C₁₋₈alkyl, C₃₋₁₂cycloalkyl and C₆₋₁₄aryl;

at each occurrence, R^x and R^y, which may be the same or different, are independently selected from hydrogen, C₁₋₈alkyl, C₃₋₁₂cycloalkyl and C₆₋₁₄aryl; or R^x and R^y together with the common atom to which they are attached, form a cyclic ring which is substituted or unsubstituted and wherein the cyclic ring optionally contains one or more hetero atoms selected from O, N or S;

‘m’ is an integer ranging from 0 to 3, both inclusive;

‘n’ is an integer ranging from 0 to 3, both inclusive;

‘q’ is an integer ranging from 1 to 4, both inclusive;

‘s’ is an integer ranging from 0 to 1, both inclusive; and

‘t’ is an integer ranging from 0 to 1, both inclusive;

with the proviso that ‘m’ and ‘n’ are not ‘0’ simultaneously.

The compounds of formula (I) may involve one or more embodiments. Embodiments of formula (I) include compounds of formula (II), compounds of formula (III) and compounds of formula (IV) as described hereinafter. It is to be understood that the embodiments below are illustrative of the present invention and are not intended to limit the claims to the specific embodiments exemplified. It is also to be understood that

the embodiments defined herein may be used independently or in conjunction with any definition of any other embodiment defined herein. Thus, the invention contemplates all possible combinations and permutations of the various independently described embodiments. For example, the invention provides compounds of formula (I) as defined above, wherein R^3 is hydrogen (according to an embodiment defined below), 'n' is 0, 1 or 2 (according to another embodiment defined below), and 's' is 1 and 't' is 0 (according to another embodiment defined below).

According to one embodiment, specifically provided are compounds of formula (I), in which ring A is C_{6-14} aryl (e.g. phenyl) or 5-14 membered heteroaryl (e.g. pyridinyl).

According to another embodiment, specifically provided are compounds of formula (I), in which ring A is phenyl or pyridinyl.

According to yet another embodiment, specifically provided are compounds of formula (I), in which ring B is C_{6-14} aryl (e.g. phenyl), 5-14 membered heteroaryl (e.g. pyridinyl) or C_{3-12} cycloalkyl (e.g. cyclopentyl or cyclohexyl).

According to yet another embodiment, specifically provided are compounds of formula (I), in which ring B is phenyl, pyridinyl, cyclopentyl or cyclohexyl.

According to yet another embodiment, specifically provided are compounds of formula (I), in which R^3 is hydrogen.

According to yet another embodiment, specifically provided are compounds of formula (I), in which R^3 is C_{1-8} alkyl (e.g. methyl or ethyl).

According to yet another embodiment, specifically provided are compounds of formula (I), in which R^3 is hydrogen, methyl or ethyl.

According to yet another embodiment, specifically provided are compounds of formula (I), in which P is $-(CR^xR^y)_qNR^xC(O)-$ or $-(CR^xR^y)_qNR^xS(O)_2-$. In this embodiment, R^x and R^y are independently selected from hydrogen and C_{1-4} alkyl (e.g. methyl or ethyl), and 'q' is 1.

According to yet another embodiment, specifically provided are compounds of formula (I), in which P is $-CH_2NHC(O)-$ or $-CH_2NHS(O)_2-$.

According to yet another embodiment, specifically provided are compounds of formula (I), in which L is $-(CR^xR^y)_qNR^xC(O)-$ or $-(CR^xR^y)_qNR^xS(O)_2-$. In this

embodiment, R^x and R^y are independently selected from hydrogen and C_{1-4} alkyl (e.g. methyl or ethyl), and 'q' is 1.

According to yet another embodiment, specifically provided are compounds of formula (I), in which L is $-CH_2NHC(O)-$ or $-CH_2NHS(O)_2-$.

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R^1 is independently selected from cyano, halogen (e.g. F, Cl or Br), C_{1-8} alkyl (e.g. methyl), halo C_{1-8} alkyl (e.g. trifluoromethyl or difluoromethyl), C_{1-8} alkoxy (e.g. methoxy), halo C_{1-8} alkoxy C_{1-8} alkyl (e.g. (2,2,2-trifluoroethoxy)methyl), C_{3-12} cycloalkyl (e.g. cyclopropyl), 3-15 membered heterocyclyl C_{1-8} alkyl (e.g. (pyrrolidin-1-yl)methyl), 5-14 membered heteroaryl (e.g. 4-methylthiophenyl or 5-isopropyl-1,3,4-oxadiazol-2-yl), 5-14 membered heteroaryl C_{1-8} alkyl (e.g. (3,5-dimethyl-1*H*-pyrazol-1-yl)methyl), $-S(O)_2NHR$, $-NHCOR$, $-CONHR$ and $-C\equiv CR$.

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R^1 is independently selected from $-S(O)_2NHR$, $-NHCOR$, $-CONHR$ and $-C\equiv CR$. In this embodiment R is independently selected from C_{1-8} alkyl (e.g. isopropyl or *tert*-butyl), C_{3-12} cycloalkyl (e.g. cyclopropyl) and C_{6-14} aryl (e.g. 3,5-difluorophenyl, 4-(trifluoromethyl)phenyl, 3-(trifluoromethyl)phenyl, 4-fluoro-3-(trifluoromethyl)phenyl, 2-fluoro-5-(trifluoromethyl)phenyl, 2-chloro-4-methylphenyl, 2,5-dichlorophenyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 2-chloro-4-(trifluoromethyl)phenyl, 3-(difluoromethyl)-4-fluorophenyl or 3-(difluoromethyl)phenyl).

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R^1 is independently cyano, F, Cl, Br, methyl, trifluoromethyl, difluoromethyl, methoxy, (2,2,2-trifluoroethoxy)methyl, cyclopropyl, (pyrrolidin-1-yl)methyl, 4-methylthiophenyl, 5-isopropyl-1,3,4-oxadiazol-2-yl, (3,5-dimethyl-1*H*-pyrazol-1-yl)methyl, $-S(O)_2NHR$, $-NHCOR$, $-CONHR$ and $-C\equiv CR$. In this embodiment R is independently selected from isopropyl, *tert*-butyl, cyclopropyl, 3,5-difluorophenyl, 4-(trifluoromethyl)phenyl, 3-(trifluoromethyl)phenyl, 4-fluoro-3-(trifluoromethyl)phenyl, 2-fluoro-5-(trifluoromethyl)phenyl, 2-chloro-4-methylphenyl, 2,5-dichlorophenyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 2-chloro-4-

(trifluoromethyl)phenyl, 3-(difluoromethyl)-4-fluorophenyl and 3-(difluoromethyl)phenyl.

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R¹ is independently cyano, F, Cl, Br, methyl, trifluoromethyl, difluoromethyl, methoxy, (2,2,2-trifluoroethoxy)methyl, cyclopropyl, (pyrrolidin-1-yl)methyl, 4-methylthiophenyl, 5-isopropyl-1,3,4-oxadiazol-2-yl, (3,5-dimethyl-1*H*-pyrazol-1-yl)methyl, *N*-cyclopropylsulfamoyl, 4-(trifluoromethyl)benzamide, 3,5-difluorobenzamide, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, (2,5-dichlorophenyl)ethynyl, (4-chloro-2-fluorophenyl)ethynyl, (3-chloro-2-fluorophenyl)ethynyl, [2-chloro-4-(trifluoromethyl)phenyl]ethynyl, -CONH-[4-(trifluoromethyl)phenyl], -CONH-[3-(trifluoromethyl)phenyl], -CONH-[3-(difluoromethyl)phenyl], -CONH-[4-fluoro-3-(trifluoromethyl)phenyl], -CONH-[2-fluoro-5-(trifluoromethyl)phenyl] or -CONH-[2-chloro-4-methylphenyl].

According to yet another embodiment, specifically provided are compounds of formula (I), in which 'n' is 0, 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (I), in which 'n' is 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R² is independently cyano, halogen (e.g. F, Cl or Br), C₁₋₈alkyl (e.g. methyl), haloC₁₋₈alkyl (e.g. trifluoromethyl), haloC₁₋₈alkoxy (e.g. trifluoromethoxy), C₃₋₁₂cycloalkyl (e.g. cyclopropyl), C₁₋₈alkoxy (e.g. methoxy), C₆₋₁₄aryl (e.g. 3-(trifluoromethoxy)phenyl), 5-14 membered heteroaryl (e.g. 3-isopropyl-1,2,4-oxadiazol-5-yl, 3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl, 3-(4-fluorophenyl)-1,2,4-oxadiazol-5-yl, 3-(3,5-dimethoxyphenyl)-1,2,4-oxadiazol-5-yl, 3-(3-fluoro-5-(trifluoromethyl)phenyl)-1,2,4-oxadiazol-5-yl, 4-methylthiophen-2-yl, 6-(trifluoromethyl)pyridin-3-yl, 1,5,6-trimethyl-1*H*-benzo[d]imidazol-2-yl or 5-(trifluoromethyl)-1*H*-benzo[d]imidazol-2-yl), -NHR, -C(O)NHR, -NHC(O)R or -C≡CR.

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R² is independently -NHR, -C(O)NHR or -NHC(O)R. In this embodiment R is independently selected from C₁₋₈alkyl (e.g. isopropyl or *tert*-butyl), C₃₋₁₂cycloalkyl (e.g. cyclopropyl, 1-[2-

(trifluoromethyl)phenyl]cyclopropyl, or 1-[4-(trifluoromethyl)phenyl]cyclopropyl), C₃₋₈cycloalkylC₁₋₈alkyl (e.g. (cyclopropyl)methyl), C₆₋₁₄aryl (e.g. 2,5-dichlorophenyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 4-(trifluoromethyl)phenyl, 3-(trifluoromethyl)phenyl, 2-(trifluoromethyl)phenyl, 3-(difluoromethyl)phenyl, 2-chloro-4-(trifluoromethyl)phenyl, 4-fluoro-3-(trifluoromethyl)phenyl, 2-fluoro-5-(trifluoromethyl)phenyl, 2-chloro-4-methylphenyl, 2-fluoro-4-methylphenyl, 5-chloro-2-methylphenyl, 3-(difluoromethyl)-4-fluorophenyl, 3-(difluoromethyl)phenyl, 4-(methylsulfonyl)phenyl or 2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl), C₆₋₁₄arylC₁₋₈alkyl (e.g. 2-(trifluoromethyl)benzyl or 4-fluoro-2-(trifluoromethyl)benzyl), and 5-14 membered heteroaryl (e.g. 6-fluorobenzo[d]thiazol-2-yl, 1*H*-benzo[d]imidazol-2-yl, 5-(trifluoromethyl)-1,3,4-thiadiazol-2-yl, 6-fluoropyridin-3-yl, 5-(trifluoromethyl)pyridin-2-yl, 3-chloropyridin-4-yl, 2-morpholinopyrimidin-5-yl, or 6-(morpholin-4-yl)pyridin-3-yl).

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R² is -C≡CR. In this embodiment R is independently selected from C₁₋₈alkyl (e.g. isopropyl or *tert*-butyl), C₃₋₁₂cycloalkyl (e.g. cyclopropyl), C₆₋₁₄aryl (2,5-dichlorophenyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 2-chloro-4-(trifluoromethyl)phenyl, or 2-(trifluoromethyl)phenyl) and 5-14 membered heteroaryl (e.g. 6-fluoropyridin-3-yl, 5-(trifluoromethyl)pyridin-2-yl, 3-chloropyridin-4-yl, 2-morpholinopyrimidin-5-yl, or 6-(morpholin-4-yl)pyridin-3-yl).

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R² is -C≡CR. In this embodiment R is independently selected from isopropyl, *tert*-butyl, cyclopropyl, 2,5-dichlorophenyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 2-chloro-4-(trifluoromethyl)phenyl, 2-(trifluoromethyl)phenyl, 6-fluoropyridin-3-yl, 5-(trifluoromethyl)pyridin-2-yl, 3-chloropyridin-4-yl, 2-morpholinopyrimidin-5-yl, and 6-(morpholin-4-yl)pyridin-3-yl.

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R² is -NHR. In this embodiment R is 3-(trifluoromethyl)phenyl.

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R^2 is $-NHC(O)R$. In this embodiment R is cyclopropyl.

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R^2 is $-C(O)NHR$. In this embodiment R is (cyclopropyl)methyl, 4-(trifluoromethyl)phenyl, 3-(trifluoromethyl)phenyl, 3-(difluoromethyl)phenyl, 4-fluoro-3-(trifluoromethyl)phenyl, 2-fluoro-5-(trifluoromethyl)phenyl, 2-chloro-4-methylphenyl, 2-fluoro-4-methylphenyl, 5-chloro-2-methylphenyl, 3-(difluoromethyl)-4-fluorophenyl, 3-(difluoromethyl)phenyl, 2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl, 2-(trifluoromethyl)benzyl, 4-fluoro-2-(trifluoromethyl)benzyl, 1-[2-(trifluoromethyl)phenyl]cyclopropyl, 1-[4-(trifluoromethyl)phenyl]cyclopropyl, 6-fluorobenzo[d]thiazol-2-yl, 1*H*-benzo[d]imidazol-2-yl, 5-(trifluoromethyl)-1,3,4-thiadiazol-2-yl or 4-(methylsulfonyl)phenyl.

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R^2 is independently cyano, F, Cl, Br, methyl, trifluoromethyl, trifluoromethoxy, methoxy, cyclopropyl, 3-(trifluoromethoxy)phenyl, 3-isopropyl-1,2,4-oxadiazol-5-yl, 3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl, 3-(4-fluorophenyl)-1,2,4-oxadiazol-5-yl, 3-(3,5-dimethoxyphenyl)-1,2,4-oxadiazol-5-yl, 3-(3-fluoro-5-(trifluoromethyl)phenyl)-1,2,4-oxadiazol-5-yl, 4-methylthiophen-2-yl, 6-(trifluoromethyl)pyridin-3-yl, 1,5,6-trimethyl-1*H*-benzo[d]imidazol-2-yl, 5-(trifluoromethyl)-1*H*-benzo[d]imidazol-2-yl, [3-(trifluoromethyl)phenyl]amine, cyclopropanecarboxamido, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, (2,5-dichlorophenyl)ethynyl, (4-chloro-2-fluorophenyl)ethynyl, (3-chloro-2-fluorophenyl)ethynyl, [2-chloro-4-(trifluoromethyl)phenyl]ethynyl, [2-(trifluoromethyl)phenyl]ethynyl, (6-fluoropyridin-3-yl)ethynyl, [5-(trifluoromethyl)pyridin-2-yl]ethynyl, (3-chloropyridin-4-yl)ethynyl, (2-morpholinopyrimidin-5-yl)ethynyl, [6-(morpholin-4-yl)pyridin-3-yl]ethynyl, $C(O)NH-$ [(cyclopropyl)methyl], $-CONH-[4-(trifluoromethyl)phenyl]$, $-CONH-[3-(trifluoromethyl)phenyl]$, $-CONH-[3-(difluoromethyl)phenyl]$, $-CONH-[4-fluoro-3-(trifluoromethyl)phenyl]$, $-CONH-[2-fluoro-5-(trifluoromethyl)phenyl]$, $-CONH-[2-$

chloro-4-methylphenyl], -CONH-[2-fluoro-4-methylphenyl], -CONH-[5-chloro-2-methylphenyl], -CONH-[3-(difluoromethyl)-4-fluorophenyl], -CONH-[3-(difluoromethyl)phenyl], -CONH-[2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl], -CONH-[2-(trifluoromethyl)benzyl], -CONH-[4-fluoro-2-(trifluoromethyl)benzyl], -CONH-{1-[2-(trifluoromethyl)phenyl]cyclopropyl}, -CONH-{1-[4-(trifluoromethyl)phenyl]cyclopropyl}, -CONH-{6-fluorobenzo[d]thiazol-2-yl}, -CONH-{1*H*-benzo[d]imidazol-2-yl}, -CONH-{5-(trifluoromethyl)-1,3,4-thiadiazol-2-yl}, or -CONH-[4-(methylsulfonyl)phenyl].

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R^2 is -C(O)NHR or -C $\equiv$ CR. In this embodiment R is cyclopropyl or cyclopropylmethyl, each of which may be optionally substituted with one or more substituents selected from halogen (e.g. F, Cl or Br), C_{1-8} alkyl (e.g. methyl or ethyl), halo C_{1-8} alkyl (e.g. trifluoromethyl) and substituted phenyl (e.g. 2-(trifluoromethyl)phenyl and 4-(trifluoromethyl)phenyl).

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R^2 is -C(O)NHR or -C $\equiv$ CR. In this embodiment R is phenyl optionally substituted with one or more substituents selected from halogen (e.g. F, Cl or Br), C_{1-8} alkyl (e.g. methyl or ethyl), halo C_{1-8} alkyl (e.g. trifluoromethyl or difluoromethyl), C_{1-8} alkoxy (e.g. methoxy or ethoxy), halo C_{1-8} alkoxy (e.g. trifluoromethoxy) and -SO₂R^x (e.g. methylsulfonyl).

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R^2 is -C(O)NHR or -C $\equiv$ CR. In this embodiment R is benzyl optionally substituted with one or more substituents selected from halogen (e.g. F, Cl or Br), C_{1-8} alkyl (e.g. methyl or ethyl) and halo C_{1-8} alkyl (e.g. trifluoromethyl).

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R^2 is -C $\equiv$ CR. In this embodiment R is pyridine or pyrimidine, each of which may be optionally substituted with one or more substituents selected from halogen (e.g. F, Cl or Br), C_{1-8} alkyl (e.g. methyl or ethyl), halo C_{1-8} alkyl (e.g. trifluoromethyl) and 5 membered heterocyclyl (e.g. morpholiny).

According to yet another embodiment, specifically provided are compounds of formula (I), in which 'm' is 0, 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (I), in which 'm' is 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of W is C₁₋₈alkyl (e.g. isopropyl or *tert*-butyl), haloC₁₋₈alkyl (e.g. trifluoromethyl or 1-fluoro-2-methylpropan-2-yl), hydroxyC₁₋₈alkyl (e.g. 1-hydroxy-2-methylpropan-2-yl) or C₃₋₁₂cycloalkyl (e.g. cyclopropyl).

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of W is C₁₋₄alkyl (e.g. isopropyl or *tert*-butyl), haloC₁₋₈alkyl (e.g. trifluoromethyl or 1-fluoro-2-methylpropan-2-yl), hydroxyC₁₋₈alkyl (e.g. 1-hydroxy-2-methylpropan-2-yl) or C₃₋₆cycloalkyl (e.g. cyclopropyl).

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of W is isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl or cyclopropyl.

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of Q is C₁₋₈alkyl (e.g. isopropyl or *tert*-butyl), haloC₁₋₈alkyl (e.g. trifluoromethyl or 1-fluoro-2-methylpropan-2-yl), C₁₋₈alkoxyC₁₋₈alkyl (e.g. 1-methoxy-2-methylpropan-2-yl), hydroxyC₁₋₈alkyl (e.g. 1-hydroxy-2-methylpropan-2-yl), C₃₋₁₂cycloalkyl (e.g. cyclopropyl, cyclobutyl or cyclopentyl), C₆₋₁₄aryl (e.g. 2-fluorophenyl), 3-15 membered heterocyclyl (e.g. tetrahydrofuranyl, tetrahydrofuran-2-yl, (S)-tetrahydrofuran-2-yl or (R)-tetrahydrofuran-2-yl) or 5-14 membered heteroaryl (e.g. isoxazolyl or 1-methyl-1*H*-imidazole-2-yl).

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of Q is isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-methoxy-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl, cyclopropyl, cyclobutyl, cyclopentyl, 2-fluorophenyl, tetrahydrofuranyl, tetrahydrofuran-2-yl, (S)-tetrahydrofuran-2-yl, (R)-tetrahydrofuran-2-yl, isoxazolyl or 1-methyl-1*H*-imidazole-2-yl.

According to yet another embodiment, specifically provided are compounds of formula (I), in which 's' is 0 and 't' is 1.

According to yet another embodiment, specifically provided are compounds of formula (I), in which 's' is 1 and 't' is 0.

According to yet another embodiment, specifically provided are compounds of formula (I), in which 's' is 1 and 't' is 1.

According to yet another embodiment, specifically provided are compounds of formula (I), in which 's' is 0 and 't' is 0.

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R is independently selected from C₁₋₈alkyl (e.g. isopropyl or *tert*-butyl), C₃₋₁₂cycloalkyl (e.g. cyclopropyl, 1-[2-(trifluoromethyl)phenyl]cyclopropyl, or 1-[4-(trifluoromethyl)phenyl]cyclopropyl), C₃₋₈cycloalkylC₁₋₈alkyl (e.g. (cyclopropyl)methyl), C₆₋₁₄aryl (e.g. 2,5-dichlorophenyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 4-(trifluoromethyl)phenyl, 3-(trifluoromethyl)phenyl, 2-(trifluoromethyl)phenyl, 3-(difluoromethyl)phenyl, 2-chloro-4-(trifluoromethyl)phenyl, 4-fluoro-3-(trifluoromethyl)phenyl, 2-fluoro-5-(trifluoromethyl)phenyl, 2-chloro-4-methylphenyl, 2-fluoro-4-methylphenyl, 5-chloro-2-methylphenyl, 3-(difluoromethyl)-4-fluorophenyl, 3-(difluoromethyl)phenyl, 4-(methylsulfonyl)phenyl or 2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl), C₆₋₁₄arylC₁₋₈alkyl (e.g. 2-(trifluoromethyl)benzyl or 4-fluoro-2-(trifluoromethyl)benzyl), and 5-14 membered heteroaryl (e.g. 6-fluorobenzo[d]thiazol-2-yl, 1*H*-benzo[d]imidazol-2-yl, or 5-(trifluoromethyl)-1,3,4-thiadiazol-2-yl).

According to yet another embodiment, specifically provided are compounds of formula (I), in which each occurrence of R is independently selected from isopropyl, *tert*-butyl, cyclopropyl, 1-[2-(trifluoromethyl)phenyl]cyclopropyl, 1-[4-(trifluoromethyl)phenyl]cyclopropyl), (cyclopropyl)methyl, 2,5-dichlorophenyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 4-(trifluoromethyl)phenyl, 3-(trifluoromethyl)phenyl, 2-(trifluoromethyl)phenyl, 3-(difluoromethyl)phenyl, 2-chloro-4-(trifluoromethyl)phenyl, 4-fluoro-3-(trifluoromethyl)phenyl, 2-fluoro-5-(trifluoromethyl)phenyl, 2-chloro-4-methylphenyl, 2-fluoro-4-methylphenyl, 5-chloro-2-methylphenyl, 3-(difluoromethyl)-4-fluorophenyl, 3-(difluoromethyl)phenyl, 4-(methylsulfonyl)phenyl, 2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl, 2-

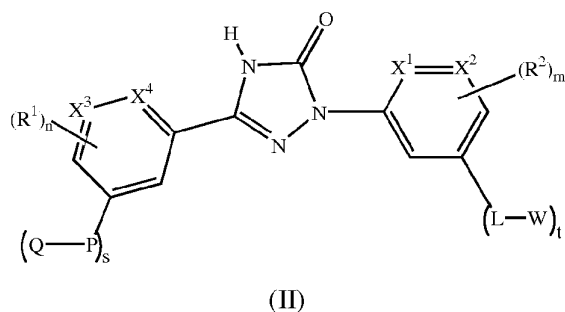
(trifluoromethyl)benzyl, 4-fluoro-2-(trifluoromethyl)benzyl, 6-fluorobenzo[d]thiazol-2-yl, 1*H*-benzo[d]imidazol-2-yl, or 5-(trifluoromethyl)-1,3,4-thiadiazol-2-yl.

According to an embodiment, specifically provided are compounds of formula (I) that exhibit an IC_{50} value of less than 500 nM, preferably less than 100 nM, more preferably less than 50 nM, with respect to mPGES-1 inhibition.

Further embodiments relating to groups R^1 , R^2 , m , n , s , t , P , L , Q and W (and groups defined therein) are described hereinafter in relation to the compounds of formula (II), formula (III) and formula (IV). It is to be understood that these embodiments are not limited to use in conjunction with formula (II), formula (III) or formula (IV), but apply independently and individually to the compounds of formula (I). For example, in an embodiment described hereinafter, the invention specifically provides compounds of formula (II), formula (III) or formula (IV) in which 'n' is 0, 1 or 2, and consequently, there is also provided a compound of formula (I) in which 'n' is 0, 1 or 2.

The invention also provides a compound of formula (II), which is an embodiment of a compound of formula (I).

Accordingly the invention provides a compound of formula (II)



or a pharmaceutically acceptable salt thereof,

wherein,

X^1 , X^2 , X^3 and X^4 are each independently selected from CH and N;

each occurrence of L is independently selected from $-CH_2NHC(O)-$ and $-CH_2NHS(O)_2-$;

each occurrence of P is independently selected from $-CH_2NHC(O)-$ and $-CH_2NHS(O)_2-$;

each occurrence of Q is independently selected from C₁₋₈alkyl, haloC₁₋₈alkyl, C₁₋₈alkoxyC₁₋₈alkyl, hydroxyC₁₋₈alkyl, carboxylC₁₋₈alkyl, C₃₋₁₂cycloalkyl, C₆₋₁₄aryl, 3-15 membered heterocyclyl, and 5-14 membered heteroaryl;

each occurrence of W is independently selected from C₁₋₈alkyl, haloC₁₋₈alkyl, C₁₋₈alkoxyC₁₋₈alkyl, hydroxyC₁₋₈alkyl, carboxylC₁₋₈alkyl, C₃₋₁₂cycloalkyl, C₆₋₁₄aryl, 3-15 membered heterocyclyl, and 5-14 membered heteroaryl;

each occurrence of R¹ is independently selected from halogen, cyano, C₁₋₈alkyl, C₁₋₈alkoxy, haloC₁₋₈alkoxyC₁₋₈alkyl, haloC₁₋₈alkyl, C₃₋₆cycloalkyl, 5 membered heterocyclylC₁₋₈alkyl, 5 membered heteroaryl, 5 membered heteroarylC₁₋₈alkyl, and -C≡CR;

each occurrence of R² is independently selected from halogen, cyano, C₁₋₈alkyl, C₁₋₈alkoxy, haloC₁₋₈alkyl, C₃₋₆cycloalkyl, 5 membered heteroaryl, -C(O)NHR, -NHC(O)R, -S(O)₂NHR and -C≡CR;

each occurrence of R is independently selected from C₁₋₈alkyl, C₃₋₁₂cycloalkyl, and C₆₋₁₄aryl;

‘m’ is an integer ranging from 0 to 3, both inclusive;

‘n’ is an integer ranging from 0 to 3, both inclusive;

‘s’ is an integer ranging from 0 to 1, both inclusive; and

‘t’ is an integer ranging from 0 to 1, both inclusive;

with the provisos that (i) ‘s’ and ‘t’ are not ‘0’ simultaneously, and (ii) ‘m’ and ‘n’ are not ‘0’ simultaneously.

The compounds of formula (II) may involve one or more embodiments. It is to be understood that the embodiments below are illustrative of the present invention and are not intended to limit the claims to the specific embodiments exemplified. It is also to be understood that the embodiments defined herein may be used independently or in conjunction with any definition of any other embodiment defined herein. Thus the invention contemplates all possible combinations and permutations of the various independently described embodiments. For example, the invention provides compounds of formula (II) as defined above wherein X¹, X², X³ and X⁴ are CH (according to an embodiment defined below), ‘m’ is 0, 1 or 2 (according to another embodiment defined below), and ‘s’ is 0 and ‘t’ is 1 (according to another embodiment defined below).

According to one embodiment, specifically provided are compounds of formula (II), in which X^1 , X^2 , X^3 and X^4 are CH.

According to another embodiment, specifically provided are compounds of formula (II), in which X^1 is N and X^2 , X^3 and X^4 are CH.

According to yet another embodiment, specifically provided are compounds of formula (II), in which X^2 is N and X^1 , X^3 and X^4 are CH.

According to yet another embodiment, specifically provided are compounds of formula (II), in which X^3 is N and X^1 , X^2 and X^4 are CH.

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of L is $-\text{CH}_2\text{NHC}(\text{O})-$.

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of P is $-\text{CH}_2\text{NHC}(\text{O})-$.

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of R^1 is independently selected from cyano, halogen (e.g. F, Cl or Br), C_{1-8} alkyl (e.g. methyl), halo C_{1-8} alkyl (e.g. trifluoromethyl or difluoromethyl), C_{1-8} alkoxy (e.g. methoxy), halo C_{1-8} alkoxy C_{1-8} alkyl (e.g. (2,2,2-trifluoroethoxy)methyl), C_{3-6} cycloalkyl (e.g. cyclopropyl), 5 membered heterocyclyl C_{1-8} alkyl (e.g. (pyrrolidin-1-yl)methyl), 5 membered heteroaryl (e.g. 4-methylthiophenyl, 5-isopropyl-1,3,4-oxadiazol-2-yl), 5 membered heteroaryl C_{1-8} alkyl (e.g. (3,5-dimethyl-1*H*-pyrazol-1-yl)methyl) and $-\text{C}\equiv\text{CR}$.

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of R^1 is independently selected from cyano, Cl, F, CHF_2 , CF_3 , OCH_3 , CH_3 , (2,2,2-trifluoroethoxy)methyl, cyclopropyl, (pyrrolidin-1-yl)methyl, 4-methylthiophenyl, 5-isopropyl-1,3,4-oxadiazol-2-yl, (3,5-dimethyl-1*H*-pyrazol-1-yl)methyl and $-\text{C}\equiv\text{CR}$. In this embodiment R is isopropyl, *tert*-butyl, cyclopropyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 3-(trifluoromethyl)phenyl or 2-chloro-4-(trifluoromethyl)phenyl.

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of R^1 is independently selected from cyano, Cl, F, CHF_2 , CF_3 , OCH_3 , CH_3 , (2,2,2-trifluoroethoxy)methyl, cyclopropyl, (pyrrolidin-1-yl)methyl, 4-methylthiophenyl, 5-isopropyl-1,3,4-oxadiazol-2-yl, (3,5-dimethyl-1*H*-

pyrazol-1-yl)methyl, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, (2,5-dichlorophenyl)ethynyl, (4-chloro-2-fluorophenyl)ethynyl, (3-chloro-2-fluorophenyl)ethynyl, (3-(trifluoromethyl)phenyl)ethynyl and (2-chloro-4-(trifluoromethyl)phenyl)ethynyl.

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of R^1 is independently cyano, Cl, F, CHF_2 , CF_3 , OCH_3 , CH_3 , (2,2,2-trifluoroethoxy)methyl, cyclopropyl, (pyrrolidin-1-yl)methyl, 4-methylthiophenyl, 5-isopropyl-1,3,4-oxadiazol-2-yl, (3,5-dimethyl-1*H*-pyrazol-1-yl)methyl, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, (2,5-dichlorophenyl)ethynyl, (4-chloro-2-fluorophenyl)ethynyl, (3-chloro-2-fluorophenyl)ethynyl, (3-(trifluoromethyl)phenyl)ethynyl or (2-chloro-4-(trifluoromethyl)phenyl)ethynyl; and 'n' is 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (II), in which 'n' is 0, 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of R^2 is independently selected from cyano, halogen (e.g. F, Cl or Br), C_{1-8} alkyl (e.g. methyl), haloC_{1-8} alkyl (e.g. trifluoromethyl), C_{1-8} alkoxy (e.g. methoxy), 5 membered heteroaryl (e.g. 3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl), $-\text{C}(\text{O})\text{NHR}$, $-\text{NHC}(\text{O})\text{R}$ and $-\text{C}\equiv\text{CR}$.

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of R^2 is independently $-\text{C}(\text{O})\text{NHR}$, $-\text{NHC}(\text{O})\text{R}$ or $-\text{C}\equiv\text{CR}$. In this embodiment R is isopropyl, *tert*-butyl, cyclopropyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 3-(trifluoromethyl)phenyl, 2-chloro-4-(trifluoromethyl)phenyl, 3-(difluoromethyl)-4-fluorophenyl or 3-(difluoromethyl)phenyl).

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of R^2 is independently cyano, Cl, F, CH_3 , CF_3 , OCH_3 , 3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl, cyclopropanecarboxamido, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, $-\text{CONH}-[3-(\text{trifluoromethyl})\text{phenyl}]$, $-\text{CONH}-[3-(\text{difluoromethyl})-4\text{-fluorophenyl}]$ or $-\text{CONH}-[3-(\text{difluoromethyl})\text{phenyl}]$.

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of R^2 is independently cyano, Cl, F, CH_3 , CF_3 ,

OCH₃, 3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl, cyclopropanecarboxamido, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, -CONH-[3-(trifluoromethyl)phenyl], -CONH-[3-(difluoromethyl)-4-fluorophenyl] or -CONH-[3-(difluoromethyl)phenyl]; and 'm' is 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (II), in which 'm' is 0, 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of W is C₁₋₈alkyl (e.g. isopropyl or *tert*-butyl), haloC₁₋₈alkyl (e.g. trifluoromethyl or 1-fluoro-2-methylpropan-2-yl), hydroxyC₁₋₈alkyl (e.g. 1-hydroxy-2-methylpropan-2-yl) or C₃₋₁₂cycloalkyl (e.g. cyclopropyl).

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of W is C₁₋₄alkyl (e.g. isopropyl or *tert*-butyl), haloC₁₋₈alkyl (e.g. trifluoromethyl or 1-fluoro-2-methylpropan-2-yl), hydroxyC₁₋₈alkyl (e.g. 1-hydroxy-2-methylpropan-2-yl) or C₃₋₆cycloalkyl (e.g. cyclopropyl).

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of W is isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl or cyclopropyl.

According to yet another embodiment, specifically provided are compounds of formula (II), in which W is isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl or cyclopropyl; 't' is 1; and 's' is 0.

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of Q is C₁₋₈alkyl (e.g. isopropyl or *tert*-butyl), haloC₁₋₈alkyl (e.g. trifluoromethyl or 1-fluoro-2-methylpropan-2-yl), C₁₋₈alkoxyC₁₋₈alkyl (e.g. 1-methoxy-2-methylpropan-2-yl), hydroxyC₁₋₈alkyl (e.g. 1-hydroxy-2-methylpropan-2-yl), C₃₋₁₂cycloalkyl (e.g. cyclopropyl, cyclobutyl or cyclopentyl), C₆₋₁₄aryl (e.g. 2-fluorophenyl), 3-15 membered heterocyclyl (e.g. tetrahydrofuranyl, tetrahydrofuran-2-yl, (S)-tetrahydrofuran-2-yl or (R)-tetrahydrofuran-2-yl) or 5-14 membered heteroaryl (e.g. isoxazolyl or 1-methyl-1*H*-imidazole-2-yl).

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of Q is isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-methoxy-2-methylpropan-2-yl, 1-hydroxy-2-

methylpropan-2-yl, cyclopropyl, cyclobutyl, cyclopentyl, 2-fluorophenyl, tetrahydrofuranyl, tetrahydrofuranyl, tetrahydrofuran-2-yl, (S)-tetrahydrofuran-2-yl, (R)-tetrahydrofuran-2-yl, isoxazolyl or 1-methyl-1*H*-imidazole-2-yl.

According to yet another embodiment, specifically provided are compounds of formula (II), in which Q is isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-methoxy-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl, cyclopropyl, cyclobutyl, cyclopentyl, 2-fluorophenyl, tetrahydrofuranyl, tetrahydrofuranyl, tetrahydrofuran-2-yl, (S)-tetrahydrofuran-2-yl, (R)-tetrahydrofuran-2-yl, isoxazolyl or 1-methyl-1*H*-imidazole-2-yl; 's' is 1; and 't' is 0.

According to yet another embodiment, specifically provided are compounds of formula (II), in which 's' is 0 and 't' is 1.

According to yet another embodiment, specifically provided are compounds of formula (II), in which 's' is 1 and 't' is 0.

According to yet another embodiment, specifically provided are compounds of formula (II), in which 's' is 1 and 't' is 1.

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of R is independently selected from C₁₋₈alkyl (e.g. isopropyl or *tert*-butyl), C₃₋₁₂cycloalkyl (e.g. cyclopropyl) and C₆₋₁₄aryl (e.g. 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 3-(trifluoromethyl)phenyl, 2-chloro-4-(trifluoromethyl)phenyl, 3-(difluoromethyl)-4-fluorophenyl or 3-(difluoromethyl)phenyl).

According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of R is phenyl optionally substituted with one or more substituents selected from Cl, F, CH₃, trifluoromethyl and difluoromethyl.

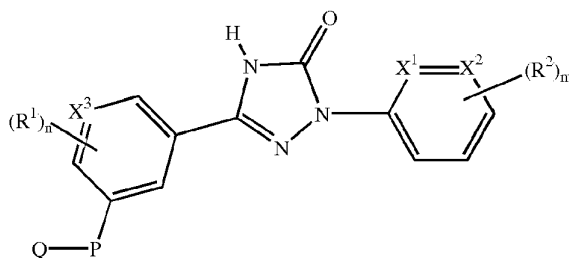
According to yet another embodiment, specifically provided are compounds of formula (II), in which each occurrence of R is independently isopropyl, *tert*-butyl, cyclopropyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 3-(trifluoromethyl)phenyl, 2-chloro-4-(trifluoromethyl)phenyl, 3-(difluoromethyl)-4-fluorophenyl or 3-(difluoromethyl)phenyl.

According to an embodiment, specifically provided are compounds of formula (II) that exhibit an IC₅₀ value of less than 500 nM, preferably less than 100 nM, more preferably less than 50 nM with respect to mPGES-1 inhibition.

Further embodiments relating to groups R^1 , R^2 , m , n , s , t , P , L , Q and W (and groups defined therein) are described herein in relation to the compounds of formula (I), formula (III) or formula (IV). It is to be understood that these embodiments are not limited to use in conjunction with formula (I), formula (III) or formula (IV), but apply independently and individually to the compounds of formula (II). For example, in an embodiment described hereinafter, the invention specifically provides compounds of formula (I), formula (III) or formula (IV) in which 'n' is 0, 1 or 2, and consequently there is also provided a compound of formula (II) in which 'n' is 0, 1 or 2.

The invention also provides a compound of formula (III), which is an embodiment of a compound of formula (I).

Accordingly the invention provides the compound of formula (III)



(III)

or a pharmaceutically acceptable salt thereof,

wherein,

X^1 , X^2 and X^3 are each independently selected from CH and N;

P is selected from $-CH_2NHC(O)-$ and $-CH_2NHS(O)_2-$;

Q is selected from C_{1-8} alkyl, halo C_{1-8} alkyl, C_{1-8} alkoxy C_{1-8} alkyl, hydroxy C_{1-8} alkyl, carboxyl C_{1-8} alkyl, C_{3-12} cycloalkyl, C_{6-14} aryl, 3-15 membered heterocyclyl, and 5-14 membered heteroaryl;

each occurrence of R^1 is independently selected from halogen, cyano, C_{1-8} alkyl, C_{1-8} alkoxy, halo C_{1-8} alkyl and C_{3-6} cycloalkyl;

each occurrence of R^2 is independently selected from halogen, cyano, C_{1-8} alkyl, C_{1-8} alkoxy, halo C_{1-8} alkyl, C_{3-6} cycloalkyl, 5 membered heteroaryl, $-C(O)NHR$, $-NHC(O)R$, $-S(O)_2NHR$ and $-C\equiv CR$;

each occurrence of R is independently selected from C₁₋₈alkyl, C₃₋₁₂cycloalkyl, and C₆₋₁₄aryl;

‘m’ is an integer ranging from 0 to 3, both inclusive; and

‘n’ is an integer ranging from 0 to 3, both inclusive;

with the proviso that ‘m’ and ‘n’ are not ‘0’ simultaneously.

The compounds of formula (III) may involve one or more embodiments. It is to be understood that the embodiments below are illustrative of the present invention and are not intended to limit the claims to the specific embodiments exemplified. It is also to be understood that the embodiments defined herein may be used independently or in conjunction with any definition of any other embodiment defined herein. Thus the invention contemplates all possible combinations and permutations of the various independently described embodiments. For example, the invention provides compounds of formula (III) as defined above wherein X¹, X² and X³ are CH (according to an embodiment defined below), ‘m’ is 0, 1 or 2 (according to another embodiment defined below) and ‘n’ is 0, 1 or 2 (according to another embodiment defined below).

According to one embodiment, specifically provided are compounds of formula (III), in which X¹, X² and X³ are CH.

According to another embodiment, specifically provided are compounds of formula (III), in which X¹ is N and X² and X³ are CH.

According to yet another embodiment, specifically provided are compounds of formula (III), in which X² is N and X¹ and X³ are CH.

According to yet another embodiment, specifically provided are compounds of formula (III), in which X³ is N and X¹ and X² are CH.

According to yet another embodiment, specifically provided are compounds of formula (III), in which P is -CH₂NHC(O)-.

According to yet another embodiment, specifically provided are compounds of formula (III), in which each occurrence of R¹ is independently selected from cyano, halogen (e.g. F, Cl or Br), C₁₋₈alkyl (e.g. methyl), haloC₁₋₈alkyl (e.g. trifluoromethyl or difluoromethyl), C₁₋₈alkoxy (e.g. methoxy) and C₃₋₆cycloalkyl (e.g. cyclopropyl).

According to yet another embodiment, specifically provided are compounds of formula (III), in which each occurrence of R^1 is independently selected from cyano, Cl, F, CHF_2 , CF_3 , OCH_3 , CH_3 and cyclopropyl.

According to yet another embodiment, specifically provided are compounds of formula (III), in which each occurrence of R^1 is independently cyano, Cl, F, CHF_2 , CF_3 , OCH_3 , CH_3 or cyclopropyl; and 'n' is 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (III), in which 'n' is 0, 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (III), in which each occurrence of R^2 is independently selected from cyano, halogen (e.g. F, Cl or Br), C_{1-8} alkyl (e.g. methyl), halo C_{1-8} alkyl (e.g. trifluoromethyl), C_{1-8} alkoxy (e.g. methoxy), 5 membered heteroaryl (e.g. 3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl), $-C(O)NHR$, $-NHC(O)R$ and $-C\equiv CR$.

According to yet another embodiment, specifically provided are compounds of formula (III), in which each occurrence of R^2 is independently $-C(O)NHR$, $-NHC(O)R$ and $-C\equiv CR$. In this embodiment R is isopropyl, *tert*-butyl, cyclopropyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 3-(trifluoromethyl)phenyl, 2-chloro-4-(trifluoromethyl)phenyl, 3-(difluoromethyl)-4-fluorophenyl or 3-(difluoromethyl)phenyl.

According to yet another embodiment, specifically provided are compounds of formula (III), in which each occurrence of R^2 is independently cyano, Cl, F, CH_3 , CF_3 , OCH_3 , 3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl, cyclopropanecarboxamido, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, $-CONH-[3-(trifluoromethyl)phenyl]$, $-CONH-[3-(difluoromethyl)-4-fluorophenyl]$ or $-CONH-[3-(difluoromethyl)phenyl]$.

According to yet another embodiment, specifically provided are compounds of formula (III), in which R^2 is independently cyano, Cl, F, CH_3 , CF_3 , OCH_3 , 3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl, cyclopropanecarboxamido, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, $-CONH-[3-(trifluoromethyl)phenyl]$, $-CONH-[3-(difluoromethyl)-4-fluorophenyl]$ or $-CONH-[3-(difluoromethyl)phenyl]$; and 'm' is 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (III), in which 'm' is 0, 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (III), in which Q is C₁₋₈alkyl (e.g. isopropyl or *tert*-butyl), haloC₁₋₈alkyl (e.g. trifluoromethyl or 1-fluoro-2-methylpropan-2-yl), C₁₋₈alkoxyC₁₋₈alkyl (e.g. 1-methoxy-2-methylpropan-2-yl), hydroxyC₁₋₈alkyl (e.g. 1-hydroxy-2-methylpropan-2-yl), C₃₋₁₂cycloalkyl (e.g. cyclopropyl, cyclobutyl or cyclopentyl), C₆₋₁₄aryl (e.g. 2-fluorophenyl), 3-15 membered heterocyclyl (e.g. tetrahydrofuranyl, tetrahydrofuran-2-yl, (S)-tetrahydrofuran-2-yl or (R)-tetrahydrofuran-2-yl) or 5-14 membered heteroaryl (e.g. isoxazolyl or 1-methyl-1*H*-imidazole-2-yl).

According to yet another embodiment, specifically provided are compounds of formula (III), in which Q is isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-methoxy-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl, cyclopropyl, cyclobutyl, cyclopentyl, 2-fluorophenyl, tetrahydrofuranyl, tetrahydrofuran-2-yl, (S)-tetrahydrofuran-2-yl, (R)-tetrahydrofuran-2-yl, isoxazolyl or 1-methyl-1*H*-imidazole-2-yl.

According to yet another embodiment, specifically provided are compounds of formula (III), in which each occurrence of R is independently selected from C₁₋₈alkyl (e.g. isopropyl or *tert*-butyl), C₃₋₁₂cycloalkyl (e.g. cyclopropyl) and C₆₋₁₄aryl (e.g. 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 3-(trifluoromethyl)phenyl, 2-chloro-4-(trifluoromethyl)phenyl, 3-(difluoromethyl)-4-fluorophenyl or 3-(difluoromethyl)phenyl).

According to yet another embodiment, specifically provided are compounds of formula (III), in which each occurrence of R is phenyl optionally substituted with one or more substituents selected from Cl, F, CH₃, trifluoromethyl and difluoromethyl.

According to yet another embodiment, specifically provided are compounds of formula (III), in which each occurrence of R is independently isopropyl, *tert*-butyl, cyclopropyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 3-(trifluoromethyl)phenyl, 2-chloro-4-(trifluoromethyl)phenyl, 3-(difluoromethyl)-4-fluorophenyl or 3-(difluoromethyl)phenyl.

According to yet another embodiment, specifically provided are compounds of formula (III), in which:

each R¹ is, independently, cyano, Cl, F, CHF₂, CF₃, OCH₃, CH₃ or cyclopropyl;

each R² is, independently, cyano, Cl, F, CH₃, CF₃ or OCH₃;

P is $-\text{CH}_2\text{NHC(O)}-$;

'm' is 1 or 2;

'n' is 0, 1 or 2; and

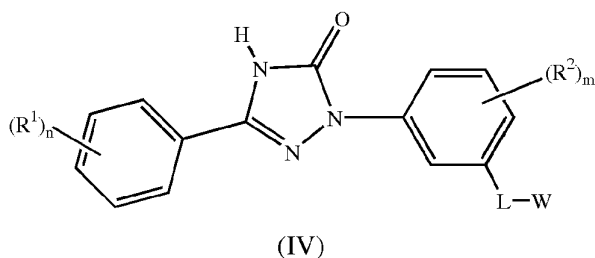
Q is isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-methoxy-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl, cyclopropyl, cyclobutyl, cyclopentyl, 2-fluorophenyl, tetrahydrofuranyl, tetrahydrofuran-2-yl, (S)-tetrahydrofuran-2-yl, (R)-tetrahydrofuran-2-yl, isoxazolyl or 1-methyl-1*H*-imidazole-2-yl.

According to an embodiment, specifically provided are compounds of formula (III) which exhibit an IC_{50} value of less than 500 nM, preferably less than 100 nM, more preferably less than 50 nM with respect to mPGES-1 inhibition.

Further embodiments relating to groups R^1 , R^2 , m, n, s, t, P, L, Q and W (and groups defined therein) are described herein in relation to the compounds of formula (I), formula (II) or formula (IV). It is to be understood that these embodiments are not limited to use in conjunction with formula (I), formula (II) or formula (IV), but apply independently and individually to the compounds of formula (III). For example, in an embodiment described hereinafter, the invention specifically provides compounds of formula (I), formula (II) or formula (IV) in which 'n' is 0, 1 or 2, and consequently there is also provided a compound of formula (III) in which 'n' is 0, 1 or 2.

The invention also provides a compound of formula (IV), which is an embodiment of a compound of formula (I).

Accordingly the invention provides the compound of formula (IV)



or a pharmaceutically acceptable salt thereof,
wherein,

L is selected from $-\text{CH}_2\text{NHC(O)}-$ and $-\text{CH}_2\text{NHS(O)}_2-$;

W is selected from C₁₋₈alkyl, haloC₁₋₈alkyl, C₁₋₈alkoxyC₁₋₈alkyl, hydroxyC₁₋₈alkyl, carboxylC₁₋₈alkyl, C₃₋₁₂cycloalkyl, C₆₋₁₄aryl, 3-15 membered heterocyclyl, and 5-14 membered heteroaryl;

each occurrence of R¹ is independently selected from halogen, cyano, C₁₋₈alkyl, C₁₋₈alkoxy, haloC₁₋₈alkoxyC₁₋₈alkyl, haloC₁₋₈alkyl, C₃₋₆cycloalkyl, 5 membered heterocyclylC₁₋₈alkyl, 5 membered heteroaryl, 5 membered heteroarylC₁₋₈alkyl, and -C≡CR;

each occurrence of R² is independently selected from halogen, cyano, C₁₋₈alkyl, C₁₋₈alkoxy and haloC₁₋₈alkyl;

each occurrence of R is independently selected from C₁₋₈alkyl, C₃₋₁₂cycloalkyl, and C₆₋₁₄aryl;

‘m’ is an integer ranging from 0 to 3, both inclusive; and

‘n’ is an integer ranging from 0 to 3, both inclusive;

with the proviso that ‘m’ and ‘n’ are not ‘0’ simultaneously.

The compounds of formula (IV) may involve one or more embodiments. It is to be understood that the embodiments below are illustrative of the present invention and are not intended to limit the claims to the specific embodiments exemplified. It is also to be understood that the embodiments defined herein may be used independently or in conjunction with any definition of any other embodiment defined herein. Thus the invention contemplates all possible combinations and permutations of the various independently described embodiments. For example, the invention provides compounds of formula (IV) as defined above wherein L is -CH₂NHC(O)- (according to an embodiment defined below), ‘m’ is 0, 1 or 2 (according to another embodiment defined below), and ‘n’ is 0, 1 or 2 (according to another embodiment defined below).

According to one embodiment, specifically provided are compounds of formula (IV), in which L is -CH₂NHC(O)-.

According to another embodiment, specifically provided are compounds of formula (IV), in which each occurrence of R¹ is independently selected from cyano, halogen (e.g. F, Cl or Br), C₁₋₈alkyl (e.g. methyl), haloC₁₋₈alkyl (e.g. trifluoromethyl or difluoromethyl), C₁₋₈alkoxy (e.g. methoxy), haloC₁₋₈alkoxyC₁₋₈alkyl (e.g. (2,2,2-trifluoroethoxy)methyl), 5 membered heterocyclylC₁₋₈alkyl (e.g. (pyrrolidin-1-

yl)methyl), 5 membered heteroaryl (e.g. 4-methylthiophenyl, 5-isopropyl-1,3,4-oxadiazol-2-yl), 5 membered heteroarylC₁₋₈alkyl (e.g. (3,5-dimethyl-1*H*-pyrazol-1-yl)methyl) and -C≡CR.

According to yet another embodiment, specifically provided are compounds of formula (IV), in which each occurrence of R¹ is independently selected from cyano, Cl, F, CHF₂, CF₃, OCH₃, CH₃, (2,2,2-trifluoroethoxy)methyl, (pyrrolidin-1-yl)methyl, 4-methylthiophenyl, 5-isopropyl-1,3,4-oxadiazol-2-yl, (3,5-dimethyl-1*H*-pyrazol-1-yl)methyl and -C≡CR. In this embodiment R is isopropyl, *tert*-butyl, cyclopropyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 3-(trifluoromethyl)phenyl or 2-chloro-4-(trifluoromethyl)phenyl.

According to yet another embodiment, specifically provided are compounds of formula (IV), in which each occurrence of R¹ is independently selected from cyano, Cl, F, CHF₂, CF₃, OCH₃, CH₃, (2,2,2-trifluoroethoxy)methyl, (pyrrolidin-1-yl)methyl, 4-methylthiophenyl, 5-isopropyl-1,3,4-oxadiazol-2-yl, (3,5-dimethyl-1*H*-pyrazol-1-yl)methyl, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, (2,5-dichlorophenyl)ethynyl, (4-chloro-2-fluorophenyl)ethynyl, (3-chloro-2-fluorophenyl)ethynyl, (3-(trifluoromethyl)phenyl)ethynyl and (2-chloro-4-(trifluoromethyl)phenyl)ethynyl.

According to yet another embodiment, specifically provided are compounds of formula (IV), in which each occurrence of R¹ is independently cyano, Cl, F, CHF₂, CF₃, OCH₃, CH₃, (2,2,2-trifluoroethoxy)methyl, (pyrrolidin-1-yl)methyl, 4-methylthiophenyl, 5-isopropyl-1,3,4-oxadiazol-2-yl, (3,5-dimethyl-1*H*-pyrazol-1-yl)methyl, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, (2,5-dichlorophenyl)ethynyl, (4-chloro-2-fluorophenyl)ethynyl, (3-chloro-2-fluorophenyl)ethynyl, (3-(trifluoromethyl)phenyl)ethynyl or (2-chloro-4-(trifluoromethyl)phenyl)ethynyl; and 'n' is 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (IV), in which 'n' is 0, 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (IV), in which each occurrence of R² is independently selected from cyano, halogen (e.g. F, Cl or Br), C₁₋₈alkyl (e.g. methyl), haloC₁₋₈alkyl (e.g. trifluoromethyl) and C₁₋₈alkoxy (e.g. methoxy).

According to yet another embodiment, specifically provided are compounds of formula (IV), in which each occurrence of R^2 is independently cyano, Cl, F, CH_3 , CF_3 or OCH_3 .

According to yet another embodiment, specifically provided are compounds of formula (IV), in which each occurrence of R^2 is independently cyano, Cl, F, CH_3 , CF_3 or OCH_3 ; and 'm' is 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (IV), in which 'm' is 0, 1 or 2.

According to yet another embodiment, specifically provided are compounds of formula (IV), in which W is C_{1-8} alkyl (e.g. isopropyl or *tert*-butyl), halo C_{1-8} alkyl (e.g. trifluoromethyl or 1-fluoro-2-methylpropan-2-yl), hydroxy C_{1-8} alkyl (e.g. 1-hydroxy-2-methylpropan-2-yl) or C_{3-12} cycloalkyl (e.g. cyclopropyl).

According to yet another embodiment, specifically provided are compounds of formula (IV), in which W is C_{1-4} alkyl (e.g. isopropyl or *tert*-butyl), halo C_{1-8} alkyl (e.g. trifluoromethyl or 1-fluoro-2-methylpropan-2-yl), hydroxy C_{1-8} alkyl (e.g. 1-hydroxy-2-methylpropan-2-yl) or C_{3-6} cycloalkyl (e.g. cyclopropyl).

According to yet another embodiment, specifically provided are compounds of formula (IV), in which W is isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl or cyclopropyl.

According to yet another embodiment, specifically provided are compounds of formula (IV), in which each occurrence of R is independently selected from C_{1-8} alkyl (e.g. isopropyl or *tert*-butyl), C_{3-12} cycloalkyl (e.g. cyclopropyl) and C_{6-14} aryl (e.g. 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 3-(trifluoromethyl)phenyl, 2-chloro-4-(trifluoromethyl)phenyl, 3-(difluoromethyl)-4-fluorophenyl or 3-(difluoromethyl)phenyl).

According to yet another embodiment, specifically provided are compounds of formula (IV), in which each occurrence of R is phenyl optionally substituted with one or more substituents selected from Cl, F, trifluoromethyl and difluoromethyl.

According to yet another embodiment, specifically provided are compounds of formula (IV), in which each occurrence of R is independently isopropyl, *tert*-butyl, cyclopropyl, 4-chloro-2-fluorophenyl, 3-chloro-2-fluorophenyl, 3-(trifluoromethyl)phenyl,

2-chloro-4-(trifluoromethyl)phenyl, 3-(difluoromethyl)-4-fluorophenyl or 3-(difluoromethyl)phenyl).

According to yet another embodiment, specifically provided are compounds of formula (IV), in which:

each occurrence of R^1 is independently Cl, F, CHF_2 , CF_3 , OCH_3 , CH_3 , 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, (2,5-dichlorophenyl)ethynyl, (4-chloro-2-fluorophenyl)ethynyl, 3-chloro-2-fluorophenyl)ethynyl, 3-(trifluoromethyl)phenyl)ethynyl or 2-chloro-4-(trifluoromethyl)phenyl)ethynyl;

each occurrence of R^2 is independently Cl, F, CH_3 , CF_3 or OCH_3 ;

L is $-\text{CH}_2\text{NHC(O)}-$;

'm' is 1 or 2;

'n' is 0, 1 or 2; and

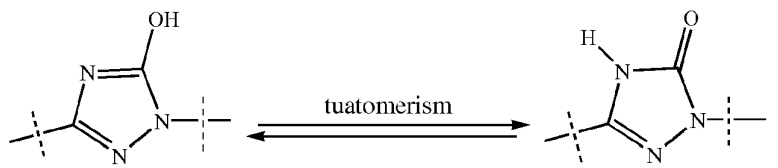
W is isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl or cyclopropyl.

According to an embodiment, specifically provided are compounds of formula (IV) that exhibit an IC_{50} value of less than 500 nM, preferably less than 100 nM, more preferably less than 50 nM with respect to mPGES-1 inhibition.

It should be understood that the formulas (I), (II), (III) and (IV), structurally encompass all geometrical isomers, stereoisomers, enantiomers and diastereomers, *N*-oxides, and pharmaceutically acceptable salts that may be contemplated from the chemical structure of the genera described herein.

Compounds of the present invention include the compounds in Examples 1-192.

According to an embodiment, the compounds of formula (I) (wherein R^3 is H), formula (II), formula (III) or formula (IV) structurally encompass all tautomeric forms whether such tautomer exists in equilibrium or predominantly in one form. Such tautomeric form may be different or the same when the compound is bound to the mPGES-1 enzyme.



The present application also provides a pharmaceutical composition that includes at least one compound described herein and at least one pharmaceutically acceptable excipient (such as a pharmaceutically acceptable carrier or diluent). Preferably, the pharmaceutical composition comprises a therapeutically effective amount of at least one compound described herein. The compounds described herein may be associated with a pharmaceutically acceptable excipient (such as a carrier or a diluent) or be diluted by a carrier, or enclosed within a carrier which can be in the form of a capsule, sachet, paper or other container.

The compounds and pharmaceutical compositions of the present invention are useful for inhibiting the activity of mPGES-1, which is related to a variety of disease states.

The present invention further provides a method of inhibiting mPGES-1 in a subject in need thereof by administering to the subject one or more compounds described herein in an amount effective to cause inhibition of such receptor.

Detailed Description of the Invention

Definitions

The terms “halogen” or “halo” means fluorine (fluoro), chlorine (chloro), bromine (bromo), or iodine (iodo).

The term “alkyl” refers to a hydrocarbon chain radical that includes solely carbon and hydrogen atoms in the backbone, containing no unsaturation, having from one to eight carbon atoms (i.e. C₁₋₈alkyl), and which is attached to the rest of the molecule by a single bond, e.g., methyl, ethyl, n-propyl, 1-methylethyl (isopropyl), n-butyl, n-pentyl, and 1,1-dimethylethyl (t-butyl). The term “C₁₋₆ alkyl” refers to an alkyl chain having 1 to 6 carbon atoms. The term “C₁₋₄alkyl” refers to an alkyl chain having 1 to 4 carbon atoms. Unless set forth or recited to the contrary, all alkyl groups described or claimed herein may be straight chain or branched, substituted or unsubstituted.

The term “alkenyl” refers to a hydrocarbon chain containing from 2 to 10 carbon atoms (i.e. C₂₋₁₀alkenyl) and including at least one carbon-carbon double bond. Non-limiting examples of alkenyl groups include ethenyl, 1-propenyl, 2-propenyl (allyl), *iso*-propenyl, 2-methyl-1-propenyl, 1-butenyl, and 2-butenyl. Unless set forth or recited to

the contrary, all alkenyl groups described or claimed herein may be straight chain or branched, substituted or unsubstituted.

The term “alkynyl” refers to a hydrocarbyl radical having at least one carbon-carbon triple bond, and having 2 to about 12 carbon atoms (with radicals having 2 to about 10 carbon atoms being preferred i.e. C₂₋₁₀alkynyl). Non-limiting examples of alkynyl groups include ethynyl, propynyl, and butynyl. Unless set forth or recited to the contrary, all alkynyl groups described or claimed herein may be straight chain or branched, substituted or unsubstituted.

The term “alkoxy” denotes an alkyl group attached via an oxygen linkage to the rest of the molecule (i.e. C₁₋₈ alkoxy). Representative examples of such groups are -OCH₃ and -OC₂H₅. Unless set forth or recited to the contrary, all alkoxy groups described or claimed herein may be straight chain or branched, substituted or unsubstituted.

The term “alkoxyalkyl” or “alkyloxyalkyl” refers to an alkoxy or alkyloxy group as defined above directly bonded to an alkyl group as defined above (i.e. C₁₋₈alkoxyC₁₋₈alkyl or C₁₋₈alkyloxyC₁₋₈alkyl). Example of such alkoxyalkyl moiety includes, but are not limited to, -CH₂OCH₃ and -CH₂OC₂H₅. Unless set forth or recited to the contrary, all alkoxyalkyl groups described herein may be straight chain or branched, substituted or unsubstituted.

The term “haloalkyl” refers to at least one halo group (selected from F, Cl, Br or I), linked to an alkyl group as defined above (i.e. haloC₁₋₈alkyl). Examples of such haloalkyl moiety include, but are not limited to, trifluoromethyl, difluoromethyl and fluoromethyl groups. Unless set forth or recited to the contrary, all haloalkyl groups described herein may be straight chain or branched, substituted or unsubstituted.

The term “haloalkoxy” refers to an alkoxy group substituted with one or more halogen atoms (i.e. haloC₁₋₈alkoxy). Examples of “haloalkoxy” include but are not limited to fluoromethoxy, difluoromethoxy, trifluoromethoxy, 2,2,2-trifluoroethoxy, pentafluoroethoxy, pentachloroethoxy, chloromethoxy, dichloromethoxy, trichloromethoxy and 1-bromoethoxy. Unless set forth or recited to the contrary, all haloalkoxy groups described herein may be straight chain or branched, substituted or unsubstituted.

The term “haloalkoxyalkyl” refers to haloalkoxy group as defined above directly bonded to an alkyl group as defined above (i.e. haloC₁₋₈alkoxyC₁₋₈alkyl). Examples of “haloC₁₋₈alkoxyC₁₋₈alkyl” include but are not limited to (2,2,2-trifluoroethoxy)methyl or (2,2-difluoroethoxy)methyl. Unless set forth or recited to the contrary, all haloalkoxyalkyl groups described herein may be straight chain or branched, substituted or unsubstituted.

The term “hydroxyalkyl” refers to an alkyl group as defined above wherein one to three hydrogen atoms on different carbon atoms is/are replaced by hydroxyl groups (i.e. hydroxyC₁₋₈alkyl). Examples of hydroxyalkyl moieties include, but are not limited to -CH₂OH, -C₂H₄OH and -CH(OH)C₂H₄OH. Unless set forth or recited to the contrary, all hydroxyalkyl groups described herein may be straight chain or branched, substituted or unsubstituted.

The term “carboxyl” means the group -COOH.

The term “carboxylalkyl” refers to C₁₋₈alkyl group as defined above wherein at least one of the hydrogen atoms of the C₁₋₈alkyl group is replaced by a carboxyl group (i.e. “carboxylC₁₋₈alkyl”). Examples of carboxylalkyl moieties include, but are not limited to carboxymethyl (-CH₂-COOH), carboxylethyl (-CH₂-CH₂-COOH), carboxylisopropyl (-C(CH₃)₂-COOH) and carboxyltertbutyl (-C(CH₃)₂CH₂-COOH). Unless set forth or recited to the contrary, all carboxylalkyl groups described herein may be straight chain or branched, substituted or unsubstituted.

The term “cycloalkyl” denotes a non-aromatic mono or multicyclic ring system of 3 to about 12 carbon atoms, (i.e. C₃₋₁₂cycloalkyl). Examples of monocyclic cycloalkyl include but are not limited to cyclopropyl, cyclobutyl, cyclopentyl, and cyclohexyl. Examples of multicyclic cycloalkyl groups include, but are not limited to, perhydronaphthyl, adamantyl and norbornyl groups, bridged cyclic groups or spirobicyclic groups, e.g., spiro(4,4)non-2-yl. The term “C₃₋₆cycloalkyl” refers to the cyclic ring having 3 to 6 carbon atoms. Unless set forth or recited to the contrary, all cycloalkyl groups described or claimed herein may be substituted or unsubstituted.

The term “cycloalkylalkyl” refers to a non-aromatic cyclic ring-containing radical having 3 to about 8 carbon atoms directly attached to an alkyl group (i.e. C₃₋₈cycloalkylC₁₋₈alkyl). The cycloalkylalkyl group may be attached to the main structure at

any carbon atom in the alkyl group that results in the creation of a stable structure. Non-limiting examples of such groups include cyclopropylmethyl, cyclobutylethyl, and cyclopentylethyl. Unless set forth or recited to the contrary, all cycloalkylalkyl groups described or claimed herein may be substituted or unsubstituted.

The term “cycloalkenyl” refers to a cyclic ring-containing radical having 3 to about 8 carbon atoms with at least one carbon-carbon double bond, (i.e. C₃₋₈cycloalkenyl). Examples of “cycloalkenyl” include but are not limited to cyclopropenyl, cyclobutenyl, and cyclopentenyl. Unless set forth or recited to the contrary, all cycloalkenyl groups described or claimed herein may be substituted or unsubstituted.

The term “cycloalkenylalkyl” refers to a non-aromatic cyclic ring-containing radical having 3 to about 8 carbon atoms with at least one carbon-carbon double bond, directly attached to an alkyl group, (i.e. C₃₋₈cycloalkenylC₁₋₈alkyl). Unless set forth or recited to the contrary, all cycloalkenylalkyl groups described or claimed herein may be substituted or unsubstituted.

The term “aryl” refers to an aromatic radical having 6 to 14 carbon atoms (i.e. C₆₋₁₄aryl), including monocyclic, bicyclic and tricyclic aromatic systems, such as phenyl, naphthyl, tetrahydronaphthyl, indanyl, and biphenyl. Unless set forth or recited to the contrary, all aryl groups described or claimed herein may be substituted or unsubstituted.

The term “aryloxy” refers to an aryl group as defined above attached via an oxygen linkage to the rest of the molecule (i.e. C₆₋₁₄aryloxy). Examples of aryloxy moieties include, but are not limited to phenoxy and naphthoxy. Unless set forth or recited to the contrary, all aryloxy groups described herein may be substituted or unsubstituted.

The term “arylalkyl” refers to an aryl group as defined above directly bonded to an alkyl group as defined above, i.e. C₆₋₁₄arylC₁₋₈alkyl, such as -CH₂C₆H₅ and -C₂H₄C₆H₅. Unless set forth or recited to the contrary, all arylalkyl groups described or claimed herein may be substituted or unsubstituted.

The term “heterocyclic ring” or “heterocyclyl” unless otherwise specified refers to substituted or unsubstituted non-aromatic 3 to 15 membered ring radical (i.e. 3 to 15 membered heterocyclyl) which consists of carbon atoms and from one to five hetero atoms selected from nitrogen, phosphorus, oxygen and sulfur. The heterocyclic ring

radical may be a mono-, bi- or tricyclic ring system, which may include fused, bridged or spiro ring systems, and the nitrogen, phosphorus, carbon, oxygen or sulfur atoms in the heterocyclic ring radical may be optionally oxidized to various oxidation states. In addition, the nitrogen atom may be optionally quaternized; also, unless otherwise constrained by the definition the heterocyclic ring or heterocyclyl may optionally contain one or more olefinic bond(s). Examples of such heterocyclic ring radicals include, but are not limited to azepinyl, azetidiny, benzodioxolyl, benzodioxanyl, chromanyl, dioxolanyl, dioxaphospholanyl, decahydroisoquinolyl, indanyl, indoliny, isoindoliny, isochromanyl, isothiazolidinyl, isoxazolidinyl, morpholiny, oxazoliny, oxazolidinyl, oxadiazolyl, 2-oxopiperazinyl, 2-oxopiperidinyl, 2-oxopyrrolidinyl, 2-oxoazepinyl, octahydroindolyl, octahydroisoindolyl, perhydroazepinyl, piperazinyl, 4-piperidonyl, pyrrolidinyl, piperidinyl, phenothiazinyl, phenoxazinyl, quinuclidinyl, tetrahydroisquinolyl, tetrahydrofuryl or tetrahydrofuranyl, tetrahydropyranyl, thiazoliny, thiazolidinyl, thiamorpholiny, thiamorpholiny sulfoxide and thiamorpholiny sulfone. Unless set forth or recited to the contrary, all heterocyclyl groups described or claimed herein may be substituted or unsubstituted.

The term "heterocyclylalkyl" refers to a heterocyclic ring radical directly bonded to an alkyl group (i.e. 3 to 15 membered heterocyclylC₁₋₈alkyl). Unless set forth or recited to the contrary, all heterocyclylalkyl groups described or claimed herein may be substituted or unsubstituted.

The term "heteroaryl" unless otherwise specified refers to substituted or unsubstituted 5 to 14 membered aromatic heterocyclic ring radical with one or more heteroatom(s) independently selected from N, O or S (i.e. 5 to 14 membered heteroaryl). The heteroaryl may be a mono-, bi- or tricyclic ring system. Examples of such heteroaryl ring radicals include, but are not limited to oxazolyl, isoxazolyl, imidazolyl, furyl, indolyl, isoindolyl, pyrrolyl, triazolyl, triazinyl, tetrazoyl, thienyl, thiazolyl, isothiazolyl, pyridyl, pyrimidinyl, pyrazinyl, pyridazinyl, pyrazolyl, benzofuranyl, benzothiazolyl, benzoxazolyl, benzimidazolyl, benzothieryl, benzopyranyl, carbazolyl, quinoliny, isoquinoliny, quinazoliny, cinnoliny, naphthyridiny, pteridinyl, puriny, quinoxaliny, quinolyl, isoquinolyl, thiadiazolyl, indoliziny, acridiny, phenazinyl and phthalazinyl.

Unless set forth or recited to the contrary, all heteroaryl groups described or claimed herein may be substituted or unsubstituted.

The term “heteroarylalkyl” refers to a heteroaryl ring radical directly bonded to an alkyl group (i.e. 5 to 14 membered heterarylC₁₋₈alkyl). Unless set forth or recited to the contrary, all heteroarylalkyl groups described or claimed herein may be substituted or unsubstituted.

Unless otherwise specified, the term “substituted” as used herein refers to substitution with any one or any combination of the following substituents: hydroxy, halogen, carboxyl, cyano, nitro, oxo (=O), thio (=S), substituted or unsubstituted alkyl, substituted or unsubstituted haloalkyl, substituted or unsubstituted hydroxyl alkyl, substituted or unsubstituted alkoxy, substituted or unsubstituted haloalkoxy, substituted or unsubstituted alkenyl, substituted or unsubstituted alkynyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted cycloalkylalkyl, substituted or unsubstituted cycloalkenyl, substituted or unsubstituted amino, substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, substituted or unsubstituted heteroaryl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted heterocyclalkyl, substituted or unsubstituted heterocyclic ring, substituted or unsubstituted guanidine, -COOR^{x'}, -C(O)R^{x'}, -C(S)R^{x'}, -C(O)NR^{x'}R^{y'}, -C(O)ONR^{x'}R^{y'}, -NR^{x'}CONR^{y'}R^{z'}, -N(R^{x'})SOR^{y'}, -N(R^{x'})SO₂R^{y'}, -(=N-N(R^{x'})R^{y'}), -NR^{x'}C(O)OR^{y'}, -NR^{x'}R^{y'}, -NR^{x'}C(O)R^{y'}, -NR^{x'}C(S)R^{y'}, -NR^{x'}C(S)NR^{y'}R^{z'}, -SONR^{x'}R^{y'}, -SO₂NR^{x'}R^{y'}, -OR^{x'}, -OC(O)NR^{y'}R^{z'}, -OC(O)OR^{y'}, -OC(O)R^{x'}, -OC(O)NR^{x'}R^{y'}, -SR^{x'}, -SOR^{x'}, -SO₂R^{x'}, and -ONO₂, wherein each occurrence of R^{x'}, R^{y'} and R^{z'} are independently selected from hydrogen, substituted or unsubstituted alkyl, substituted or unsubstituted alkoxy, substituted or unsubstituted alkenyl, substituted or unsubstituted alkynyl, substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted cycloalkylalkyl, substituted or unsubstituted cycloalkenyl, substituted or unsubstituted amino, substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, substituted heterocyclalkyl, substituted or unsubstituted heteroarylalkyl, and substituted or unsubstituted heterocyclic ring. The substituents in the aforementioned “substituted” groups cannot be further substituted. For example, when the substituent on

“substituted alkyl” is “substituted aryl”, the substituent on “substituted aryl” can be unsubstituted alkenyl but cannot be “substituted alkenyl”.

The term “pharmaceutically acceptable salt” includes salts prepared from pharmaceutically acceptable bases or acids including inorganic or organic bases and inorganic or organic acids. Examples of such salts include, but are not limited to, acetate, benzenesulfonate, benzoate, bicarbonate, bisulfate, bitartrate, borate, bromide, camsylate, carbonate, chloride, clavulanate, citrate, dihydrochloride, edetate, edisylate, estolate, esylate, fumarate, gluceptate, gluconate, glutamate, glycolylarsanilate, hexylresorcinate, hydrabamine, hydrobromide, hydrochloride, hydroxynaphthoate, iodide, isothionate, lactate, lactobionate, laurate, malate, maleate, mandelate, mesylate, methylbromide, methylnitrate, methylsulfate, mucate, napsylate, nitrate, *N*-methylglucamine ammonium salt, oleate, oxalate, pamoate (embonate), palmitate, pantothenate, phosphate, diphosphate, polygalacturonate, salicylate, stearate, sulfate, subacetate, succinate, tannate, tartrate, teoclate, tosylate, triethiodide and valerate. Examples of salts derived from inorganic bases include, but are not limited to, aluminum, ammonium, calcium, copper, ferric, ferrous, lithium, magnesium, manganic, mangamous, potassium, sodium, and zinc.

The term “treating” or “treatment” of a state, disorder or condition includes: (a) preventing or delaying the appearance of clinical symptoms of the state, disorder or condition developing in a subject that may be afflicted with or predisposed to the state, disorder or condition but does not yet experience or display clinical or subclinical symptoms of the state, disorder or condition; (b) inhibiting the state, disorder or condition, i.e., arresting or reducing the development of the disease or at least one clinical or subclinical symptom thereof; or (c) relieving the disease, i.e., causing regression of the state, disorder or condition or at least one of its clinical or subclinical symptoms.

The term “subject” includes mammals (especially humans) and other animals, such as domestic animals (e.g., household pets including cats and dogs) and non-domestic animals (such as wildlife).

A “therapeutically effective amount” means the amount of a compound that, when administered to a subject for treating a state, disorder or condition, is sufficient to effect such treatment. The “therapeutically effective amount” may vary depending on the

compound, the disease and its severity and the age, weight, physical condition and responsiveness of the subject to be treated.

The sensation of pain can be triggered by any number of physical or chemical stimuli and the sensory neurons which mediate the response to this harmful stimulus are termed as “nociceptors”. Nociceptors are primary sensory afferent (C and A δ fibers) neurons that are activated by a wide variety of noxious stimuli including chemical, mechanical, thermal, and proton (pH<6) modalities. Nociceptors are the nerves which sense and respond to parts of the body which suffer from damage. They signal tissue irritation, impending injury, or actual injury. When activated, they transmit pain signals (via the peripheral nerves as well as the spinal cord) to the brain.

The term “chronic pain” usually refers to pain which persists for 3 months or longer and can lead to significant changes in a patient's personality, lifestyle, functional ability and overall quality of life. Chronic pain can be classified as either nociceptive or neuropathic. Nociceptive pain includes tissue injury-induced pain and inflammatory pain such as that associated with arthritis. Neuropathic pain is caused by damage to the sensory nerves of the peripheral or central nervous system and is maintained by aberrant somatosensory processing. The pain is typically well localized, constant, and often with an aching or throbbing quality. Visceral pain is the subtype of nociceptive pain that involves the internal organs. It tends to be episodic and poorly localized. Nociceptive pain is usually time limited, meaning when the tissue damage heals, the pain typically resolves (arthritis is a notable exception in that it is not time limited).

Pharmaceutical Compositions

The compounds of the invention are typically administered in the form of a pharmaceutical composition. Such compositions can be prepared using procedures known in the pharmaceutical art and comprise at least one compound of the invention. The pharmaceutical composition of the present patent application comprises one or more compounds described herein and one or more pharmaceutically acceptable excipients. Typically, the pharmaceutically acceptable excipients are approved by regulatory authorities or are generally regarded as safe for human or animal use. The pharmaceutically acceptable excipients include, but are not limited to, carriers, diluents,

glidants and lubricants, preservatives, buffering agents, chelating agents, polymers, gelling agents, viscosifying agents, and solvents.

Examples of suitable carriers include, but are not limited to, water, salt solutions, alcohols, polyethylene glycols, peanut oil, olive oil, gelatin, lactose, terra alba, sucrose, dextrin, magnesium carbonate, sugar, amylose, magnesium stearate, talc, gelatin, agar, pectin, acacia, stearic acid, lower alkyl ethers of cellulose, silicic acid, fatty acids, fatty acid amines, fatty acid monoglycerides and diglycerides, fatty acid esters, and polyoxyethylene.

The pharmaceutical composition may also include one or more pharmaceutically acceptable auxiliary agents, wetting agents, suspending agents, preserving agents, buffers, sweetening agents, flavoring agents, colorants or any combination of the foregoing.

The pharmaceutical compositions may be in conventional forms, for example, capsules, tablets, solutions, suspensions, injectables or products for topical application. Further, the pharmaceutical composition of the present invention may be formulated so as to provide a desired release profile.

Administration of the compounds of the invention, in pure form or in an appropriate pharmaceutical composition, can be carried out using any of the accepted routes of administration of pharmaceutical compositions. The route of administration may be any route which effectively transports the active compound of the patent application to the appropriate or desired site of action. Suitable routes of administration include, but are not limited to, oral, nasal, buccal, dermal, intradermal, transdermal, parenteral, rectal, subcutaneous, intravenous, intraurethral, intramuscular, or topical.

Solid oral formulations include, but are not limited to, tablets, capsules (soft or hard gelatin), dragees (containing the active ingredient in powder or pellet form), troches and lozenges.

Liquid formulations include, but are not limited to, syrups, emulsions, and sterile injectable liquids, such as suspensions or solutions.

Topical dosage forms of the compounds include ointments, pastes, creams, lotions, powders, solutions, eye or ear drops, and impregnated dressings, and may contain

appropriate conventional additives such as preservatives, solvents to assist drug penetration.

The pharmaceutical compositions of the present patent application may be prepared by conventional techniques, e.g., as described in *Remington: The Science and Practice of Pharmacy*, 20th Ed., 2003 (Lippincott Williams & Wilkins).

Suitable doses of the compounds for use in treating the diseases and disorders described herein can be determined by those skilled in the relevant art. Therapeutic doses are generally identified through a dose ranging study in humans based on preliminary evidence derived from animal studies. Doses are generally sufficient to result in a desired therapeutic benefit without causing unwanted side effects. Mode of administration, dosage forms, and suitable pharmaceutical excipients can also be well used and adjusted by those skilled in the art. All changes and modifications are envisioned within the scope of the present patent application.

In another embodiment, the present invention relates to a pharmaceutical composition comprising a compound as described herein, a second therapeutic agent, and optionally a pharmaceutically-acceptable excipient. In one embodiment, the pharmaceutical composition includes a compound as described herein and a second therapeutic agent, wherein each of the compound described herein and the second therapeutic agent is formulated in admixture with a pharmaceutically-acceptable excipient.

Methods of Treatment

Compounds of the present invention are particularly useful because they may inhibit the activity of prostaglandin E synthases (and particularly microsomal prostaglandin E synthase-1 (mPGES-1)), i.e., they prevent, inhibit, or suppress the action of mPGES-1 or a complex of which the mPGES-1 enzyme forms a part, and/or may elicit mPGES-1 modulating effect. Compounds of the invention are thus useful in the treatment of those conditions treatable by inhibition of a PGES, and particularly mPGES-1.

Compounds of the invention are thus expected to be useful in the treatment of inflammation. The term "inflammation" will be understood by those skilled in the art to include any condition characterized by a localized or a systemic protective response,

which may be elicited by physical trauma, infection, chronic diseases, such as those mentioned hereinbefore, and/or chemical and/or physiological reactions to external stimuli (e.g. as part of an allergic response). Any such response, which may serve to destroy, dilute or sequester both the injurious agent and the injured tissue, may be manifest by, for example, heat, swelling, pain, redness, dilation of blood vessels and/or increased blood flow.

The term "inflammation" is also understood to include any inflammatory disease, disorder or condition per se, any condition that has an inflammatory component associated with it, and/or any condition characterized by inflammation as a symptom, including, *inter alia*, acute, chronic, ulcerative, specific, allergic, infection by pathogens, immune reactions due to hypersensitivity, entering foreign bodies, physical injury, and necrotic inflammation, and other forms of inflammation known to those skilled in the art. The term thus also includes, for the purposes of this invention, inflammatory pain, pain generally and/or fever.

The compounds of the present invention may also be useful in the treatment of asthma, chronic obstructive pulmonary disease, pulmonary fibrosis, inflammatory bowel disease, irritable bowel syndrome, inflammatory pain, chronic pain, acute pain, fever, migraine, headache, low back pain, fibromyalgia, myofascial disorders, viral infections (e.g. influenza, common cold, herpes zoster, hepatitis C and AIDS), bacterial infections, fungal infections, dysmenorrhea, burns, surgical or dental procedures, malignancies (e.g. breast cancer, colon cancer, and prostate cancer), hyperprostaglandin E syndrome, classic Bartter syndrome, atherosclerosis, gout, arthritis, osteoarthritis, juvenile arthritis, rheumatoid arthritis, juvenile onset rheumatoid arthritis, rheumatic fever, ankylosing spondylitis, Hodgkin's disease, systemic lupus erythematosus, vasculitis, pancreatitis, nephritis, bursitis, conjunctivitis, iritis, scleritis, uveitis, wound healing, dermatitis, eczema, psoriasis, stroke, diabetes mellitus, neurodegenerative disorders such as Alzheimer's disease and multiple sclerosis, autoimmune diseases, allergic disorders, rhinitis, ulcers, mild to moderately active ulcerative colitis, familial adenomatous polyposis, coronary heart disease, sarcoidosis and any other disease with an inflammatory component.

Compounds of the invention may also have effects that are not linked to inflammatory mechanisms, such as in the reduction of bone loss in a subject. Conditions that may be mentioned in this regard include osteoporosis, osteoarthritis, Paget's disease and/or periodontal diseases.

By virtue of the mPGES-1 inhibitory activity of compounds of the present invention, the compounds are useful for the relief of pain, fever and inflammation of a variety of conditions including rheumatic fever, symptoms associated with influenza or other viral infections, common cold, low back and neck pain, dysmenorrhea, headache, migraine (acute and prophylactic treatment), toothache, sprains and strains, myositis, neuralgia, synovitis, arthritis, including rheumatoid arthritis, juvenile rheumatoid arthritis, degenerative joint diseases (osteoarthritis), acute gout and ankylosing spondylitis, acute, subacute and chronic musculoskeletal pain syndromes such as bursitis, burns, injuries, and pain following surgical (post-operative pain) and dental procedures as well as the preemptive treatment of surgical pain. The pain may be mild pain, moderate pain, severe pain, musculoskeletal pain, complex regional pain syndrome, neuropathic pain, back pain such as acute visceral pain, neuropathies, acute trauma, chemotherapy – induced mononeuropathy pain states, polyneuropathy pain states (such as diabetic peripheral neuropathy & chemotherapy induced neuropathy), autonomic neuropathy pain states, peripheral nervous system (PNS) lesion or central nervous system (CNS) lesion or disease related pain states, polyradiculopathies of cervical, lumbar or sciatica type, cauda equina syndrome, piriformis syndrome, paraplegia, quadriplegia, pain states related to various Polyneuritis conditions underlying various infections, chemical injuries, radiation exposure, underlying disease or deficiency conditions (such as beriberi, vitamin deficiencies, hypothyroidism, porphyria, cancer, HIV, autoimmune disease such as multiple sclerosis and spinal-cord injury, fibromyalgia, nerve injury, ischaemia, neurodegeneration, stroke, post stroke pain, inflammatory disorders, oesophagitis, gastroesophageal reflux disorder (GERD), irritable bowel syndrome, inflammatory bowel disease, pelvic hypersensitivity, urinary incontinence, cystitis, stomach duodenal ulcer, muscle pain, pain due to colicky and referred pain. Compounds of the present invention may also be useful for the treatment or prevention of endometriosis, hemophilic arthropathy and Parkinson's disease.

Compounds of the present invention will also inhibit prostanoid-induced smooth muscle contraction by preventing the synthesis of contractile prostanoids and hence may be of use in the treatment of dysmenorrhea, premature labor and asthma.

In addition, the compounds of the present invention may inhibit cellular neoplastic transformations and metastatic tumor growth and hence can be used in the treatment of cancer, and pain associated with cancer. Furthermore, the present invention provides preferred embodiments of the methods and uses as described herein, in which cancer includes Acute Lymphoblastic Leukemia, Acute Myeloid Leukemia, Adolescents Cancer, Adrenocortical Carcinoma, Anal Cancer, Appendix Cancer, Astrocytomas, Atypical Teratoid, Basal Cell Carcinoma, Bile Duct Cancer, Extrahepatic, Bladder Cancer, Bone Cancer, Brain Stem Glioma, Brain Tumor, Breast Cancer, Bronchial Tumors, Burkitt Lymphoma, Carcinoid Tumor, Carcinoma of Unknown Primary, Cardiac (Heart) Tumors, Central Nervous System tumors, Cervical Cancer, Childhood Cancers, Chordoma, Chronic Lymphocytic Leukemia, Chronic Myelogenous Leukemia, Chronic Myeloproliferative Disorders, Colon Cancer, Colorectal Cancer, Craniopharyngioma, Cutaneous T-Cell Lymphoma, Duct Bile Extrahepatic cancer, Ductal Carcinoma In Situ, Embryonal Tumors, Central Nervous System cancer, Endometrial Cancer, Ependymoma, Esophageal Cancer, Esthesioneuroblastoma, Ewing Sarcoma, Extracranial Germ Cell Tumor, Extragonadal Germ Cell Tumor, Extrahepatic Bile Duct Cancer, Eye Cancer, Fibrous Histiocytoma of Bone, Malignant, and Osteosarcoma, Gall bladder Cancer, Gastric (Stomach) Cancer, Gastrointestinal Carcinoid Tumor, Gastrointestinal Stromal Tumors, Germ Cell Tumor, Gestational Trophoblastic Tumor, Glioma, Hairy Cell Leukemia, Head and Neck Cancer, Heart Cancer, Hepatocellular (Liver) Cancer, Histiocytosis, Langerhans Cell, Hodgkin Lymphoma, Hypopharyngeal Cancer, Intraocular Melanoma, Islet Cell Tumors, Pancreatic Neuroendocrine Tumors, Kaposi Sarcoma, Kidney cancer, Langerhans Cell Histiocytosis, Laryngeal Cancer, Acute Lymphoblastic Leukemia, Acute Myeloid Leukemia, Chronic Lymphocytic Leukemia, Chronic Myelogenous Leukemia, Hairy Cell Leukemia, Lip and Oral Cavity Cancer, Liver Cancer, Lobular Carcinoma In Situ, Lung Cancer, AIDS-Related Lymphoma, Cutaneous T-Cell Lymphoma, Hodgkin Lymphoma, Non-Hodgkin Lymphoma, Primary Central Nervous System (CNS) Lymphoma, Macroglobulinemia, Waldenström, Male

Breast Cancer, Malignant Fibrous Histiocytoma of Bone and Osteosarcoma, Melanoma, Merkel Cell Carcinoma, Mesothelioma, Malignant, Metastatic Squamous Neck Cancer with Occult Primary, Midline Tract Carcinoma Involving NUT Gene, Mouth Cancer, Multiple Endocrine Neoplasia Syndromes, Multiple Myeloma/Plasma Cell Neoplasm, Mycosis Fungoides, Myelodysplastic Syndromes, Myelodysplastic/Myeloproliferative Neoplasms, Myelogenous Leukemia, Chronic, Myeloid Leukemia Acute, Multiple Myeloma, Chronic Myeloproliferative Disorders, Nasal Cavity and Paranasal Sinus Cancer, Nasopharyngeal Cancer, Neuroblastoma, Non-Hodgkin Lymphoma, Non-Small Cell Lung Cancer, Oral Cancer, Oral Cavity Cancer, Lip and, Oropharyngeal Cancer, Osteosarcoma and Malignant Fibrous Histiocytoma of Bone, Ovarian Cancer, Pancreatic Cancer, Papillomatosis, Paraganglioma, Paranasal Sinus and Nasal Cavity Cancer, Parathyroid Cancer, Penile Cancer, Pharyngeal Cancer, Pheochromocytoma, Pituitary Tumor, Plasma Cell Neoplasm/Multiple Myeloma, Pleuropulmonary Blastoma, Pregnancy and Breast Cancer, Primary Central Nervous System (CNS) Lymphoma, Prostate Cancer, Rectal Cancer, Renal Cell (Kidney) Cancer, Renal Pelvis and Ureter, Transitional Cell Cancer, Retinoblastoma, Rhabdomyosarcoma, Salivary Gland Cancer, Ewing Sarcoma, Kaposi Sarcoma, Osteosarcoma, Rhabdomyosarcoma, Soft Tissue Sarcoma, Uterine Sarcoma, Sézary Syndrome, Skin Cancer, Small Cell Lung Cancer, Small Intestine Cancer, Soft Tissue Sarcoma, Squamous Cell Carcinoma, Squamous Neck Cancer with Occult Primary, Metastatic, Stomach (Gastric) Cancer, T-Cell Lymphoma, Cutaneous, Testicular Cancer, Throat Cancer, Thymoma and Thymic Carcinoma, Thyroid Cancer, Transitional Cell Cancer of the Renal Pelvis and Ureter, Trophoblastic Tumor, Gestational, Unknown Primary, Carcinoma of, Ureter and Renal Pelvis, Transitional Cell Cancer, Urethral Cancer, Uterine Cancer, Endometrial, Uterine Sarcoma, Vaginal Cancer, Vulvar Cancer, Waldenström, Macroglobulinemia, Wilms Tumor and Women's Cancers.

The compounds of the present invention may be useful in the treatment of disease, disorder, syndrome or condition selected from the group consisting of inflammation, asthma, chronic obstructive pulmonary disease, pulmonary fibrosis, inflammatory bowel disease, irritable bowel syndrome, pain, inflammatory pain, chronic pain, acute pain, fever, migraine, headache, low back pain, fibromyalgia, myofascial

disorders, viral infections, influenza, common cold, herpes zoster, hepatitis C, AIDS, bacterial infections, fungal infections, dysmenorrhea, burns, surgical or dental procedures, malignancies hyperprostaglandin E syndrome, classic Bartter syndrome, synovitis, atherosclerosis, gout, arthritis, osteoarthritis, juvenile arthritis, rheumatoid arthritis, juvenile onset rheumatoid arthritis, rheumatic fever, ankylosing spondylitis, Hodgkin's disease, systemic lupus erythematosus, vasculitis, pancreatitis, nephritis, bursitis, conjunctivitis, iritis, scleritis, uveitis, wound healing, dermatitis, eczema, psoriasis, stroke, diabetes mellitus, cancer, neurodegenerative disorders such as Alzheimer's disease, Parkinson's disease, Amyotrophic lateral sclerosis and multiple sclerosis, autoimmune diseases, allergic disorders, rhinitis, ulcers, mild to moderately active ulcerative colitis, familial adenomatous polyposis, coronary heart disease, and sarcoidosis.

The compounds of the present invention may be useful in the treatment of pain, chronic pain, acute pain, rheumatoid arthritic pain or osteoarthritic pain.

The compounds of the present invention may be useful in the treatment of inflammation, neurodegenerative disorders such as Parkinson's disease, Alzheimer's disease and amyotrophic lateral sclerosis.

The compounds of the present invention may be useful in the treatment prevention or management of the cancer.

Compounds of the present invention are indicated both in the therapeutic and/or prophylactic treatment of the above-mentioned conditions. For the above-mentioned therapeutic uses the dosage administered will, of course, vary with the compound employed, the mode of administration, the treatment desired and the disorder indicated. The daily dosage of the compound of the invention may be in the range from 0.05 mg/kg to 100 mg/kg.

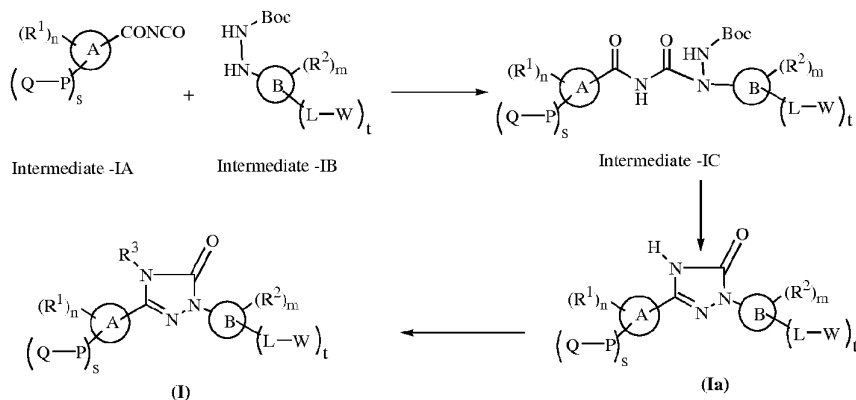
General Methods of Preparation

The compounds described herein, including compounds of general formula (I), (II), (III), and (IV) and specific examples can be prepared using techniques known to one skilled in the art through the reaction sequences depicted in Synthetic Scheme-I, Synthetic Scheme-II, Synthetic Scheme-III, Synthetic Scheme-IV, Synthetic Scheme-V,

Synthetic Scheme-VI, Synthetic Scheme-VII, Synthetic Scheme-VIII, Synthetic Scheme-IX, Synthetic Scheme-X and Synthetic Scheme-XI as well as by other methods. Furthermore, in the following schemes, where specific acids, bases, reagents, coupling agents, solvents, etc. are mentioned, it is understood that other suitable acids, bases, reagents, coupling agents etc. may be used and are included within the scope of the present invention. Modifications to reaction conditions, for example, temperature, duration of the reaction or combinations thereof, are envisioned as part of the present invention. The compounds obtained by using the general reaction sequences may be of insufficient purity. These compounds can be purified by using any of the methods for purification of organic compounds known to persons skilled in the art, for example, crystallization or silica gel or alumina column chromatography using different solvents in suitable ratios. All possible geometrical isomers and stereo isomers are envisioned within the scope of this invention.

A general approach for the preparation of a compound of formula (I) or (Ia) is depicted in the Synthetic Scheme-I (wherein ring A, ring B, R^1 , R^2 , R^3 , L, P, Q, W, m, n, s and t are as defined with respect to a compound of formula (I)).

Synthetic Scheme-I



An Isocyanate compound of formula (Intermediate-IA) may be obtained from a corresponding acid, amide or acid halide derivative.

A compound of formula (Intermediate-IA) can be treated (or coupled) with a compound of formula (Intermediate-IB) to form a compound of formula (Intermediate-

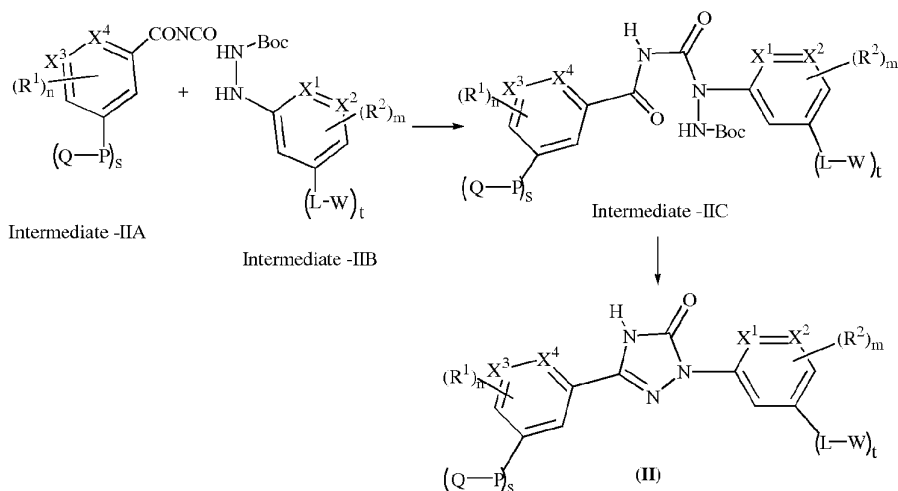
IC). The compound of formula (Intermediate-IA) can be reacted with the compound of formula (Intermediate-IB) in solvent such as DCM, toluene or EDC. According to the process, the compound of formula Intermediate-IC may be isolated or not isolated.

A compound of formula (Intermediate-IC) can be deprotected (e.g., treated with an acid) to obtain a compound of formula (Ia). The acid used in the conversion of the compound of formula (Intermediate-IC) may be organic acid such as TFA, CSA or methane sulphonic acid.

A compound of formula (I) (wherein R^3 is not hydrogen) can be prepared from a compound of formula (Ia) by general alkylation methods by using C_{1-8} alkyl halide/ C_{3-12} cycloalkyl halide/ C_{6-14} aryl halide in the presence of one or more inorganic bases such as NaH, K_2CO_3 or $CsCO_3$.

In another approach, the compound of formula (II) can be prepared following the synthetic steps depicted in Synthetic Scheme-II (wherein R^1 , R^2 , L, P, Q, W, X^1 , X^2 , X^3 , X^4 , m, n, s and t are as defined with respect to a compound of formula (II)).

Synthetic Scheme-II

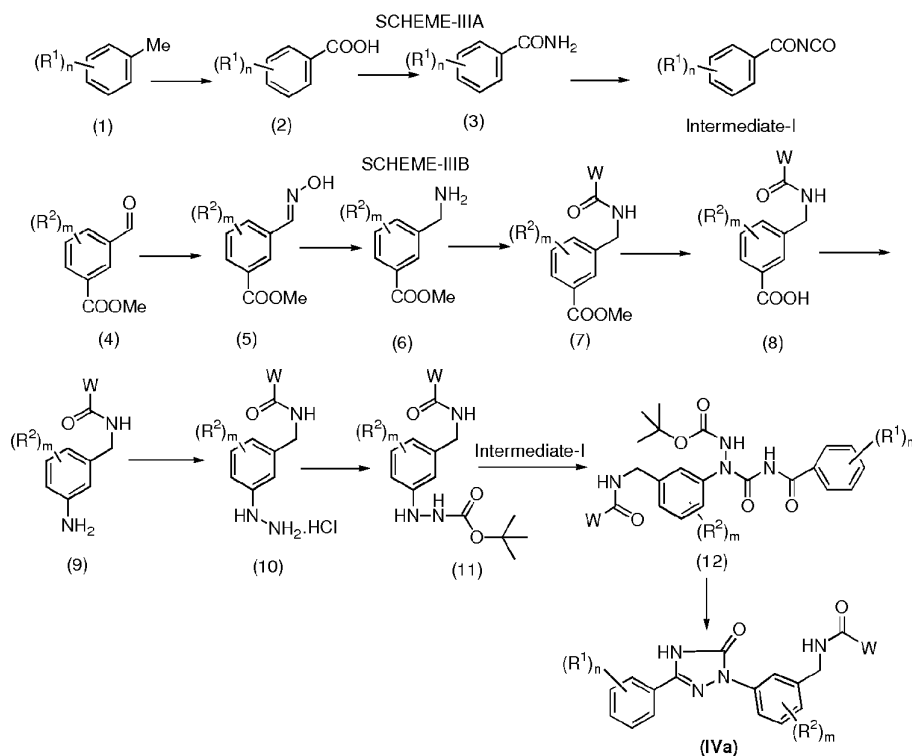


An Isocyanate compound of formula (Intermediate-IIA) may be obtained from a corresponding acid, amide or acid halide derivative. A compound of formula (Intermediate-IIA) can be treated with (or coupled to) a compound of formula (Intermediate-IIB) to form a compound of formula (Intermediate-IIC). A compound of formula (Intermediate-IIC) can be treated with an acid such as an organic acid (e.g. TFA,

CSA or methane sulphonic acid) at room temperature to obtain a compound of formula (II). According to the process, the compound of formula (Intermediate-IIA) can be reacted with the compound of formula (Intermediate-IIB) in a solvent such as DCM, toluene or EDC. According to the process, the compound of formula Intermediate-IIC may be isolated or not isolated.

An approach for the preparation of compound of formula (IVa) is schematically represented in Synthetic Scheme-III (wherein R^1 , R^2 , W, m, and n are as defined with respect to a compound of formula (IV)).

Synthetic Scheme-III



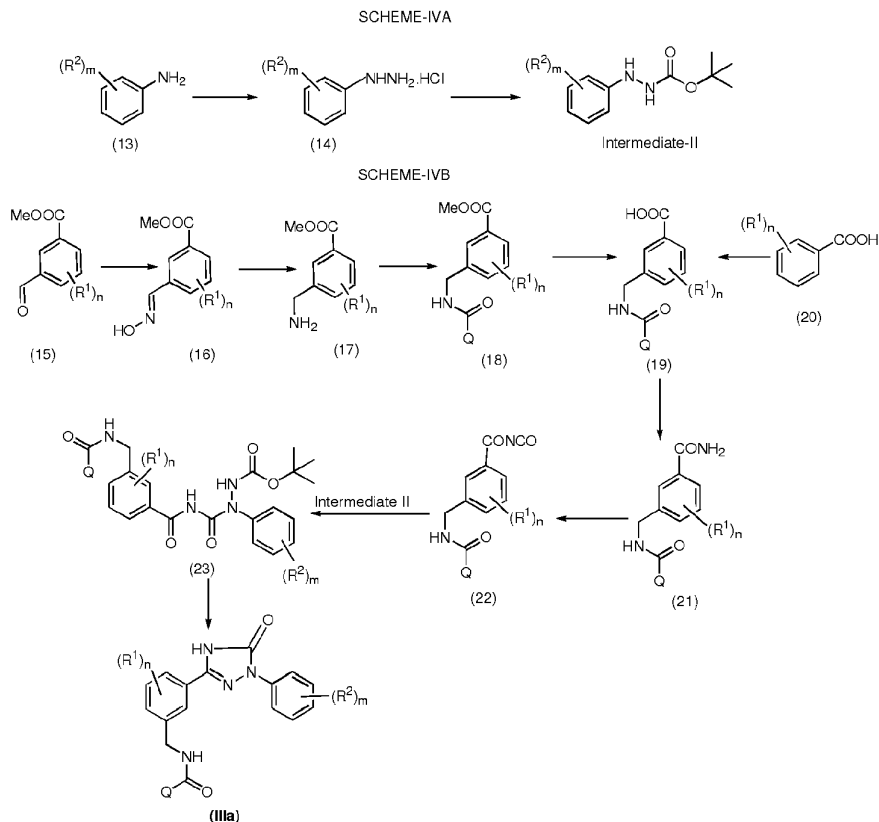
A compound of formula (1) (Scheme-III A) can be converted to a compound of formula (2) by using general oxidation methods known in the art using oxidizing agents such as KMnO_4 , CrO_3 , $\text{K}_2\text{Cr}_2\text{O}_7$ etc. A compound of formula (2) can be converted to a compound of formula (3) by using reaction conditions such as oxalyl chloride/ NH_3 , thionyl chloride/ NH_3 or EDCI/ NH_4Cl . A compound of formula (3) can be treated with

oxalyl chloride in solvents such as EDC or toluene to obtain Intermediate-I (Scheme-III A).

A compound of formula (5) (Scheme-III B) can be prepared from a compound of formula (4) according to known procedures in the literature and can be further converted to a compound of formula (6) by known reducing agents such as Fe/HCl or Zn/HCl. A compound of formula (6) can be converted to a compound of formula (7) by reacting it with aromatic or aliphatic carbonyl chloride using solvent such as dry THF. A compound of formula (7) can be converted to a compound of formula (8) by hydrolysis, for example with NaOH, KOH or LiOH. A compound of formula (8) can be treated with NaN₃ and conc. H₂SO₄ (e.g., at about 50°C) to give a compound of formula (9). A compound of formula (9) can be converted to a compound of formula (10) by reaction with an aqueous solution of sodium nitrite and a solution of stannous chloride in conc. HCl. A compound of formula (10) can be converted to a compound of formula (11) by using an aqueous solution of inorganic bases such as Na₂CO₃ or K₂CO₃ and BOC anhydride in a solvent such as THF or DMF. A compound of formula (11) can be treated with Intermediate-I to obtain a compound of formula (12). The compound of formula (11) can be reacted with the Intermediate-I in an aprotic solvent such as DCM, toluene or EDC to obtain the compound of formula (12). A compound of formula (12) can be treated with organic acid to obtain compound of formula (IVa) (Scheme-III B). The organic acid used in the conversion of the compound of formula (12) may be TFA, CSA or methane sulphonic acid.

An approach for the preparation of compound of formula (IIIa) is schematically represented in Synthetic Scheme-IV (wherein R¹, R², Q, m, and n are as defined with respect to a compound of formula (III)).

Synthetic Scheme-IV



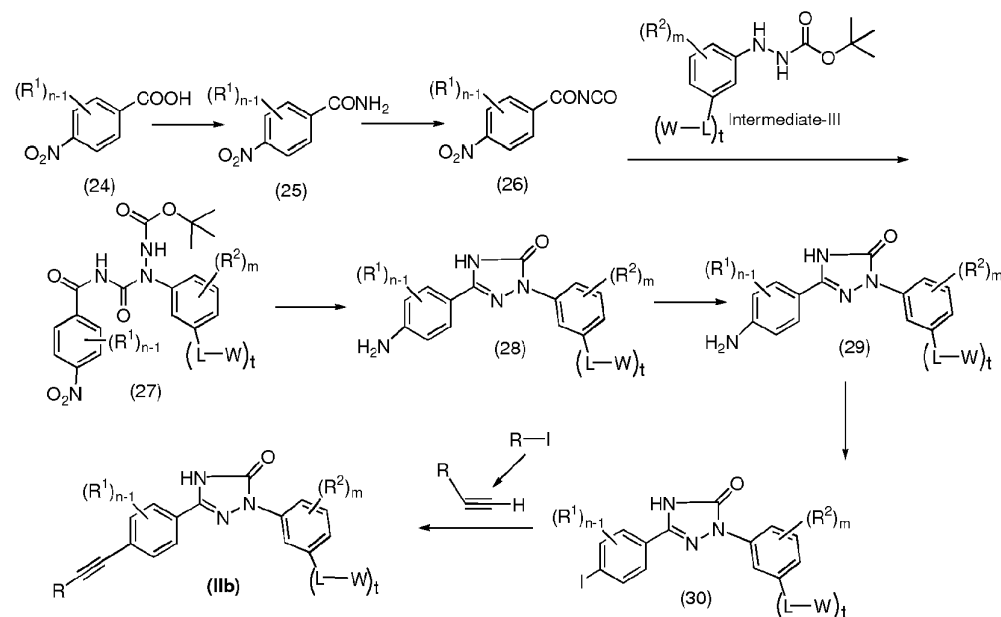
A compound of formula **(13)** can be converted to a compound of formula **(14)** by reaction with an aqueous solution of sodium nitrite and a solution of stannous chloride in conc. HCl, which can be further converted to Intermediate-II by using BOC protection methods known in the art of synthesis in the presence of inorganic bases such as Na₂CO₃, K₂CO₃ by using aprotic solvents such as THF, DMF at 0-100°C. A compound of formula **(19)** can be prepared from a compound of formula **(15)** by following the reaction steps as given in scheme IIB for a compound of formula **(8)**. Also, a compound of formula **(19)** can be directly prepared from a compound of formula **(20)** by using 2,2,2-trifluoro-*N*-(hydroxymethyl)acetamide and sulphuric acid. A compound of formula **(19)** can be converted to a compound of formula **(21)** by using reaction conditions such as oxalyl chloride/NH₃, thionyl chloride/NH₃ or EDCI/NH₄Cl, which can be further treated with oxalyl chloride in a solvent such as EDC or toluene to give a compound of formula **(22)**.

A compound of formula **(22)** can be treated with Intermediate-II to obtain a compound of formula **(23)**. The compound of formula **(22)** can be treated with the

Intermediate-II in an aprotic solvent, such as DCM, toluene or EDC to obtain a compound of formula (23). A compound of formula (23) can be treated with organic acid to obtain compound of formula (IIIa). The organic acid used in the conversion of a compound of formula (23) may be TFA, CSA or methane sulphonic acid. The compound of formula (IIIa) can be converted to a free amine for further amide derivatization by procedures known in the art of synthesis.

An approach for the preparation of compound of formula (IIb) is schematically represented in Synthetic Scheme-V (wherein L, W, R¹, R², R, t, m, and n are as defined with respect to a compound of formula (II)).

Synthetic Scheme-V

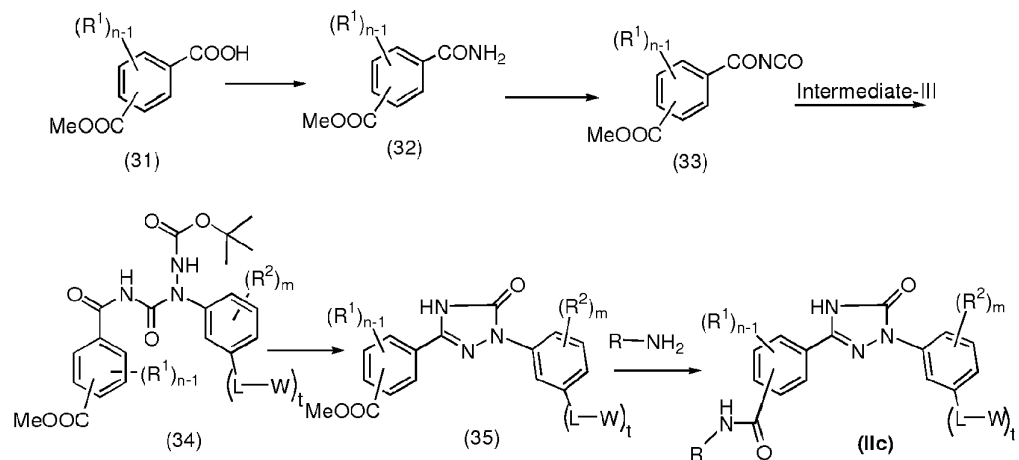


A compound of formula (24) can be converted to a compound of formula (25) by using reaction conditions such as oxalyl chloride/NH₃, thionyl chloride/NH₃ or EDCI/NH₄Cl, which can be further treated with oxalyl chloride in a solvent such as EDC or toluene to give a compound of formula (26). A compound of formula (26) can be treated with Intermediate-III in an aprotic solvent, such as DCM, toluene or EDC to obtain a compound of formula (27), which then further treated with an organic acid, such as TFA, CSA or methane sulphonic acid at room temperature give a compound of

formula (28). A compound of formula (28) can be converted to a compound of formula (29) by reduction methods known in the art of synthesis such as Pd/C, Fe-HCl or Raney Nickel in a solvent such as MeOH or EtOH, which can be further converted to a compound of formula (30) by iodination using KI followed by diazotization using PTSA and NaNO₂. A compound of formula (30) can be converted to a compound of formula (IIb) by a sequence of transformations such as transition metal catalyzed reactions for example Sonogashira coupling.

An approach for the preparation of compound of formula (IIc) is schematically represented in Synthetic Scheme-VI (wherein L, W, R¹, R², R, m, and n are as defined with respect to a compound of formula (II)).

Synthetic Scheme-VI



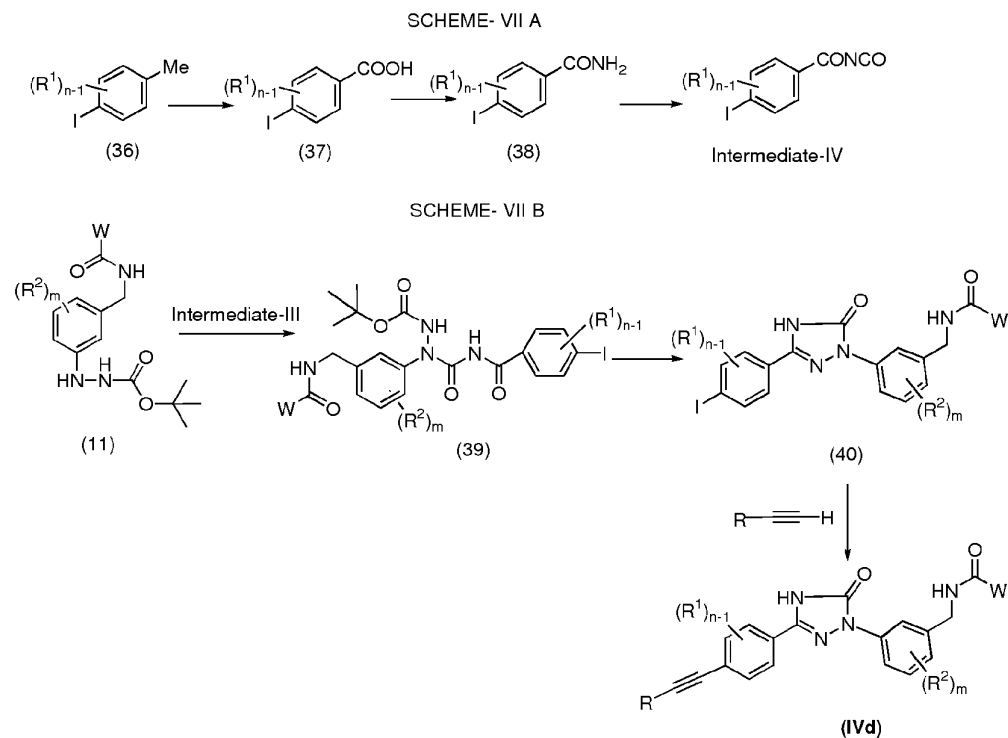
A compound of formula (31) can be converted to a compound of formula (32) by using standard conditions such as oxalyl chloride/NH₃, thionyl chloride/NH₃ or EDCI/NH₄Cl, which can be further treated with oxalyl chloride in a solvent, such as EDC or toluene to give a compound of formula (33). A compound of formula (33) can be treated with (or coupled to) Intermediate-III in an aprotic solvent, such as DCM, toluene or EDC to obtain a compound of formula (34), which when further treated with organic acid, such as TFA, CSA or methane sulphonic acid at room temperature give a compound of formula (35). A compound of formula (IIc) can be prepared from a compound of

formula (35) by using trimethyl aluminum and solvents such as toluene, xylene or EDC at a temperature of 0-100°C.

Alternatively, the methyl ester of a compound of formula (35) can be converted to its acid derivative under hydrolysis condition, which can be further treated with the amines of the formula R-NH₂ to give compound of formula (IIc) using standard coupling conditions such as EDCI, HBTU, TBTU, DCC etc.

An approach for the preparation of a compound of formula (IVd) is schematically represented in Synthetic Scheme-VII (wherein R¹, R², R, W, m, and n are as defined with respect to a compound of formula (IV)).

Synthetic Scheme-VII

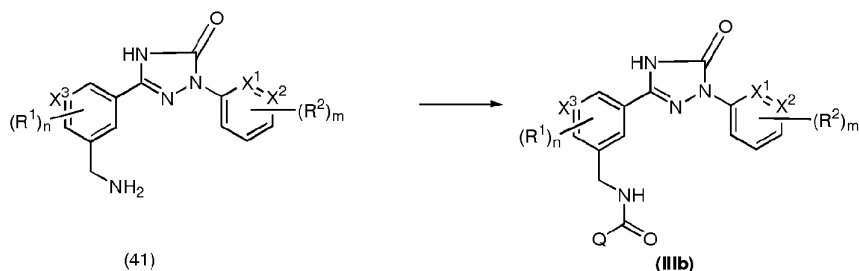


A compound of formula (36) can be converted to Intermediate-IV by following the same reaction steps as for Intermediate-I described in scheme IIIA. A compound of formula (11) can be treated with Intermediate-IV in an aprotic solvent, such as DCM, toluene or EDC to obtain a compound of formula (39), which when further treated with organic acid, such as TFA, CSA or methane sulphonic acid at room temperature give a

compound of formula (40). A compound of formula (40) can be converted to a compound of formula (IVd) by a sequence of transformations such as transition metal catalyzed reactions for example Sonogashira coupling.

In another approach, the compound of formula (IIIb) can be prepared following the synthetic steps depicted in Synthetic Scheme-VIII (wherein X^1 , X^2 , X^3 , R^1 , R^2 , Q , m and n are as defined with respect to a compound of formula (III)).

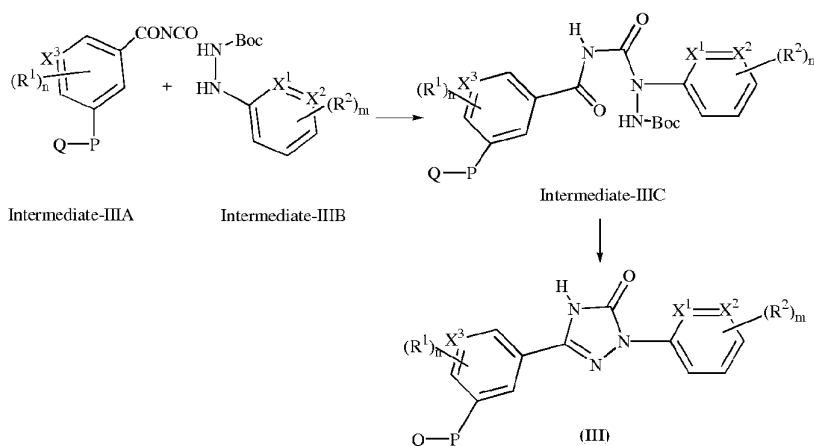
Synthetic Scheme-VIII



A compound of formula (41) can be reacted with $Q-C(O)LG$ (wherein LG represents a suitable leaving group (e.g., OH or Cl or Br or O -alkyl)) under suitable reaction conditions to obtain a compound of formula (IIIb). When LG represents Cl the reaction can be performed in a suitable solvent such as DMF, DCM or THF in the temperature range of $0-120^{\circ}C$, optionally in the presence of a suitable base such as DIPEA or Et_3N . Furthermore, when LG represents O -alkyl the reaction can be performed with a suitable reagent such as trimethylaluminium or a strong base such as sodium hydride (NaH) in a suitable solvent such as toluene or DMF. When LG represents OH , the reaction can be performed with a suitable coupling reagent known in the art, for example, 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide (EDCI) in a suitable solvent such as DMF or tetrahydrofuran (THF) in the temperature range of $0-120^{\circ}C$, optionally in the presence of a suitable base such as DIPEA (diisopropylethyl amine) or Et_3N . Alternatively, the reaction can be performed using a suitable reagent such as isobutyl chloroformate, oxalyl chloride or thionyl chloride in a suitable solvent such as DMF, DCM or THF, in the presence of a suitable base such as DIPEA or Et_3N .

In another approach, the compound of formula (III) can be prepared following the synthetic steps depicted in Synthetic Scheme-IX (wherein R^1 , R^2 , P, Q, X^1 , X^2 , X^3 , m and n are as defined with respect to a compound of formula (III)).

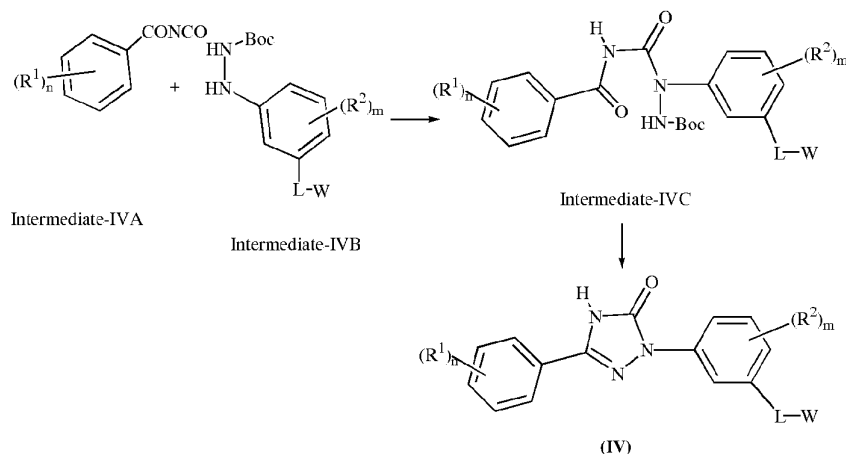
Synthetic Scheme-IX



An Isocyanate compound of formula (Intermediate-III A) may be obtained from a corresponding acid, amide or acid halide derivative. A compound of formula (Intermediate-III A) can be reacted with a compound of formula (Intermediate-III B) to form a compound of formula (Intermediate-III C) and the compound of formula (Intermediate-III C) can be converted to a compound of formula (III). According to the process, the compound of formula (Intermediate-III A) can be reacted with the compound of formula (Intermediate-III B) in a solvent such as DCM, toluene or EDC. According to the process, the compound of formula (Intermediate-III C) may be isolated or not isolated. According to the process, the compound of formula (Intermediate-III C) is converted to compound of formula (III) using an acid. The acid used in the process can be an organic acid such as TFA, CSA or methane sulphonic acid.

In another approach, the compound of formula (IV) can be prepared following the synthetic steps depicted in Synthetic Scheme-X (wherein R^1 , R^2 , L, W, m and n are as defined with respect to a compound of formula (IV)).

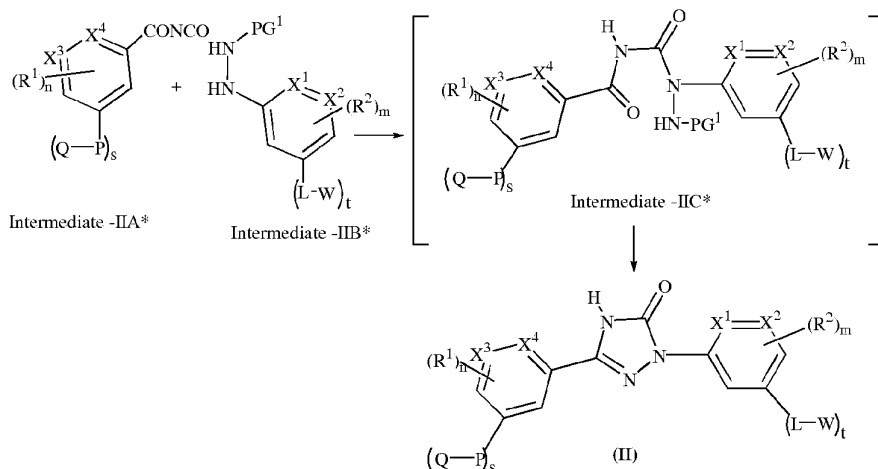
Synthetic Scheme-X



An Isocyanate compound of formula (Intermediate-IVA) may be obtained from a corresponding acid, amide or acid halide derivative. A compound of formula (Intermediate-IVA) can be reacted with a compound of formula (Intermediate-IVB) to form a compound of formula (Intermediate-IVC) and the compound of formula (Intermediate-IVC) can be converted to a compound of formula (IV). According to the process, the compound of formula (Intermediate-IVA) can be reacted with the compound of formula (Intermediate-IVB) in solvent such as DCM, toluene or EDC. According to the process, the compound of formula (Intermediate-IVC) may be isolated or not isolated. According to the process, the compound of formula (Intermediate-IVC) is converted to a compound of formula (IV) using an acid. The acid used in the process can be an organic acid such as TFA, CSA or methane sulphonic acid.

In another approach, the compound of formula (II) can be prepared following the synthetic steps depicted in Synthetic Scheme-XI (wherein R^1 , R^2 , L, P, Q, W, X^1 , X^2 , X^3 , X^4 , m, n, s and t are as defined with respect to a compound of formula (II) and PG^1 represents an amine protecting group such as, but not limited to, *tert*-Butoxycarbonyl (Boc) or carboxybenzyl (Cbz) or benzyl.

Synthetic Scheme-XI



An Isocyanate compound of formula (Intermediate-IIA*) may be obtained from a corresponding acid, amide or acid halide derivative. A compound of formula (Intermediate-IIA*) can be reacted with a compound of formula (Intermediate-IIB*) to form a compound of formula (Intermediate-IIC*) and the compound of formula (Intermediate-IIC*) can be converted to the compound of formula (II). According to the process, the compound of formula (Intermediate-IIA*) can be reacted with the compound of formula (Intermediate-IIB*) in a solvent such as DCM, toluene or EDC. According to the process, the compound of formula Intermediate-IIC* may be isolated or not isolated. According to the process, the compound of formula (Intermediate-IIC*) is converted to compound of formula (II) using an acid. The acid used in the process can be an organic acid such as TFA, CSA or methane sulphonic acid.

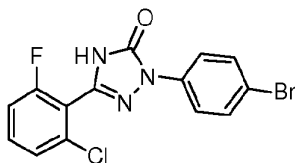
Q, W or other substituents (e.g. R^1 or R^2) if present in the formula (I), (II), (III) or (IV) may be transformed into another chemical group at a chemically compatible stage of the synthetic sequence, in the presence of a suitable reagent by following the procedures known in the art of organic synthesis to obtain the final compound of formula (I), (II), (III) or (IV).

Experimental

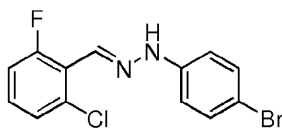
Unless otherwise stated, work-up includes distribution of the reaction mixture between the organic and aqueous phase indicated within parentheses, separation of layers and drying the organic layer over sodium sulphate (Na_2SO_4), filtration and evaporation of the solvent under reduced pressure. Purification, unless otherwise mentioned, includes purification by silica gel chromatographic techniques, in suitable solvents of a suitable polarity as the mobile phase. Abbreviations used in the description of the chemistry and in the examples that follow are: AIBN: Azobisisobutyronitrile; NBS: *N*-Bromosuccinimide; CCl_4 : Carbon tetrachloride; CSA: Camphor sulphonic acid; TFA: Trifluoro acetic acid; NaHCO_3 : Sodium bicarbonate; PCl_5 : Phosphorous pentachloride; POCl_3 : Phosphorous oxychloride; NaO^tBu : Sodium *O*-*tert* butyl; K_2CO_3 : Potassium carbonate; DIPEA: *N,N*-Diisopropylethylamine; LDA: Lithium diisopropylamide; TEA: Triethylamine; TBAF: Tetra-*n*-butylammonium fluoride; DCC: *N,N'*-Dicyclohexylcarbodiimide; HOBT: 1-Hydroxybenzotriazole; TBTU: *O*-(Benzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium tetrafluoroborate; HBTU: *O*-(Benzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate; EDCI: 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide; BOP: (Benzotriazol-1-yl)oxytris(dimethylamino)phosphonium hexafluorophosphate; Boc/BOC: *tert*-Butoxycarbonyl; BOC anhydride: Di-*tert*-butyl dicarbonate; Cbz: Benzyloxycarbonyl; DAST: Diethylaminosulfur trifluoride; PTSA: *p*-Toluenesulfonic acid; DBU: 1,8-Diazabicyclo[5.4.0]undec-7-ene; DCM or MDC: Dichloromethane; DEE: Diethylether; DMSO: Di-methyl sulfoxide; THF: Tetrahydrofuran; EDC: Ethylene dichloride; EtOAc or EA: Ethyl acetate; CHCl_3 : Chloroform; MeOH: Methanol; RT: Room temperature; h: hours.

Intermediate-1

2-(4-Bromophenyl)-5-(2-chloro-6-fluorophenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one

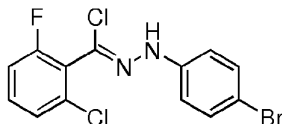


Step-1:- Preparation of (2E)-1-(4-bromophenyl)-2-(2-chloro-6-fluorobenzylidene) hydrazine



To a solution of 2-chloro-6-fluorobenzaldehyde (1.0 g, 6.32 mmol) in ethanol was added (4-bromophenyl)hydrazine (1.7 g, 7.59 mmol) and aq. solution of NaHCO_3 (0.637 g, 4.59 mmol). The reaction mixture was stirred at RT for 5-6 h. The reaction mass was quenched in water and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated. The obtained crude product was purified by column chromatography on silica gel, eluting with 4.6 % EA: pet. ether to afford 0.650 g of the desired product. ^1H NMR (300 MHz, $\text{DMSO}-d_6$): δ 6.98 (d, $J = 7.8$ Hz, 2H), 7.23-7.40 (m, 5H), 8.07 (s, 1H), 10.87 (br s, 1H). MS (m/z): 329.17 ($\text{M}+\text{H}$) $^+$.

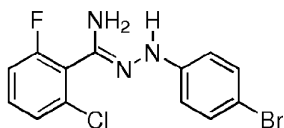
Step-2:- Preparation of N-(4-bromophenyl)-2-chloro-6-fluorobenzenecarbohydrazonoyl chloride



To a solution of (2E)-1-(4-bromophenyl)-2-(2-chloro-6-fluorobenzylidene) hydrazine (0.300 g, 1.32 mmol) in benzene (10 mL) was added PCl_5 (0.330 g, 1.58 mmol). The reaction mixture was stirred at RT for 18 h. The reaction mass was quenched in water, neutralized with NaHCO_3 and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 0.110 g of the desired

product. ^1H NMR (300 MHz, DMSO d_6): δ 7.17 (d, J = 8.7 Hz, 2H), 7.41 (d, J = 8.7 Hz, 3H), 7.45-7.61 (m, 2H), 10.26 (s, 1H). MS (m/z): 361.34 (M-H) $^-$.

Step-3:- Preparation of *N'*-(4-bromophenyl)-2-chloro-6-fluorobenzenecarbohydrazonamide



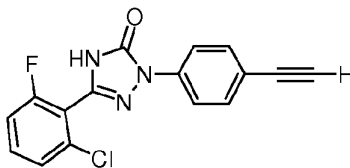
To a cold solution of *N'*-(4-bromophenyl)-2-chloro-6-fluorobenzenecarbohydrazonoyl chloride (0.070 g, 0.267 mmol) in dry THF was added aq. ammonia. The reaction mass was stirred at 0-5°C for 2-3 h. The reaction mass was quenched in water, neutralized with dilute acetic acid and extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 0.040 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 6.29 (br s, 2H), 6.80 (d, J = 8.4 Hz, 2H), 7.24 (d, J = 9.3 Hz, 2H), 7.28-7.50 (m, 3H), 8.33 (s, 1H).

Step-4:- Preparation of 2-(4-bromophenyl)-5-(2-chloro-6-fluorophenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one

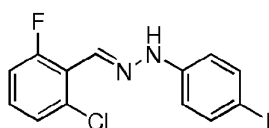
To a cold solution of *N'*-(4-bromophenyl)-2-chloro-6-fluorobenzenecarbohydrazonamide (0.050 g, 0.146 mmol) in CHCl_3 was added pyridine (0.5 mL) and phosgene (1.0 mL) at 0-5°C. The reaction mass was stirred at 0-5°C for 2-3 h. The reaction mass was quenched in water, neutralized with dilute acetic acid and extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated. The obtained crude product was purified with column chromatography on silica gel eluting with 1.0 % MeOH: DCM to afford 0.020 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.50 (t, J = 8.7 Hz, 1H), 7.56-7.71 (m, 4H), 7.89 (d, J = 9.3 Hz, 2H), 12.69 (s, 1H). MS (m/z): 368.15 (M) $^+$.

Intermediate-2

5-(2-Chloro-6-fluorophenyl)-2-(4-ethynylphenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one

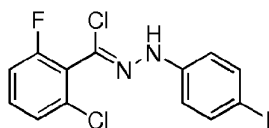


Step-1:- Preparation of (1*E*)-1-(2-chloro-6-fluorobenzylidene)-2-(4-iodophenyl)hydrazine



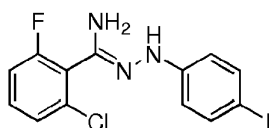
To a solution of (4-iodophenyl)hydrazine (1.0 g, 4.27 mmol) in ethanol (10 mL) was slowly added solution of 2-chloro-6-fluorobenzaldehyde (0.675 g, 4.27 mmol) in ethanol (10 mL). The reaction mass was stirred at RT for 5-6 h. The reaction mass was quenched in water and extracted with ethyl acetate. The organic layer was concentrated to afford 1.2 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 6.87 (d, J = 8.4 Hz, 2H), 7.26-7.36 (m, 3H), 7.53 (d, J = 9.0 Hz, 2H), 8.06 (s, 1H), 10.84 (s, 1H). MS (m/z): 374.87 (M) $^+$.

Step-2:- Preparation of 2-chloro-6-fluoro-*N*-(4-iodophenyl)benzenecarbohydrazonoyl chloride



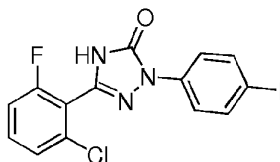
The title compound was prepared according to the procedure described in step-2 of Intermediate-1 using (1*E*)-1-(2-chloro-6-fluorobenzylidene)-2-(4-iodophenyl)hydrazine (1.2 g, 3.20 mmol), PCl_5 (0.990 g, 4.8 mmol) and benzene (20 mL) to afford 1.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.00 (d, J = 8.7 Hz, 2H), 7.41 (d, J = 8.7 Hz, 1H), 7.48-7.60 (m, 4H), 10.22 (s, 1H).

Step-3:- Preparation of 2-chloro-6-fluoro-*N*-(4-iodophenyl)benzenecarbohydrazonamide



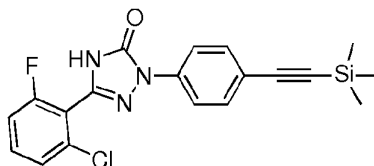
The title compound was prepared according to the procedure described in step-3 of Intermediate-1 by using 2-chloro-6-fluoro-*N*-(4-iodophenyl)benzenecarbohydrazonoyl chloride (1.00 g, 2.44 mmol), aq. ammonia (2.0 mL) and dry THF (10 mL) to afford 0.900 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 6.28 (br s, 2H), 6.70 (d, *J* = 8.4 Hz, 2H), 7.27 (t, *J* = 8.7 Hz, 1H), 7.36-7.42 (m, 3H), 7.45-7.51 (m, 1H), 8.33 (s, 1H).

Step-4:- Preparation of 5-(2-chloro-6-fluorophenyl)-2-(4-iodophenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one



The title a compound was prepared according to the procedure described in step-4 of Intermediate-1 using 2-chloro-6-fluoro-*N'*-(4-iodophenyl)benzene carbohydrazonamide (0.900 g, 2.31 mmol), pyridine (0.65 mL, 5.79 mmol), phosgene (2.50 mL, 4.63 mmol) and CHCl₃ (20 mL). The obtained crude product was purified with column chromatography on silica gel eluting with 1.0 % MeOH: DCM to afford 0.400 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 7.47 (t, *J* = 9.0 Hz, 2H), 7.54-7.82 (m, 5H), 12.62 (s, 1H). MS (*m/z*): 414.25 (M-H)⁺.

Step-5:- Preparation of 5-(2-chloro-6-fluorophenyl)-2-{4-[(trimethylsilyl)ethynyl]phenyl}-2,4-dihydro-3*H*-1,2,4-triazol-3-one



To a solution of 5-(2-chloro-6-fluorophenyl)-2-(4-iodophenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one (0.200 g, 0.483 mmol) in DMSO (2 mL) was added ethynyl(trimethyl)silane (0.071 g, 0.724 mmol), copper iodide (0.005 g, 0.007 mmol), bis(triphenylphosphine) palladium(II) chloride (0.200 g, mmol) and TEA (2.0 mL). The reaction mass was stirred at RT for 24 h. The reaction mass was quenched in water and neutralized with dilute

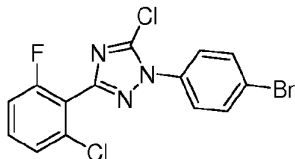
acetic acid and extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 0.200 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 2.31 (s, 9H), 7.44-7.47 (m, 3H), 7.54-7.71 (m, 3H), 7.94-8.01 (m, 1H), 77.95 (br s, 1H). MS (m/z): 384.29 (M-H) $^-$.

Step-6:- Preparation of 5-(2-chloro-6-fluorophenyl)-2-(4-ethynylphenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one

To a solution of 5-(2-chloro-6-fluorophenyl)-2-{4-[(trimethylsilyl)ethynyl] phenyl}-2,4-dihydro-3H-1,2,4-triazol-3-one (0.200 g, 0.570 mmol) in DCM was added TBAF (0.362 g, 1.11 mmol). The reaction mass was stirred at RT for 2-3 h. The reaction mass was quenched in water and filtered through celite bed and concentrated. The obtained crude product was purified with column chromatography on silica gel eluting with 2.0 % MeOH: DCM to afford 0.400 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 4.21 (s, 1H), 7.50 (t, J = 9.3 Hz, 1H), 7.58 (d, J = 8.4 Hz, 3H), 7.67-7.71 (m, 1H), 7.96 (d, J = 8.1 Hz, 2H), 12.69 (br s, 1H).

Intermediate-3

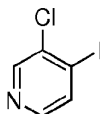
1-(4-bromophenyl)-5-chloro-3-(2-chloro-6-fluorophenyl)-1H-1,2,4-triazole



To 2-(4-bromophenyl)-5-(2-chloro-6-fluorophenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one (Intermediate-1, 0.100 g, 0.271 mmol) was added POCl₃. The reaction mass was refluxed for 48 h. The reaction mass was quenched in water, basified with NaHCO₃ and extracted with ethyl acetate and concentrated. The obtained crude product was purified with column chromatography on silica gel eluting with 0.5 % EA: DCM to afford 0.110 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.44 (t, J = 9.3 Hz, 1H), 7.53 (d, J = 7.8 Hz, 1H), 7.60-7.68 (m, 1H), 7.72 (d, J = 9.0 Hz, 2H), 7.85 (d, J = 8.7 Hz, 2H). MS (m/z): 388.22 (M+H) $^+$.

Intermediate-4

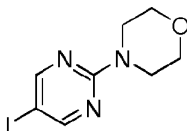
3-Chloro-4-iodopyridine



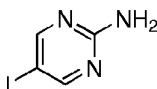
To a cold solution of 3-chloro pyridine (1.0 g, 8.88 mmol) in THF (30.0 mL) was added LDA (5.9 mL, 8.88 mmol) at -75°C . The reaction mixture was stirred at -75°C for 4 h. Iodine (2.2 g, 8.88 mmol) was added and continued stirring at -75°C for 1 h. The reaction mixture was quenched in water at -70°C , extracted with ethyl acetate and concentrated to afford 0.500 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 8.01 (d, $J = 5.1$ Hz, 1H), 8.14 (d, $J = 4.8$ Hz, 1H), 8.63 (s, 1H).

Intermediate-5

4-(5-Iodopyrimidin-2-yl)morpholine

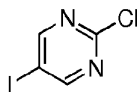


Step-1:- Preparation of 5-iodopyrimidin-2-amine



To a solution of pyrimidin-2-amine (1.0 g, 0.010 mol) in DMSO (10 mL) was added iodine (3.2 g, 0.012 mol). The reaction mixture was stirred at 120°C for 1 h. The reaction mass was quenched in water and excess of iodine was neutralised with sodium metabisulphate. The reaction mass was extracted with ethyl acetate and concentrated to afford 0.400 g of the desired product.

Step-2:- Preparation of 2-chloro-5-iodopyrimidine



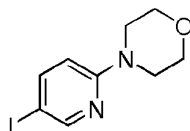
To a solution of 5-iodopyrimidin-2-amine (10.0 g, 0.045 mol) in acetonitrile (150 mL) was added CuCl_2 (11.57 g, 0.067 mol) and *tert*-butyl nitrite (6.99 g, 0.067 mol). The reaction mass was heated at 70°C for 5-6 h. The reaction mass was diluted with ether and the solid obtained was filtered off. The obtained product was purified with column chromatography on silica gel eluting with DCM to afford 1.700 g of the desired product.

Step-3:- Preparation of 4-(5-iodopyrimidin-2-yl)morpholine

The mixture of 2-chloro-5-iodopyrimidine (0.200 g, 0.836 mmol) in morpholine (3.0 mL) was refluxed for 2-3 h. The reaction mass was quenched in water and the solid obtained was filtered off. The obtained solid was dried to afford 0.180 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.63(s, 8H), 8.52 (s, 2H); MS (m/z): 292.25 (M+H) $^+$.

Intermediate-6

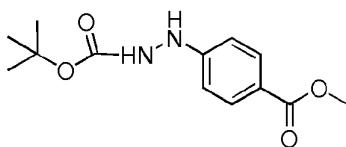
4-(5-Iodopyridin-2-yl)morpholine



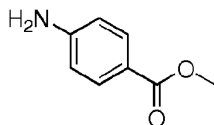
A solution of 2-chloro-5-iodo pyridine (0.200 g, 0.836 mmol) in morpholine (3.0 mL) was refluxed for 12-15 h. The reaction mass was quenched in ice and the solid obtained was filtered off to afford 0.170 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.40 (t, J = 4.8 Hz, 4H), 3.67 (t, J = 4.5 Hz, 4H), 6.74 (d, J = 8.7 Hz, 1H), 7.77-7.81 (m, 1H), 8.28 (s, 1H).

Intermediate-7

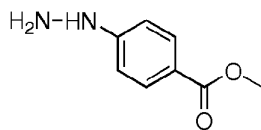
tert-Butyl 2-[4-(methoxycarbonyl)phenyl]hydrazinecarboxylate



Step-1:- Preparation of methyl 4-aminobenzoate



To a solution of 4-amino benzoic acid (20.0 g, 0.14 mol) in methanol (400 mL) was added conc. sulphuric acid (40 mL). The reaction mass was stirred at RT for 6-7 h. The reaction mass was quenched with water and basified with NaHCO_3 and extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 15.0 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.72 (s, 3H), 5.96 (br s, 2H), 6.55 (d, J = 8.1 Hz, 2H), 7.64 (d, J = 8.4 Hz, 2H) MS (m/z): 152.21 (M+H) $^+$.

Step-2:- Preparation of methyl 4-hydrazinylbenzoate

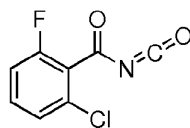
To a cold solution of methyl 4-aminobenzoate (15.0 g, 0.099 mol) in conc. HCl was added aq. solution of sodium nitrite (7.5 g, 0.109 mmol) at 0-5°C. The reaction mass was stirred at RT for 1-2 h. The reaction mass was cooled to 0°C and stannous chloride (0.049 g, 0.210 mmol) was added and further stirred at RT for 2-3 h. The reaction mass was filtered to afford 17.0 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.77 (s, 3H), 7.03 (d, *J* = 8.4 Hz, 2H), 7.82 (d, *J* = 8.4 Hz, 2H), 10.69 (br s, 2H); MS (m/z): 167.28 (M+H)⁺.

Step-3:- Preparation of *tert*-butyl 2-[4-(methoxycarbonyl)phenyl]hydrazinecarboxylate

To a solution of methyl 4-hydrazinylbenzoate (10.0 g, 0.049 mol) in DCM (150 mL) was added TEA (10 mL) and BOC anhydride (10.7 g, 0.049 mmol). The reaction mass was stirred at RT for 12 h. The reaction mass was quenched in water and extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated. The obtained solid was washed with pentane to afford 6.5 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.40 (s, 9H), 3.74 (s, 3H), 6.65 (d, *J* = 8.4 Hz, 2H), 7.74 (d, *J* = 8.7 Hz, 2H), 8.31 (s, 1H), 8.93 (br s, 1H).

Intermediate-8

2-Chloro-6-fluorobenzoyl isocyanate

Step-1:- Preparation 2-chloro-6-fluorobenzamide

A solution of 2-chloro-6-fluorobenzonitrile (8.0 g, 51.61 mmol) in conc. sulphuric acid (50 mL) was heated at 60-70°C for 6-7 h. The reaction mass was quenched in water and

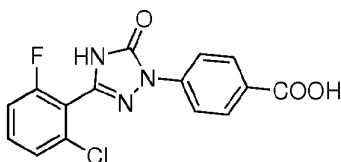
extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 5.5 g of desired product.

Step-2:- Preparation of 2-chloro-6-fluorobenzoyl isocyanate

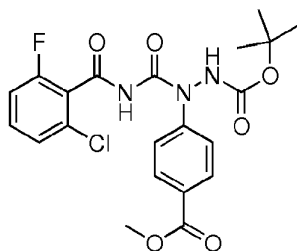
To a solution of 2-chloro-6-fluorobenzamide (2.0 g, 0.011 mmol) in EDC (10 mL) was added oxalyl chloride (2.18 g, 0.017 mmol). The reaction mass was refluxed for 24 h. Excess of solvent was removed under vacuum to afford 2.0 g of desired product.

Intermediate-9

4-[3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoic acid

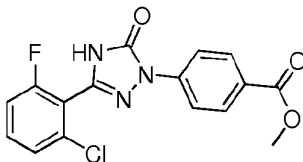


Step-1:- Preparation of *tert*-butyl 2-[(2-chloro-6-fluorobenzoyl)carbamoyl]-2-[4-(methoxycarbonyl)phenyl]hydrazinecarboxylate



To a solution of *tert*-butyl 2-[4-(methoxycarbonyl)phenyl]hydrazinecarboxylate (Intermediate-7, 2.0 g, 7.54 mmol) in DCM (15 mL) was added 2-chloro-6-fluorobenzoyl isocyanate (Intermediate-8, 1.50 g, 7.54 mmol). The reaction mass was stirred at RT for 2 h. Excess of solvent was removed under vacuum to afford 3.0 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.42 (s, 9H), 3.83 (s, 3H), 7.24-7.46 (m, 5H), 7.94 (d, *J* = 7.2 Hz, 2H), 9.91 (br s, 1H), 11.34-11.39 (br m, 1H); MS (*m/z*): 464.06 (M-H)⁺.

Step-2:- Preparation of methyl 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoate



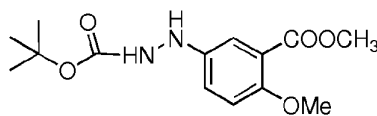
To a solution of *tert*-butyl 2-[(2-chloro-6-fluorobenzoyl)carbamoyl]-2-[4-(methoxycarbonyl)phenyl]hydrazinecarboxylate (3.0 g, 6.45 mmol) in DCM (30 mL) was added trifluoro acetic acid (2.0 mL). The reaction mass was stirred at RT for 2-3 h. Excess of solvent was removed at low temperature. The reaction mass was quenched in ice and filtered off to afford 2.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.86 (s, 3H), 7.51-7.60 (m, 2H), 7.70 (br s, 1H), 8.09 (br s, 4H), 12.79 (br s, 1H); MS (m/z): 348.55 (M+H) $^+$.

Step-3:- Preparation of 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoic acid

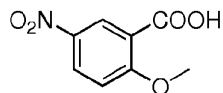
To a solution of methyl 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoate (2.0 g, 5.76 mmol) in THF: water (5 mL:10 mL) was added sodium hydroxide (0.691 g, 17.27 mmol). The reaction mass was stirred at RT for 3 h. The reaction mass was diluted with water and washed the aqueous layer with diethyl ether and toluene. The aqueous layer was cooled to 15°C and acidified with dilute HCl. The reaction mass was filtered off to afford 1.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.50 (t, J = 9.0 Hz, 1H), 7.58 (d, J = 7.8 Hz, 1H), 7.67-7.74 (m, 1H), 8.06 (s, 4H), 12.77 (br s, 1H).

Intermediate-10

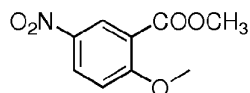
tert-Butyl 2-[4-methoxy-3-(methoxycarbonyl)phenyl]hydrazinecarboxylate



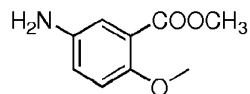
Step-1:- Preparation of 2-methoxy-5-nitrobenzoic acid



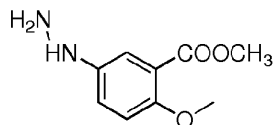
To a solution of 2-chloro-5-nitrobenzoic acid (3.0 g, 0.14 mol) in methanol (500 mL) was added sodium methoxide (28.1 g, 0.520 mol). The reaction mass was refluxed for 15 h. Excess of solvent was removed under vacuum and the reaction mass was diluted with water and acidified with dilute HCl to obtain solid which was filtered off to afford 25.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.96 (s, 3H), 7.36 (d, J = 9.3 Hz, 1H), 8.37 (d, J = 9.0 Hz, 1H), 8.45 (s, 1H), 13.32 (br s, 1H); MS (m/z): 198.25 (M+H) $^+$.

Step-2:- Preparation of methyl 2-methoxy-5-nitrobenzoate

To a solution of 2-methoxy-5-nitrobenzoic acid (25.0 g, 0.12 mol) in DMF (100 mL) was added K_2CO_3 (26.3 g, 0.19 mol). The reaction mass was stirred at 80°C for 1 h followed by addition of methyl iodide (12.3 mL, 0.19 mol). The reaction mass was further stirred at 80°C for 5-6 h. The reaction mass was filtered and the obtained filtrate was concentrated. The residue was further diluted with water and obtained solid was filtered off to afford 20.0 g of desired product. 1H NMR (300 MHz, DMSO d_6): δ 3.83 (s, 3H), 3.97 (s, 3H), 7.38 (d, $J = 9.3$ Hz, 1H), 8.41 (dd, $J = 2.4$ Hz & 2.4 Hz, 1H), 8.47 (s, 1H); MS (m/z): 212.44 (M+H) $^+$.

Step-3:- Preparation of methyl 5-amino-2-methoxybenzoate

A solution of methyl 2-methoxy-5-nitrobenzoate (20.0g, 0.097 mol) in methanol (300 mL) and 10% Pd/C (5.0 g) was stirred under hydrogen atmosphere under 70-80 psi pressure in Parr apparatus for 4-5 h. The reaction mass was filtered and the obtained filtrate was concentrated to afford 15.0 g of desired product. 1H NMR (300 MHz, DMSO d_6): δ 3.67 (s, 3H), 3.74 (s, 3H), 4.96 (br s, 2H), 6.75 (dd, $J = 2.4$ Hz, 1H), 6.85 (d, $J = 8.7$ Hz, 1H), 6.92 (s, 1H); MS (m/z): 182.25 (M+H) $^+$.

Step-4:- Preparation of methyl 5-hydrazinyl-2-methoxybenzoate

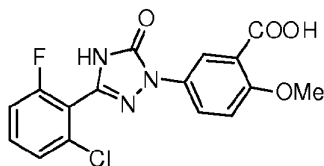
The title compound was prepared according to the procedure described in step-2 of Intermediate-7 by using methyl 5-amino-2-methoxybenzoate (15.0 g, 0.082 mol), stannous chloride (37.25 g, 0.16 mmol), sodium nitrite (6.28 g, 0.091 mmol), conc.HCl (400 mL) and water (100 mL) to afford 14.0 g of desired product. 1H NMR (300 MHz, DMSO d_6): δ 3.75 (s, 6H), 7.09 (m, 1H), 7.20-7.23 (m, 1H), 7.34 (s, 1H), 10.20 (br s, 2H); MS (m/z): 197.27 (M+H) $^+$.

Step-5:- Preparation of *tert*-butyl 2-[4-(methoxycarbonyl)phenyl]hydrazinecarboxylate

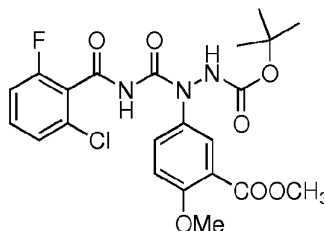
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using methyl 5-hydrazinyl-2-methoxybenzoate (10.0 g, 0.051 mol), DCM (150 mL), TEA (10 mL) and BOC anhydride (11.2 g, 0.051 mmol) to afford 2.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.43 (s, 9H), 3.71 (s, 3H), 3.79 (s, 3H), 6.81-6.85 (m, 1H), 6.97-6.99 (m, 2H), 7.45 (s, 1H), 8.78 (s, 1H) ; MS (m/z): 295.19 (M-H) $^-$.

Intermediate-11

5-[3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-2-methoxybenzoic acid

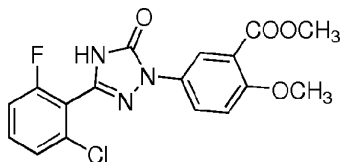


Step-1:- Preparation *tert*-butyl 2-[(2-chloro-6-fluorobenzoyl)carbamoyl]-2-[4-methoxy-3-(methoxycarbonyl)phenyl]hydrazinecarboxylate



The title compound was prepared according to the procedure described in step-1 of Intermediate-9 by using 2-chloro-6-fluorobenzoyl isocyanate (Intermediate-8, 2.0 g, 6.75 mmol), *tert*-butyl 2-[4-methoxy-3-(methoxycarbonyl)phenyl]hydrazinecarboxylate (Intermediate-10, 1.61 g, 8.18 mmol) and DCM (30 mL) to afford 3.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.40 (s, 9H), 3.81 (s, 6H), 7.13-7.83 (m, 6H), 9.6 (br s, 1H), 11.13 (br s, 1H); MS (m/z): 494.05 (M-H) $^-$.

Step-2:- Preparation of methyl 5-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-2-methoxybenzoate



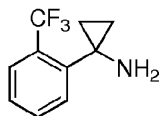
The title compound was prepared according to the procedure described in step-2 of Intermediate-9 by using *tert*-butyl 2-[(2-chloro-6-fluorobenzoyl)carbamoyl]-2-[4-methoxy-3-(methoxycarbonyl)phenyl]hydrazinecarboxylate (3.0 g, 6.06 mmol), DCM (30 mL) and trifluoro acetic acid (3.0 mL) to afford 2.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.80 (s, 3H), 3.84 (s, 3H), 7.28 (d, $J = 9.3$ Hz, 1H), 7.49 (t, $J = 8.7$ Hz, 1H), 7.57 (d, $J = 8.4$ Hz, 1H), 7.65-7.73 (m, 1H), 8.06 (d, $J = 9.0$ Hz, 1H), 8.18 (s, 1H), 12.62 (s, 1H); MS (m/z): 376.27 (M-H).

Step-3:- Preparation of 5-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl]-2-methoxybenzoic acid

The title compound was prepared according to the procedure described in step-3 of Intermediate-9 by using of methyl 5-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl]-2-methoxybenzoate (1.5 g, 3.97 mol) and sodium hydroxide (0.320 g, 7.95 mmol) to afford 0.700 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.84 (s, 3H), 7.24 (t, $J = 9.3$ Hz, 1H), 7.48 (t, $J = 8.7$ Hz, 1H), 7.57 (d, $J = 8.7$ Hz, 1H), 7.65-7.72 (m, 1H), 8.00 (d, $J = 8.7$ Hz, 1H), 8.17 (s, 1H), 12.60 (s, 1H), 12.85 (br s, 1H).

Intermediate-12

1-[2-(Trifluoromethyl)phenyl]cyclopropanamine

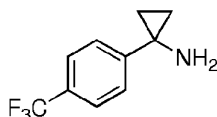


To a cold solution of 2-(trifluoromethyl)benzonitrile (1.0 g, 0.58 mmol) in diethyl ether (20 mL) was added titanium isopropoxide (2.0 g, 0.70 mmol) at -70°C and ethyl magnesium bromide (4.30 mL, 12.86 mmol). The reaction mass was stirred at RT for 2-3 h. Followed by addition of boron trifluoride solution (1.5 mL) and continued stirring for 2 h at RT. The reaction mass was quenched in 1N HCl and basified with NaOH solution. The obtained crude product was further purified by column chromatography on silica gel eluting with 1% EtOAc: DCM eluent to afford 0.250 g of desired product. ^1H NMR

(300 MHz, DMSO d_6): δ 0.82-0.84 (s, 2H), 0.91-0.93 (br s, 2H), 2.24 (br s, 2H), 7.41 (t, J = 6.6 Hz, 1H), 7.58-7.67 (m, 3H).

Intermediate-13

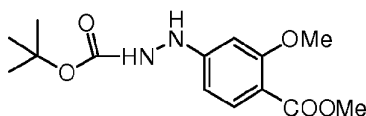
1-[4-(Trifluoromethyl)phenyl]cyclopropanamine



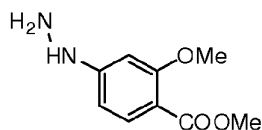
The title compound was prepared according to the procedure described in Intermediate-12 by using of 4-(trifluoromethyl)benzonitrile (1.0 g, 0.58 mmol), diethyl ether (20 mL), titanium isopropoxide (2.0 g, 0.70 mmol), ethyl magnesium bromide (4.30 mL, 12.86 mmol) and boron trifluoride solution (1.5 mL) to afford 0.210 g of desired product. ^1H NMR (CDCl_3): δ 1.05 (s, 2H), 1.60 (s, 2H), 1.78 (s, 2H), 7.36 (d, J = 7.8 Hz, 2H), 7.55 (d, J = 8.1 Hz, 2H).

Intermediate-14

tert-Butyl 2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate



Step-1:- Preparation of methyl 4-hydrazinyl-2-methoxybenzoate



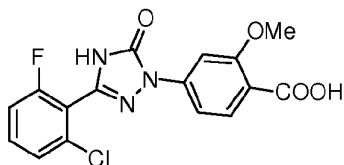
The title compound was prepared according to the procedure described in step-2 of Intermediate-7 by using methyl 4-amino-2-methoxybenzoate (10.0 g, 0.055 mol), stannous chloride (31.00 g, 0.138 mmol), sodium nitrite (4.57 g, 0.066 mmol), 6 N HCl (200 mL) and water (100 mL) to afford 14.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.73 (s, 3H), 3.80 (s, 3H), 6.51 (d, J = 8.4 Hz, 1H), 6.69 (s, 1H), 7.68 (d, J = 8.4 Hz, 1H), 8.75 (br s, 1H), 10.28 (br hump, 2H); MS (m/z): 197.01 ($\text{M}+\text{H}$) $^+$.

Step-2:- Preparation of *tert*-butyl 2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazine carboxylate

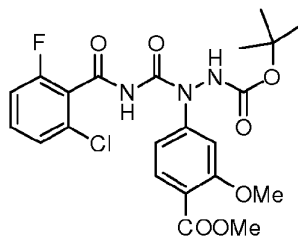
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using methyl 4-hydrazinyl-2-methoxybenzoate (5.0 g, 0.025 mol), DCM (70 mL), TEA (5.0 mL) and BOC anhydride (6.11 g, 0.028 mmol) to afford 3.50 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.40 (s, 9H), 3.68 (s, 3H), 3.72 (s, 3H), 6.26 (s, 2H), 7.59 (d, J = 8.7 Hz, 1H), 8.26 (s, 1H), 8.94 (s, 1H); MS (m/z): 297.02 (M+H) $^+$.

Intermediate-15

4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxybenzoic acid

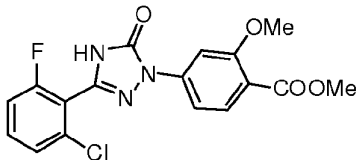


Step-1:- Preparation of *tert*-butyl 2-((2-chloro-6-fluorobenzoyl)carbamoyl)-2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate



The title compound was prepared according to the procedure described in step-1 of Intermediate-9 by using 2-chloro-6-fluorobenzoyl isocyanate (Intermediate-8, 2.0 g, 6.75 mmol), *tert*-butyl 2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate (Intermediate-14, 2.0 g, 10.10 mmol) and DCM (30 mL) to afford 3.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.42 (s, 9H), 3.76 (s, 6H), 6.99 (d, J = 8.1 Hz, 1H), 7.11 (s, 1H), 7.2-7.50 (m, 4H), 9.88 (s, 1H), 11.32 (s, 1H); MS (m/z): 493.92 (M-H) $^-$.

Step-2:- Preparation of methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxybenzoate



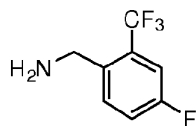
The title compound was prepared according to the procedure described in step-2 of Intermediate-9 by using *tert*-butyl 2-((2-chloro-6-fluorobenzoyl)carbamoyl)-2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate (3.0 g, 6.06 mmol), DCM (30 mL) and trifluoro acetic acid (2.0 mL) to afford 2.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.81 (s, 3H), 3.85 (s, 3H), 7.34-7.73 (m, 6H), 12.80 (s, 1H); MS (m/z): 376.16 (M-H) $^-$.

Step-3:- Preparation of 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxybenzoic acid

The title compound was prepared according to the procedure described in step-3 of Intermediate-9 by using of methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxybenzoate (1.0 g, 2.65 mol) and sodium hydroxide (0.210 g, 5.30 mmol) to afford 0.600 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.84 (s, 3H), 7.59 (t, J = 7.8 Hz, 2H), 7.67-7.75 (m, 3H), 7.81 (d, J = 8.4 Hz, 1H), 12.58 (br, 1H), 12.80 (s, 1H); MS (m/z): 362.21 (M-H) $^-$.

Intermediate-16

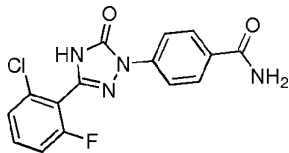
(4-Fluoro-2-(trifluoromethyl)phenyl)methanamine



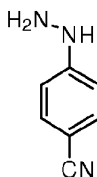
To a solution of 4-fluoro-2-(trifluoromethyl)benzonitrile (2.0 g) in ethanol (10.0 mL) was added Raney Ni (catalytic amount). The reaction mixture was subjected for hydrogenation in Parr apparatus under 50 psi for 2-3 h. The reaction mass was filtered through celite and the filtrate was concentrated to afford 0.400 g of desired product. ^1H NMR (CDCl_3): δ 3.98 (s, 2H), 7.24 (t, J = 9.0 Hz, 1H), 7.34 (d, J = 8.7 Hz, 1H), 7.54 (t, J = 7.8 Hz, 1H); MS $[\text{M}+\text{H}]^+$: 194.03.

Intermediate-17

4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)benzamide

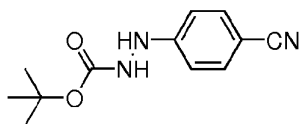


Step-1:- Preparation of 4-hydrazinylbenzonitrile



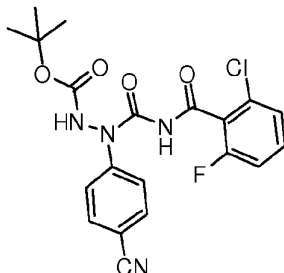
To a cold solution of 4-cyanoaniline (6.0 g, 0.050 mol) in conc. HCl was added aq. solution of sodium nitrite (3.85 g, 0.055 mmol) at -15°C . The reaction mass was stirred at $0-10^{\circ}\text{C}$ for 15 minutes and filtered off to remove insolubles. The filtrate was added to stannous chloride in conc. HCl (24.0 g, 0.166 mmol). The reaction mass was stirred at -15°C for 30 minutes. The reaction mass was filtered to afford 5.2 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.04 (d, 2H), 7.70 (d, 2H), 9.17 (br s, 1H), 10.66 (br s, 2H).

Step-2:- Preparation of *tert*-butyl 2-(4-cyanophenyl)hydrazinecarboxylate



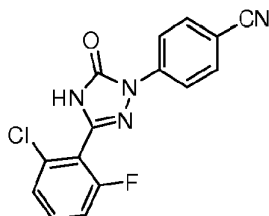
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using 4-hydrazinylbenzonitrile (10.0 g, 0.059 mmol), BOC anhydride (14.5 g, 0.065 mmol), TEA and DCM to afford 12.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.41 (s, 9H), 6.70 (d, $J = 8.4$ Hz, 2H), 7.53 (d, $J = 8.4$ Hz, 2H), 8.48 (br s, 1H), 9.00 (br s, 1H); $(\text{M}+\text{H})^+$. 233.94.

Step-3:- Preparation of *tert*-butyl 2-((2-chloro-6-fluorobenzoyl)carbamoyl)-2-(4-cyanophenyl)hydrazinecarboxylate



To a solution of *tert*-butyl 2-(4-cyanophenyl)hydrazinecarboxylate (2.5 g, 0.01 mmol) in DCM (15 mL) was added 2-chloro-6-fluorobenzoyl isocyanate (Intermediate-8, 2.56 g, 0.012 mmol). The reaction mass was stirred at RT for 2 h. Excess of solvent was removed under vacuum to afford 4.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.42 (s, 9H), 7.25-7.38 (m, 2H), 7.47-7.60 (m, 3H), 7.82-7.85 (m, 2H), 9.96 (br s, 1H), 11.43 (br s, 1H); MS (m/z): 431.02 (M+H) $^+$.

Step-4:- Preparation of 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzonitrile



The title compound was prepared according to the procedure described in step-2 of Intermediate-9 by using *tert*-butyl 2-((2-chloro-6-fluorobenzoyl)carbamoyl)-2-cyanophenylhydrazinecarboxylate (4.0 g, 0.009 mol), DCM (30 mL), trifluoro acetic acid (5.0 mL) to afford 2.2 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.50 (t, 1H), 7.58 (d, 1H), 7.70 (q, 1H), 7.95 (d, J = 8.7 Hz, 2H), 7.53 (d, J = 8.4 Hz, 2H), 12.80 (br s, 1H); MS (m/z): 331.80 (M+H) $^+$.

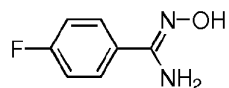
Step-5:- Preparation of 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzamide

A solution of 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzonitrile (1.0 g, 0.003 mol) in conc. sulphuric acid (10 mL) was heated to 70°C for 15 h. The reaction mixture was quenched in ice water. The reaction mass was basified till pH ~ 6 -6.5 with dilute NaOH. The obtained solid was filtered off, washed with water and

suck dried to afford 0.300 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.36 (br s, 1H), 7.84 (t, 1H), 7.57 (d, 1H), 7.67 (d, 1H), 7.98 (m, 5H), 12.70 (br s, 1H); MS (m/z): 333.11 (M+H) $^+$.

Intermediate-18

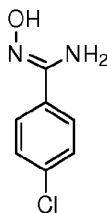
4-Fluoro-*N'*-hydroxybenzimidamide



To a solution of 4-fluorobenzonitrile (0.500 g, 4.13 mmol) in ethanol (3 mL) was added hydroxyl amine HCl (0.427g, 6.19 mmol) and potassium carbonate (1.14g, 8.26 mmol). The reaction mass was stirred at RT for 15-17 h. Excess of solvent was removed under vacuum. The obtained residue was diluted with water, acidified with dilute HCl and extracted with DCM. The organic layer was separated, dried over anhydrous sodium sulphate and concentrated to afford 0.450 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 5.84 (br s, 2H), 7.17-7.27 (m, 2H), 7.68-7.73 (m, 2H), 9.64 (br s, 1H); MS (m/z): 155.13 (M+H) $^+$.

Intermediate-19

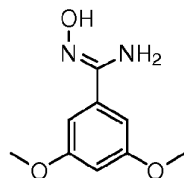
4-Chloro-*N'*-hydroxybenzimidamide



To a solution of 4-chloro benzonitrile (1.000 g, 7.26 mmol) in ethanol (20 mL) was added hydroxyl amine HCl (0.752 g, 10.90 mmol) and potassium carbonate (3.00 g, 21.80 mmol). The reaction mass was refluxed for 10-12 h. Excess of solvent was removed under vacuum and the residue was diluted with water, acidified with dilute HCl. Precipitate obtained was filtered off and sucked dried to afford 0.500 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 5.88 (br s, 2H), 7.42 (d, J = 8.1 Hz, 2H), 7.68 (d, J = 8.4 Hz, 2H), 9.70 (br s, 1H); MS (m/z): 171.13 (M+H) $^+$.

Intermediate-20

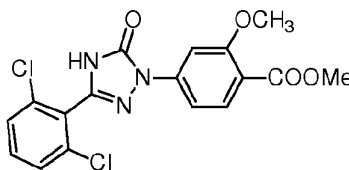
N'-Hydroxy-3,5-dimethoxybenzimidamide



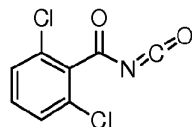
The title compound was prepared according to the procedure described in Intermediate-19 by using 3,5-dimethoxy benzonitrile (1.00 g, 6.13 mmol), hydroxyl amine HCl (0.634 g, 9.23 mmol) and potassium carbonate (2.53g, 18.40 mmol), ethanol (20 mL) to afford 0.400 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.75 (s, 6H), 5.79 (s, 2H), 6.49 (s, 1H), 6.84 (s, 2H), 9.62 (br s, 1H); MS (m/z): 197.11 (M+H) $^+$.

Intermediate-21

Methyl 4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxybenzoate

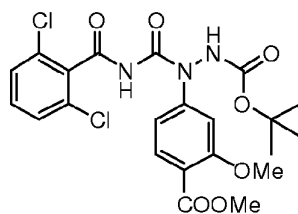


Step-1:- Preparation of 2,6-dichlorobenzoyl isocyanate



The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 2,6-dichlorobenzamide (1.00 g, 5.26 mmol), oxalyl chloride (0.795 g, 6.31 mmol) and toluene (10 mL) to afford 1.00 g of desired product.

Step-2:- Preparation of *tert*-butyl 2-((2,6-dichlorobenzoyl)carbamoyl)-2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate



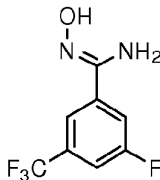
The title compound was prepared according to the procedure described in step-1 of Intermediate-9 by using 2,6-dichlorobenzoyl isocyanate (1.0 g, 3.38 mmol), *tert*-butyl 2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate (Intermediate-14, 0.805 g, 3.72 mmol) and DCM (20 mL) to afford 1.5 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.42 (s, 9H), 3.76 (s, 6H), 6.99 (d, J = 6.0 Hz, 1H), 7.10 (s, 1H), 7.60-7.37 (m, 3H), 7.67 (d, J = 7.8 Hz, 1H), 9.86 (br s, 1H), 11.30 (br s, 1H).

Step-3:- Preparation of methyl 4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate

The title compound was prepared according to the procedure described in step-2 of Intermediate-9 by using *tert*-butyl 2-((2,6-dichlorobenzoyl)carbamoyl)-2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate (1.5 g, 2.2 mmol), DCM (50 mL) and trifluoro acetic acid (5.0 mL) to afford 0.900 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.78 (s, 3H), 3.84 (s, 3H), 7.62- 7.30 (m, 5H), 7.82 (d, J = 8.7 Hz, 1H), 12.76 (br s, 1H); MS (m/z): 393.85 ($M+H$) $^+$.

Intermediate-22

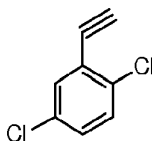
3-Fluoro-*N'*-hydroxy-5-(trifluoromethyl)benzimidamide



The title compound was prepared according to the procedure described in Intermediate-19 by using 3-fluoro-5-(trifluoromethyl)benzonitrile (2.00 g, 10.0 mmol), hydroxyl amine .HCl (1.09 g, 15 mmol) and potassium carbonate (2.2 g, 15 mmol), ethanol (20 mL) to afford 0.900 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 6.10 (s, 2H), 7.69 (d, J = 7.8 Hz, 1H), 7.79 (d, J = 10.2 Hz, 1H), 10.03 (s, 1H); MS (m/z): 223.17 ($M+H$) $^+$.

Intermediate-23

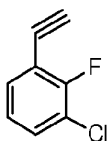
1,4-Dichloro-2-ethynylbenzene



The title compound was prepared according to the procedure described in step-5 and step-6 of Intermediate-2 by using 1,4-dichloro-2-iodobenzene (1.0 g, 3.6 mmol), ethynyl(trimethyl)silane (0.541 g, 5.5 mmol), copper iodide (0.027 g, 0.14 mmol), bis(triphenylphosphine) palladium(II) chloride (0.050 g, 0.072 mmol), TBAF (catalytic) and DCM to afford 0.550 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 4.73 (s, 1H), 7.50 (d, J = 8.7 Hz, 1H), 7.58 (d, J = 8.7 Hz, 1H), 7.71 (s, 1H).

Intermediate-24

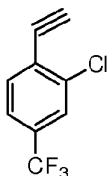
1-Chloro-3-ethynyl-2-fluorobenzene



The title compound was prepared according to the procedure described in step-5 and step-6 of Intermediate-2 by using 1-chloro-2-fluoro-3-iodobenzene (1.0 g, 3.9 mmol), ethynyl(trimethyl)silane (0.541 g, 5.5 mmol), copper iodide (0.027 g, 0.14 mmol), bis(triphenylphosphine) palladium(II) chloride (0.050 g, 0.072 mmol), TBAF (catalytic) and DCM to afford 0.500 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 4.66 (s, 1H), 7.25 (t, J = 8.4 Hz, 1H), 7.54 (t, J = 7.2 Hz, 1H), 7.65 (t, J = 7.8 Hz, 1H).

Intermediate-25

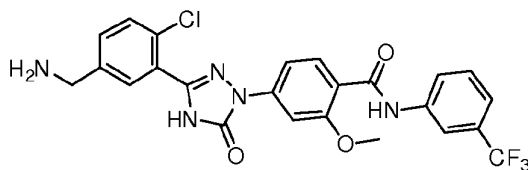
2-Chloro-1-ethynyl-4-(trifluoromethyl)benzene



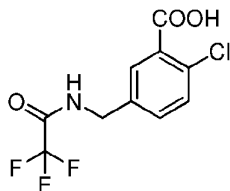
The title compound was prepared according to the procedure described in step-5 and step-6 of Intermediate-2 by using 2-chloro-1-iodo-4-(trifluoromethyl)benzene (1.0 g, 3.2 mmol), ethynyl(trimethyl)silane (0.541 g, 5.5 mmol), copper iodide (0.027 g, 0.14 mmol), bis(triphenylphosphine) palladium(II) chloride (0.050 g, 0.072 mmol), TBAF (catalytic) and DCM to afford 0.525 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 4.88 (s, 1H), 7.73 (d, J = 8.4 Hz, 1H), 7.83 (d, J = 7.8 Hz, 1H), 8.01 (s, 1H).

Intermediate-26

4-(3-(5-(Aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide

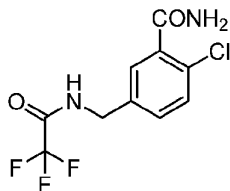


Step 1: Preparation of 2-chloro-5-[[[(trifluoroacetyl)amino]methyl]benzoic acid



To a solution of 2-chlorobenzoic acid (0.500 g, 3.49 mmol) in conc. H_2SO_4 was added 2,2,2-trifluoro-*N*-(hydroxymethyl)acetamide (0.547 g, 3.49 mmol). The mixture was stirred at RT for 16 h. The reaction mixture was poured into ice-water and stirred for 2 h. The precipitate obtained was collected by filtration, dried and re-crystallized from toluene/butan-2-one (7:1) to afford 0.800 g of the title product. ^1H NMR (300 MHz, $\text{DMSO } d_6$): δ 4.42 (d, $J = 6.0$ Hz, 2H), 7.43 (d, $J = 9.9$ Hz, 1H), 7.54 (d, $J = 8.4$ Hz, 1H), 7.71 (s, 1H), 10.06 (br s, 1H), 13.47 (br s, 1H); MS (m/z): 280.18 (M-H) $^-$.

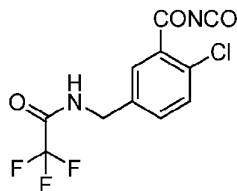
Step 2: Preparation of 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzamide



To a cold solution of 2-chloro-5-[[[(trifluoroacetyl)amino]methyl]benzoic acid (1.50 g, 5.33 mmol) in THF : DCM (20:10 mL) was added oxalyl chloride (0.6 mL, 6.40 mmol) and DMF (2-3 drop) at 0°C . The reaction mixture was stirred at RT for 2 h and concentrated. A solution of the concentrated mass in THF (15 mL) was treated with ammonia gas (purged through reaction mass) at 0°C and the reaction mixture was stirred at RT for 1 h. The reaction mixture was diluted with ethyl acetate. The reaction mixture was washed with water, dilute HCl and brine. The organic layer was separated, dried, filtered and concentrated to afford 0.800 g of the desired product. ^1H NMR (300 MHz,

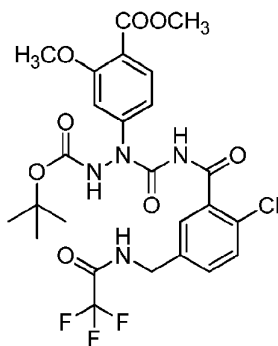
DMSO d_6): δ 4.39 (s, 2H), 7.30- 7.34 (m, 2H), 7.46 (d, J = 7.8 Hz, 1H), 7.63 (s, 1H), 7.90 (s, 1H), 10.06 (s, 1H).

Step 3: Preparation of 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate



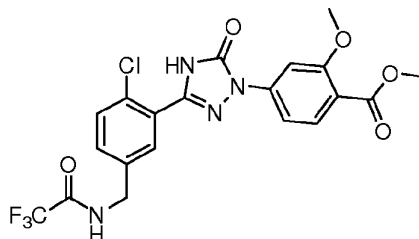
The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzamide (0.700 g, 2.5 mmol), oxalyl chloride (0.3 mL, 3.0 mmol) and EDC (30 mL) to afford 0.700 g of the desired product.

Step 4: Preparation of *tert*-butyl 2-((2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl)carbamoyl)-2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate



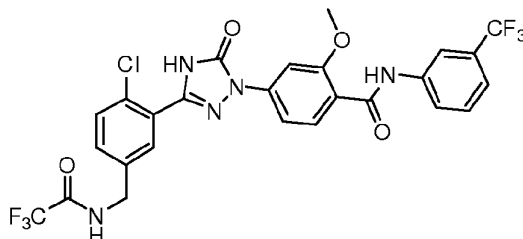
The title compound was prepared according to the procedure described in step-1 of Intermediate-9 by using 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate (0.700 g, 2.2 mmol), *tert*-butyl 2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate (Intermediate-14, 0.675 mL, 2.20 mmol) and DCM (30 mL) to afford 1.2 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.39 (s, 9H), 3.09 (m, 6H), 4.48 (s, 2H), 6.22- 6.26 (m, 2H), 7.13- 7.76 (m, 4H), 8.94 (s, 1H), 10.08 (s, 1H), 11.02 (s, 1H); MS (m/z): (M) $^+$.602.72.

Step 5: Preparation of methyl 4-(3-(2-chloro-5-((2,2,2-trifluoroacetamido)methyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate



The title compound was prepared according to the procedure described in step-2 of Intermediate-9 by using *tert*-butyl 2-((2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl)carbamoyl)-2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate (1.20 g, 1.99 mmol), TFA (2 mL) and DCM (20 mL) to afford 0.280 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.78 (s, 3H), 3.85 (s, 3H), 4.46 (d, J = 6.0 Hz, 2H), 7.51 (t, J = 8.1 Hz, 1H), 7.65- 7.68 (m, 3H), 7.78- 7.84 (m, 2H), 10.09 (m, 1H), 12.70 (br s, 1H); MS (m/z): (M) $^+$.484.95.

Step 6: Preparation of 4-(3-(2-chloro-5-((2,2,2-trifluoroacetamido)methyl)phenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide



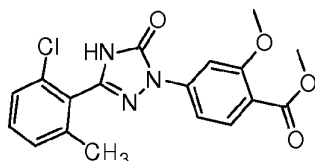
To a solution of methyl 4-(3-(2-chloro-5-((2,2,2-trifluoroacetamido) methyl)phenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxybenzoate (0.200 g, 0.413 mmol) in dry toluene (3 mL) was added 3-(trifluoromethyl)aniline (0.100 g, 0.619 mmol) followed by addition of trimethyl aluminium (2M solution in toluene) (2 mL). The reaction mass was refluxed for 3-4 h. The reaction mass was quenched in water and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 0.170 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.95 (s, 3H), 4.46 (d, J = 6.0 Hz, 2H), 7.56 (t, J = 6.0 Hz, 2H), 7.59- 7.85 (m, 6H), 7.96 (d, J = 7.8 Hz, 1H), 8.26 (s, 1H), 10.09 (s, 1H), 10.37 (s, 1H), 12.65 (s, 1H); MS (m/z): (M) $^+$.614.08.

Step 7: Preparation of 4-(3-(5-(aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide

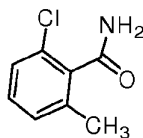
To a solution of 4-(3-(2-chloro-6-((2,2,2-trifluoroacetamido)methyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide (0.150 g, 0.24 mmol) in THF was added aq. KOH (0.028 g, 0.48 mmol). The mixture was stirred at RT for 3-4 h. The reaction mixture was poured into water and extracted in DCM. Organic layer was dried and concentrated to afford 0.080 g of the title product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.92-3.96 (m, 5H), 7.43- 7.61 (m, 5H), 7.70- 7.85 (m, 3H), 7.97 (m, 2H), 8.26 (s, 1H), 10.35 (m, 1H).

Intermediate-27

Methyl 4-(3-(2-chloro-6-methylphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate

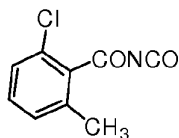


Step 1: Preparation of 2-chloro-6-methylbenzamide



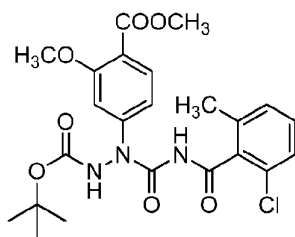
A solution of 2-chloro-6-methylbenzonitrile (2.0 g) in conc. sulphuric acid (10 mL) was heated at 100°C for 3-4 h. The reaction mass was quenched in ice-water and the solid obtained was filtered. The precipitate was suck dried to afford 1.5 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 2.28 (s, 3H), 7.21- 7.27 (m, 3H), 7.65 (s, 1H), 7.91 (s, 1H).

Step 2: Preparation of 2-chloro-6-methylbenzoyl isocyanate



The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 2-chloro-6-methylbenzamide (1.0 g, 5.9 mmol), oxalyl chloride (0.894 g, 7.1 mmol) and EDC (20 mL) to afford 1.00 g of the desired product.

Step 3: Preparation of *tert*-butyl 2-((2-chloro-6-methylbenzoyl)carbamoyl)-2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate



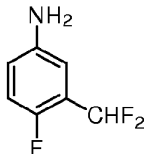
The title compound was prepared according to the procedure described in step-1 of Intermediate-9 by using 2-chloro-6-methylbenzoyl isocyanate (0.793 g, 4.06 mmol), *tert*-butyl 2-(3-methoxy-4-(methoxycarbonyl)phenyl) hydrazinecarboxylate (Intermediate - 14, 1.2 g, 4.06 mmol) and DCM (30 mL) to afford 1.5 g of the desired product.

Step 4: Preparation of methyl 4-(3-(2-chloro-6-methylphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate

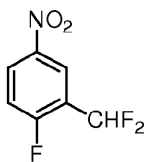
The title compound was prepared according to the procedure described in step-2 of Intermediate-9 by using *tert*-butyl 2-((2-chloro-6-methylbenzoyl)carbamoyl)-2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate (2.0 g), TFA (5 mL) and DCM (30 mL) to afford 0.700 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 2.31 (s, 3H), 3.78 (s, 3H), 3.84 (s, 3H), 7.40 (s, 1H), 7.49 (s, 2H), 7.64 (d, *J* = 8.4 Hz, 1H), 7.75 (s, 1H), 7.81 (d, *J* = 8.7 Hz, 1H), 12.54 (s, 1H); MS (*m/z*): 373.95 (M+H)⁺.

Intermediate-28

3-(Difluoromethyl)-4-fluoroaniline



Step 1: Preparation of 2-(difluoromethyl)-1-fluoro-4-nitrobenzene



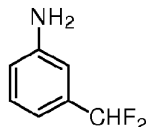
A solution of 2-fluoro-5-nitrobenzaldehyde (3.0 g, 17 mmol) in DCM (50 mL) was added DAST (3.42 g, 21 mmol) and stirred at RT for 18 h under nitrogen atmosphere. The reaction mass was quenched in ice-water and extracted with DCM. The organic layer was dried and concentrated to afford 2.5 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.31 (s, 1H), 7.69 (t, $J = 8.7$ Hz, 1H), 8.46-8.50 (m, 2H).

Step 2: Preparation of 3-(difluoromethyl)-4-fluoroaniline

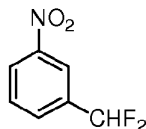
A solution of 2-(difluoromethyl)-1-fluoro-4-nitrobenzene (1.2 g, 6.2 mmol) in methanol (20 mL) was added iron powder (4.8 g, 24.8 mmol) followed by conc. HCl (5 mL) drop-wise. The reaction mass was stirred at RT for 1-2 h. The reaction mass was quenched in ice-water, basified with NaHCO_3 and extracted with DCM. The organic layer was dried and concentrated to afford 0.800 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 5.25 (s, 2H), 6.66-6.73 (m, 2H), 6.86- 7.04 (m, 2H).

Intermediate-29

3-(Difluoromethyl)aniline



Step 1: Preparation of 1-(difluoromethyl)-3-nitrobenzene



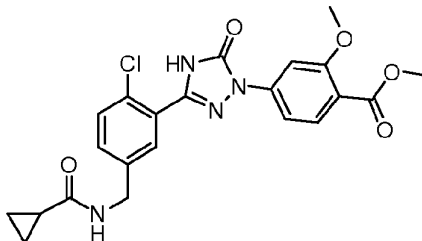
The title compound was prepared according to the procedure described in step-1 of Intermediate-28 by using 3-nitrobenzaldehyde (2.0 g, 13 mmol), DAST (2.55 g, 15 mmol) and DCM (30 mL) to afford 1.5 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.21 (s, 1H), 7.83 (t, $J = 8.1$ Hz, 1H), 8.05 (m, 1H), 8.40 (br s, 2H).

Step 2: Preparation of 3-(difluoromethyl)aniline

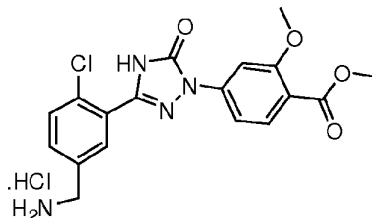
The title compound was prepared according to the procedure described in step-2 of Intermediate-28 by using 1-(difluoromethyl)-3-nitrobenzene (1.0 g, 5.0 mmol), Iron powder (3.0 g, 15.0 mmol), conc. HCl (5 mL) and methanol (20 mL) to afford 0.700 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 5.36 (br s, 2H), 6.62- 6.82 (m, 4H), 7.11 (t, $J = 7.8$ Hz, 1H); MS (m/z): 144.05 (M+H) $^+$.

Intermediate-30

Methyl 4-(3-(2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate



Step 1: Preparation of methyl 4-(3-(5-(aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate hydrochloride



A solution of methyl 4-(3-(2-chloro-5-((2,2,2-trifluoroacetamido)methyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-5 of Intermediate-26, 0.500 g) in methanol (20 mL) was added conc. HCl (5 mL) dropwise. The reaction mass was refluxed for 24 h. The reaction mass was concentrated to afford 0.450 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.78 (s, 3H), 3.85 (s, 3H), 4.10 (d, *J* = 5.4 Hz, 2H), 7.63 (d, *J* = 9.0 Hz, 1H), 7.74- 7.84 (m, 4H), 7.94 (s, 1H), 8.57 (s, 2H), 12.81 (s, 2H); MS (*m/z*): 388.96 (M+H)⁺.

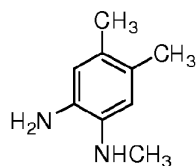
Step 2: Preparation of methyl 4-(3-(2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate

A solution of methyl 4-(3-(5-(aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate hydrochloride (0.500 g, 1.29 mmol) in THF (20 mL) was added DIPEA (2 mL) and cyclopropyl carbonyl chloride (0.201 g, 1.9 mmol) under nitrogen atmosphere. The reaction mass was stirred at RT for 2-4 h. The reaction mass was quenched in water, extracted with DCM and concentrated to afford 0.400 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.69 (m, 4H), 1.60 (s, 1H), 3.79 (s,

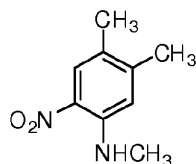
3H), 3.86 (s, 3H), 4.33 (d, $J = 5.4$ Hz, 2H), 7.41- 7.54 (m, 3H), 7.61 (d, $J = 8.4$ Hz, 1H), 7.73 (s, 1H), 7.79 (d, 1H), 8.71 (m, H); MS (m/z): 456.97 (M+H)⁺.

Intermediate-31

*N*¹,4,5-Trimethylbenzene-1,2-diamine



Step 1: Preparation of *N*¹,4,5-trimethyl-2-nitroaniline



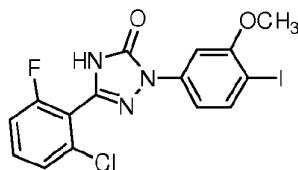
A solution of 4,5-dimethyl-2-nitroaniline (2.00 g, 12 mmol) in toluene (25 mL) was added NaOH (1.93 g, 48 mmol). The reaction mass was stirred at 100°C for 1 h, followed by addition of dimethyl sulphate (4.6 g, 36 mmol) and reaction mass was stirred at RT for 24 h. The reaction mass was quenched in water, extracted with DCM and concentrated to afford 1.5 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 2.11 (s, 3H), 2.24 (d, $J = 7.8$ Hz, 3H), 2.92 (s, 3H), 6.77 (m, 1H), 7.79 (m, 1H), 8.06 (s, 1H); MS (m/z): 180.92 (M+H)⁺.

Step 2: Preparation of *N*¹,4,5-trimethylbenzene-1,2-diamine

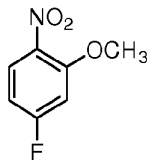
The title compound was prepared according to the procedure described in step-2 of Intermediate-28 by using *N*¹,4,5-trimethyl-2-nitroaniline (0.350 g, 1.9 mmol), methanol (20 mL), Iron powder (1.75 g, 9.75 mmol), conc. HCl (5 mL) to afford 0.120 g of desired product.

Intermediate-32

3-(2-Chloro-6-fluorophenyl)-1-(4-iodo-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one

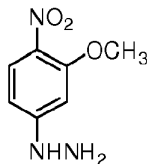


Step-1:- Preparation of 4-fluoro-2-methoxy-1-nitrobenzene



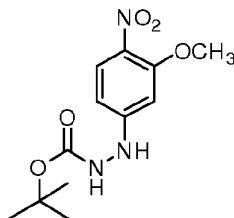
To a solution of 4-fluoro-2-hydroxy-1-nitrobenzene (5.0 g, 31.8 mmol) in DMF (10 mL) was added K_2CO_3 (13.1 g, 95.4 mmol). The reaction mixture was stirred at RT for 1 h followed by addition of methyl iodide (9.93 g, 69.9 mmol) and the reaction mixture was stirred at 60°C for 2 h. The reaction mass was concentrated and quenched in water. The reaction mass was basified with saturated sodium bicarbonate solution and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 4.5 g of desired product. 1H NMR (300 MHz, DMSO d_6): δ 3.93 (s, 3H), 6.97 (t, $J = 7.8$ Hz, 1H), 7.31 (d, $J = 8.7$ Hz, 1H), 8.02 (s, 1H).

Step-2:- Preparation of (3-methoxy-4-nitrophenyl)hydrazine



To a solution of 4-fluoro-2-methoxy-1-nitrobenzene (2.5 g, 14.60 mmol) in ethanol (25 mL) was added hydrazine hydrate (2.1 g, 43.8 mmol). The reaction mixture was refluxed for 3-4 h. The reaction mixture was concentrated to afford 2.0 g of desired product. 1H NMR (300 MHz, DMSO d_6): δ 3.84 (s, 3H), 4.46 (s, 2H), 6.31 (d, $J = 9.3$ Hz, 1H), 6.50 (s, 1H), 7.83 (d, $J = 9.3$ Hz, 1H), 8.25 (s, 1H); MS (m/z): (184.01 M) $^+$.

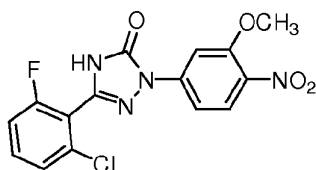
Step-3:- Preparation of *tert*-butyl 2-(3-methoxy-4-nitrophenyl)hydrazinecarboxylate



The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using (3-methoxy-4-nitrophenyl)hydrazine (0.500 g, 2.68 mmol), BOC anhydride (0.703 g, 3.22 mmol), sodium carbonate (0.426 g, 4.02 mmol) and DCM to afford 0.200 g of desired product. 1H NMR (300 MHz, DMSO d_6): δ 1.36 (s, 9H), 3.81

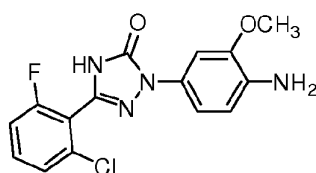
(s, 3H), 6.25- 6.29 (m, 2H), 7.87 (d, $J = 8.7$ Hz, 1H), 7.78 (s, 1H), 9.12 (s, 1H); MS (m/z): 283.88 (M)⁺.

Step-4:- Preparation of 3-(2-chloro-6-fluorophenyl)-1-(3-methoxy-4-nitrophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



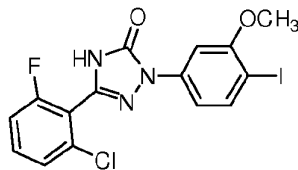
The title compound was prepared according to the procedure described in step-1 and step-2 of Intermediate-9 by using *tert*-butyl 2-(3-methoxy-4-nitrophenyl)hydrazinecarboxylate (1.0 g) and 2-chloro-6-fluorobenzoyl isocyanate (Intermediate-8, 1.0 g), TFA (5 mL) and DCM (40 mL) to afford 0.900 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.96 (s, 3H), 7.51 (t, $J = 8.7$ Hz, 1H), 7.59 (d, $J = 7.8$ Hz, 1H), 7.68- 7.73 (m, 2H), 7.88 (s, 1H), 8.09 (d, $J = 8.7$ Hz, 1H), 12.93 (s, 1H); MS (m/z): 363 (M)⁺.

Step-5:- Preparation of 1-(4-amino-3-methoxyphenyl)-3-(2-chloro-6-fluorophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



The title compound was prepared according to the procedure described in step-2 of Intermediate- 28 by using 3-(2-chloro-6-fluorophenyl)-1-(3-methoxy-4-nitrophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (0.100 g), iron powder (catalytic), conc.HCl (5-6 drops) and methanol (5 mL) to afford 0.070 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.77 (s, 3H), 4.81 (s, 2H), 6.66 (d, $J = 8.7$ Hz, 1H), 7.17 (d, $J = 8.1$ Hz, 1H), 7.26 (s, 1H), 7.50- 7.57 (m, 2H), 7.68 (m, 1H), 12.39 (s, 1H).

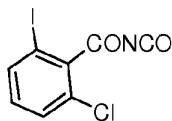
Step-6:- Preparation of 3-(2-chloro-6-fluorophenyl)-1-(4-iodo-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



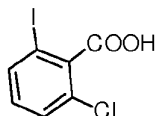
To a solution of 1-(4-amino-3-methoxyphenyl)-3-(2-chloro-6-fluorophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (0.280 g, 0.83 mmol) in acetonitrile (5.0 ml) was added PTSA (0.477 g, 2.51 mmol). The reaction mass was stirred at RT for 1h followed by addition of potassium iodide (0.347 g, 2.0 mmol) and sodium nitrite (0.115 g, 1.67 mmol) at 0-5°C and further stirred for 2h. Excess solvent was removed under vacuum and the residue reaction mass was quenched in water and extracted with ethyl acetate. The organic layer was washed with sodium metabisulphate and concentrated to obtain a crude product, which was further purified by column chromatography on silica gel eluting with 10% EtOAc: pet. ether to afford 0.200 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.85 (s, 3H), 7.40 (d, *J* = 8.7 Hz, 1H), 7.46- 7.74 (m, 4H), 7.83 (d, *J* = 8.1 Hz, 1H), 12.71 (br s, 1H); MS (*m/z*): 445.84 (M+H)⁺.

Intermediate-33

2-Chloro-6-iodobenzoyl isocyanate



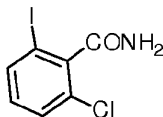
Step-1:- Preparation of 2-chloro-6-iodobenzoic acid



To the cold solution of 2-amino-6-chlorobenzoic acid (2.0 g, 11.6 mmol) in conc. HCl (10 mL) was added aq. solution of sodium nitrite (0.8 g, 11.6 mmol) at 0°C followed by addition of solution of potassium iodide (2.88 g, 17.4 mmol) in water and conc. sulphuric acid (1 mL). The reaction mixture was refluxed for 2 h. The reaction mass was washed with sodium metabisulphate solution and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 2.0 g

of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.08 (t, J = 7.8 Hz, 1H), 7.42 (d, J = 8.4 Hz, 1H), 7.76 (d, J = 7.8 Hz, 1H), 8.37 (br s, 1H); MS (m/z): 280.59 (M) $^-$.

Step-2:- Preparation of 2-chloro-6-iodobenzamide



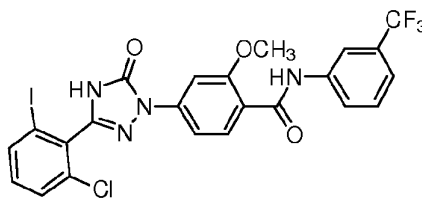
The title compound was prepared according to the procedure described in step-2 of Intermediate-26 by using 2-chloro-6-iodobenzoic acid (2.7 g, 9.6 mmol), oxalyl chloride (1.4 g, 11.5 mmol), ammonia gas, THF (20 mL) and DCM (10 mL) to afford 2.3 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.11 (t, J = 7.8 Hz, 1H), 7.49 (d, J = 8.4 Hz, 1H), 7.73 (s, 1H), 7.80 (d, J = 7.8 Hz, 1H), 8.00 (s, 1H); MS (m/z): 281.98 (M+H) $^+$.

Step-3:- Preparation of 2-chloro-6-iodobenzoyl isocyanate

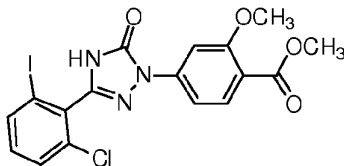
The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 2-chloro-6-iodobenzamide (1.0 g, 3.5 mmol), oxalyl chloride (0.538 g, 4.2 mmol) and EDC (20 mL) to afford 1.00 g of the desired product.

Intermediate-34

4-(3-(2-Chloro-6-iodophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxy-N-(3-(trifluoromethyl)phenyl)benzamide



Step-1:- Preparation of methyl 4-(3-(2-chloro-6-iodophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxybenzoate



The title compound was prepared according to the procedure described in step-1 and step-2 of Intermediate-9 by using *tert*-butyl 2-(3-methoxy-4-(methoxycarbonyl)phenyl)

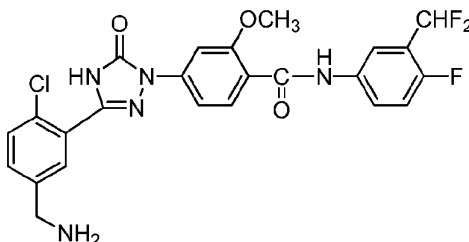
hydrazinecarboxylate (Intermediate-14, 1.0g) and 2-chloro-6-iodobenzoyl isocyanate (Intermediate-33, 1.0 g), TFA (5 mL) and DCM (40 mL) to afford 0.900 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.78 (s, 3H), 3.85 (s, 3H), 3.37 (t, J = 7.8 Hz, 1H), 7.63 (d, J = 9.0 Hz, 1H), 7.70- 7.81 (m, 3H), 8.00 (d, J = 7.8 Hz, 1H), 12.65 (br s, 1H); MS (m/z): 285.75 (M+H) $^+$.

Step-2:- Preparation of 4-(3-(2-chloro-6-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide

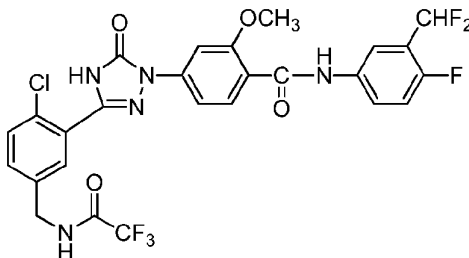
The title compound was prepared by following the procedure as described for step-6 of Intermediate-26 by using methyl 4-(3-(2-chloro-6-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (0.100 g, 0.206 mmol), 3-(trifluoromethyl)aniline (0.041 g, 0.247 mmol), trimethyl aluminium (2M solution in toluene) (0.5 mL), dry toluene (10.0 mL) to afford 0.075 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.95 (s, 3H), 7.19- 8.03 (m, 9H), 8.25 (s, 1H), 10.36 (s, 1H), 12.65 (s, 1H); MS (m/z): 614.98 (M+H) $^+$.

Intermediate-35

4-(3-(5-(Aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(difluoromethyl)-4-fluorophenyl)-2-methoxybenzamide



Step-1:- Preparation of 4-(3-(2-chloro-5-((2,2,2-trifluoroacetamido)methyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(difluoromethyl)-4-fluorophenyl)-2-methoxybenzamide



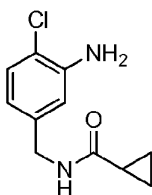
The title compound was prepared by following the procedure as described for step-6 of Intermediate-26 by using methyl 4-(3-(2-chloro-6-((2,2,2-trifluoroacetamido) methyl) phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Step-5 of Intermediate-26, 0.100 g, 0.20 mmol), 3-(difluoromethyl)-4-fluoroaniline (0.050 g, 0.30 mmol), trimethyl aluminium (2M solution in toluene) (1 mL) to afford 0.070 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.95 (s, 3H), 4.47 (s, 2H), 6.71 (m, 1H), 7.04 (m, 1H), 7.34- 7.40 (m, 1H), 7.50- 7.54 (m, 1H), 7.65- 7.70 (m, 3H), 7.79- 7.87 (m, 3H), 8.12 (s, 1H), 10.09 (m, 1H), 10.28 (s, 1H), 12.67 (brs, 1H); MS (*m/z*): 614.04 (M+H)⁺.

Step-2:- Preparation of 4-(3-(5-(aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(difluoromethyl)-4-fluorophenyl)-2-methoxybenzamide

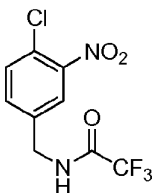
The solution of 4-(3-(2-chloro-5-((2,2,2-trifluoroacetamido)methyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(difluoromethyl)-4-fluorophenyl)-2-methoxybenzamide (0.070 g, 0.11 mmol) in 20% aq. KOH (10 mL) and aq. NH₃ (2 mL) was stirred for 24 h at RT. Excess of solvent was removed under vacuum and filtered off to afford 0.015 g of desired product.

Intermediate-36

N-(3-Amino-4-chlorobenzyl)cyclopropanecarboxamide



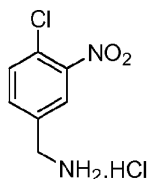
Step-1:- Preparation of *N*-(4-chloro-3-nitrobenzyl)-2,2,2-trifluoroacetamide



To a solution of 1-chloro-2-nitrobenzene (10.0 g, 0.063 mmol) in conc. sulphuric acid (150 mL) was added *N*-hydroxy methyl trifluoro acetamide (9.98 g, 0.069 mmol). The reaction mixture was heated at 70-80°C for 24 h. The reaction mass was quenched in ice cold water, neutralised with sodium hydroxide and extracted with DCM. The organic

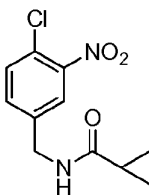
layer was concentrated and the obtained crude product was purified by column chromatography on silica gel eluting with 4% EtOAc: pet. ether to afford 5.5 g of desired product. ^1H NMR (300 MHz, DMSO d_6): 4.48 (d, $J = 3.9$ Hz, 2H), 7.62- 7.64 (m, 1H), 7.72- 7.78 (m, 1H), 7.97- 8.01 (m, 1H), 10.09 (brs, 1H) .

Step-2:- Preparation of (4-chloro-3-nitrophenyl)methanamine hydrochloride



To a solution of *N*-(4-chloro-3-nitrobenzyl)-2,2,2-trifluoroacetamide (0.800 g) in methanol (20 mL) was added conc. HCl (2.0 ml). The reaction mass was refluxed for 18 h. Excess solvent was removed under vacuum to afford 0.700 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.18 (s, 1H), 7.35 (s, 1H), 7.52 (s, 1H), 7.86 (s, 2H), 8.28 (s, 1H), 8.72 (br s, 2H) .

Step-3:- Preparation of *N*-(4-chloro-3-nitrobenzyl)cyclopropanecarboxamide



The title compound was prepared by following the procedure as described in step-2 of Intermediate-30 by using (4-chloro-3-nitrophenyl)methanamine hydrochloride (0.700 g, 3.14 mmol), cyclopropylcarbonyl chloride (0.490 g, 4.71 mmol), DIPEA (3.0 mL) and THF (15 mL) to afford 0.400 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.69 (d, $J = 6.0$ Hz, 2H), 1.55- 1.61 (m, 1H), 4.33 (d, $J = 6.0$ Hz), 7.55 (d, $J = 7.8$ Hz, 1H), 7.70 (d, $J = 8.4$ Hz, 1H), 7.87 (s, 1H), 8.68 (br s, 1H); MS (m/z): 255.09 ($M+H$) $^+$.

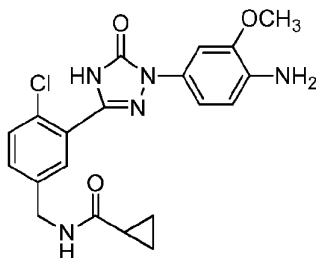
Step-4:- Preparation of *N*-(3-amino-4-chlorobenzyl)cyclopropanecarboxamide

The title compound was prepared by following the procedure as described in step-2 of Intermediate-28 by using *N*-(4-chloro-3-nitrobenzyl)cyclopropanecarboxamide (0.400 g), methanol (10 mL), iron powder (catalytic) and conc.HCl (5 mL). The reaction mass was refluxed for 2 h. The reaction mass was quenched in water and basified with NaHCO_3

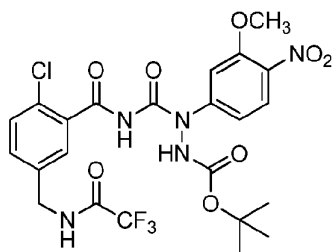
and extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 0.200 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.65 (d, J = 8.4 Hz, 4H), 1.57 (m, 1H), 4.11 (d, J = 5.4 Hz, 2H), 5.34 (s, 2H), 6.41 (d, J = 7.2 Hz, 1H), 7.10 (d, J = 7.8 Hz, 1H), 8.50 (br s, 1H); MS (m/z): 225.07 ($M+H$) $^+$.

Intermediate-37

N-(3-(1-(4-Amino-3-methoxyphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-4-chlorobenzyl)cyclopropanecarboxamide

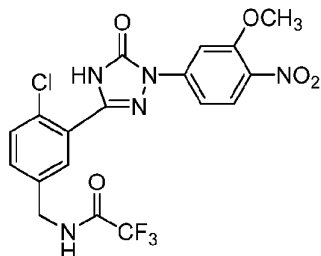


Step-1:- Preparation of *tert*-butyl 2-((2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl)carbamoyl)-2-(3-methoxy-4-nitrophenyl)hydrazinecarboxylate



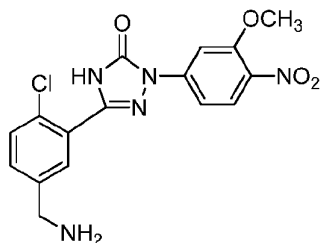
The title compound was prepared according to the procedure described in step-1 of Intermediate-9 by using 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate (step-3 of Intermediate-26, 1.400 g, 4.57 mmol), *tert*-butyl 2-(3-methoxy-4-nitrophenyl)hydrazinecarboxylate (step-3 of Intermediate-32, 1.30 g, 4.57 mmol) and DCM 30 mL) to afford 2.0 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.33- 1.46 (m, 9H), 3.87 (s, 3H), 4.40 (s, 2H), 7.06 (d, J = 8.4 Hz, 1H), 7.32- 7.36 (m, 3H), 7.45- 7.52 (m, 1H), 7.96 (d, J = 8.7 Hz, 1H), 9.92 (s, 1H), 10.07 (br s, 1H), 11.13 (br s, 1H); MS (m/z): 589.69 (M) $^+$.

Step-2:- Preparation of *N*-(4-chloro-3-(1-(3-methoxy-4-nitrophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



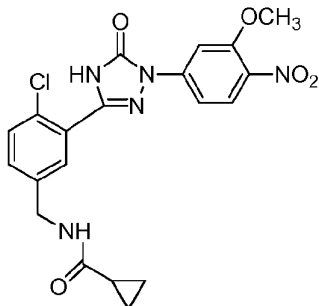
The title compound was prepared according to the procedure described in step-2 of Intermediate-9 by using *tert*-butyl 2-((2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl)carbamoyl)-2-(3-methoxy-4-nitrophenyl)hydrazinecarboxylate (2.0 g, 4.23 mmol), TFA (30 mL) and DCM (5 mL) to afford 1.300 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.96 (s, 3H), 4.47 (d, $J = 5.4$ Hz, 2H), 7.54 (d, $J = 8.7$ Hz, 1H), 7.67- 7.74 (m, 3H), 7.93 (s, 1H), 8.10 (d, $J = 9.0$ Hz, 1H), 10.11 (m, 1H), 12.86 (br s, 1H); MS (m/z): 471.88 (M+H) $^+$.

Step-3:- Preparation of 3-(5-(aminomethyl)-2-chlorophenyl)-1-(3-methoxy-4-nitrophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



The solution of *N*-(4-chloro-3-(1-(3-methoxy-4-nitrophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (0.500 g, 1.06 mmol) in 20% aq. KOH (20 mL) was stirred for 2-3 h at RT. Excess of solvent was removed under vacuum and filtered off remaining reaction mass to afford 0.400 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.80 (br s, 2H), 3.72 (s, 2H), 3.91 (s, 3H), 7.27- 7.39 (m, 2H), 7.78 (d, $J = 9.9$ Hz, 2H), 8.98 (d, $J = 8.7$ Hz, 1H), 8.25 (s, 1H); MS (m/z): 375.91 (M) $^+$.

Step-4:- Preparation of *N*-(4-chloro-3-(1-(3-methoxy-4-nitrophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)cyclopropanecarboxamide



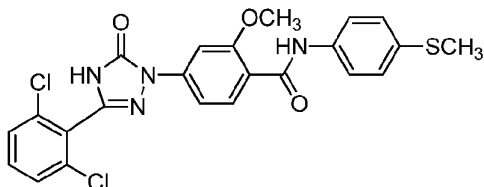
The title compound was prepared according to the procedure as described in step-2 Intermediate-30 by using 3-(5-(aminomethyl)-2-chlorophenyl)-1-(3-methoxy-4-nitrophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (0.500 g, 1.33 mmol), cyclopropylcarbonyl chloride (0.207 g, 1.99 mmol), DIPEA (2.0 mL) and THF (30 mL) to afford 0.300 g of crude product which was triturated with methanol: DEE to afford 0.300 g of pure product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.69 (s, 4H), 1.60 (m, 1H), 3.97 (s, 3H), 4.33 (d, *J* = 5.4 Hz, 2H), 7.42- 7.55 (m 3H), 7.70 (d, *J* = 8.4 Hz, 1H), 7.87 (s, 1H), 8.12 (d, *J* = 8.7 Hz, 1H), 8.69- 8.71 (m, 1H), 12.83 (s, 1H); MS (*m/z*): 442.14 (M-H)⁻.

Step-5:- Preparation of *N*-(3-(1-(4-amino-3-methoxyphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-4-chlorobenzyl)cyclopropanecarboxamide

The title compound was prepared according to the procedure as described in step-2 Intermediate-28 by using *N*-(4-chloro-3-(1-(3-methoxy-4-nitrophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)cyclopropanecarboxamide (0.500 g, 1.12 mmol), iron powder (catalytic amount), methanol (20 mL), conc. HCl (5.0 mL) to afford 0.350 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.69 (s, 4H), 1.22 (br s, 2H), 1.60 (m, 1H), 3.97 (s, 3H), 4.33 (d, *J* = 5.4 Hz, 2H), 7.45- 7.58 (m, 3H), 7.70 (d, *J* = 9.3 Hz, 1H), 7.87 (s, 1H), 8.13 (d, *J* = 9.0 Hz, 1H), 8.71 (m, 1H), 12.30 (br s, 1H); MS (*m/z*): 414.09 (M+H)⁺.

Intermediate-38

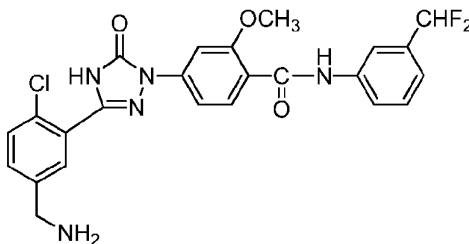
4-(3-(2,6-Dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(4-(methylthio)phenyl)benzamide



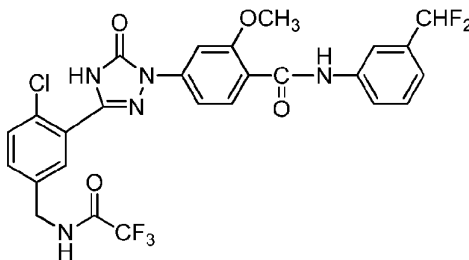
The title compound was prepared according to the procedure as described for step-6 of Intermediate-26 by using methyl 4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-21, 0.150 g, 0.380 mmol), 4-(methylthio)aniline (0.079 g, 0.570 mmol), trimethyl aluminium (2M solution in toluene) (1 mL) and dry toluene (5.0 mL) to afford 0.080 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 2.46 (s, 3H), 3.94 (s, 3H), 7.26 (d, J = 8.4 Hz, 2H), 7.69- 7.72 (m, 6H), 7.78 (d, J = 6.9 Hz, 1H), 7.82 (s, 1H), 10.06 (s, 1H), 12.76 (br s, 1H); MS (m/z): 501.0 (M) $^+$.

Intermediate-39

4-(3-(5-(Aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-N-(3-(difluoromethyl)phenyl)-2-methoxybenzamide



Step-1:- Preparation of 4-(3-(2-chloro-5-((2,2,2-trifluoroacetamido)methyl)phenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-N-(3-(difluoromethyl)phenyl)-2-methoxybenzamide



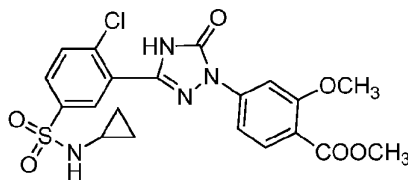
The title compound was prepared according to the procedure described in step-6 of Intermediate-26 by using methyl 4-(3-(2-chloro-5-((2,2,2-trifluoroacetamido)methyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-5 of Intermediate-26, 0.100g, 0.20 mmol), 3-(difluoromethyl)aniline (Intermediate-29, 0.045 g, 0.30 mmol), trimethyl aluminium (2M solution in toluene) (1.0 mL) and dry toluene (5.0 mL) to afford 0.080 g of crude product which was triturated with methanol: DCM: ether to afford 0.080 g of pure product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.95 (s, 3H), 4.47 (d, *J* = 5.7 Hz, 2H), 7.05 (m, 1H), 7.30 (m, 1H), 7.50 (d, *J* = 7.8 Hz, 2H), 7.70 (m, 3H), 7.80 (d, *J* = 6.3 Hz, 3H), 8.08 (s, 1H), 10.11 (br s, 1H), 10.28 (s, 1H), 12.72 (br s, 1H); MS (*m/z*): 596.12 (M+H)⁺.

Step-2:- Preparation of 4-(3-(5-(aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(difluoromethyl)phenyl)-2-methoxybenzamide

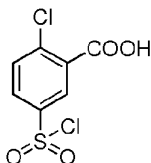
The solution of 4-(3-(2-chloro-5-((2,2,2-trifluoroacetamido)methyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(difluoromethyl)phenyl)-2-methoxybenzamide (0.070 g, 0.11 mmol) in 20% aq. KOH (10 mL) and aq. NH₃ (2 mL) was stirred for 24 h at RT. Excess of solvent was removed under vacuum and filtered off to afford 0.015 g of desired product.

Intermediate-40

Methyl 4-(3-(2-chloro-5-(*N*-cyclopropylsulfamoyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate



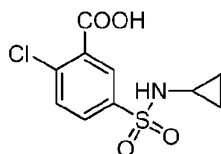
Step-1:- Preparation of 2-chloro-5-(chlorosulfonyl)benzoic acid



To a solution of 2-chlorobenzoic acid (5.0 g, 31.93 mmol) was slowly added chlorosulphonic acid (25 mL). The reaction mixture was heated at 100°C for 5-6 h,

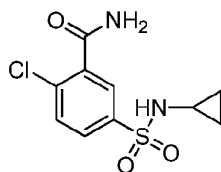
followed by stirring at RT for 24 h. The reaction mass was quenched in ice cold water slowly and the solid obtained was filtered off, washed with water and dried to afford 4.0 g of pure product. ^1H NMR (300 MHz, DMSO d_6): δ 7.50 (d, J = 7.8 Hz, 1H), 7.68 (d, J = 8.1 Hz, 1H), 7.96 (d, J = 1.5 Hz, 1H), 12.69-12.94 (br s, 1H).

Step-2:- Preparation of 2-chloro-5-(*N*-cyclopropylsulfamoyl)benzoic acid



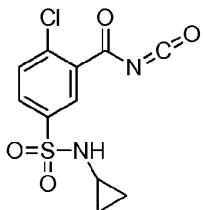
To a solution of 2-chloro-5-(chlorosulfonyl)benzoic acid (4.0 g, 15.68 mmol) in DCM (60 mL) was added cyclopropyl amine (1.8 g, 31.57 mmol) and the reaction mixture was stirred at RT for 16 h. The reaction mass was concentrated and the residue was diluted with ice cold water and acidified with dilute HCl. The crude solid was filtered off, washed with water and dried to afford 3.0 g of pure product. ^1H NMR (300 MHz, DMSO d_6): δ 0.37 (s, 2H), 0.49 (d, J = 5.4 Hz, 2H), 2.12-2.13 (m, 1H), 7.82 (d, J = 8.4 Hz, 1H), 7.91 (d, J = 8.4 Hz, 1H), 8.13 (s, 1H), 8.18 (s, 1H), 13.92 (br s, 1H).

Step-3:- Preparation of 2-chloro-5-(*N*-cyclopropylsulfamoyl)benzamide



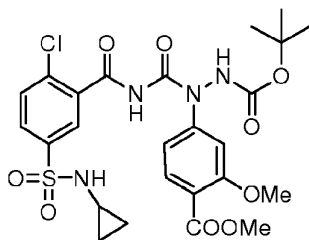
The title compound was prepared by following procedure as described for step-2 of Intermediate -26 by using 2-chloro-5-(*N*-cyclopropylsulfamoyl)benzoic acid (3.0 g, 10.0 mmol), DCM (30 mL), oxalyl chloride, ammonia gas (1.2 mL, 13.0 mmol), DMF (2-3 drop) and THF (40 mL) to afford 2.500 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.41 (m, 2H), 0.47-0.51 (m, 2H), 2.12 (m, 2H), 7.74- 7.82 (m, 4H), 8.11 (br s, 2H); MS (m/z): 273.00 (M-H) $^+$.

Step-4:- Preparation of 2-chloro-5-(*N*-cyclopropylsulfamoyl)benzoyl isocyanate



To a solution of 2-chloro-5-(*N*-cyclopropylsulfamoyl)benzamide (1.50 g, 5.46 mmol) in EDC (20 mL) was added oxalyl chloride (0.59 mL, 6.55 mmol). The reaction mass was refluxed for 1-2 h. Excess of solvent was removed under vacuum to afford 1.5 g of desired product.

Step-5:- Preparation of *tert*-butyl 2-((2-chloro-5-(*N*-cyclopropylsulfamoyl)benzoyl)carbamoyl)-2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate



To a solution of *tert*-butyl 2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate (Intermediate 14, 1.5 g, 5.08 mmol) in DCM (20 mL) was added 2-chloro-5-(*N*-cyclopropylsulfamoyl)benzoyl isocyanate (1.6 g, 5.32 mmol). The reaction mass was stirred at RT for 2 h. Excess of solvent was removed under vacuum to afford 0.500 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.50 (m, 4H), 1.41 (s, 9H), 2.11 (m, 1H), 3.68-3.75 (m, 6H), 7.59 (d, J = 8.7 Hz, 1H), 7.78- 7.87 (m, 4H), 8.10 (m, 2H), 8.27 (s, 1H), 8.95 (s, 1H).

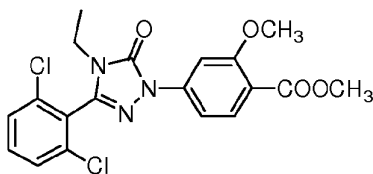
Step-6:- Preparation of methyl 4-(3-(2-chloro-5-(*N*-cyclopropylsulfamoyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate

The title compound was prepared according to the procedure described in step-2 of Intermediate-9 by using *tert*-butyl 2-((2-chloro-5-(*N*-cyclopropylsulfamoyl)benzoyl)carbamoyl)-2-(3-methoxy-4-(methoxycarbonyl)phenyl)hydrazinecarboxylate (0.500 g, 0.83 mmol), TFA (2 mL) and DCM (10 mL) to afford 0.100 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.40 (m, 2H), 0.52 (d, J = 5.4 Hz, 2H), 2.18

(m, 1H), 3.77 (s, 3H), 3.85 (s, 3H), 7.65 (d, $J = 9.0$ Hz, 1H), 7.82 (d, $J = 8.1$ Hz, 2H), 7.93 (s, 2H), 8.18 (s, 2H), 12.8 (br s, 1H); MS (m/z): 478.97 (M+H)⁺.

Intermediate-41

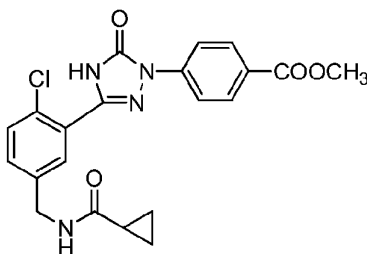
Methyl 4-(3-(2,6-dichlorophenyl)-4-ethyl-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxybenzoate



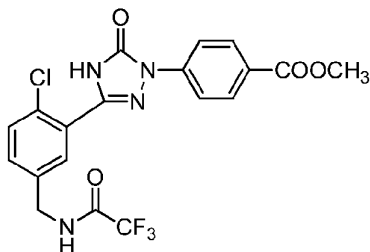
To a solution of methyl 4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-21, 0.100 g, 0.25 mmol) in dry DMF (3 mL) was added NaH (0.012 g, 0.30 mmol) at 0°C and the reaction mixture was stirred for 20-30 minutes. Ethyl bromide (0.028 g, 0.25 mmol) was added and the reaction mixture was further stirred at 60°C for 5-6 h. The reaction mass was quenched with water and extracted in ethyl acetate. The organic layer was dried and concentrated to afford 0.040 g of the crude product which was further purified by column chromatography in basic alumina eluting with 0.5-1.0 % methanol : DCM to afford 0.040 g of pure product.¹H NMR (300 MHz, DMSO d_6): δ 1.13 (t, 3H), 3.54-3.56 (m, 2H), 3.78 (s, 3H), 3.86 (s, 3H), 7.65 (d, $J = 8.4$ Hz, 1H), 7.77 (m, 4H), 7.84 (d, $J = 8.4$ Hz, 1H); MS (m/z): 421.95 (M+H)⁺.

Intermediate-42

Methyl 4-(3-(2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)benzoate

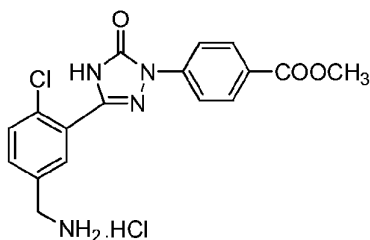


Step-1:- Preparation of methyl 4-(3-(2-chloro-5-((2,2,2-trifluoroacetamido)methyl)phenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)benzoate



The title compound was prepared according to the procedure described in step-1 and step-2 of Intermediate-9 by using 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate (step-3 of Intermediate-26, 1.300 g, 4.2 mmol) and *tert*-butyl 2-[4-(methoxycarbonyl)phenyl]hydrazinecarboxylate (Intermediate-7, 1.150g, 3.8 mmol) in TFA (5-7 mL) and DCM (40 mL). The reaction mass was quenched with water and extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 1.300 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.84 (s, 3H), 4.44 (d, J = 5.7 Hz, 2H), 7.49 (t, 1H), 7.66 (d, J = 8.7 Hz, 2H), 8.04-8.13 (m, 4H), 10.07 (m, 1H), 12.69 (s, 1H); MS (m/z): 453.14 (M-H) $^-$.

Step-2:- Preparation of methyl 4-(3-(5-(aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)benzoate hydrochloride



The title compound was prepared according to the procedure described in step-1 of Intermediate-30 by using methyl 4-(3-(2-chloro-5-((2,2,2-trifluoroacetamido)methyl)phenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)benzoate (1.00 g, 2.19 mmol), methanol (20 mL) and HCl (10 mL) to afford 1.00 g of desired product.

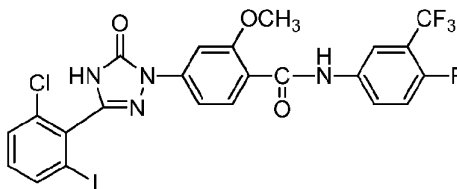
Step-3:- Preparation of methyl 4-(3-(2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)benzoate

The title compound was prepared by following the procedure as described for step- 2 of Intermediate-30 by using methyl 4-(3-(5-(aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)benzoate hydrochloride (0.250 g, 0.5 mmol), cyclopropane

carbonyl chloride (0.108 g, 1.0 mmol), DIPEA (2.0 mL), THF (10 mL) to afford 0.200 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.05 (s, 2H), 1.60 (m, 2H), 3.14 (m, 1H), 3.87 (s, 3H), 4.33 (d, $J = 5.7$ Hz, 2H), 7.41- 7.61 (m, 4H), 8.11-8.17 (m, 3H), 8.69 (m, 1H), 12.65 (br s, 1H); MS (m/z): 427.14 (M+H) $^+$.

Intermediate-43

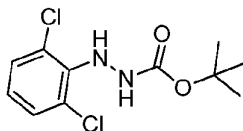
4-(3-(2-Chloro-6-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(4-fluoro-3-(trifluoromethyl)phenyl)-2-methoxybenzamide



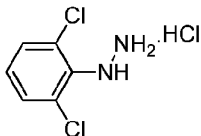
The title compound was prepared by following the procedure as described for step- 6 of Intermediate-26 by using methyl 4-(3-(2-chloro-6-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-1 of Intermediate-34, 0.100 g, 0.20 mmol), 4-fluoro-3-(trifluoromethyl)aniline (0.044 g, 0.34 mmol), trimethyl aluminium (2M solution in toluene) (1.0 mL) and dry toluene (8.0 mL) to afford 0.020 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.94 (s, 3H), 3.78 (t, 1H), 7.52 (t, 2H), 7.71-7.82 (m, 3H), 8.03 (d, $J = 6.6$ Hz, 2H), 10.38 (s, 1H), 12.69 (br s, 1H).

Intermediate-44

tert-Butyl 2-(2,6-dichlorophenyl)hydrazinecarboxylate



Step-1:- Preparation of (2,6-dichlorophenyl)hydrazine hydrochloride



To a cold solution of 2,6-dichloroaniline (10.0 g, 61.72 mmol) in conc. HCl was added aqueous solution of sodium nitrite (5.11 g, 74.00 mmol) at -15°C. The reaction mass was

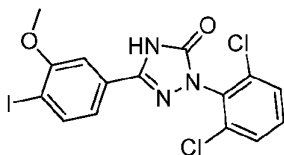
stirred at 0-10°C for 15 minutes. The reaction mass was filtered off to remove insolubles and stannous chloride in conc. HCl (34.7 g, 150 mmol) was added to the filtrate. The reaction mass was further stirred at -15°C for 30 min and filtered to afford 15.00 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.18- 7.28 (m, 1H), 7.39- 7.52 (m, 2H), 10.30 (br s, 3H); MS (m/z): 176.90 (M) $^+$.

Step-2:- Preparation of *tert*-butyl 2-(2,6-dichlorophenyl)hydrazinecarboxylate

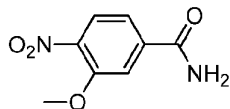
To the cold solution of (2,6-dichlorophenyl)hydrazine hydrochloride (15.0 g, 84.0 mmol) in THF (50 mL) was added aqueous solution of K_2CO_3 (23.3 g, 169 mmol) and BOC anhydride solution in THF (20.3 g, 93.00 mmol) at 0°C. The reaction mass was stirred at RT for 12 h. The reaction mass was quenched in water and extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated. The obtained solid was washed with pentane to afford 6.5 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.34 (s, 9H), 6.85 (m, 1H), 7.06 (s, 1H), 7.27 (d, J = 7.8 Hz, 2H), 9.01 (s, 1H); MS (m/z): 276.88 (M-H) $^-$.

Intermediate-45

1-(2,6-Dichlorophenyl)-3-(4-iodo-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



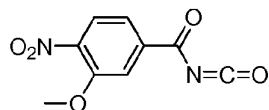
Step-1:- Preparation of 3-methoxy-4-nitrobenzamide



To a cold solution of 3-methoxy-4-nitrobenzoic acid (2.0 g, 10.10 mmol) in THF (25 mL) was added oxalyl chloride (1.1 mL, 12.1 mmol) and DMF (2-3 drop) at 0°C. The reaction mixture was stirred at RT for 2 h and concentrated. The solution of the concentrated mass in THF (15 mL) was treated with ammonia gas (purged through reaction mass) at 0°C and the reaction mixture was stirred at RT for 1 h. The reaction mass was diluted with ethyl acetate, washed with water, dilute HCl and brine. The organic layer was separated, dried, filtered and concentrated. The concentrate was used for the next step without

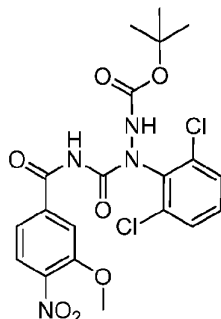
further purification. ^1H NMR (300 MHz, DMSO d_6): δ 3.97 (s, 3H), 7.56 (d, J = 8.1 Hz, 1H), 7.72- 7.75 (m, 2H), 7.94 (d, J = 8.4 Hz, 1H), 8.25 (s, 1H); MS (m/z): 195.98 (M) $^+$.

Step-2:- Preparation of 3-methoxy-4-nitrobenzoyl isocyanate



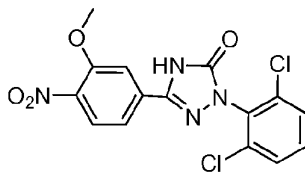
To a solution of 3-methoxy-4-nitrobenzamide (1.00 g, 5.10 mmol) in EDC (10 mL) was added oxalyl chloride (0.64 g, 6.12 mmol). The reaction mass was refluxed for 24 h. Excess of solvent was removed under vacuum to afford 1.0 g of desired product.

Step-3:- Preparation of *tert*-butyl 2-(2,6-dichlorophenyl)-2-((3-methoxy-4-nitrobenzoyl)carbamoyl)hydrazinecarboxylate



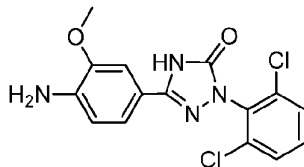
To a solution of *tert*-butyl 2-(2,6-dichlorophenyl) hydrazinecarboxylate (Intermediate-44, 1.2 g, 4.5 mmol) in DCM (20 mL) was added 3-methoxy-4-nitrobenzoyl isocyanate (1.0 g, 4.5 mmol). The reaction mass was stirred at RT for 2 h. Excess of solvent was removed under vacuum to afford 1.5 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.35 (s, 9H), 3.97 (s, 3H), 7.89 (m, 1H), 7.27- 7.35 (m, 2H), 7.55 (d, 1H), 7.62- 7.75 (m, 1H), 7.94 (d, 1H), 8.26 (s, 1H), 9.03 (s, 1H); MS (m/z): 496.78 (M-H) $^+$.

Step-4:- Preparation of 1-(2,6-dichlorophenyl)-3-(3-methoxy-4-nitrophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



To a solution of *tert*-butyl 2-(2,6-dichlorophenyl)-2-((3-methoxy-4-nitrobenzoyl)carbamoyl)hydrazinecarboxylate (1.0 g, 2.002 mmol) in DCM (20 mL) was added trifluoro acetic acid (2.0 mL). The reaction mass was stirred at RT for 2-3 h. Excess of solvent was removed at low temperature. The reaction mass was quenched in ice and filter off to afford 0.200 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.98 (s, 3H), 7.56- 7.62 (m, 2H), 7.69- 7.75 (m, 3H), 8.02 (d, J = 8.4 Hz, 1H), 11-12 (br s, 1H); MS (m/z): 382.07 ($M+H$) $^+$.

Step-5:- Preparation of 3-(4-amino-3-methoxyphenyl)-1-(2,6-dichlorophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



To a solution of 1-(2,6-dichlorophenyl)-3-(3-methoxy-4-nitrophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (0.200 g, 0.52 mmol) in methanol (10 mL) was added iron powder (0.060 g, 1.84 mmol) and conc. HCl (2 mL). The reaction mixture was stirred at RT for 2 h. The reaction mass was quenched in water, basified with saturated sodium bicarbonate solution and extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 0.150 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.80 (s, 3H), 5.34 (s, 2H), 6.66 (d, J = 7.8 Hz, 1H), 7.18- 7.21 (m, 2H), 7.58 (m, 1H), 7.69 (d, J = 7.8 Hz, 2H), 12.27 (s, 1H); MS (m/z): 351.05 (M) $^+$.

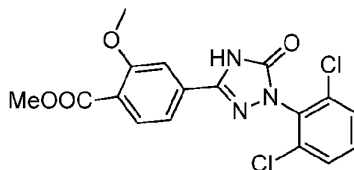
Step-6:- Preparation of 1-(2,6-dichlorophenyl)-3-(4-iodo-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one

To a solution of 3-(4-amino-3-methoxyphenyl)-1-(2,6-dichlorophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (0.300 g, 0.85 mmol) in acetonitrile (20 mL) was added PTSA (0.488 g, 2.56 mmol). The reaction mixture was stirred at RT for 2 h, followed by addition of aqueous solution of NaNO₂ (0.118 g, 1.170 mmol) and KI (0.284 g, 1.70 mmol). The reaction mixture was stirred at RT for 2 h. The reaction mass was concentrated and quenched in water. The reaction mass was washed with sodium metabisulphite solution and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 0.220 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.89

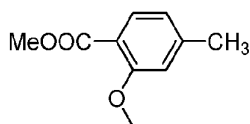
(s, 3H), 7.23 (d, $J = 7.8$ Hz, 1H), 7.40 (s, 1H), 7.61 (m, 1H), 7.71- 7.73 (m, 2H), 7.92 (d, $J = 8.4$ Hz, 1H), 12.74 (s, 1H); MS (m/z): 461.93 (M)⁺.

Intermediate-46

Methyl 4-(1-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-3-yl)-2-methoxybenzoate

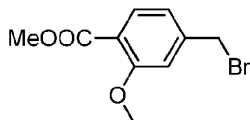


Step-1:- Preparation of methyl 2-methoxy-4-methylbenzoate



To a solution of 2-hydroxy-4-methylbenzoic acid (10.00 g, 65 mmol) in DMF (10 mL) was added K_2CO_3 (13.6 g, 98 mmol). The reaction mixture was stirred at 60°C for 1 h. Methyl iodide (14.0 g, 98 mmol) was added to the reaction mixture and was stirred at 60°C for 18 h. The reaction mass was quenched in water and concentrated. The reaction mass was extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 10.00 g of desired product. ¹H NMR (300 MHz, DMSO d_6): δ 2.34 (s, 3H), 3.74 (s, 3H), 3.79 (s, 3H), 6.81 (d, $J = 7.2$ Hz, 1H), 6.97 (s, 1H), 7.56 (d, $J = 7.8$ Hz, 1H); MS (m/z): 180.95 (M+H)⁺.

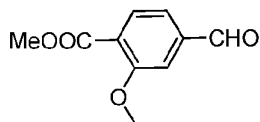
Step-2:- Preparation of methyl 4-(bromomethyl)-2-methoxybenzoate



To a solution of methyl 2-methoxy-4-methylbenzoate (10.00 g, 55 mmol) in CCl_4 (150 mL) were added NBS (11.8 g, 66 mmol) and AIBN (0.100 g). The reaction mixture was refluxed for 18 h. The reaction mass was diluted with diethyl ether and washed with 25% NaOH solution. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 10.00 g of desired product. ¹H NMR (300 MHz, DMSO d_6): δ 3.77

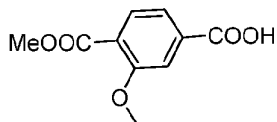
(m, 6H), 4.68 (s, 2H), 7.06 (d, $J = 7.8$ Hz, 1H), 7.23 (s, 1H), 7.61 (d, $J = 7.8$ Hz, 1H); MS (m/z): 259.13 (M)⁺.

Step-3:- Preparation of methyl 4-formyl-2-methoxybenzoate



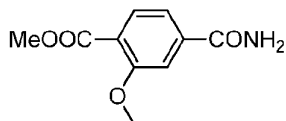
To a solution of sodium methoxide (0.260 g, 4.82 mmol) in methanol (200 mL) was added nitromethane (4.03 g, 11.58 mmol) under nitrogen atmosphere and the reaction mixture was stirred and reflux for 30 minutes. Methyl 4-(bromomethyl)-2-methoxybenzoate (10.00 g, 9.65 mmol) in methanol was added and the reaction mixture was refluxed for 7-8 h. The reaction mass was concentrated and diluted with CHCl₃. The diluted reaction mass was washed with 2N NaOH solution. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 6.00 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.83 (m, 6H), 7.36 (d, $J = 6.6$ Hz, 2H), 7.79 (d, $J = 7.8$ Hz, 1H), 10.04 (s, 1H).

Step-4:- Preparation of 3-methoxy-4-(methoxycarbonyl)benzoic acid



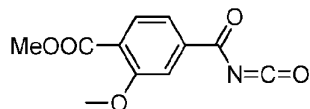
To a cold solution of methyl 4-formyl-2-methoxybenzoate (6.0 g, 7.73 mmol) in acetone (50 mL) was added aqueous sulphamic acid solution (4.5 g, 11.59 mmol) and aqueous sodium chlorite solution (4.04 g, 11.59 mmol) at 0-5°C. The reaction mixture was stirred at RT for 3-4 h. Excess of solvent was removed under vacuum. The obtained residue was diluted with water and basified with saturated sodium bicarbonate solution. The aqueous layer was washed with diethyl ether to remove organic impurities and the obtained aqueous layer was again acidified with conc.HCl. The solid obtained was filtered and suck dried to afford 2.00 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.80 (s, 3H), 3.87 (s, 3H), 7.55- 7.59 (m, 2H), 7.70 (d, $J = 7.8$ Hz, 1H), 13.37 (s, 1H); MS (m/z): 210.97 (M+H)⁺.

Step-5:- Preparation of methyl 4-carbamoyl-2-methoxybenzoate



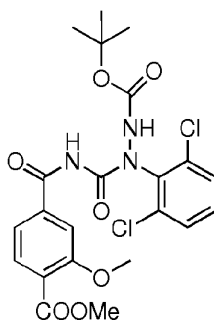
The title compound was prepared according to the procedure described in step-2 of Intermediate-26 by using 3-methoxy-4-(methoxycarbonyl)benzoic acid (2.00 g, 9.52 mmol), oxalyl chloride (1.02 g, 11.42 mmol), ammonia gas and THF (25 mL) to afford 1.40 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.79 (s, 3H), 3.87 (s, 3H), 7.49 (d, J = 8.4 Hz, 1H), 7.57 (s, 2H), 7.68 (d, J = 7.8 Hz, 1H), 8.14 (s, 1H); MS (m/z): 209.99 (M) $^+$.

Step-6:- Preparation of methyl 4-(isocyanatocarbonyl)-2-methoxybenzoate



The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using methyl 4-carbamoyl-2-methoxybenzoate (1.4 g, 6.69 mmol), oxalyl chloride (0.72 mL, 8.03 mmol) and EDC (10 mL) to afford 1.00 g of the desired product.

Step-7:- Preparation of *tert*-butyl 2-(2,6-dichlorophenyl)-2-((3-methoxy-4-(methoxycarbonyl)benzoyl)carbamoyl)hydrazinecarboxylate



The title compound was prepared according to the procedure described in step-1 of Intermediate-9 by using methyl 4-(isocyanatocarbonyl)-2-methoxybenzoate (0.84 g, 3.61 mmol), *tert*-butyl 2-(2,6-dichlorophenyl)hydrazinecarboxylate (Intermediate-44, 1.0 g, 3.61 mmol) and DCM (20 mL) to afford 1.50 g of the desired product. ^1H NMR (300

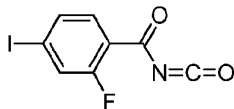
MHz, DMSO d_6): δ 1.35 (s, 9H), 3.80 (s, 3H), 3.87 (s, 3H), 7.28- 7.42 (m, 6H), 9.57 (s, 1H), 10.85 (s, 1H); MS (m/z): 511.73 (M)⁺.

Step-8:- Preparation of methyl 4-(1-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-methoxybenzoate

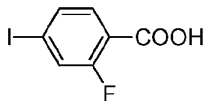
The title compound was prepared according to the procedure described in step-2 of Intermediate-9 by using *tert*-butyl 2-(2,6-dichlorophenyl)-2-((3-methoxy-4-(methoxycarbonyl) benzoyl)carbamoyl)hydrazinecarboxylate (1.5 g, 2.9 mmol), DCM (20 mL) and trifluoro acetic acid (2.0 mL) to afford 0.600 g of desired product. ¹H NMR (300 MHz, DMSO d_6): δ 3.80 (s, 3H), 3.80 (s, 3H), 7.49- 7.71 (m, 3H), 7.74- 7.79 (m, 3H), 12.85 (s, 1H); MS (m/z): 393.95 (M)⁺.

Intermediate-47

2-Fluoro-4-iodobenzoyl isocyanate

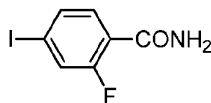


Step-1:- Preparation of 2-fluoro-4-iodobenzoic acid



A solution of sodium dichromate (25.2 g, 84.0 mmol) in acetic acid (100 mL) was stirred for 10 min, followed by addition of 2-fluoro-4-iodo-1-methylbenzene (10.0 g, 42 mmol). The reaction mixture was stirred for 5 min followed by slow addition of conc.H₂SO₄ (60 mL) for 1-2 h. The reaction mixture stirred at 90-100°C for 7-8 h. The reaction mixture was cooled and quenched with ice water. The reaction mass was extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 5.0 g of the desired product. ¹H NMR (300 MHz, DMSO d_6): δ 7.58 (t, J = 8.4 Hz, 1H), 7.67 (d, J = 7.5 Hz, 1H), 7.75 (d, J = 8.4 Hz, 1H), 13 (br s, 1H); MS (m/z): 264.99 (M-H)⁻.

Step-2:- Preparation of 2-fluoro-4-iodobenzamide



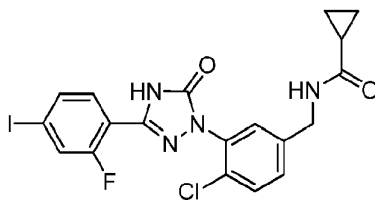
The title compound was prepared according to the procedure described in step-2 of Intermediate-26 by using 2-fluoro-4-iodobenzoic acid (5.00 g, 18.0 mmol), oxalyl chloride (2.0 mL, 22.0 mmol), ammonia gas, DCM (100 mL) and THF (100 mL) to afford 2.00 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.40 (t, J = 7.8 Hz, 1H), 7.63- 7.76 (m, 1H); MS (m/z): 266.19 (M+H) $^+$.

Step-3:- Preparation of 2-fluoro-4-iodobenzoyl isocyanate

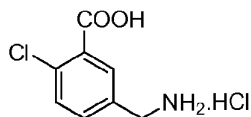
The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 2-fluoro-4-iodobenzamide (0.500 g, 1.8 mmol), oxalyl chloride (0.285 g, 2.2 mmol) and EDC (10 mL) to afford 0.350 g of the desired product.

Intermediate-48

N-(4-Chloro-3-(3-(2-fluoro-4-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide

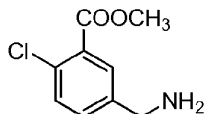


Step-1:- Preparation of 5-(aminomethyl)-2-chlorobenzoic acid hydrochloride



To a solution of 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoic acid (step-1 of Intermediate-26, 10.0 g, 35.50 mmol) in methanol (100 mL) was added conc. HCl (10 mL). The reaction mixture was refluxed for 18 h. The reaction mixture was concentrated and crude reaction mixture was used for next step.

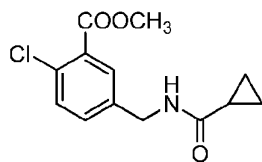
Step-2:- Preparation of methyl 5-(aminomethyl)-2-chlorobenzoate



To a solution of 5-(aminomethyl)-2-chlorobenzoic acid hydrochloride (17.0 g, 76.5 mmol) in methanol (200 mL) was added conc. H_2SO_4 (15 mL). The reaction mixture was refluxed for 15 h. The reaction mixture was concentrated and water was added. The aq.

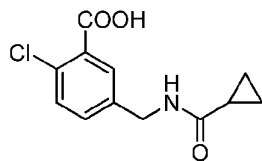
solution was basified with sodium bicarbonate. The reaction mixture was extracted with 10% MeOH: DCM and concentrated to afford 15.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 2.1 (br s, 2H), 3.73 (s, 2H), 3.85 (s, 3H), 7.50 (s, 2H), 7.77 (s, 1H); MS (m/z): 200.05 (M+H) $^+$.

Step-3:- Preparation of methyl 2-chloro-5-(cyclopropanecarboxamidomethyl)benzoate



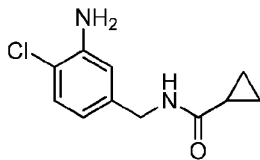
To a solution of methyl 5-(aminomethyl)-2-chlorobenzoate (15.0 g, 75 mmol) in dry THF (150 mL) was added DIPEA (15 mL) under nitrogen atmosphere. Cyclopropanecarbonyl chloride was added to the reaction mixture at 10°C and (11.81 g, 113 mmol). The reaction mixture was stirred at RT for 5-6 h. The reaction mixture was quenched with ice water and extracted with ethyl acetate. The organic layer was washed with dilute HCl, followed with sodium bicarbonate solution and concentrated to afford 10.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.69 (m, 4H), 1.59 (m, 1H), 3.86 (s, 3H), 4.29 (d, J = 6.0 Hz, 2H), 7.43 (d, J = 6.6 Hz, 1H), 7.53 (d, J = 8.4 Hz, 1H), 7.67 (s, 1H), 8.64 (br s, 1H); MS (m/z): 268.16 (M+H) $^+$.

Step-4:- Preparation of 2-chloro-5-(cyclopropanecarboxamidomethyl)benzoic acid



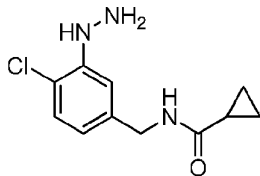
To a solution of methyl 2-chloro-5-(cyclopropanecarboxamidomethyl)benzoate (10.0 g, 30 mmol) in THF: methanol (10:20 mL) was added aqueous solution of NaOH (6.0 g, 140 mmol). The reaction mixture was refluxed for 3-5 h. The reaction mixture was concentrated, diluted with water and acidified with dilute HCl. The reaction mixture was extracted with 10% MeOH: DCM and concentrated to afford 4.0 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.67 (m, 4H), 1.59 (m, 1H), 4.28 (d, J = 6.0 Hz, 2H), 7.37- 7.50 (m, 2H), 7.65 (s, 1H), 8.64 (m, 1H), 13.40 (br s, 1H).

Step-5:- Preparation of *N*-(3-amino-4-chlorobenzyl)cyclopropanecarboxamide



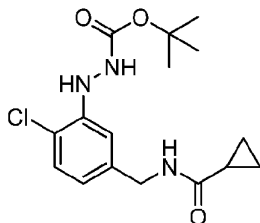
To a solution of 2-chloro-5-(cyclopropanecarboxamidomethyl)benzoic acid (5.0 g, 19 mmol) in conc. H_2SO_4 (30 mL) was added NaN_3 (1.4 g, 21 mmol) lotwise at 50°C . The reaction mixture was stirred at 50°C for 18 h. The reaction mass was quenched with ammonia and ice water. The solid obtained was filtered and suck dried to afford 3.0 g of desired product. ^1H NMR (300 MHz, $\text{DMSO}-d_6$): δ 0.66 (m, 4H), 1.56 (m, 1H), 4.12 (d, $J = 6.0$ Hz, 2H), 5.27 (s, 2H), 6.41 (d, $J = 6.9$ Hz, 1H), 6.61 (s, 1H), 7.10 (d, $J = 8.4$ Hz, 1H), 8.48 (m, 1H); MS (m/z): 225.22 ($\text{M}+\text{H}^+$).

Step-6:- Preparation of *N*-(4-chloro-3-hydrazinylbenzyl)cyclopropanecarboxamide



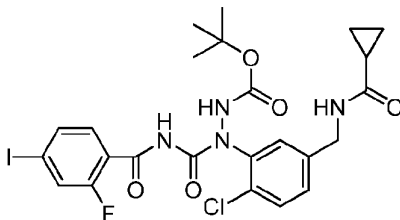
To a cold solution of *N*-(3-amino-4-chlorobenzyl)cyclopropanecarboxamide (2.00 g, 8.90 mmol) in conc. HCl (25 mL) was added aqueous solution of sodium nitrite (0.676 g, 9.7 mmol) at -20°C to -25°C . The reaction mass was stirred at -20°C to -25°C for 30 min, followed by addition of stannous chloride solution (5.0 g, 22.2 mmol) in conc. HCl slowly. The reaction mass was stirred at -20°C to -25°C for 1 h. The reaction mass was basified with aqueous NaOH at below 0°C . The reaction mass was diluted with water, extracted with 10% MeOH : DCM and concentrated to afford 1.6 g of desired product. ^1H NMR (300 MHz, $\text{DMSO}-d_6$): δ 0.66 (m, 4H), 1.58 (m, 1H), 4.10 (s, 2H), 4.18 (d, $J = 5.4$ Hz, 2H), 6.47 (m, 2H), 7.11 (m, 2H), 8.51 (br s, 1H); MS (m/z): 240.05 ($\text{M}+\text{H}^+$).

Step-7:- Preparation of *tert*-butyl 2-(2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl)hydrazinecarboxylate



To a cold solution of *N*-(4-chloro-3-hydrazinylbenzyl)cyclopropanecarboxamide (1.60 g, 6.69 mmol) in acetonitrile (20 mL) was added aqueous solution of Na_2CO_3 (1.06 g, 10.04 mmol) and BOC anhydride (1.06 g, 7.36 mmol). The reaction mass was stirred at RT for 12 h. The reaction mass was quenched in water and extracted with ethyl acetate and concentrated to afford 1.00 g of the crude product which was further purified by column chromatography using basic alumina eluting with 0.3% MeOH: DCM to afford 0.900 g of pure product. ^1H NMR (300 MHz, $\text{DMSO}-d_6$): δ 0.63 (m, 4H), 1.40 (s, 9H), 1.54 (m, 1H), 4.15 (d, $J = 6.0$ Hz, 2H), 6.59 (m, 2H), 7.17 (d, $J = 7.8$ Hz, 1H), 7.31 (s, 1H), 8.51 (m, 1H), 8.87 (s, 1H); MS (m/z): 338.41 ($\text{M}-\text{H}$) $^-$.

Step-8:- Preparation of *tert*-butyl 2-(2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-2-((2-fluoro-4-iodobenzoyl)carbamoyl)hydrazinecarboxylate



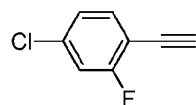
The title compound was prepared according to the procedure described in step-1 of Intermediate-9 by using 2-fluoro-4-iodobenzoyl isocyanate (Intermediate-47, 0.500 g, 1.71 mmol), *tert*-butyl 2-(2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl)hydrazine carboxylate (0.582 g, 1.71 mmol) and DCM (20 mL) to afford 0.200 g of the desired product. ^1H NMR (300 MHz, $\text{DMSO}-d_6$): δ 0.69 (m, 4H), 1.42 (s, 9H), 1.56 (m, 1H), 4.17 (d, $J = 5.7$ Hz, 2H), 6.58- 6.63 (m, 2H), 7.18 (d, $J = 8.1$ Hz, 1H), 7.30 (m, 3H), 7.63- 7.79 (m, 2H), 8.53 (m, 1H), 8.90 (s, 1H); MS (m/z): 630.85 ($\text{M}+\text{H}$) $^+$.

Step-9:- Preparation of *N*-(4-chloro-3-(3-(2-fluoro-4-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide

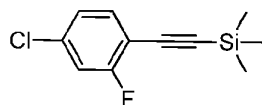
To a solution of *tert*-butyl 2-(2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-2-((2-fluoro-4-iodobenzoyl)carbamoyl)hydrazinecarboxylate (0.200 g, 0.31 mmol) in DCM (10 mL) was added trifluoro acetic acid (1.0 mL). The reaction mass was stirred at RT for 12 h and refluxed for 15 h. The reaction mass was cooled to RT, basified with NaHCO₃ (pH =8), extracted with DCM and concentrated to afford 0.110 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆) : δ 0.68 (m, 4H), 1.59 (m, 1H), 4.32 (d, *J* = 5.7 Hz, 2H), 7.38 (d, *J* = 8.4 Hz, 1H), 7.45 (s, 1H), 7.51- 7.63 (m, 2H), 7.75 (d, *J* = 8.1 Hz, 1H), 7.88 (d, *J* = 10.2 Hz, 1H), 8.68 (m, 1H), 12.43 (br s, 1H); MS (*m/z*): 513.01 (M+H)⁺.

Intermediate-49

4-Chloro-1-ethynyl-2-fluorobenzene



Step-1:- Preparation of ((4-chloro-2-fluorophenyl)ethynyl)trimethylsilane



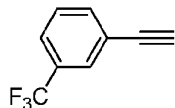
To a solution of 4-chloro-2-fluoro-1-iodobenzene (2.0 g, 7.8 mmol) in DMSO (10 mL) was added ethynyl(trimethyl)silane (1.14 g, 11.0 mmol), copper iodide (0.029 g, 0.15 mmol), bis(triphenylphosphine) palladium(II) chloride (0.218 g, 0.3 mmol) and TEA (1 mL). The reaction mass was stirred at RT for 4-5 h. The reaction mass quenched in water, passed through celite bed and extracted with ethyl acetate and concentrated to afford 1.0 g of desired product.

Step-2:- Preparation of 4-chloro-1-ethynyl-2-fluorobenzene

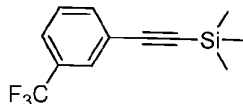
To a solution of ((4-chloro-2-fluorophenyl)ethynyl)trimethylsilane (1.0 g, 4.42 mmol) in DCM (20 mL) was added TBAF (catalytic) and reaction mixture was stirred for 1-2 h at RT. The reaction mass was quenched in water, extracted with DCM and concentrated to afford 0.900 g of the crude product which was further purified by column chromatography eluting with pet. ether to afford to afford 0.500 g of pure product. ¹H NMR (300 MHz, DMSO *d*₆): δ 4.60 (s, 1H), 7.32 (d, *J* = 8.1 Hz, 1H), 7.56- 7.61 (m, 2H).

Intermediate-50

1-Ethynyl-3-(trifluoromethyl)benzene



Step-1:- Preparation of trimethyl((3-(trifluoromethyl)phenyl)ethynyl)silane



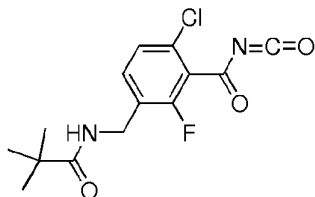
To a solution of 1-iodo-3-(trifluoromethyl)benzene (1.0 g, 3.6 mmol) in ethyl acetate (20 mL) was added ethynyl(trimethyl)silane (0.540 g, 5.5 mmol), copper iodide (0.042 g, 0.22 mmol), bis(triphenylphosphine) palladium(II) chloride (0.129 g, 0.18 mmol) and TEA (2.2 mL, 14.7 mmol). The reaction mass stirred at 50-60°C for 24 h. The reaction mass was quenched in water, extracted with ethyl acetate and concentrated to afford 0.500 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.26 (s, 9H), 7.44 (t, *J* = 7.8 Hz, 1H), 7.55 (d, *J* = 7.8 Hz, 1H), 7.62 (d, *J* = 7.2 Hz, 1H), 7.72 (s, 1H).

Step-2:- Preparation of 1-ethynyl-3-(trifluoromethyl)benzene

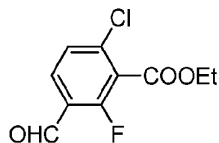
To a solution of trimethyl((3-(trifluoromethyl)phenyl)ethynyl)silane (0.500 g, 2.06 mmol) in ethanol (10 mL) was added potassium carbonate (0.253g, 1.8 mmol) and the reaction mixture was stirred at RT for 18 h. The reaction mass was quenched in water, neutralized with dil. HCl and extracted with hexane and concentrated to afford 0.200 g of the crude product which was further purified by column chromatography using silica (60-120 mesh) eluting with pet. ether to afford 0.500 g of pure product. ¹H NMR (300 MHz, DMSO *d*₆): δ 4.41 (s, 1H), 7.64 (d, *J* = 7.5 Hz, 1H), 7.77- 7.81 (m, 3H).

Intermediate-51

6-Chloro-2-fluoro-3-(pivalamidomethyl)benzoyl isocyanate

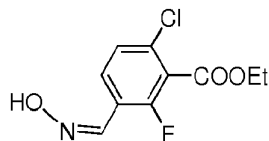


Step-1:- Preparation of ethyl 6-chloro-2-fluoro-3-formylbenzoate



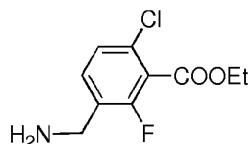
To a cold solution of ethyl 2-chloro-6-fluorobenzoate (0.500 g, 24.7 mmol) in THF (35 mL) was added LDA (6.6g, 61.8 mmol) at -78°C . The reaction mixture was stirred for 2 h at same temperature, followed by addition of DMF (2.7 g, 37 mmol). The reaction mixture was further stirred for 2 h at same temperature. The reaction mass was quenched in dilute HCl, extracted with ethyl acetate and concentrated to afford crude product which was further purified by column chromatography eluting with EtOAc: pet.ether to afford 2.00 g of pure product. ^1H NMR (300 MHz, DMSO d_6): δ 1.31 (t, 3H), 4.42 (q, 2H), 7.94- 7.99 (m, 2H), 10.14 (s, 1H).

Step-2:- Preparation of ethyl 6-chloro-2-fluoro-3-((hydroxyimino)methyl)benzoate



To a solution of ethyl 6-chloro-2-fluoro-3-formylbenzoate (1.1 g, 4.0 mmol) in MeOH (5 mL) was added (aq. solution) hydroxyl amine (4 mL) and the reaction mixture was heated at 60°C for 2-3 h. The reaction mass was concentrated and extracted with EtOAc. The organic layer was washed with brine and concentrated to afford 0.900 g of the product. ^1H NMR (300 MHz, DMSO d_6): δ 1.31 (t, $J = 6.9$ Hz, 3H), 4.40 (q, $J = 6.6$ Hz, 2H), 7.47 (d, $J = 9.0$ Hz, 1H), 7.86 (t, $J = 8.1$ Hz, 1H), 8.20 (s, 1H), 11.89 (br s, 1H).

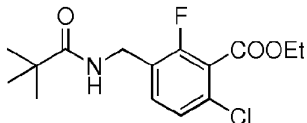
Step-3:- Preparation of ethyl 3-(aminomethyl)-6-chloro-2-fluorobenzoate



To a solution of ethyl 6-chloro-2-fluoro-3-((hydroxyimino)methyl)benzoate (0.800 g, 3.26 mmol) in MeOH (5 mL) was added Zn (0.850 g, 13.06 mmol), followed by dropwise addition of conc. HCl (catalytic) and the reaction mixture was heated at 60°C for 2-3 h. Excess of solvent was removed under vacuum and the residue was diluted with water and basified with NaHCO_3 , extracted with EtOAc and concentrated to afford 0.600

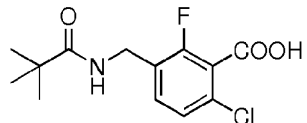
g of the product. ^1H NMR (300 MHz, DMSO d_6): δ 1.31 (t, $J = 7.2$ Hz, 3H), 3.76 (s, 2H), 4.38 (q, $J = 6.9$ Hz, 2H), 7.41 (d, $J = 8.1$ Hz, 1H), 7.65 (t, $J = 8.1$ Hz, 1H).

Step-4:- Preparation of ethyl 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoate



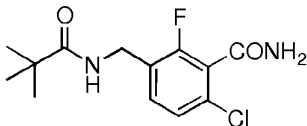
To a solution of ethyl 3-(aminomethyl)-6-chloro-2-fluorobenzoate (0.540 g, 2.5 mmol) in THF (5 mL) was added pivaloyl chloride (0.335 g, 2.79 mmol) and TEA (0.1 mL) and the reaction mixture was stirred for 2-3 h at RT. Excess of solvent was removed under vacuum and the residue was diluted with water, extracted with EtOAc and concentrated to afford 0.500 g of the product. ^1H NMR (300 MHz, DMSO d_6): δ 1.11 (s, 9H), 1.31 (t, $J = 7.5$ Hz, 3H), 4.26 (d, $J = 5.1$ Hz, 2H), 4.39 (q, $J = 6.6$ Hz, 2H), 7.33- 7.43 (m, 2H), 8.13 (m, 1H).

Step-5:- Preparation of 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoic acid



The title compound was prepared according to the procedure described in step-4 of Intermediate-48 by using ethyl 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoate (0.800 g) THF: methanol (2:2 mL) and aqueous solution of NaOH (1.0 g) to afford 0.500 g of the product. ^1H NMR (300 MHz, DMSO d_6): δ 1.11 (s, 9H), 4.26 (d, $J = 5.4$ Hz, 2H), 7.26- 7.38 (m, 2H), 8.14 (m, 1H), 14.07 (br s, 1H).

Step-6:- Preparation of 6-chloro-2-fluoro-3-(pivalamidomethyl)benzamide



The title compound was prepared according to the procedure described in step-2 of Intermediate-26 by using 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoic acid (0.400 g, 1.39 mmol), THF (25 mL), oxalyl chloride (0.14 mL, 1.67 mmol) and ammonia gas to afford 0.220 g of the product. ^1H NMR (300 MHz, DMSO d_6): δ 1.12 (s, 9H), 4.25 (d, $J =$

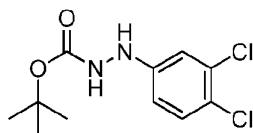
5.4 Hz, 2H), 7.20 (d, $J = 8.4$ Hz, 1H), 7.30 (d, $J = 8.4$ Hz, 1H), 7.82 (s, 1H), 8.12 (br s, 2H); MS (m/z): 287.01 ($M+H^+$).

Step-7:- Preparation of 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoyl isocyanate

To a solution of 6-chloro-2-fluoro-3-(pivalamidomethyl)benzamide (0.300 g, 1.04 mmol) in EDC (10 mL) was added oxalyl chloride (0.11 mL, 1.25 mmol). The reaction mass was heated at 60-70°C for 4-5 h. Excess of solvent was removed under vacuum to afford 1.0 g of desired product.

Intermediate-52

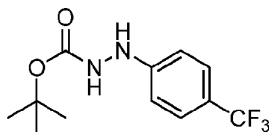
tert-Butyl 2-(3,4-dichlorophenyl)hydrazinecarboxylate



The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using (3,4-dichlorophenyl)hydrazine hydrochloride (1.00 g, 4.6 mmol), BOC anhydride (1.22 g, 5.6 mmol), Na_2CO_3 (1.24 g, 11.7 mmol), acetonitrile (20 mL) and water (10 mL) to afford 0.700 g of desired product. 1H NMR (300 MHz, $DMSO-d_6$): δ 1.40 (s, 9H), 6.62 (d, $J = 8.7$ Hz, 1H), 6.77 (s, 1H), 7.34 (d, $J = 8.7$ Hz, 1H), 8.02 (s, 1H), 8.91 (s, 9H).

Intermediate-53

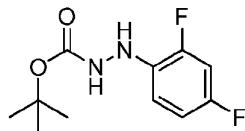
tert-Butyl 2-(4-(trifluoromethyl)phenyl)hydrazinecarboxylate



The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using 4-(trifluoromethyl)phenylhydrazine hydrochloride (1.00 g, 5.6 mmol), BOC anhydride (1.46 g, 6.8 mmol), Na_2CO_3 (0.900 g, 8.5 mmol), acetonitrile (20 mL) and water (10 mL) to afford 0.600 g of desired product. 1H NMR (300 MHz, $DMSO-d_6$): δ 1.41 (s, 9H), 6.74 (d, $J = 8.7$ Hz, 2H), 7.45 (d, $J = 9.0$ Hz, 2H), 8.23 (s, 1H), 8.94 (br s, 1H).

Intermediate-54

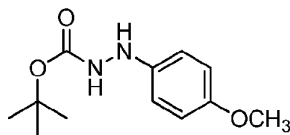
tert-Butyl 2-(2,4-difluorophenyl)hydrazinecarboxylate



The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using (2,4-difluorophenyl)hydrazine hydrochloride (1.00 g, 5.54 mmol), BOC anhydride (1.81 g, 8.31 mmol), Na_2CO_3 (1.46 g, 13.85 mmol), acetonitrile (20 mL) and water (10 mL) to afford 0.500 g of desired product. ^1H NMR (300 MHz, $\text{DMSO } d_6$): δ 1.40 (s, 9H), 6.70- 6.72 (m, 1H), 6.90 (t, $J = 8.1$ Hz, 1H), 7.07- 7.14 (m, 1H), 7.47 (s, 1H), 8.83 (br s, 1H).

Intermediate-55

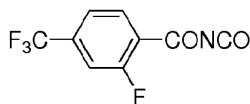
tert-Butyl 2-(4-methoxyphenyl)hydrazinecarboxylate



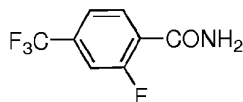
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using (4-methoxyphenyl)hydrazine hydrochloride (1.00 g, 5.72 mmol), BOC anhydride (1.8 g, 8.59 mmol), Na_2CO_3 (1.51 g, 14.31 mmol), acetonitrile (20 mL) and water (10 mL) to afford 0.400 g of desired product. ^1H NMR (300 MHz, $\text{DMSO } d_6$): δ 1.39 (s, 9H), 3.64 (s, 3H), 6.60 (d, $J = 7.8$ Hz, 2H), 7.74 (d, $J = 8.4$ Hz, 2H), 7.18 (br s, 1H), 8.68 (br s, 1H).

Intermediate-56

2-Fluoro-4-(trifluoromethyl)benzoyl isocyanate



Step-1:- Preparation of 2-fluoro-4-(trifluoromethyl)benzamide



The title compound was prepared according to the procedure described in step-2 of Intermediate-26 by using 2-fluoro-4-(trifluoromethyl)benzoic acid (1.5 g), THF (25 mL), oxalyl chloride (0.5 mL) and ammonia gas to afford 1.35 g of the product. ^1H NMR (300

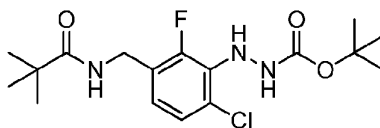
MHz, DMSO d_6): δ 7.65 (d, J = 8.4 Hz, 1H), 7.77- 7.84 (m, 3H), 7.96 (br s, 1H); MS (m/z): 208.06 (M^+).

Step-2:- Preparation of 2-fluoro-4-(trifluoromethyl)benzoyl isocyanate

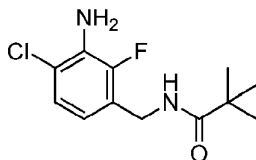
The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 2-fluoro-4-(trifluoromethyl)benzamide (0.200 g, 0.96 mmol), EDC (10 mL) and oxalyl chloride (0.1 mL, 1.15 mmol) to afford 0.200 g of the product.

Intermediate-57

tert-Butyl 2-(6-chloro-2-fluoro-3-(pivalamidomethyl)phenyl)hydrazinecarboxylate

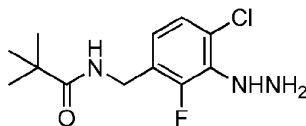


Step-1:- Preparation of *N*-(3-amino-4-chloro-2-fluorobenzyl)pivalamide



To a solution of 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoic acid (step-5 of Intermediate-51, 1.0 g, 3.48 mmol) in conc. sulphuric acid (10.0 mL) was added sodium azide (0.271 g, 4.18 mmol). The reaction mass was stirred at 50-60°C for 18 h. The reaction mass was quenched in ammonia solution (cold) and solid obtained was filtered and suck dried to afford 0.570 g of the product. ^1H NMR (300 MHz, DMSO d_6): δ 1.11 (s, 9H), 4.21 (d, J = 5.7 Hz, 2H), 5.30 (s, 2H), 6.39 (t, J = 7.8 Hz, 1H), 7.00 (d, J = 8.4 Hz, 1H), 7.98 (m, 1H); MS (m/z): 259.12 ($M+H^+$).

Step-2:- Preparation of *N*-(4-chloro-2-fluoro-3-hydrazinylbenzyl)pivalamide



The title compound was prepared according to the procedure described in step-2 of Intermediate-7 by using *N*-(3-amino-4-chloro-2-fluorobenzyl)pivalamide (0.500 g, 1.93 mmol), sodium nitrite (0.200 g, 2.90 mmol), stannous chloride.H₂O (4.84 g, 1.09 mmol), conc. HCl (20.0 g) and water (3 mL) to afford 0.300 g of the desired product. ^1H NMR

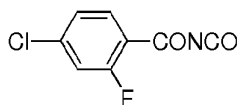
(300 MHz, DMSO d_6): δ 1.12 (s, 9H), 4.23(d, J = 5.4 Hz, 2H), 4.34 (br s, 2H), 5.91 (s, 1H), 6.65 (t, J = 7.8 Hz, 1H), 8.02 (m, 1H) ; MS (m/z): 274.08 (M+H⁺).

Step-3:- Preparation of *tert*-butyl 2-(6-chloro-2-fluoro-3-(pivalamidomethyl)phenyl)hydrazinecarboxylate

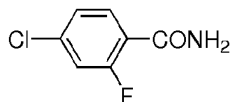
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using *N*-(4-chloro-2-fluoro-3-hydrazinylbenzyl)pivalamide (0.300 g, 1.09 mmol), BOC anhydride (0.360 g, 1.64 mmol), Na₂CO₃ (0.231 g, 2.18 mmol), acetonitrile (10 mL) and water (5 mL) to afford 0.150 g of desired product. ¹H NMR (300 MHz, DMSO d_6): δ 1.11 (s, 9H), 1.37 (s, 9H), 4.20 (d, J = 5.7 Hz, 2H), 6.64 (m, 1H), 7.06 (br s, 2H), 8.00 (br s, 1H), 8.98 (s, 1H).

Intermediate-58

4-Chloro-2-fluorobenzoyl isocyanate

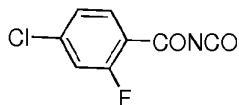


Step-1:- Preparation of 4-chloro-2-fluorobenzamide



The title compound was prepared according to the procedure described in step-2 of Intermediate-26 by using 4-chloro-2-fluoro benzoic acid (1.5 g), THF (25 mL), oxalyl chloride (0.5 mL) and ammonia gas to afford 1.45 g of the product. ¹H NMR (300 MHz, DMSO d_6): δ 7.36 (d, J = 6.3 Hz, 1H), 7.53 (dd, J = 8.7 Hz, 1H), 7.64- 7.76 (m, 3H).

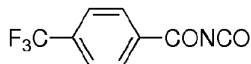
Step-2:- Preparation of 4-chloro-2-fluorobenzoyl isocyanate



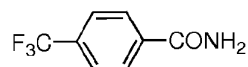
The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 4-chloro-2-fluorobenzamide (0.200 g, 1.26 mmol), EDC (10 mL) and oxalyl chloride (0.14 mL, 1.15 mmol) to afford 0.200 g of the product.

Intermediate -59

4-(Trifluoromethyl)benzoyl isocyanate



Step-1:- Preparation of 4-(trifluoromethyl)benzamide



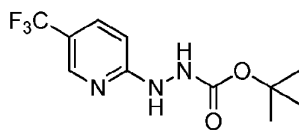
To a cold solution of 4-(trifluoromethyl)benzonitrile (0.500 g, 2.92 mmol) in DMSO (6.0 mL) was added H₂O₂ (50%) (5 mL) at 0°C, followed by portion-wise addition of K₂CO₃ (0.121 g, 0.87 mmol). The reaction mass was allowed to attain RT and stirred for 1 h. The reaction mass was quenched in ice water and extracted with DCM and concentrated to afford 0.300 g of product. ¹H NMR (400 MHz, DMSO *d*₆): δ 7.60 (br s, 1H), 7.82 (d, *J* = 7.5 Hz, 2H), 8.05 (d, *J* = 6.6 Hz, 2H), 8.17 (br s, 1H); MS (*m/z*): 190.11 (M+H⁺).

Step-2:- Preparation of 4-(trifluoromethyl)benzoyl isocyanate

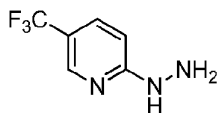
The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 4-(trifluoromethyl)benzamide (0.200 g, 1.05 mmol), EDC (10 mL) and oxalyl chloride (0.15 mL, 1.58 mmol) to afford 0.200 g of the product.

Intermediate-60

tert-Butyl 2-(5-(trifluoromethyl)pyridin-2-yl)hydrazinecarboxylate



Step-1:- Preparation of 2-hydrazinyl-5-(trifluoromethyl)pyridine



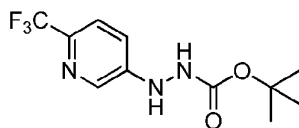
To a solution of 2-chloro-5-(trifluoromethyl)pyridine (1.5 g, 8.26 mmol) in ethanol (20.0 mL) was added hydrazinehydrate (1.21 g, 24.79 mmol). The reaction mass was refluxed for 5-6 h. The reaction mass was concentrated and quenched in ice water and solid obtained was filtered and suck dried to afford 0.700 g of the product. ¹H NMR (300 MHz, DMSO *d*₆): δ 4.32 (br s, 2H), 6.78 (d, *J* = 8.8 Hz, 1H), 7.67 (d, *J* = 8.4 Hz, 1H), 8.25 (br s, 2H); MS (*m/z*): 178.15 (M+H⁺).

Step-2:- Preparation of *tert*-butyl 2-(5-(trifluoromethyl)pyridin-2-yl)hydrazinecarboxylate

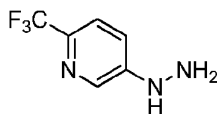
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using 2-hydrazinyl-5-(trifluoromethyl)pyridine (0.700 g, 3.95 mmol), BOC anhydride (1.3 g, 5.9 mmol), Na_2CO_3 (0.838 g, 7.90 mmol), acetonitrile (15 mL) and water (7 mL) to afford 0.500 g of desired product. ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ 1.40 (s, 9H), 6.62 (d, $J = 8.8$ Hz, 1H), 7.83 (dd, $J = 8.0$ Hz, 1H), 8.37 (s, 1H), 8.95 (s, 1H), 9.01 (br s, 1H); MS (m/z): 278.00 ($\text{M}+\text{H}^+$).

Intermediate-61

tert-Butyl 2-(6-(trifluoromethyl)pyridin-3-yl)hydrazinecarboxylate



Step-1:- Preparation of 5-hydrazinyl-2-(trifluoromethyl)pyridine



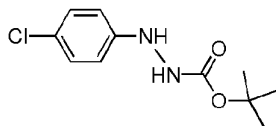
To a cold solution of 6-(trifluoromethyl)pyridin-3-amine (1.5 g, 9.25 mmol) in 4N HCl (60.0 mL) was added aqueous solution of NaNO_2 (0.766 g, 11.11 mmol) at 0°C . The reaction mass was stirred at same temperature for 30 minutes. The reaction mass was added to a solution of $\text{SnCl}_2\cdot\text{H}_2\text{O}$ in 4N HCl at 80°C and further continued stirring for 5-6 h at same temperature. The reaction mass was cooled, basified and extracted with DCM. The organic layer was dried and concentrated to afford 0.700 g of the product. ^1H NMR (300 MHz, $\text{DMSO}-d_6$): δ 4.29 (br s, 2H), 7.18 (dd, $J = 8.6$ Hz, 1H), 7.51 (d, $J = 8.8$ Hz, 1H), 7.68 (s, 1H), 8.13 (d, $J = 2.4$ Hz, 1H); MS (m/z): 178.15 ($\text{M}+\text{H}^+$).

Step-2:- Preparation of *tert*-butyl 2-(6-(trifluoromethyl)pyridin-3-yl)hydrazinecarboxylate

The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using 5-hydrazinyl-2-(trifluoromethyl)pyridine (0.700 g, 3.95 mmol), BOC anhydride (1.3 g, 5.9 mmol), Na_2CO_3 (0.838 g, 7.90 mmol), acetonitrile (15 mL) and water (7 mL) to afford 0.400 g of desired product. ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ 1.31 (s, 9H), 7.10 (dd, $J = 8.0$ Hz, 1H), 7.64 (d, $J = 8.6$ Hz, 1H), 8.09 (d, $J = 2.3$ Hz, 1H), 8.55 (br s, 1H), 9.08 (br s, 1H); MS (m/z): 278.05 ($\text{M}+\text{H}^+$).

Intermediate-62

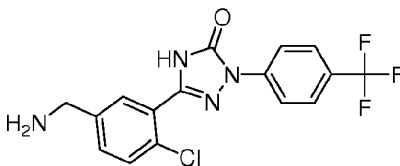
tert-Butyl 2-(4-chlorophenyl)hydrazinecarboxylate



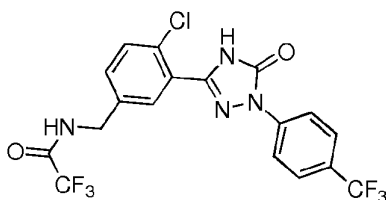
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using (4-chlorophenyl)hydrazine hydrochloride (1.5 g, 10.5 mmol), BOC anhydride (2.7 g, 12.6 mmol), Na₂CO₃ (2.23 g, 21.03 mmol), acetonitrile (20 mL) and water (10 mL) to afford 0.900 g of desired product.

Intermediate-63

3-(5-(Aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



Step-1:- Preparation of *N*-(4-chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



To a solution of *tert*-butyl 2-(4-(trifluoromethyl)phenyl)hydrazinecarboxylate (Intermediate-53, 1.00 g, 3.63 mmol) in DCM (20 mL), 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate (step-3 of Intermediate-26, 1.30 g, 4.03 mmol) was added and the reaction mass was stirred at RT for 2 h. After completion of reaction, excess of solvent was removed under reduced pressure to obtain 0.700 g of the crude product. To a solution of obtained crude product in DCM (5.0 mL), TFA (5.0 mL) was added and the reaction mass was stirred at RT for 2-3 h. Excess of the solvent was removed from the reaction mass under reduced pressure and the reaction mass was quenched in ice and filtered to obtain 0.700 g of the desired title product. ¹H NMR (300 MHz, DMSO *d*₆): δ 4.45 (d, 1H), 7.50 (d, *J* = 4.8 Hz, 1H), 7.65- 7.68 (m, 2H), 8.85 (d, *J*

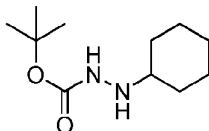
= 6.6Hz, 2H), 8.18 (d, J = 6.3Hz, 2H), 10.05 (s, 1H), 12.69 (s, 1H); MS (m/z): 463.17 ($M-H$).

Step-2:- Preparation of 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one

To a solution of *N*-(4-chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (0.400 g, 0.861 mmol) in THF (5.0 mL) was added KOH aqueous solution (0.096 g, 1.72 mmol) and the reaction mass was stirred at RT for 3-4 h. The reaction mass was quenched in water and extracted with ethyl acetate and concentrated to afford 0.200 g of the desired product.

Intermediate-64

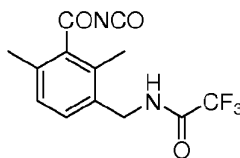
tert-Butyl 2-Cyclohexyl hydrazinecarboxylate



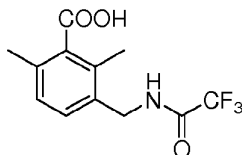
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using cyclohexyl hydrazine hydrochloride (1.00 g, 6.6 mmol), BOC anhydride (1.73 g, 7.9 mmol), Na_2CO_3 (1.75 g, 16.6 mmol), acetonitrile (20 mL) and water (10 mL) to afford 0.500 g of desired product. 1H NMR (300 MHz, DMSO d_6): δ 0.95-1.02 (m, 2H), 1.07-1.14 (m, 2H), 1.16 (s, 9H), 1.39-1.51 (m, 2H), 1.67-1.98 (m, 2H), 2.61 (m, 1H), 8.16 (br s, 1H), 8.75 (br s, 1H). MS (m/z): 214.79 (M^+).

Intermediate-65

3-((2,2,2-Trifluoroacetamido)methyl)-2,6-dimethylbenzoyl isocyanate

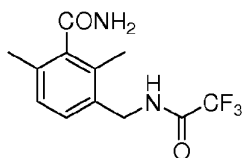


Step-1:- Preparation of 3-((2,2,2-trifluoroacetamido)methyl)-2,6-dimethylbenzoic acid



To a solution of 2,6-dimethyl benzoic acid (6.8 g, 0.045 mmol) in conc.H₂SO₄ (100 mL), 2,2,2-trifluoro-*N*-(hydroxymethyl)acetamide (7.1 g, 0.049 mmol) was added and the reaction mixture was stirred at RT for 24 h. After completion of the reaction, the reaction mass was quenched in ice water and stirred for 15 minutes. The obtained solid product was filtered and dried to afford 7.0 g of the desired title product. ¹H NMR (300 MHz, DMSO *d*₆): δ 2.26 (s, 6H), 4.36 (t, *J* = 12.3 Hz, 2H), 7.09 (m, 1H), 7.15 (m, 1H), 9.89-9.94 (m, 1H).

Step-2:- Preparation of 3-((2,2,2-trifluoroacetamido)methyl)-2,6-dimethylbenzamide



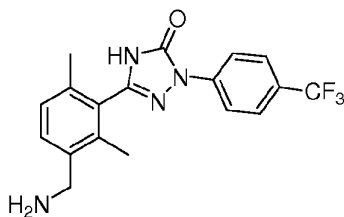
The title compound was prepared according to the procedure described in step-2 of Intermediate-26 by using 3-((2,2,2-trifluoroacetamido)methyl)-2,6-dimethylbenzoic acid (1.0 g), oxalyl chloride (1.0 mL), THF (50 mL) and ammonia gas to afford 0.800 g of the desired product.

Step-3:- Preparation of 3-((2,2,2-trifluoroacetamido)methyl)-2,6-dimethylbenzoyl isocyanate

The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 3-((2,2,2-trifluoroacetamido)methyl)-2,6-dimethylbenzamide (1.0 g, 3.6 mmol), oxalyl chloride and EDC (10 mL) to afford 0.900 g of the desired product.

Intermediate-66

5-(3-(Aminomethyl)-2,6-dimethylphenyl)-2-(4-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one

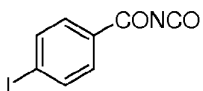


To a solution of *N*-(3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)-2,2,2-trifluoroacetamide (Example-106, 0.800 g, 1.74 mmol)

in THF (10 mL), aqueous solution of KOH was added and the reaction mass was stirred at RT for 3-4 h. After completion of the reaction, the reaction mass was quenched with water and extracted with ethyl acetate. The organic layer was separated, dried over anhydrous sodium sulphate and concentrated to afford 0.200 g of the desired product. MS (m/z): 362.86 (M+H)⁺.

Intermediate-67

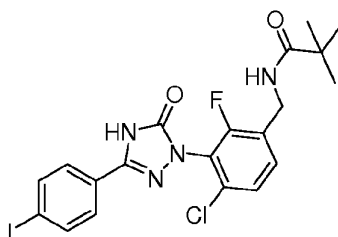
4-Iodobenzoyl isocyanate



The title compound was prepared according to the procedures described in step-2 and step-3 of Intermediate-26 by using 4-iodo benzoic acid (2.0 g), oxalyl chloride (1.0 ml), ammonia gas, DCM (20 mL), THF (20 mL) to afford 1.5 g of the desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 7.17 (br s, 1H), 7.64 (d, 2H), 8.01 (s, 2H); MS (m/z): 461.14 (M+H)⁺.

Intermediate-68

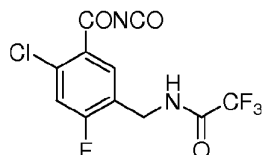
N-(4-Chloro-2-fluoro-3-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide



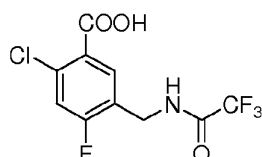
The title compound was prepared according to the procedure described in Example-83 by using *tert*-butyl 2-(6-chloro-2-fluoro-3-(pivalamidomethyl)phenyl)hydrazinecarboxylate (Intermediate-57, 0.250g), 4-iodobenzoyl isocyanate (Intermediate-67, 0.250 g), DCM (10 mL) and trifluoro acetic acid (0.5 mL) to afford 0.150 g of the desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 1.19 (s, 9H), 4.30 (d, *J* = 3.3 Hz, 2H), 7.39 (m, 1H), 7.45 (d, 1H), 7.62 (d, 2H), 7.90 (d, *J* = 3.6 Hz, 2H), 8.19 (t, 1H), 13.0 (br s, 1H); MS (m/z): 528.98 (M+H)⁺.

Intermediate-69

5-((2,2,2-Trifluoroacetamido)methyl)-2-chloro-4-fluorobenzoyl isocyanate

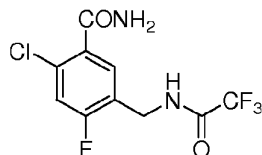


Step-1:- Preparation of 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzoic acid



The title compound was prepared according to the procedure described in step-1 of Intermediate-26 by using 2-chloro-4-fluorobenzoic acid (5.0 g, 0.002 mmol), 2,2,2-trifluoro-*N*-(hydroxymethyl)acetamide (4.0 g, 0.002 mmol) and conc.H₂SO₄ (50 mL) to afford 4.0 g of the desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 4.44 (d, *J* = 12.0 Hz, 2H), 7.59 (m, 1H), 7.92 (m, 1H), 10.0 (br s, 1H), 13.5 (br s, 1H).

Step-2:- Preparation of 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzamide



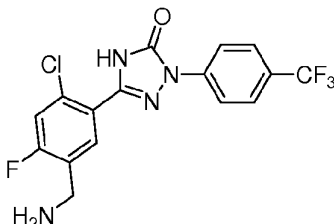
The title compound was prepared according to the procedure described in step-2 of Intermediate-26 by using 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzoic acid (1.0 g), oxalyl chloride (1.0 mL), THF (50 mL) and ammonia gas to afford 0.800 g of the desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 4.53 (m, 2H), 7.43 (m, 1H), 7.44 (m, 2H), 7.48 (m, 1H), 10.0 (br s, 1H); MS (*m/z*): 297.10 (*M*-H)⁻.

Step-3:- Preparation of 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzoyl isocyanate

The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzamide (1.0 g, 3.6 mmol), oxalyl chloride (1.0mL) and EDC (10 mL) to afford 0.750 g of the desired product.

Intermediate-70

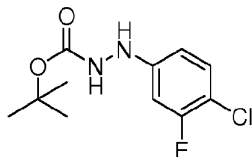
5-(5-(Aminomethyl)-2-chloro-4-fluorophenyl)-2-(4-(trifluoromethyl)phenyl)-2H-1,2,4-triazol-3(4H)-one



The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-2-fluoro-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1H-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (Example-113, 0.100 g, 0.207 mmol), KOH (0.034 g, 0.622 mol), water (5 mL) and THF (5.0 mL) to afford 0.080 g of the desired product.

Intermediate-71

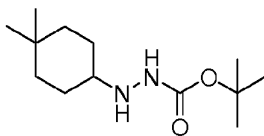
tert-Butyl 2-(4-chloro-3-fluorophenyl)hydrazinecarboxylate



The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using 4-chloro-3-fluorophenylhydrazine hydrochloride (1.00 g), BOC anhydride (1.5g), Na₂CO₃ (0.900 g, 8.5 mmol), acetonitrile (20 mL) and water (10 mL) to afford 0.600 g of the desired product.

Intermediate-72

4,4-Dimethyl *tert*-butyl 2-cyclohexyl hydrazinecarboxylate

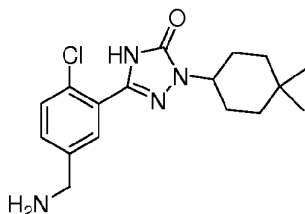


The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using 4,4-dimethylcyclohexylhydrazine hydrochloride (1.00 g), BOC

anhydride (1.5g), Na₂CO₃ (0.900 g, 8.5 mmol), acetonitrile (20 mL) and water (10 mL) to afford 0.600 g of the desired product.

Intermediate-73

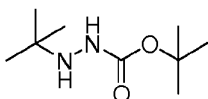
5-(5-(Aminomethyl)-2-chlorophenyl)-2-(4,4-dimethylcyclohexyl)-2*H*-1,2,4-triazol-3(4*H*)-one



The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-3-(4,5-dihydro-1-(4,4-dimethylcyclohexyl)-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (Example-117, 0.150 g, 0.348 mmol), KOH (0.039 g, 0.696 mol), water (2 mL) and THF (5.0 mL) to afford 0.100 g of the desired product.

Intermediate-74

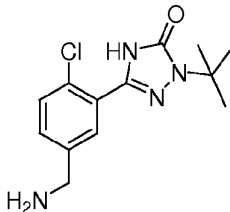
tert-Butyl 2-*tert*-butylhydrazinecarboxylate



The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using *tert*-butylhydrazine (1.00 g), BOC anhydride (1.5g), Na₂CO₃ (0.900 g, 8.5 mmol), acetonitrile (20 mL) and water (10 mL) to afford 0.600 g of the desired product.

Intermediate-75

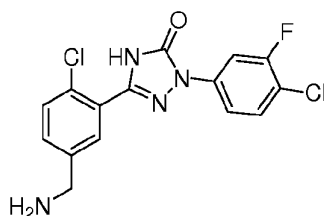
2-*tert*-Butyl-5-(5-(aminomethyl)-2-chlorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one



The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(3-(1-*tert*-butyl-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-4-chlorobenzyl)-2,2,2-trifluoroacetamide (Example-120, 0.400 g, 1.062 mmol), KOH (0.119 g, 2.12 mmol), water (5 mL) and THF (5.0 mL) to afford 0.300 g of the desired product.

Intermediate-76

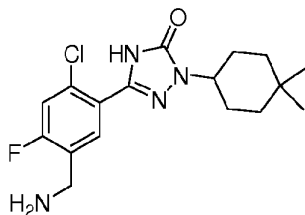
5-(5-(Aminomethyl)-2-chlorophenyl)-2-(4-chloro-3-fluorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one



The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-3-(1-(4-chloro-3-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (Example-116, 0.400 g, 0.890 mmol), KOH (0.099 g, 1.78 mmol), water (2 mL) and THF (4.0 mL) to afford 0.300 g of the desired product.

Intermediate-77

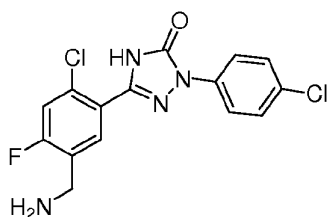
5-(5-(Aminomethyl)-2-chloro-4-fluorophenyl)-2-(4,4-dimethylcyclohexyl)-2*H*-1,2,4-triazol-3(4*H*)-one



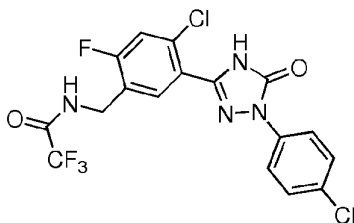
The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-2-fluoro-5-(4,5-dihydro-1-(4,4-dimethylcyclohexyl)-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (Example-124, 1.0 g), KOH (1.0 g), water (5 mL) and THF (40.0 mL) to afford 0.700 g of the desired product.

Intermediate-78

5-(5-(Aminomethyl)-2-chloro-4-fluorophenyl)-2-(4-chlorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one



Step-1:- Preparation of *N*-(4-chloro-5-(1-(4-chlorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)-2,2,2-trifluoroacetamide



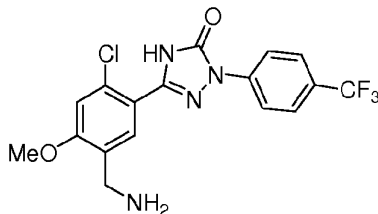
The title compound was prepared according to the procedure described in Example-83 by using 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzoyl isocyanate (Intermediate-69, 0.400 g), *tert*-butyl 2-(4-chlorophenyl)hydrazinecarboxylate (Intermediate-62, 0.400 g), DCM (20 mL) and trifluoro acetic acid (5.0 mL) to afford 0.250 g of the desired product.

Step-2:- Preparation of 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(4-chlorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one

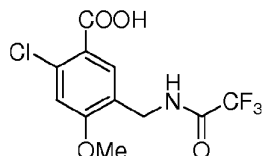
The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-5-(1-(4-chlorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)-2,2,2-trifluoroacetamide (0.160 g), KOH (0.050 g), water (2 mL) and THF (10.0 mL) to afford 0.070 g of the desired product.

Intermediate-79

5-(5-(Aminomethyl)-2-chloro-4-methoxyphenyl)-2-(4-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one

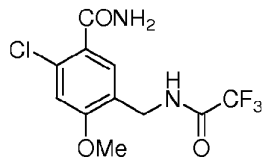


Step-1:- Preparation of 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-methoxybenzoic acid



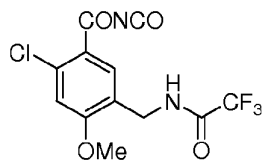
The title compound was prepared according to the procedure described in step-1 of Intermediate-26 by using 2-chloro-4-methoxybenzoic acid (0.060 g), 2,2,2-trifluoro-*N*-(hydroxymethyl)acetamide (0.050 g) and H₂SO₄ (2 mL) to afford 0.040 g of the desired product.

Step-2:- Preparation of 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-methoxybenzamide



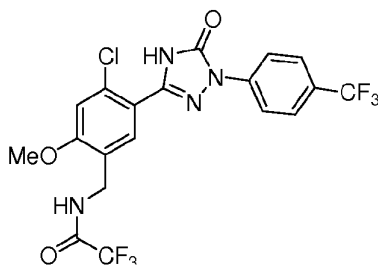
The title compound was prepared according to the procedure described in step-2 of Intermediate-26 by using 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-methoxybenzoic acid (1.0 g), oxalyl chloride (1.0 mL), THF (50 mL) and ammonia gas to afford 0.800 g of the desired product.

Step-3:- Preparation of 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-methoxybenzoyl isocyanate



The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-methoxybenzamide (1.0 g, 3.6 mmol), oxalyl chloride (1.0 mL) and EDC (10 mL) to afford 0.750 g of the desired product.

Step-4:- Preparation of *N*-(4-chloro-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-methoxybenzyl)-2,2,2-trifluoroacetamide



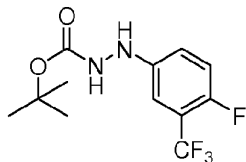
The title compound was prepared according to the procedure described in Example-83 by using 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-methoxybenzoyl isocyanate (0.500 g), *tert*-butyl 2-(4-(trifluoromethyl)phenyl)hydrazinecarboxylate (Intermediate-53, 0.450 g), DCM (20 mL), and trifluoro acetic acid (5.0 mL) to afford 0.180 g of the desired product.

Step-5:- Preparation of 5-(5-(aminomethyl)-2-chloro-4-methoxyphenyl)-2-(4-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one

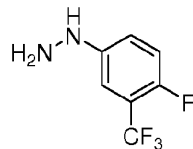
The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-methoxybenzyl)-2,2,2-trifluoroacetamide (0.200 g), water (5 mL), KOH (0.200 g) and THF (10.0 mL) to afford 0.150 g of the desired product.

Intermediate-80

tert-Butyl 2-[4-fluoro-3-(trifluoromethyl)phenyl]hydrazinecarboxylate



Step1:- Preparation of 1-(4-fluoro-3-(trifluoromethyl)phenyl)hydrazine



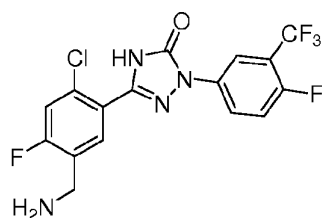
The title compound was prepared according to the procedure described in step-1 of Intermediate-61 by using 4-fluoro-3-(trifluoromethyl)aniline (3.0 g, 0.016 mmol), NaNO_2 (1.73 g, 0.025 mmol), $\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$ (9.39 g, 0.041 mmol) and conc. HCl (100 mL) to afford 2.0 g of the desired product.

Step-2:- Preparation of *tert*-butyl 2-[4-fluoro-3-(trifluoromethyl)phenyl]hydrazinecarboxylate

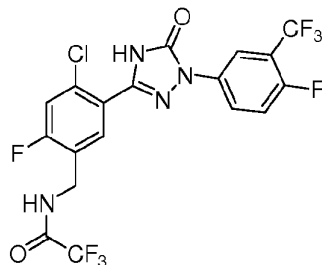
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using 1-(4-fluoro-3-(trifluoromethyl)phenyl)hydrazine (1.00 g), BOC anhydride (1.5g), Na_2CO_3 (0.900 g, 8.5 mmol), acetonitrile (20 mL) and water (10 mL) to afford 0.600 g of the desired product. ^1H NMR (300 MHz, CDCl_3): δ 1.40 (s, 9H), 6.86-6.94 (m, 2H), 7.26- 7.32 (m, 1H), 7.98 (s, 1H), 8.95 (s, 1H); MS (m/z): 436.55 (M^+H^+).

Intermediate-81

5-(5-(Aminomethyl)-2-chloro-4-fluorophenyl)-2-(4-fluoro-3-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one



Step-1:- Preparation of *N*-(4-chloro-2-fluoro-5-(1-(4-fluoro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



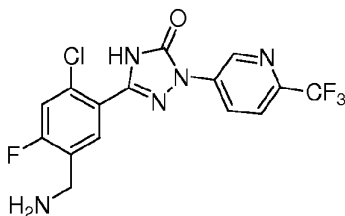
The title compound was prepared according to the procedure described in Example-83 by using 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzoyl isocyanate (Intermediate-69, 0.400 g), *tert*-butyl 2-[4-fluoro-3-(trifluoromethyl)phenyl]hydrazinecarboxylate (Intermediate-80, 0.400 g), DCM (20 mL) and TFA (5.0 mL) to afford 0.200 g of the desired product.

Step-2:- Preparation of 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(4-fluoro-3-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one

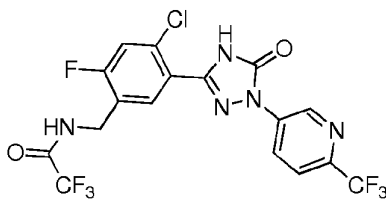
The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-3-(1-(4-fluoro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (0.400 g), KOH (0.100 g) and THF (10.0 mL) to afford 0.150 g of desired product.

Intermediate-82

5-(5-(Aminomethyl)-2-chloro-4-fluorophenyl)-2-(6-(trifluoromethyl)pyridin-3-yl)-2*H*-1,2,4-triazol-3(4*H*)-one



Step-1:- Preparation of *N*-(4-chloro-2-fluoro-5-(1-(6-(trifluoromethyl)pyridin-3-yl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



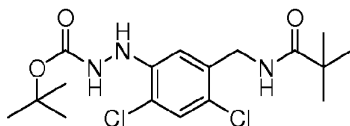
The title compound was prepared according to the procedure described in Example-83 by using 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzoyl isocyanate (Intermediate-69, 0.350 g), *tert*-butyl 2-[6-(trifluoromethyl)pyridin-3-yl]hydrazinecarboxylate (Intermediate-61, 0.350 g), DCM (20 mL) and TFA (5.0 mL) to afford 0.250 g of the desired product.

Step-2:- Preparation of 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(6-(trifluoromethyl)pyridin-3-yl)-2*H*-1,2,4-triazol-3(4*H*)-one

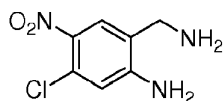
The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-2-fluoro-5-(1-(6-(trifluoromethyl)pyridin-3-yl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (0.500 g), KOH (0.200 g), water (5 mL) and THF (10.0 mL) to afford 0.150 g of desired product.

Intermediate-83

tert-Butyl 2-(2,4-dichloro-5-[(2,2-dimethylpropanoyl)amino]methyl}phenyl)hydrazinecarboxylate

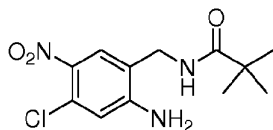


Step-1:- Preparation of 2-(aminomethyl)-5-chloro-4-nitrobenzenamine



To a solution of 2-amino-4-chloro-5-nitrobenzamide (3.0 g, 13.7 mmol) in dry THF (50 mL), borane dimethyl sulphide complex (1.56 g, 20.6 mmol) was added at RT and the reaction mass was refluxed for 24 h. The reaction mass was cooled to RT and dilute HCl was added till it became acidic. Further, the reaction mixture was stirred for 1 h and then the reaction mixture was basified with dilute NaOH and extracted with ethyl acetate. The organic layer was separated, dried over anhydrous sodium sulphate and concentrated to afford 2.0 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.58 (s, 2H), 6.72 (br s, 3H), 8.04 (br s, 1H); MS (m/z): 200.28 (M-H).

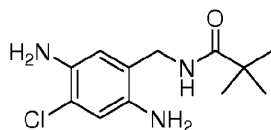
Step-2:- Preparation of *N*-(2-amino-4-chloro-5-nitrobenzyl)pivalamide



To a solution of 2-(aminomethyl)-5-chloro-4-nitrobenzenamine (2.5 g, 12.25 mmol) in THF (35 mL), TEA (3.0 mL) and pivaloyl chloride (1.96 mL, 14.7 mmol) were added and the reaction mass was stirred at RT for 5 h. The reaction mass was quenched with

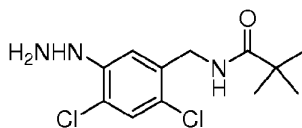
water and extracted with ethyl acetate. The organic layer was washed with dilute HCl and dilute sodium bicarbonate solution, separated, dried over anhydrous sodium sulphate and concentrated to afford 2.0 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.12 (s, 9H), 4.30 (d, $J = 5.4$ Hz, 2H), 7.81 (d, $J = 9.9$ Hz, 1H), 7.96 (d, $J = 6.3$ Hz, 1H), 8.21 (m, 1H).

Step3:- Preparation of *N*-(2,5-diamino-4-chlorobenzyl)pivalamide



To a solution of *N*-(2-amino-4-chloro-5-nitrobenzyl)pivalamide (2.0 g, 6.9 mmol) in methanol (20 mL), Raney Ni (2.0 g) and hydrazine hydrate (5.0 mL) were added and the reaction mass was stirred at RT for 3 h. After the completion of the reaction, the reaction mass was filtered through celite pad and obtained filtrate was concentrated to afford 1.5 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.11 (s, 9H), 3.97 (br d, 2H), 4.52 (br d, 2H), 5.44 (br s, 2H), 6.50 (d, $J = 11.4$ Hz, 2H), 7.88 (m, 1H).

Step-4:- Preparation of *N*-(2,4-dichloro-5-hydrazinylbenzyl)-2,2-dimethylpropanamide



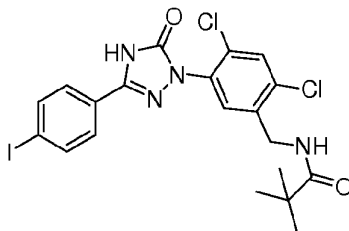
The title compound was prepared according to the procedure described in step-1 of Intermediate-61 by using *N*-(2,5-diamino-4-chlorobenzyl)pivalamide (1.5 g, 5.8 mmol), NaNO_2 (0.481 g, 6.9 mmol), $\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$ (3.20 g, 14.5 mmol), conc. HCl (50 mL) to afford 1.0 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.11 (s, 9H), 4.44 (m, 2H), 6.53 (m, 1H), 7.18 (m, 1H), 7.99 (m, 1H), 10.23 (s, 1H).

Step-5:- Preparation of *tert*-butyl 2-(2,4-dichloro-5-[(2,2-dimethylpropanoyl)amino]methyl)phenyl)hydrazinecarboxylate

The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using *N*-(2,4-dichloro-5-hydrazinylbenzyl)-2,2-dimethylpropanamide (1.0 g, 3.6 mmol), BOC anhydride (0.958 g, 4.39 mmol), Na_2CO_3 (0.776 g, 7.32 mmol), acetonitrile (10 mL) and water (5 mL) to afford 0.500 g of the desired product.

Intermediate-84

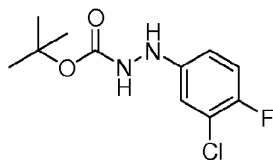
N-(2,4-Dichloro-5-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-83 by using 4-iodobenzoyl isocyanate (Intermediate-67, 0.300 g, 0.800 mmol), *tert*-butyl 2-(2,4-dichloro-5-[(2,2-dimethylpropanoyl)amino]methyl)phenylhydrazine carboxylate (Intermediate-83, 0.397g, 1.6 mmol), DCM (20 mL), trifluoro acetic acid (5.0 mL) to afford 0.150 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.09 (s, 9H), 4.31 (d, J = 4.2 Hz, 2H), 7.40- 7.58 (m, 2H), 7.61- 7.78 (m, 2H), 7.93-8.02 (br d, 2H), 8.17 (m, 1H).

Intermediate-85

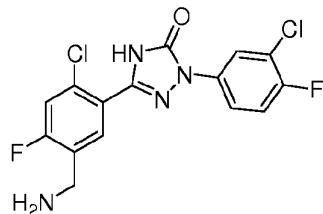
tert-Butyl 2-(3-chloro-4-fluorophenyl)hydrazinecarboxylate



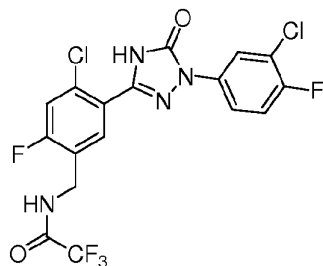
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using 1-(3-chloro-4-fluorophenyl)hydrazine (3.0 g, 0.015 mmol), BOC anhydride (3.6 g, 0.16 mmol), Na_2CO_3 (2.40 g, 0.022 mmol), acetonitrile (30 mL) and water (5 mL) to afford 1.3 g of the desired product. ^1H NMR (300 MHz, CDCl_3): δ 1.38 (s, 9H), 6.60 (m, 1H), 6.67 (m, 1H), 7.13- 7.20 (m, 1H), 7.77 (s, 1H), 8.85 (s, 1H).

Intermediate-86

5-(5-(Aminomethyl)-2-chloro-4-fluorophenyl)-2-(3-chloro-4-fluorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one



Step-1:- Preparation of *N*-(4-chloro-5-(1-(3-chloro-4-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)-2,2,2-trifluoroacetamide



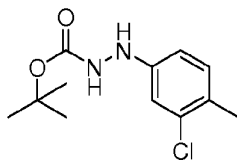
The title compound was prepared according to the procedure described in Example-83 by using 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzoyl isocyanate (Intermediate-69, 0.300 g), *tert*-butyl 2-(3-chloro-4-fluorophenyl)hydrazinecarboxylate (Intermediate-85, 0.300 g), DCM (20 mL) and trifluoro acetic acid (5.0 mL) to afford 0.200 g of the desired product.

Step-2:- Preparation of 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(3-chloro-4-fluorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one

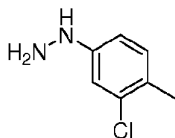
The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-5-(1-(3-chloro-4-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)-2,2,2-trifluoroacetamide (0.300 g), KOH (0.100 g), water (2 ml) and THF (10.0 mL) to afford 0.100 g of the desired product.

Intermediate-87

tert-Butyl 2-(3-chloro-4-methylphenyl)hydrazinecarboxylate



Step-1:- Preparation of 1-(3-chloro-4-methylphenyl)hydrazine



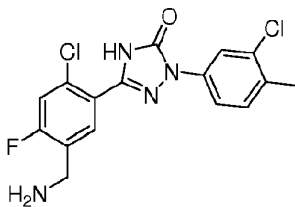
The title compound was prepared according to the procedure described in step-1 of Intermediate-61 by using 3-chloro-4-methylaniline (6.0 g, 0.0420 mmol), NaNO_2 (8.56 g, 0.050 mmol), $\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$ (23.7 g, 0.105 mmol) and conc. HCl (100 mL) to afford 4.5 g of the desired product.

Step-2:- Preparation of *tert*-butyl 2-(3-chloro-4-methylphenyl)hydrazine carboxylate

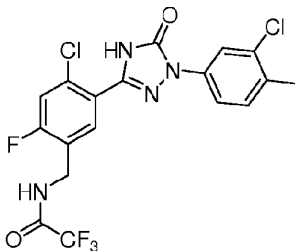
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using 1-(3-chloro-4-methylphenyl)hydrazine (4.5 g, 0.028 mmol), BOC anhydride (6.8 g, 0.032 mmol), Na_2CO_3 (4.50 g, 0.041 mmol), acetonitrile (30 mL) and water (5 mL) to afford 1.300 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.40 (s, 9H), 2.17 (s, 3H), 6.53 (d, $J = 10.4$ Hz, 1H), 6.63 (s, 1H), 7.08 (d, $J = 10.82$ Hz, 1H), 7.67 (s, 1H), 8.80 (s, 1H).

Intermediate-88

5-(5-(Aminomethyl)-2-chloro-4-fluorophenyl)-2-(3-chloro-4-methylphenyl)-2H-1,2,4-triazol-3(4H)-one



Step-1:- Preparation of *N*-(4-chloro-5-(1-(3-chloro-4-methylphenyl)-4,5-dihydro-5-oxo-1H-1,2,4-triazol-3-yl)-2-fluorobenzyl)-2,2,2-trifluoroacetamide



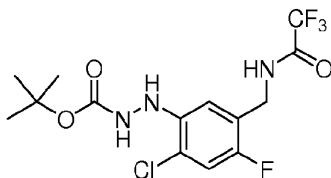
The title compound was prepared according to the procedure described in Example-83 by using 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzoyl isocyanate (Intermediate-69, 0.300 g), *tert*-butyl 2-(3-chloro-4-methylphenyl)hydrazinecarboxylate (Intermediate-87, 0.300 g), DCM (20 mL) and trifluoro acetic acid (5.0 mL) to afford 0.200 g of the desired product.

Step-2:- Preparation of 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(3-chloro-4-fluorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one

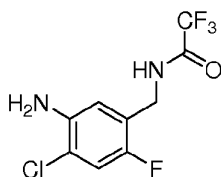
The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-5-(1-(3-chloro-4-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)-2,2,2-trifluoroacetamide (0.300 g), KOH (0.100 g), water (2 mL) and THF (10.0 mL) to afford 0.200 g of the desired product.

Intermediate-89

tert-Butyl 2-(2-chloro-4-fluoro-5-{[(trifluoroacetyl)amino]methyl}phenyl)hydrazinecarboxylate

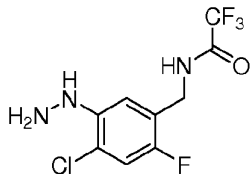


Step-1:- Preparation of *N*-(5-amino-4-chloro-2-fluorobenzyl)-2,2,2-trifluoroacetamide



The title compound was prepared according to the procedure described in step-1 of Intermediate-57 by using 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzoic acid (3.0 g), conc.H₂SO₄ and sodium azide (0.793 g, 0.012 mmol) to afford 2.1 g of the desired product.

Step-2:-Preparation of *N*-(4-chloro-2-fluoro-5-hydrazinylbenzyl)-2,2,2-trifluoroacetamide



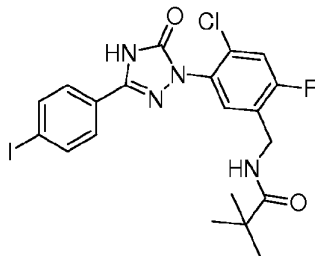
The title compound was prepared according to the procedure described in step-1 of Intermediate-61 by using *N*-(5-amino-4-chloro-2-fluorobenzyl)-2,2,2-trifluoroacetamide (2.0 g, 0.007 mmol), NaNO_2 (0.612 g, 0.008 mmol), $\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$ (4.10 g, 0.018 mmol), and conc. HCl (100 mL) to afford 1.0 g of the desired product.

Step-3:- Preparation of *tert*-butyl 2-(2-chloro-4-fluoro-5-[(trifluoroacetyl)amino]methyl) phenyl)hydrazinecarboxylate

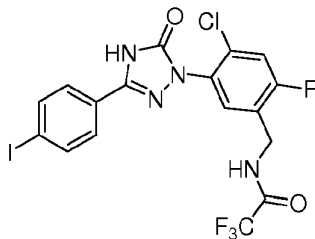
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using *N*-(4-chloro-2-fluoro-5-hydrazinylbenzyl)-2,2,2-trifluoroacetamide (1.00 g, 3.5 mmol), BOC anhydride (1.14 g, 5.26 mmol), Na_2CO_3 (0.743 g, 7.01 mmol), acetonitrile (20 mL) and water (10 mL) to afford 0.600 g of the desired product.

Intermediate-90

N-(4-Chloro-2-fluoro-5-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide

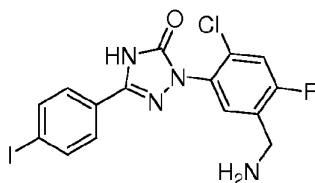


Step-1:- Preparation of *N*-(4-chloro-2-fluoro-5-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)-2,2,2-trifluoroacetamide



The title compound was prepared according to the procedure described in Example-83 by using 4-iodobenzoyl isocyanate (Intermediate-67, 0.700 g, 1.8 mmol), *tert*-butyl 2-(2-chloro-4-fluoro-5-[(trifluoroacetyl)amino]methyl)phenyl)hydrazine carboxylate (Intermediate-89, 0.992 g, 3.63 mmol), DCM (50 mL), trifluoro acetic acid (5.0 mL) to afford 0.600 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 4.47 (s, 2H), 7.61 (d, J = 2.1 Hz, 3H), 7.86- 7.90 (br d, 3H), 10.05 (br s, 1H).

Step-2:- Preparation of 2-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-5-(4-iodophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one



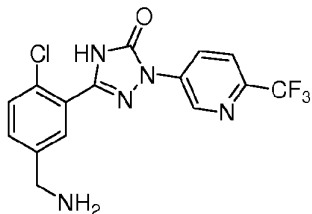
The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-2-fluoro-5-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)-2,2,2-trifluoroacetamide (0.550 g, 1.01 mmol), KOH (0.285 g, 5.09 mmol), water (5 mL) and THF (15.0 mL) to afford 0.400 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.59 (m, 2H), 3.71 (s, 1H), 7.38 (d, J = 9.6 Hz, 1H), 7.54 (d, J = 7.8 Hz, 1H), 7.60- 7.69 (m, 4H).

Step-3:- Preparation of *N*-(4-chloro-2-fluoro-5-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide

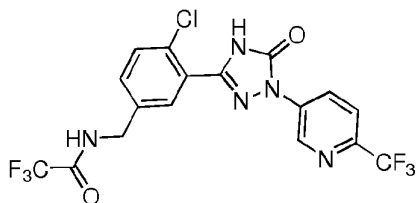
The title compound was prepared according to the procedure described in step-2 of Intermediate-83 by using 2-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-5-(4-iodophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (0.400 g, 0.900 mmol), pivaloyl chloride (0.2 mL, 1.35 mmol), DIPEA (2 mL), and THF (10 mL) to afford 0.300 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.10 (s, 9H), 4.30 (d, J = 5.4 Hz, 2H), 7.42- 7.44 (m, 1H), 7.62- 7.66 (m, 3H), 7.88- 7.91 (br d, 1H), 8.14 (m, 1H); MS (m/z): 529.42 (M^+H^+).

Intermediate-91

5-(5-(Aminomethyl)-2-chlorophenyl)-2-(6-(trifluoromethyl)pyridin-3-yl)-2*H*-1,2,4-triazol-3(4*H*)-one



Step-1:- Preparation *N*-(4-chloro-3-(1-(6-(trifluoromethyl)pyridin-3-yl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



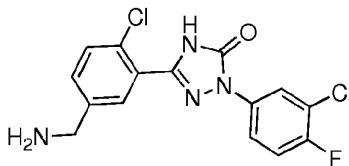
The title compound was prepared according to the procedure described in Example-83 by using 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate (step-3 of Intermediate-26, 0.500 g, 1.55 mmol), *tert*-butyl 2-[6-(trifluoromethyl)pyridin-3-yl]hydrazinecarboxylate (Intermediate-61, 0.431 g, 1.55 mmol), DCM (50 mL) and trifluoro acetic acid (5.0 mL) to afford 0.500 g of the desired product.

Step-2:- Preparation of 5-(5-(aminomethyl)-2-chlorophenyl)-2-(6-(trifluoromethyl)pyridin-3-yl)-2*H*-1,2,4-triazol-3(4*H*)-one

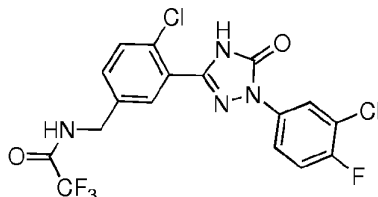
The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-3-(1-(6-(trifluoromethyl)pyridin-3-yl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (0.400 g), KOH (0.400 g), THF (5.0 mL), and water (5 mL) to afford 0.350 g of the desired product.

Intermediate-92

5-(5-(Aminomethyl)-2-chlorophenyl)-2-(3-chloro-4-fluorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one



Step-1:- Preparation of *N*-(4-chloro-3-(1-(3-chloro-4-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



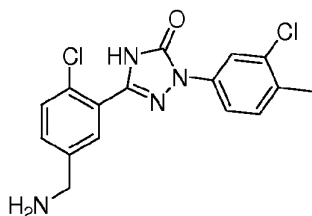
The title compound was prepared according to the procedure described in Example-83 by using 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate (step-3 of Intermediate-26, 0.500 g, 1.55 mmol), *tert*-butyl 2-(3-chloro-4-fluorophenyl)hydrazinecarboxylate (Intermediate-85, 0.455 g, 1.55 mmol), DCM (20 mL) and trifluoro acetic acid (5.0 mL) to afford 0.200 g of the desired product.

Step-2:- Preparation of 5-(5-(aminomethyl)-2-chlorophenyl)-2-(3-chloro-4-fluorophenyl)-2H-1,2,4-triazol-3(4H)-one

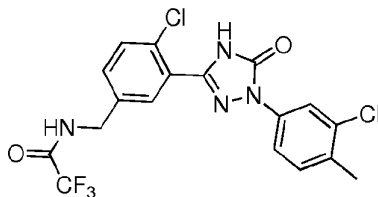
The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-3-(1-(3-chloro-4-fluorophenyl)-4,5-dihydro-5-oxo-1H-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (0.300 g), KOH (0.300 g), water (5 mL) and THF (10.0 mL) to afford 0.150 g of the desired product.

Intermediate-93

5-(5-(Aminomethyl)-2-chlorophenyl)-2-(3-chloro-4-methylphenyl)-2H-1,2,4-triazol-3(4H)-one



Step-1:- Preparation of *N*-(4-chloro-3-(1-(3-chloro-4-methylphenyl)-4,5-dihydro-5-oxo-1H-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



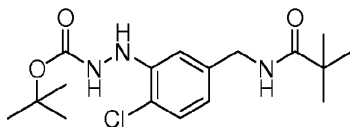
The title compound was prepared according to the procedure described in Example-83 by using 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate (step-3 of Intermediate-26, 0.500 g, 1.55 mmol), *tert*-butyl 2-(3-chloro-4-methylphenyl)hydrazinecarboxylate (Intermediate-87, 0.450 g, 1.55 mmol), DCM (20 mL) and trifluoro acetic acid (5.0 mL) to afford 0.180 g of the desired product.

Step-2:- Preparation of 5-(5-(aminomethyl)-2-chlorophenyl)-2-(3-chloro-4-methylphenyl)-2*H*-1,2,4-triazol-3(4*H*)-one

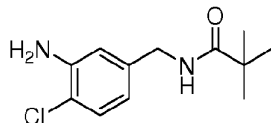
The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-3-(1-(3-chloro-4-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (0.400 g), KOH (0.400 g), water (5 mL) and THF (10.0 mL) to afford 0.350 g of the desired product.

Intermediate-94

tert-Butyl 2-(2-chloro-5-{[(2,2-dimethylpropanoyl)amino]methyl}phenyl)hydrazinecarboxylate

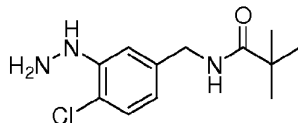


Step-1:- Preparation of *N*-(3-amino-4-chlorobenzyl)pivalamide



The title compound was prepared according to the procedure described in step-2 of Intermediate-83 by using 5-(aminomethyl)-2-chlorobenzenamine (2.0 g), pivaloyl chloride (0.2 mL), TEA (2 mL) and THF(10 mL) to afford 2.0 g of the desired product.

Step-2:- Preparation of *N*-(4-chloro-3-hydrazinylbenzyl)-2,2-dimethylpropanamide



The title compound was prepared according to the procedure described in step-1 of Intermediate-61 by using *N*-(3-amino-4-chlorobenzyl)pivalamide (1.80 g), NaNO₂ (0.62

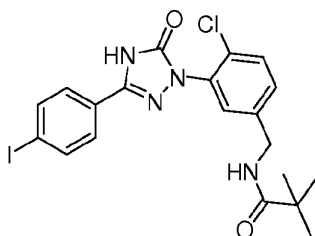
g, 0.009 mmol), $\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$ (4.2 g, 0.018 mmol) and conc. HCl (30 mL) to afford 1.5 g of the desired product.

Step-3:- Preparation of *tert*-butyl 2-(2-chloro-5-{[(2,2-dimethylpropanoyl)amino]methyl}phenyl)hydrazinecarboxylate

The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using *N*-(4-chloro-3-hydrazinylbenzyl)-2,2-dimethylpropanamide (1.5 g, 0.005 mmol), BOC anhydride (1.4 g, 0.006 mmol), Na_2CO_3 (0.922 g, 0.008 mmol), acetonitrile (30 mL) and water (5 mL) to afford 1.0 g of the desired product.

Intermediate-95

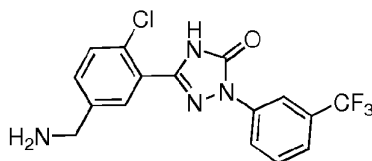
N-(4-Chloro-3-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl) pivalamide



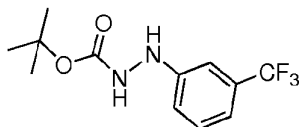
To a solution of *tert*-butyl 2-(2-chloro-5-{[(2,2-dimethylpropanoyl)amino]methyl}phenyl)hydrazine carboxylate (Intermediate-94, 0.450 g, 1.55 mmol) in DCM (20 mL) was added solution of 4-iodobenzoyl isocyanate (Intermediate-67, 0.500 g, 1.55 mmol) in DCM (10 mL) and the reaction mixture was stirred for 20 h at room temperature, followed by addition of trifluoro acetic acid (TFA, 3 mL) and further stirred for 20 h at same temperature. The reaction mass was quenched in water, extracted with DCM and concentrated to afford crude product which was purified by column chromatography eluting with MeOH: DCM to afford 0.180 g of the desired product.

Intermediate-96

5-(5-(Aminomethyl)-2-chlorophenyl)-2-(3-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one

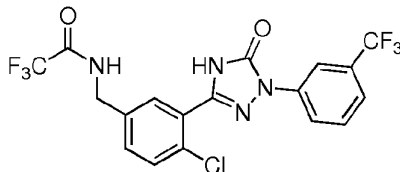


Step1:- Preparation of *tert*-butyl 2-[3-(trifluoromethyl)phenyl]hydrazinecarboxylate



The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using [3-(trifluoromethyl)phenyl]hydrazine hydrochloride (1.00 g, 0.0056 mol), BOC anhydride (1.4 g, 0.0067 mol), Na_2CO_3 (0.800 g, 0.008 mol), acetonitrile (10 mL) and water (2 mL) to afford 0.500 g of the desired product.

Step2:- Preparation of *N*-(4-chloro-3-(1-(3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



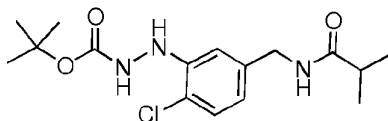
The title compound was prepared according to the procedure described in Example-83 by using 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate (step-3 of Intermediate-26, 0.500 g, 1.55 mmol), *tert*-butyl 2-[3-(trifluoromethyl)phenyl]hydrazinecarboxylate (0.428 g, 1.55 mmol), DCM (20 mL) and trifluoro acetic acid (5.0 mL) to afford 0.400 g of the desired product.

Step-3:- Preparation of 5-(5-(aminomethyl)-2-chlorophenyl)-2-(3-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one

The title compound was prepared according to the procedure described in Intermediate-66 by using *N*-(4-chloro-3-(1-(3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (0.400 g), KOH (0.200 g), water (2 mL) and THF (5.0 mL) to afford 0.200 g of the desired product.

Intermediate-97

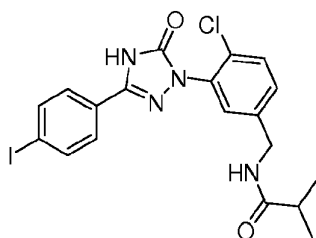
tert-Butyl 2-(2-chloro-5-[(2-methylpropanoyl)amino]methyl}phenyl)hydrazinecarboxylate



The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using *N*-(4-chloro-3-hydrazinylbenzyl)-2-methylpropanamide (2.5 g, 0.010 mol), BOC anhydride (2.5 g, 0.011 mol), Na₂CO₃ (1.6 g, 0.015 mol), acetonitrile (20 mL) and water (5 mL) to afford 1.500 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.03 (d, *J* = 6.6 Hz, 6H), 1.41 (s, 9H), 2.36-2.40 (m, 1H), 4.09 (d, *J* = 6.3 Hz, 1H), 4.15 (d, *J* = 5.4 Hz, 2H), 7.18 (d, *J* = 7.8 Hz, 2H), 7.32 (s, 1H), 8.24 (m, 2H); MS (*m/z*): 340.8 (M-H)⁺.

Intermediate-98

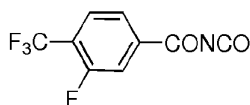
N-(4-Chloro-3-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide



The title compound was prepared according to the procedure described in Example-83 by using 4-iodobenzoyl isocyanate (Intermediate-67, 1.500 g), *tert*-butyl 2-(2-chloro-5-[(2-methylpropanoyl)amino]methyl}phenyl)hydrazinecarboxylate (Intermediate-97, 1.0 g), DCM (20 mL), and trifluoro acetic acid (5.0 mL) to afford 0.500 g of the desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 1.02 (d, *J* = 6.9 Hz, 6H), 2.44 (m, 1H), 4.28 (d, *J* = 8.7 Hz, 2H), 7.28 (br s, 1H), 7.35 (d, *J* = 7.8 Hz, 1H), 7.44 (s, 1H), 7.61 (d, *J* = 8.7 Hz, 1H), 7.91 (d, *J* = 8.1 Hz, 2H), 8.38 (m, 1H), 12.61 (m, 1H); MS (*m/z*): 495.7 (M-H)⁺.

Intermediate-99

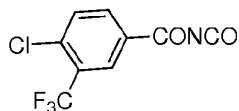
3-Fluoro-4-(trifluoromethyl)benzoyl isocyanate



The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 3-fluoro-4-(trifluoromethyl)benzamide (0.200 g), oxalyl chloride (0.025 mL) and EDC (15 mL) to afford 0.150 g of the desired product.

Intermediate-100

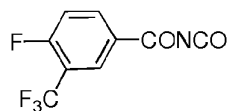
4-Chloro-3-(trifluoromethyl)benzoyl isocyanate



The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 4-chloro-3-(trifluoromethyl)benzamide (0.200 g), oxalyl chloride (0.012 mL) and EDC (15 mL) to afford 0.150 g of the desired product.

Intermediate-101

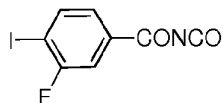
4-Fluoro-3-(trifluoromethyl)benzoyl isocyanate



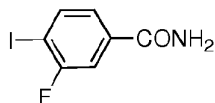
The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 4-fluoro-3-(trifluoromethyl)benzamide (0.200 g), oxalyl chloride (0.012 mL) and EDC (15 mL) to afford 0.150 g of the desired product.

Intermediate-102

3-Fluoro-4-iodobenzoyl isocyanate



Step-1:- Preparation of 3-fluoro-4-iodobenzamide



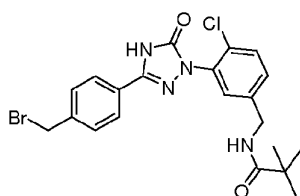
To a solution of 3-fluoro-4-iodobenzonitrile (2.7 g, 0.010 mmol) in DMSO (6.0 mL), K_2CO_3 (0.450 g, 0.003 mmol) and 30% H_2O_2 (2.4 mL) were added at 0-10°C and the reaction mass was stirred at RT for 2 h. After completion of the reaction, the reaction mass was quenched in ice cold water. The obtained solid product was filtered off to afford 2.0 g of the desired title product. 1H NMR (400 MHz, DMSO d_6): δ 7.48- 7.51 (m, 2H), 7.66 (br s, 1H), 7.69- 7.70 (m, 1H), 8.09 (brs, 1H).

Step-2:- Preparation of 3-fluoro-4-iodobenzoyl isocyanate

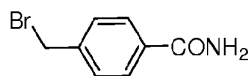
The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 3-fluoro-4-iodobenzamide (2.0 g), oxalyl chloride (0.120 mL) and EDC (25 mL) to afford 1.0 g of the desired product.

Intermediate-103

N-(3-(3-(4-(bromomethyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)-4-chlorobenzyl)pivalamide

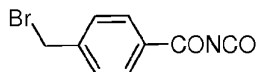


Step-1:-Preparation of 4-(bromomethyl)benzamide



Stirred the solution of 4-(bromomethyl)benzonitrile (1.5 g) in concn.H₂SO₄ at 100°C for 3-4 h. After completion of the reaction, the reaction mass was quenched in ice and filtered to afford 0.900 g of desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 4.71 (s, 2H), 7.26 (br s, 1H), 7.49 (d, *J* = 7.8 Hz, 2H), 7.82 (d, *J* = 7.8 Hz, 2H), 7.89 (br s, 1H); MS (*m/z*): 214.36 (*M*+H)⁺.

Step-2:-Preparation of 4-(bromomethyl)benzoyl isocyanate



The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 4-(bromomethyl)benzamide (0.200 g), oxalyl chloride (0.012 mL) and EDC (15 mL) to afford 0.150 g of the desired product.

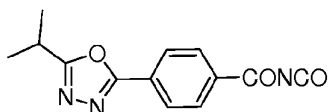
Step-3:- Preparation of *N*-(3-(3-(4-(bromomethyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)-4-chlorobenzyl)pivalamide

The title compound was prepared according to the procedure described in Example-83 by using 4-(bromomethyl)benzoyl isocyanate (0.150 g), *tert*-Butyl 2-(2-chloro-5-[(2,2-dimethylpropanoyl)amino]methyl)phenyl) hydrazinecarboxylate (Intermediate-94, 0.150 g), DCM (20 mL), and trifluoro acetic acid (5.0 mL) to afford 0.100 g of the desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 1.12 (s, 9H), 4.28 (br d, 2H), 4.75 (s, 2H),

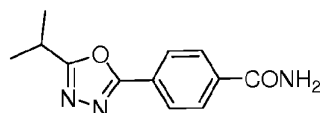
7.34 (br d, 2H), 7.43 (br s, 1H), 7.60 (br d, 2H), 7.81 (br d, 2H), 8.17 (br s, 1H), 12.57 (s, 1H).

Intermediate-104

4-(5-isopropyl-1,3,4-oxadiazol-2-yl)benzoyl isocyanate



Step1:- Preparation of 4-(5-isopropyl-1,3,4-oxadiazol-2-yl)benzamide



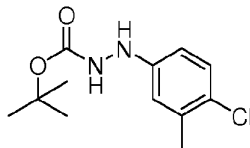
The title compound was prepared according to the procedure described in step1 of Intermediate-59 by using 4-(5-isopropyl-1,3,4-oxadiazol-2-yl)benzonitrile (0.500 g, 2.34 mmol), 50% H₂O₂ (5.0 mL), K₂CO₃ (0.097 g, 0.070 mmol) to afford 0.250 g of desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 1.3(s, 3H), 1.35 (s, 3H), 3.33(m, 1H), 7.56 (br s, 1H), 8.06 (s, 4H), 8.16 (br s, 1H); MS (m/z): 232.30 (M+H)⁺.

Step2:- Preparation of 4-(5-isopropyl-1,3,4-oxadiazol-2-yl)benzoyl isocyanate

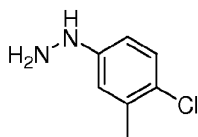
The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 4-(5-isopropyl-1,3,4-oxadiazol-2-yl)benzamide (0.200 g), oxalyl chloride (0.012 mL) and EDC (15 mL) to afford 0.150 g of the desired product.

Intermediate-105

tert-Butyl 2-(4-chloro-3-methylphenyl)hydrazinecarboxylate



Step-1:- Preparation of 1-(4-chloro-3-methylphenyl)hydrazine



The title compound was prepared according to the procedure described in step-1 of Intermediate-61 by using *N*-4-chloro-3-methylbenzenamine (2.0 g, 0.014 mmol), NaNO₂

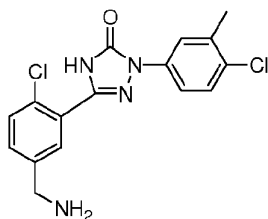
(01.16 g, 0.0169 mmol), $\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$ (7.93 g, 0.0352 mmol) and conc. HCl (30 mL) to afford 1.5 g of the desired product.

Step-2:- Preparation of *tert*-butyl 2-(4-chloro-3-methylphenyl)hydrazinecarboxylate

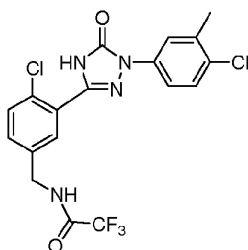
The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using 1-(4-chloro-3-methylphenyl)hydrazine (1.22 g, 0.011 mmol), BOC anhydride (2.0 g, 0.009 mmol), Na_2CO_3 (0.922 g, 0.008 mmol), acetonitrile (30 mL) and water (5 mL) to afford 1.0 g of the desired product. ^1H NMR (300 MHz, $\text{DMSO } d_6$): δ 1.40 (s, 9H), 2.21 (s, 3H), 6.49 (d, $J = 8.4$ Hz, 1H), 6.58 (s, 1H), 7.13 (d, $J = 8.7$ Hz, 1H), 7.65 (s, 1H), 8.78 (s, 1H).

Intermediate-106

5-(5-(aminomethyl)-2-chlorophenyl)-2-(4-chloro-3-methylphenyl)-2*H*-1,2,4-triazol-3(4*H*)-one



Step-1:-Preparation of *N*-(4-chloro-3-(1-(4-chloro-3-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



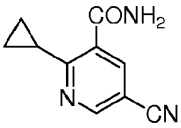
The title compound was prepared according to the procedure described in Example-83 by using 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate (step-3 of Intermediate-26, 1.0 g, 0.003 mmol), *tert*-butyl 2-(4-chloro-3-methylphenyl)hydrazinecarboxylate (Intermediate-105, 1.500 g), DCM (20 mL), and trifluoro acetic acid (5.0 mL) to afford 0.600 g of the desired product. ^1H NMR (300 MHz, $\text{DMSO } d_6$): δ 2.38 (s, 3H), 4.45 (d, $J = 5.4$ Hz, 2H), 7.53 (d, $J = 9.3$ Hz, 2H), 7.67

(s, 2H), 7.82 (d, $J = 8.1$ Hz, 1H), 7.92 (s, 1H), 10.08 (t, 1H), 12.61 (s, 1H); MS (m/z): 445.31 (M+H)⁺.

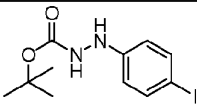
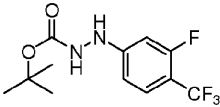
Step-2:- Preparation of 3-(5-(aminomethyl)-2-chlorophenyl)-1-(3-methoxy-4-nitrophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one

The solution of *N*-(4-chloro-3-(1-(4-chloro-3-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (0.550 g, 1.23 mmol) in 20% aq. KOH (20 mL) was stirred for 2-3 h at RT. Excess of solvent was removed under vacuum and filtered off remaining reaction mass to afford 0.400 g of desired product. MS (m/z): 349.34 (M)⁺.

Intermediate-107 was prepared according to the procedure described for step-2 of Intermediate-26 by using 5-cyano-2-cyclopropylpyridine-3-carboxylic acid, oxalyl chloride, ammonia gas, DMF, THF and DCM.

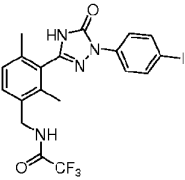
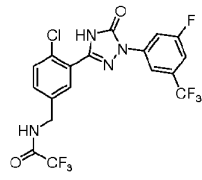
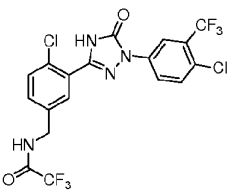
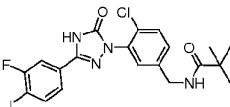
No. & Chemical Structure	Chemical name and Characterization data
Intermediate-107 	5-cyano-2-cyclopropylpyridine-3-carboxamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 107 (m, 4H), 2.49 (s, 1H), 7.82 (s, 1H), 8.13 (s, 1H), 8.20 (s, 2H), 8.87 (s, 1H); MS (m/z): 188.04 (M+H) ⁺ .

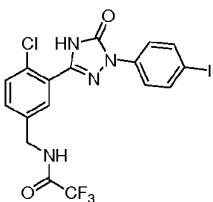
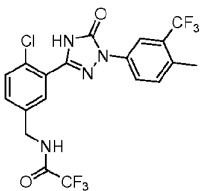
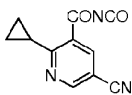
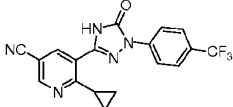
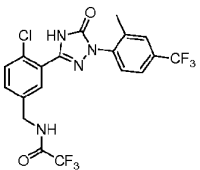
The Intermediate-108 to Intermediate-115 were prepared by following the procedure described for step-1 of Intermediate-61 by using corresponding amine, NaNO₂, SnCl₂.H₂O, 4N HCl and followed by following the procedure described for step-3 of Intermediate-7 using BOC anhydride, Na₂CO₃ in acetonitrile and water.

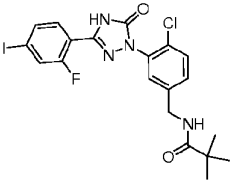
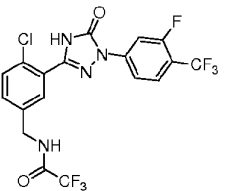
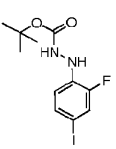
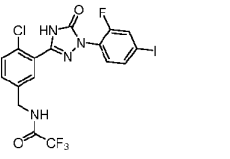
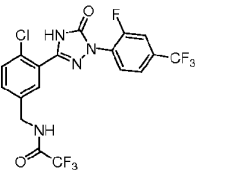
No.	Structure	Chemical name and Characterization data
Intermediate-108		<i>tert</i> -butyl 2-(4-iodophenyl)hydrazinecarboxylate. MS (m/z): 335.15 (M+H) ⁺ .
Intermediate-109		<i>tert</i> -butyl 2-[3-fluoro-4-(trifluoromethyl)phenyl]hydrazinecarboxylate. MS (m/z): 295.21 (M+H) ⁺ .

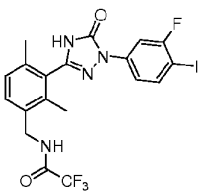
No.	Structure	Chemical name and Characterization data
Intermediate -110		<i>tert</i> -butyl 2-[3-fluoro-5-(trifluoromethyl)phenyl]hydrazinecarboxylate. MS (m/z): 295.24 (M+H) ⁺ .
Intermediate -111		<i>tert</i> -butyl 2-[4-chloro-3-(trifluoromethyl)phenyl]hydrazinecarboxylate. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.40 (s, 9H), 6.90 (d, <i>J</i> = 8.7 Hz, 1H), 7.01 (s, 1H), 7.45 (d, <i>J</i> = 9.0 Hz, 1H), 8.20 (s, 1H), 8.99 (br s, 1H).
Intermediate -112		<i>tert</i> -butyl 2-[4-methyl-3-(trifluoromethyl)phenyl]hydrazinecarboxylate. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.40 (s, 9H), 2.28 (s, 3H), 6.80 (d, <i>J</i> = 8.4 Hz, 1H), 6.91 (s, 1H), 7.18 (d, <i>J</i> = 8.4 Hz, 1H), 7.84 (s, 1H), 8.88 (br s, 1H).
Intermediate -113		<i>tert</i> -butyl 2-[2-methyl-4-(trifluoromethyl)phenyl]hydrazinecarboxylate. MS (m/z): 291.12 (M+H) ⁺ .
Intermediate -114		<i>tert</i> -butyl 2-[2-fluoro-4-(trifluoromethyl)phenyl]hydrazinecarboxylate. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.42 (s, 9H), 6.83 (t, 1H), 7.39 (d, <i>J</i> = 8.4 Hz, 1H), 7.44-7.48 (m, 1H), 8.28 (s, 1H), 9.03 (s, 1H).
Intermediate -115		<i>tert</i> -butyl 2-(3-fluoro-4-iodophenyl)hydrazinecarboxylate. MS (m/z): 351.14 (M-H) ⁻ .

The Intermediate-116 to Intermediate-128 was prepared by following the procedure described for Example-83 by using corresponding starting material mentioned in the table below, DCM and TFA.

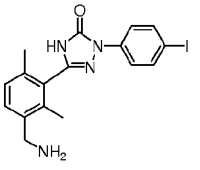
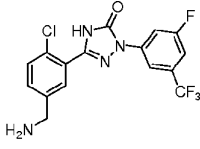
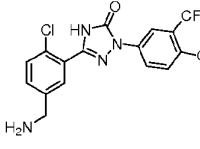
Starting material used	Intermediate No. and Structure	Intermediate chemical name and characterization data
Intermediate-65 + Intermediate-67	Intermediate-116 	<i>N</i> -(3-(4,5-dihydro-1-(4-iodophenyl)-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)-2,2,2-trifluoroacetamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 2.19 (s, 3H), 2.21 (s, 3H), 4.41 (d, <i>J</i> = 4.8 Hz, 2H), 7.20 (d, <i>J</i> = 7.8 Hz, 1H), 7.29 (d, <i>J</i> = 7.8 Hz, 1H), 7.53-7.82 (m, 4H), 9.97 (s, 1H), 12.23 (s, 1H); MS (<i>m/z</i>): 517.301 (M+H) ⁺ .
Step 3 product of Intermediate-26 + Intermediate-110	Intermediate-117 	<i>N</i> -(4-chloro-3-(1-(3-fluoro-5-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 4.46 (d, 2H), 7.51 (m, 1H), 7.60 (d, <i>J</i> = 9.3 Hz, 1H), 7.66-7.69 (m, 2H), 8.08 (d, <i>J</i> = 13.8 Hz, 1H), 8.16 (s, 1H), 10.09 (m, 1H), 12.84 (m, 1H)
Step-3 product of Intermediate-26 + Intermediate-111	Intermediate-118 	<i>N</i> -(4-chloro-3-(1-(4-chloro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 4.46 (d, <i>J</i> = 6.0 Hz, 2H), 7.49 (d, <i>J</i> = 10.2 Hz, 1H), 7.67 (d, <i>J</i> = 8.4 Hz, 2H), 7.85 (d, <i>J</i> = 9.0 Hz, 1H), 8.24 (d, <i>J</i> = 8.7 Hz, 1H), 8.44 (s, 1H), 10.09 (t, 1H), 12.80 (s, 1H); MS (<i>m/z</i>): 499.31 (M+H) ⁺ .
Intermediate-94 + Intermediate-102	Intermediate-119 	<i>N</i> -(4-chloro-3-(3-(3-fluoro-4-iodophenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.12 (s, 9H), 4.28 (d, <i>J</i> = 8.4 Hz, 2H), 7.34 (d, <i>J</i> = 7.8 Hz, 1H), 7.43-7.49 (m, 2H), 7.59-7.67 (m, 2H), 8.02 (t, <i>J</i> = 7.2 Hz, 1H), 8.19 (t, 1H), 12.67 (s, 1H); MS (<i>m/z</i>): 529.05 (M+H) ⁺ .

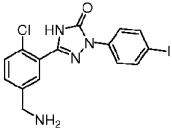
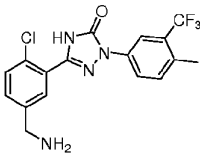
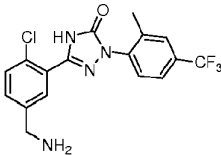
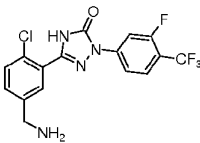
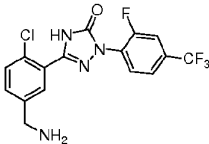
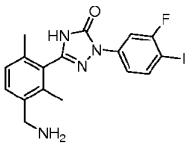
Starting material used	Intermediate No. and Structure	Intermediate chemical name and characterization data
Step-3 product of Intermediate-26 + Intermediate-67	Intermediate-120 	<i>N</i> -(4-chloro-3-(4,5-dihydro-1-(4-iodophenyl)-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 4.44 (br s, 2H), 7.46-7.73 (m, 4H), 7.80 (s, 2H), 10.11 (s, 1H); MS (m/z): 523.09 (M+H) ⁺ .
Step-3 product of Intermediate-26 + Intermediate-112	Intermediate-121 	<i>N</i> -(4-chloro-3-(1-(3-(trifluoromethyl)-4-methylphenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 2.45 (s, 3H), 4.46 (d, <i>J</i> = 6.6 Hz, 2H), 7.49 (d, <i>J</i> = 7.8 Hz, 1H), 7.57 (d, <i>J</i> = 7.8 Hz, 1H), 7.68 (s, 2H), 8.10 (d, <i>J</i> = 8.7 Hz, 1H), 8.28 (s, 1H), 10.09 (br s, 1H), 12.69 (br s, 1H); MS (m/z): 476.91 (M-H) ⁻ .
Intermediate-53 + 	Intermediate-122 	6-cyclopropyl-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)pyridine-3-carbonitrile. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.18 (m, 2H), 1.39 (m, 2H), 3.01 (m, 1H), 7.85 (d, <i>J</i> = 9.0 Hz, 2H), 8.20 (d, <i>J</i> = 7.8 Hz, 2H), 8.45 (s, 1H), 8.99 (s, 1H), 12.82 (br s, 1H); MS (m/z): 372.24 (M+H) ⁺ .
Step-3 product of Intermediate-26 + Intermediate-113	Intermediate-123 	<i>N</i> -(4-chloro-3-(1-(4-(trifluoromethyl)-2-methylphenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 2.39 (s, 3H), 4.44 (br d, 2H), 7.33-7.90 (m, 6H), 10.07 (br s, 1H), 12.51 (s, 1H); MS (m/z): 479.10 (M+H) ⁺ .
Intermediate-	Intermediate-124	<i>N</i> -(4-chloro-3-(3-(2-fluoro-4-iodophenyl)-4,5-

Starting material used	Intermediate No. and Structure	Intermediate chemical name and characterization data
47 + Intermediate-94		dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.11 (s, 9H), 4.28 (d, <i>J</i> = 5.7 Hz, 2H), 7.34 (d, <i>J</i> = 8.4 Hz, 1H), 7.42 (s, 1H), 7.53-7.61 (m, 2H), 7.75 (d, <i>J</i> = 7.2 Hz, 1H), 7.89 (d, <i>J</i> = 9.3 Hz, 1H), 8.18 (t, 1H), 12.43 (s, 1H); MS (<i>m/z</i>): 529.0110 (<i>M</i> +H) ⁺ .
Step-3 product of Intermediate-26 + Intermediate-109		<i>N</i> -(4-chloro-3-(1-(3-fluoro-4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 4.46 (d, <i>J</i> = 6.0 Hz, 2H), 7.50 (d, <i>J</i> = 8.7 Hz, 1H), 7.66 (s, 2H), 7.92-8.07 (m, 3H), 10.10 (t, 1H), 12.84 (m, 1H).
Step-3 product of Intermediate-26 + 		<i>N</i> -(4-chloro-3-(1-(2-fluoro-4-iodophenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 4.44 (d, <i>J</i> = 5.1 Hz, 2H), 7.39-7.48 (m, 2H), 7.63 (br s, 2H), 7.74 (t, <i>J</i> = 8.4 Hz, 1H), 7.90 (d, <i>J</i> = 8.7 Hz, 1H), 10.08 (br s, 1H), 12.52 (s, 1H); MS (<i>m/z</i>): 540.98 (<i>M</i> +H) ⁺ .
Step-3 product of Intermediate-26 + Intermediate-114		<i>N</i> -(4-chloro-3-(1-(2-fluoro-4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 4.44 (m, 2H), 7.34 (s, 1H), 7.49 (m, 2H), 7.75 (d, <i>J</i> = 7.8 Hz, 1H), 7.79-7.98 (m, 2H), 10.10 (br s, 1H), 12.64 (m, 1H); MS (<i>m/z</i>): 483.12 (<i>M</i> +H) ⁺ .

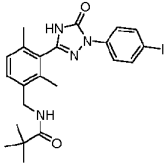
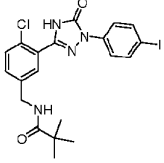
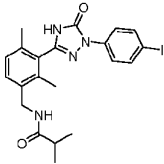
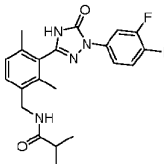
Starting material used	Intermediate No. and Structure	Intermediate chemical name and characterization data
Intermediate-65 + Intermediate-115	Intermediate-128 	<i>N</i> -(3-(1-(3-fluoro-4-iodophenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)-2,2,2-trifluoroacetamide. MS (m/z): 535.01 (M+H) ⁺ .

The Intermediate-129 to Intermediate-137 were prepared by following the procedure described in step-2 of Intermediate-106 by using corresponding starting material mentioned in the table below, KOH and water.

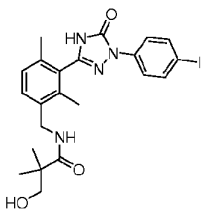
Starting material used	Intermediate No. and Structure	Intermediate chemical name and characterization data.
Intermediate-116	Intermediate-129 	5-(3-(aminomethyl)-2,6-dimethylphenyl)-2-(4-iodophenyl)-2 <i>H</i> -1,2,4-triazol-3(4 <i>H</i>)-one. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 2.10 (s, 3H), 2.12 (s, 3H), 3.13 (br s, 2H), 3.66 (s, 2H), 6.98 (d, <i>J</i> = 8.4 Hz, 1H), 7.21 (d, <i>J</i> = 8.1 Hz, 1H), 7.57 (d, <i>J</i> = 8.1 Hz, 2H), 7.90 (d, <i>J</i> = 8.7 Hz, 2H).
Intermediate-117	Intermediate-130 	5-(5-(aminomethyl)-2-chlorophenyl)-2-(3-fluoro-5-(trifluoromethyl)phenyl)-2 <i>H</i> -1,2,4-triazol-3(4 <i>H</i>)-one
Intermediate-118	Intermediate-131 	5-(5-(aminomethyl)-2-chlorophenyl)-2-(4-chloro-3-(trifluoromethyl)phenyl)-2 <i>H</i> -1,2,4-triazol-3(4 <i>H</i>)-one
Intermediate-120	Intermediate-132	5-(5-(aminomethyl)-2-chlorophenyl)-2-(4-iodophenyl)-2 <i>H</i> -1,2,4-triazol-3(4 <i>H</i>)-one. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 3.36 (br s, 2H), 3.80 (br s,

Starting material used	Intermediate No. and Structure	Intermediate chemical name and characterization data.
		2H), 7.41 (m, 1H), 7.47 (m, 1H), 7.6-7.72 (m, 2H), 7.77 (s, 1H), 7.87-7.90 (m, 2H).
Intermediate-121		5-(5-(aminomethyl)-2-chlorophenyl)-2-(3-(trifluoromethyl)-4-methylphenyl)-2H-1,2,4-triazol-3(4H)-one. MS (m/z): 381.01 (M-H) ⁻ .
Intermediate-123		5-(5-(aminomethyl)-2-chlorophenyl)-2-(4-(trifluoromethyl)-2-methylphenyl)-2H-1,2,4-triazol-3(4H)-one. MS (m/z): 383.15 (M+H) ⁺ .
Intermediate-125		5-(5-(aminomethyl)-2-chlorophenyl)-2-(3-fluoro-4-(trifluoromethyl)phenyl)-2H-1,2,4-triazol-3(4H)-one. MS (m/z): 387.21 (M+H) ⁺ .
Intermediate-127		5-(5-(aminomethyl)-2-chlorophenyl)-2-(2-fluoro-4-(trifluoromethyl)phenyl)-2H-1,2,4-triazol-3(4H)-one. MS (m/z): 387.06 (M+H) ⁺ .
Intermediate-128		5-(3-(aminomethyl)-2,6-dimethylphenyl)-2-(3-fluoro-4-iodophenyl)-2H-1,2,4-triazol-3(4H)-one. MS (m/z): 439.24 (M+H) ⁺ .

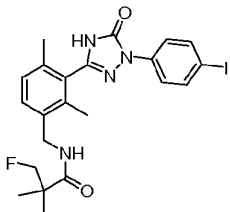
The Intermediate-138 to Intermediate-141 were prepared by following the procedure described in Example-108 by using corresponding starting material mentioned in the table below, TEA and THF.

Starting material used	Intermediate No. and Structure	Intermediate chemical name and characterization data
Intermediate-129 + Pivaloyl chloride	Intermediate-138 	<i>N</i> -(3-(4,5-dihydro-1-(4-iodophenyl)-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.19 (s, 9H), 2.15 (s, 3H), 2.19 (s, 3H), 4.22 (d, 2H), 7.17-7.21 (m, 2H), 7.79 (m, 4H), 8.00 (m, 1H), 12.00 (s, 1H).
Intermediate-132 + Pivaloyl chloride	Intermediate-139 	<i>N</i> -(4-chloro-3-(4,5-dihydro-1-(4-iodophenyl)-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.13 (s, 9H), 4.29 (d, <i>J</i> = 5.4 Hz, 2H), 7.41 (d, <i>J</i> = 8.4 Hz, 1H), 7.59 (s, 2H), 7.80 (q, <i>J</i> = 6.3 Hz, 4H), 8.19 (m, 1H), 12.59 (s, 1H); MS (<i>m/z</i>): 511.16 (<i>M</i> + <i>H</i>) ⁺ .
Intermediate-129 + Isopropyl chloride	Intermediate-140 	<i>N</i> -(3-(4,5-dihydro-1-(4-iodophenyl)-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)isobutyramide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.23 (s, 3H), 1.35 (s, 3H), 2.06 (s, 3H), 2.20 (s, 3H), 4.23 (br s, 2H), 7.08 (d, <i>J</i> = 7.8 Hz, 1H), 7.20 (d, <i>J</i> = 7.5 Hz, 1H), 7.74 (d, <i>J</i> = 8.7 Hz, 2H), 7.85 (d, <i>J</i> = 8.1 Hz, 2H), 8.17 (t, 1H).
Intermediate-137 + Isopropyl chloride	Intermediate-141 	<i>N</i> -(3-(1-(3-fluoro-4-iodophenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)isobutyramide. MS (<i>m/z</i>): 509.02 (<i>M</i> + <i>H</i>) ⁺ .

The Intermediate-142 was prepared by following the procedure described in Example-107 by using Intermediate-129, THF, DMF, 3-methoxy-2,2-dimethylpropanoic acid, TBTU and TEA.

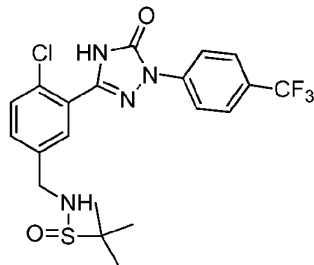
Starting material used	Intermediate No. and Structure	Intermediate chemical name and characterization data
Intermediate-129	Intermediate-142 	<i>N</i> -(3-(4,5-dihydro-1-(4-iodophenyl)-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)-3-hydroxy-2,2-dimethylpropanamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.08 (m, 6H), 2.15 (s, 3H), 2.19 (s, 3H), 3.41 (m, 2H), 4.25 (d, 2H), 4.92 (m, 1H), 6.93 (d, <i>J</i> = 6.9 Hz, 1H), 7.15 (d, <i>J</i> = 8.7 Hz, 1H), 7.28 (d, <i>J</i> = 8.1 Hz, 1H), 7.79 (s, 3H), 7.92 (t, 1H), 12.25 (s, 1H); MS (<i>m/z</i>): 521.18 (M+H) ⁺ .

The Intermediate-143 was prepared by following the procedure described in step-2 of Example-134 by using Intermediate-142, DAST and THF.

Starting material used	Intermediate No and Structure	Intermediate chemical name and characterization data
Intermediate-142	Intermediate-143 	<i>N</i> -(3-(4,5-dihydro-1-(4-iodophenyl)-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)-3-fluoro-2,2-dimethylpropanamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.08 (m, 6H), 2.15 (s, 3H), 2.19 (s, 3H), 3.34 (m, 2H), 4.25 (d, 2H), 6.91 (m, 1H), 7.15 (m, 1H), 7.25-7.27 (m, 1H), 7.79 (s, 3H), 8.08 (s, 1H), 12.24 (s, 1H); MS (<i>m/z</i>): 521.03 (M+H) ⁺ .

Intermediate-144

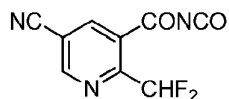
N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2-methylpropane-2-sulfonamide



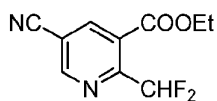
The title compound was prepared according to the procedure described in Example-108 by using 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-63, 0.450 g, 1.22 mmol), 2-methylpropane-2-sulfinic chloride (0.205 g, 1.46 mmol), TEA (2.0 mL), DCM (10 mL) to afford 0.250 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.23 (s, 9H), 4.24 (m, 2H), 7.63-7.87 (m, 4H), 7.86 (d, *J* = 8.7 Hz, 2H), 8.19 (d, *J* = 9.0 Hz, 2H); MS (*m/z*): 472.095 (M+H)⁺.

Intermediate-145

5-Cyano-2-(difluoromethyl)nicotinoyl isocyanate

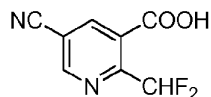


Step-1 :- Preparation of ethyl 5-cyano-2-(difluoromethyl)nicotinate



The title compound was prepared according to the procedure described in step-2 of Example-134 by using ethyl 5-cyano-2-formylnicotinate (6.0 g, 0.029 mmol), DAST (11.8 mL, 0.088 mmol), DCM (200 mL) to afford 3.5 g of desired product. MS (*m/z*): 227.06 (M+H)⁺.

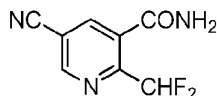
Step-2 :-Preparation of 5-cyano-2-(difluoromethyl)nicotinic acid



To a solution of ethyl 5-cyano-2-(difluoromethyl)nicotinate (1.2 g, 5.3 mmol) in THF : water (20 mL : 5 mL) was added LiOH.H₂O (0.267 g, 6.3 mmol) at 0°C and the reaction mass was stirred for 40-60 minutes. The reaction mass was acidified with dil. HCl and extracted with ethyl acetate. The organic layer was separated, dried over anhydrous

sodium sulphate and concentrated to afford 0.700 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.37-7.72 (t, J = 53.7 Hz, 1H), 8.24 (s, 1H), 9.30 (s, 1H), 14.25 (br s, 1H); MS (m/z): 199.34 (M+H) $^+$.

Step-3 :-Preparation of 5-cyano-2-(difluoromethyl)nicotinamide



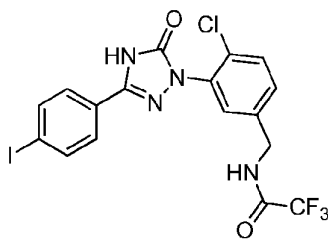
To a solution of 5-cyano-2-(difluoromethyl)nicotinic acid (0.600 g, 3.0 mmol) in dry DMF (20 mL), TBTU (10.0 g, 3.33 mmol) was added at 0°C followed by slow addition of DIPEA (0.7 mL, 3.93 mmol). The reaction mass was stirred at 0°C for 1 h and ammonium chloride (3.33 g, 60.7 mmol) was added. The reaction mass was stirred at RT for 30-48 h. After completion of reaction quenched the reaction mass with DCM, filtered off solid to afford 0.500 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 8.16 (m, 1H), 8.63 (s, 1H), 9.22 (s, 1H).

Step-4:- Preparation of 5-cyano-2-(difluoromethyl)nicotinoyl isocyanate

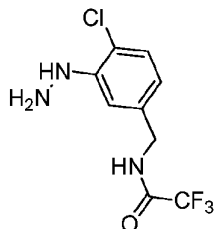
The title compound was prepared according to the procedure described in step-2 of Intermediate-8 by using 5-cyano-2-(difluoromethyl)nicotinamide (1.0 g), oxalyl chloride (3.2 g, 25.3 mmol) and DCM (20 mL) to afford 1.0 g of the desired product.

Intermediate-146

N-(4-Chloro-3-(3-(4-iodophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)benzyl)-2,2,2-trifluoroacetamide

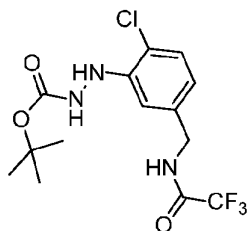


Step-1:- Preparation *N*-(4-chloro-3-hydrazinylbenzyl)-2,2,2-trifluoroacetamide



To a cold solution of *N*-(3-amino-4-chlorobenzyl)-2,2,2-trifluoroacetamide (0.500 g, 0.017 mmol) in 6N HCl (50.0 mL), aqueous solution of NaNO₂ (1.29 g, 0.018 mmol) was added at 0°C and the reaction mass was stirred at 0-5°C for 30 min. The reaction mass was added to a solution of SnCl₂·H₂O (22.18 g, 0.98 mmol) in 6N HCl at 0-5°C and further continued stirring for 5-6 h at same temperature. The reaction mass was cooled, basified and extracted with DCM. The organic layer was separated, dried and concentrated to afford 0.700 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 4.22 (br s, 2H), 4.29 (d, *J* = 6.0 Hz, 2H), 6.42-6.67 (m, 2H), 7.14-7.16 (m, 2H), 9.98 (br s, 1H).

Step-2:- Preparation *tert*-butyl 2-(2-chloro-5-((2,2,2-trifluoroacetamido)methyl)phenyl)hydrazinecarboxylate



The title compound was prepared according to the procedure described in step-3 of Intermediate-7 by using *N*-(4-chloro-3-hydrazinylbenzyl)-2,2,2-trifluoroacetamide (0.350 g, 1.3 mmol), BOC anhydride (0.316 g, 1.4 mmol), Na₂CO₃ (1.19 g, 1.9 mmol), acetonitrile (20 mL) and water (10 mL) to afford 0.500 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.41 (s, 9H), 4.28 (d, *J* = 5.7 Hz, 2H), 6.61-6.65 (m, 2H), 7.22 (d, *J* = 8.1 Hz, 1H), 7.41 (s, 1H), 9.91 (s, 1H), 9.97 (s, 1H).

Step-3:- Preparation of *N*-(4-chloro-3-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide

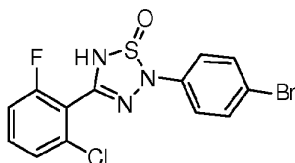
The title compound was prepared according to the procedure described in Example-83 by using 4-iodobenzoyl isocyanate (Intermediate-67, 1.500 g), *tert*-butyl 2-(2-chloro-5-

((2,2,2-trifluoroacetamido)methyl)phenyl) hydrazinecarboxylate (1.0 g), DCM (20 mL), and trifluoro acetic acid (5.0 mL) to afford 0.500 g of the desired product. ^1H NMR (400 MHz, DMSO d_6): δ 4.44 (d, J = 6.0 Hz, 2H), 7.41 (d, J = 7.8 Hz, 1H), 7.52 (s, 1H), 7.60-7.67 (m, 2H), 7.81 (d, J = 8.4 Hz, 1H), 7.90 (d, J = 8.7 Hz, 2H), 10.09 (br s, 1H), 12.63 (s, 1H).

Examples

Example-1

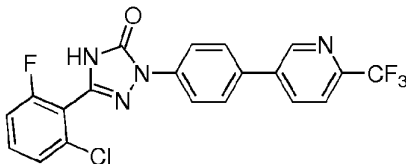
2-(4-Bromophenyl)-4-(2-chloro-6-fluorophenyl)-2,5-dihydro-1,2,3,5-thiadiazole-1-oxide



The title compound was prepared according to the procedure described in step-4 of Intermediate-1 using *N'*-(4-bromophenyl)-2-chloro-6-fluorobenzenecarbo hydrazonamide (step-3 of Intermediate-1, 0.100 g, 0.292 mmol), CHCl_3 (10 mL), pyridine (2.0 mL) and thionyl chloride (1.0 mL). The obtained product was purified with column chromatography on silica gel eluting with 0.5 % MeOH: DCM to afford 0.050 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.45 (t, J = 6.9 Hz, 3H), 7.51-7.68 (m, 4H), 12.05 (br s, 1H); MS (m/z): 388.02 (M^+).

Example-2

5-(2-Chloro-6-fluorophenyl)-2-{4-[6-(trifluoromethyl)pyridin-3-yl]phenyl}-2,4-dihydro-3*H*-1,2,4-triazol-3-one

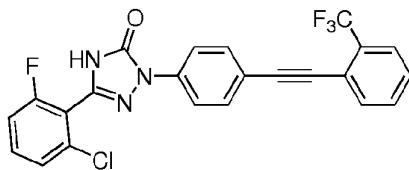


To a solution of 2-(4-bromophenyl)-5-(2-chloro-6-fluorophenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one (Intermediate-1, 0.100 g, 0.271 mmol) in DMSO (3.0 mL) was added [6-(trifluoromethyl)pyridin-3-yl]boronic acid (0.078 g, 0.40 mmol), K_2CO_3 (0.112 g, 0.81

mmol) and tetrakis(triphenyl phosphine) palladium (0) (0.062 g, 0.054 mmol). The reaction mass was stirred at 110°C for 24-48 h. The reaction mass was quenched in water and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated. The obtained product was purified with column chromatography on silica gel eluting with 4-5 % EA: DCM to afford 0.030 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 7.38-7.48 (m, 1H), 7.61-7.63 (m, 1H), 7.88-7.99 (m, 4H), 8.01-8.17 (m, 2H), 8.41 (d, *J* = 7.8 Hz, 1H), 9.14 (s, 1H), 12.60 (br s, 1H). MS (*m/z*): 435.25 (*M*+H)⁺.

Example-3

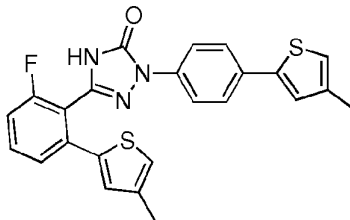
5-(2-Chloro-6-fluorophenyl)-2-(4-{[2-(trifluoromethyl)phenyl]ethynyl}phenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one



To a solution of 5-(2-chloro-6-fluorophenyl)-2-(4-ethynylphenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one (Intermediate-2, 0.100 g, 0.319 mmol) in DMSO was added 1-iodo-2-(trifluoromethyl)benzene (0.13 g, 0.479 mmol), TBAF (0.301 g, 0.958 mmol) and bis(triphenylphosphine)palladium(II) chloride (0.050 g, 0.071 mmol). The reaction mass was stirred at 110°C for 5-6 h. The reaction mass was quenched in water and extracted with DCM and concentrated. The obtained product was purified with column chromatography on silica gel eluting with 2.0 % EA: DCM to afford 0.040 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 7.50 (t, *J* = 9.3 Hz, 1H), 7.57-7.73 (m, 6H), 7.82 (br s, 2H), 8.02 (d, *J* = 8.7 Hz, 2H), 12.72 (br s, 1H). MS (*m/z*): 458.26 (*M*+H)⁺.

Example-4

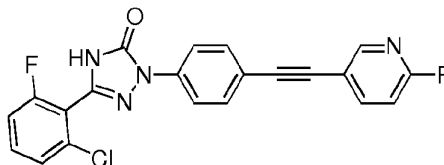
5-[2-Fluoro-6-(4-methylthiophen-2-yl)phenyl]-2-[4-(4-methylthiophen-2-yl)phenyl]-2,4-dihydro-3*H*-1,2,4-triazol-3-one



The title compound was prepared according to the procedure described in Example-2 using 2-(4-bromophenyl)-5-(2-chloro-6-fluorophenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one (Intermediate-1, 0.100 g, 0.271 mmol), (4-methylthiophen-2-yl)boronic acid (0.059 g, 0.407 mmol), K_2CO_3 (0.112 g, 0.81 mmol), tetrakis(triphenyl phosphine) palladium (0) (0.062 g, 0.054 mmol) and DMSO (3.0 mL) to afford 0.020 g of desired product. 1H NMR (300 MHz, $CDCl_3$): δ 2.29 (s, 6H), 6.86 (s, 1H), 6.91 (s, 2H), 7.13 (s, 1H), 7.21 (d, $J = 9.3$ Hz, 1H), 7.39 (d, $J = 7.8$ Hz, 1H), 7.48-7.55 (m, 1H), 7.61 (d, $J = 8.1$ Hz, 2H), 7.92 (d, $J = 9.0$ Hz, 2H), 9.27 (br s, 1H). MS (m/z): 384.88 (M-H) $^-$.

Example-5

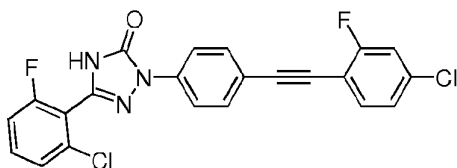
5-(2-Chloro-6-fluorophenyl)-2-{4-[(6-fluoropyridin-3-yl)ethynyl]phenyl}-2,4-dihydro-3H-1,2,4-triazol-3-one



The title compound was prepared according to the procedure described in Example-3 using 5-(2-chloro-6-fluorophenyl)-2-(4-ethynylphenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one (Intermediate-2, 0.100 g, 0.319 mmol), 2-fluoro-5-iodopyridine (0.106 g, 0.479 mmol), TBAF (0.201 g, 0.638 mmol), bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.028 mmol) and DMSO (3.0 mL). The obtained product was purified with column chromatography on silica gel eluting with 2.0 % MeOH: DCM to afford 0.030 g of the desired product. 1H NMR (300 MHz, $DMSO-d_6$): δ 7.30 (d, $J = 7.5$ Hz, 1H), 7.51 (t, $J = 8.7$ Hz, 1H), 7.58 (d, $J = 8.4$ Hz, 1H), 7.70 (d, $J = 8.1$ Hz, 3H), 8.03 (d, $J = 8.4$ Hz, 2H), 8.20 (t, $J = 7.2$ Hz, 1H), 8.49 (s, 1H), 12.73 (br s, 1H). MS (m/z): 407.33 (M-H) $^-$.

Example-6

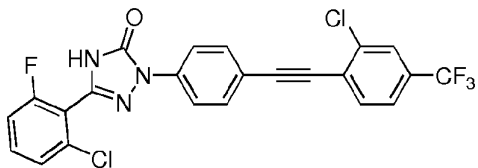
5-(2-Chloro-6-fluorophenyl)-2-{4-[(4-chloro-2-fluorophenyl)ethynyl]phenyl}-2,4-dihydro-3H-1,2,4-triazol-3-one



The title compound was prepared according to the procedure described in Example-3 using 5-(2-chloro-6-fluorophenyl)-2-(4-ethynylphenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one (Intermediate-2, 0.100 g, 0.319 mmol), 4-chloro-2-fluoro-1-iodobenzene (0.123 g, 0.479 mmol), TBAF (0.201 g, 0.638 mmol), bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.028 mmol) and DMSO (3.0 ml). The obtained crude product was purified with column chromatography on silica gel eluting with 1.0 % MeOH: DCM to afford 0.045 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.38 (d, J = 8.4 Hz, 1H), 7.50 (t, J = 8.7 Hz, 1H), 7.57-7.70 (m, 6H), 8.02 (d, J = 8.7 Hz, 2H), 12.74 (br s, 1H). MS (m/z): 442.38 (M^+).

Example-7

5-(2-Chloro-6-fluorophenyl)-2-(4-{[2-chloro-4-(trifluoromethyl)phenyl]ethynyl}phenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one

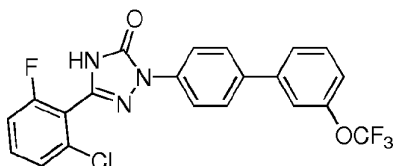


The title compound was prepared according to the procedure described in Example-3 using 5-(2-chloro-6-fluorophenyl)-2-(4-ethynylphenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one (Intermediate-2, 0.100 g, 0.319 mmol), 2-chloro-1-iodo-4-(trifluoromethyl)benzene (0.147 g, 0.479 mmol), TBAF (0.201 g, 0.638 mmol), bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.028 mmol) and DMSO (3.0 mL). The obtained product was purified with column chromatography on silica gel eluting with 1.0 % MeOH: DCM to afford 0.040 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.51 (d, J = 8.7 Hz, 1H), 7.59 (d, J = 7.8 Hz, 1H), 7.67-7.87 (m, 3H),

7.84 (t, $J = 8.4$ Hz, 2H), 8.05 (d, $J = 8.4$ Hz, 2H), 8.11 (s, 1H), 12.75 (br s, 1H). MS (m/z): 492.22 (M⁺).

Example-8

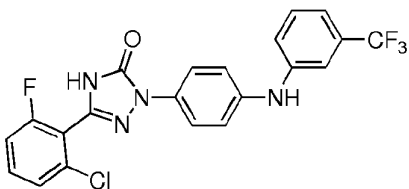
5-(2-Chloro-6-fluorophenyl)-2-[3'-(trifluoromethoxy)biphenyl-4-yl]-2,4-dihydro-3H-1,2,4-triazol-3-one



The title compound was prepared according to the procedure described in Example-2 using 5-(2-chloro-6-fluorophenyl)-2-(4-iodophenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one (step-4 of Intermediate-2, 0.100 g, 0.242 mmol), [3-(trifluoromethoxy)phenyl]boronic acid (0.075 g, 0.363 mmol), K₂CO₃ (0.100 g, 0.726 mmol), tetrakis(triphenyl) phosphine palladium (0) (0.055 g, 0.048 mmol) and DMSO (3.0 mL) to afford 0.025 g of desired product. ¹H NMR (300 MHz, CDCl₃): δ 7.19 (d, $J = 8.4$ Hz, 1H), 7.35 (s, 1H), 7.39 (d, $J = 7.8$ Hz, 1H), 7.46 (d, $J = 6.6$ Hz, 1H), 7.53 (t, $J = 7.8$ Hz, 2H), 7.65 (d, $J = 8.4$ Hz, 2H), 8.12 (d, $J = 8.4$ Hz, 2H), 10.01 (br s, 1H).

Example-9

5-(2-Chloro-6-fluorophenyl)-2-(4-{[3-(trifluoromethyl)phenyl]amino}phenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one

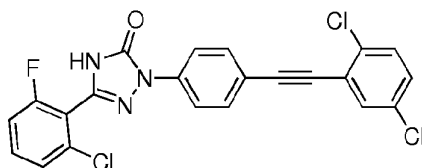


To a solution of 5-(2-chloro-6-fluorophenyl)-2-(4-iodophenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one (step-4 of Intermediate-2, 0.100 g, 0.242 mmol) in toluene (5.0 mL) was added 3-(trifluoromethyl)aniline (0.076 g, 0.292 mmol), NaOtBu (0.035 g, 0.360 mmol), tris(dibenzylideneacetone)dipalladium(0) (0.022 g, 0.024 mmol) and 9,9-dimethyl-4,5-bis(diphenylphosphino)xanthene (0.021 g, 0.036 mmol). The reaction mixture was refluxed for 12 h. The reaction mass was quenched in water and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated.

The obtained product was purified with column chromatography on silica gel eluting with 10 % EA: DCM to afford 0.030 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.09 (d, J = 7.5 Hz, 1H), 7.20 (s, 1H), 7.24 (d, J = 5.7 Hz, 2H), 7.31 (d, J = 8.1 Hz, 1H), 7.41-7.59 (m, 3H), 7.69 (d, J = 7.8 Hz, 1H), 7.81 (d, J = 8.7 Hz, 2H), 8.64 (s, 1H), 12.51 (s, 1H). MS (m/z): 449.31 ($M+H$) $^+$.

Example-10

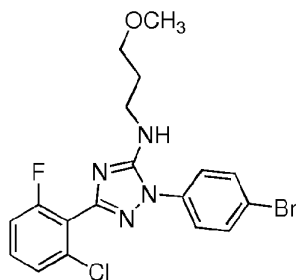
5-(2-Chloro-6-fluorophenyl)-2-{4-[(2,5-dichlorophenyl)ethynyl]phenyl}-2,4-dihydro-3H-1,2,4-triazol-3-one



The title compound was prepared according to the procedure described in Example-3 using 5-(2-chloro-6-fluorophenyl)-2-(4-ethynylphenyl)-2,4-dihydro-3H-1,2,4-triazol-3-one (Intermediate-2, 0.100 g, 0.319 mmol), 1,4-dichloro-2-iodobenzene (0.130 g, 0.470 mmol), TBAF (0.201 g, 0.638 mmol), bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.028 mmol) and DMSO (3.0 mL). The obtained product was purified with column chromatography on silica gel eluting with 1.0 % MeOH: DCM to afford 0.040 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.50-7.72 (m, 7H), 7.82 (s, 1H), 8.05 (d, J = 8.7 Hz, 2H), 12.75 (br s, 1H). MS (m/z): 458.31 (M).

Example-11

1-(4-Bromophenyl)-3-(2-chloro-6-fluorophenyl)-N-(3-methoxypropyl)-1H-1,2,4-triazol-5-amine

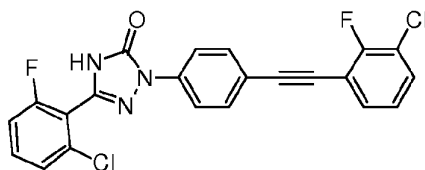


To a solution of 1-(4-bromophenyl)-5-chloro-3-(2-chloro-6-fluorophenyl)-1H-1,2,4-triazole (Intermediate-3, 0.100 g, 0.257 mmol) in DMF (3.0 mL) was added 3-

methoxypropan-1-amine (0.035 g, 0.386 mmol) and DIPEA (1.0 mL). The reaction mass was stirred at 80°C for 5-6 h. The reaction mass was quenched in water and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 0.040 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.92 (d, *J* = 5.4 Hz, 2H), 2.34 (t, *J* = 7.5 Hz, 2H), 3.28 (s, 3H), 3.56-3.64 (m, 2H), 5.69 (m, 1H), 7.06-7.15 (m, 1H), 7.22-7.49 (m, 2H), 7.54 (d, *J* = 10.2 Hz, 2H), 7.64 (d, *J* = 8.4 Hz, 2H). MS (*m/z*): 441.19 (M+H)⁺.

Example-12

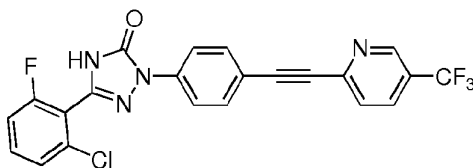
1-(4-((3-Chloro-2-fluorophenyl)ethynyl)phenyl)-3-(2-chloro-6-fluorophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



The title compound was prepared according to the procedure described in Example-3 using 5-(2-chloro-6-fluorophenyl)-2-(4-ethynylphenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one (Intermediate-2, 0.100 g, 0.319 mmol), 1-chloro-2-fluoro-3-iodobenzene (0.123 g, 0.470 mmol), TBAF (0.201 g, 0.638 mmol), bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.028 mmol) and DMSO (3.0 mL). The obtained product was purified with column chromatography on silica gel eluting with 1.0 % MeOH: DCM to afford 0.040 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 7.29 (t, *J* = 7.8 Hz, 1H), 7.50 (t, *J* = 8.7 Hz, 1H), 7.57-7.72 (m, 6H), 8.04 (d, *J* = 9.0 Hz, 2H), 12.74 (br s, 1H). MS (*m/z*): 442.35 (M⁺).

Example-13

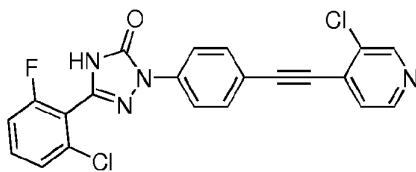
5-(2-Chloro-6-fluorophenyl)-2-(4-{[5-(trifluoromethyl)pyridin-2-yl]ethynyl}phenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one



The title compound was prepared according to the procedure described in Example-3 using 5-(2-chloro-6-fluorophenyl)-2-(4-ethynylphenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one (Intermediate-2, 0.100 g, 0.319 mmol), 2-bromo-5-(trifluoromethyl)pyridine (0.108 g, 0.479 mmol), TBAF (0.201 g, 0.638 mmol), bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.028 mmol) and DMSO (3.0 mL). The obtained product was purified with column chromatography on silica gel eluting with 1.0 % MeOH: DCM to afford 0.030 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 7.51 (t, *J* = 8.7 Hz, 1H), 7.59 (d, *J* = 7.8 Hz, 1H), 7.70 (t, *J* = 6.3 Hz, 1H), 7.79 (d, *J* = 8.7 Hz, 2H), 7.89 (d, *J* = 8.1 Hz, 1H), 8.06 (d, *J* = 8.4 Hz, 2H), 8.29 (d, *J* = 7.8 Hz, 1H), 9.00 (s, 1H), 12.75 (br s, 1H). MS (*m/z*): 459.37 (*M*⁺).

Example-14

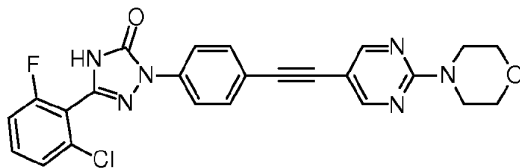
5-(2-Chloro-6-fluorophenyl)-2-{4-[(3-chloropyridin-4-yl)ethynyl]phenyl}-2,4-dihydro-3*H*-1,2,4-triazol-3-one



The title compound was prepared according to the procedure described in Example-3 using 5-(2-chloro-6-fluorophenyl)-2-(4-ethynylphenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one (Intermediate-2, 0.100 g, 0.319 mmol), 3-chloro-4-iodopyridine (Intermediate-4, 0.114 g, 0.429 mmol), TBAF (0.201 g, 0.638 mmol), bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.028 mmol) and DMSO (3.0 mL). The obtained product was purified with column chromatography on silica gel eluting with 1.0 % MeOH: DCM to afford 0.035 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 7.49 (t, *J* = 8.7 Hz, 1H), 7.56 (d, *J* = 11.7 Hz, 2H), 7.60-7.77 (m, 3H), 8.08 (d, *J* = 8.4 Hz, 2H), 8.57 (d, *J* = 5.1 Hz, 1H), 8.78 (s, 1H), 12.76 (br s, 1H). MS (*m/z*): 425.43 (*M*⁺).

Example-15

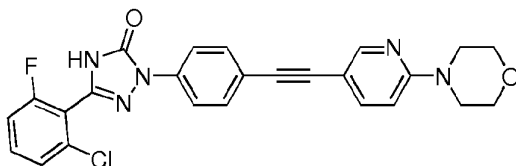
3-(2-Chloro-6-fluorophenyl)-1-(4-((2-morpholinopyrimidin-5-yl)ethynyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



The title compound was prepared according to the procedure described in Example-3 using 5-(2-chloro-6-fluorophenyl)-2-(4-ethynylphenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one (Intermediate-2, 0.100 g, 0.319 mmol), 4-(5-iodopyrimidin-2-yl)morpholine (Intermediate-5, 0.139 g, 0.470 mmol), TBAF (0.201 g, 0.638 mmol), bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.028 mmol) and DMSO (3.0 mL). The obtained product was purified with column chromatography on silica gel eluting with 2.0 % MeOH: DCM to afford 0.035 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.67 (br s, 4H), 3.76 (br s, 4H), 7.47-7.71 (m, 5H), 7.98 (d, *J* = 9.0 Hz, 2H), 8.58 (s, 2H), 12.69 (s, 1H).

Example-16

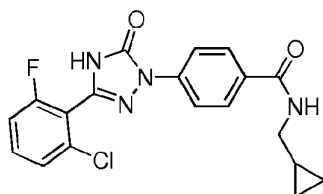
5-(2-Chloro-6-fluorophenyl)-2-(4-{[6-(morpholin-4-yl)pyridin-3-yl]ethynyl}phenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one



The title compound was prepared according to the procedure described in Example-3 using 5-(2-chloro-6-fluorophenyl)-2-(4-ethynylphenyl)-2,4-dihydro-3*H*-1,2,4-triazol-3-one (Intermediate-2, 0.100 g, 0.319 mmol), 4-(5-iodopyridin-2-yl)morpholine (Intermediate-6, 0.139 g, 0.470 mmol), TBAF (0.201 g, 0.638 mmol), bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.028 mmol) and DMSO (3.0 mL). The obtained product was purified with column chromatography on silica gel eluting with 2.0 % MeOH: DCM to afford 0.020 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.58-3.61 (m, 4H), 3.82-3.85 (br s, 4H), 6.64 (d, *J* = 8.7 Hz, 1H), 7.19 (t, *J* = 8.7 Hz, 1H), 7.36-7.47 (m, 2H), 7.58 (d, *J* = 8.4 Hz, 2H), 7.63-7.67 (m, 1H), 8.03 (d, *J* = 9.0 Hz, 2H), 8.38 (s, 1H), 10.14 (br s, 1H); MS (*m/z*): 473.64 (M-H)⁺.

Example-17

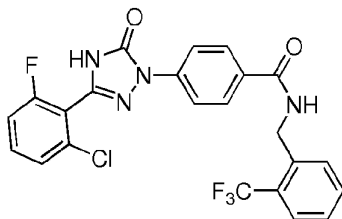
4-[3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-*N*-(cyclopropylmethyl)benzamide



To a solution of 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoic acid (Intermediate-9, 0.100 g, 0.290 mmol) in THF was added di-isopropyl ethyl amine (1.0 mL) and TBTU (0.193 g, 0.590 mmol). The reaction mass was stirred at RT for 1 h. 1-cyclopropylmethanamine (0.048 g, 0.440 mmol) was added to the reaction mass and further stirred at RT for 15 h. The reaction mass was quenched in water and extracted with ethyl acetate. The organic layer was washed with NaHCO₃ solution and water, dried over anhydrous sodium sulphate and concentrated to afford 0.025 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.23 (br s, 1H), 0.44 (d, *J* = 7.5 Hz, 1H), 0.83-0.85 (m, 1H), 1.06-1.024 (m, 1H), 1.31 (m, 1H), 3.14 (t, *J* = 5.7 Hz, 2H), 7.50 (t, *J* = 9.0 Hz, 1H), 7.58 (d, *J* = 7.2 Hz, 1H), 7.67-7.72 (m, 1H), 7.95-8.00 (m, 4H), 8.59 (m, 1H), 12.70 (br s, 1H); MS (*m/z*): 387.35 (M+H)⁺.

Example-18

4-[3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-*N*-[2-(trifluoromethyl)benzyl]benzamide

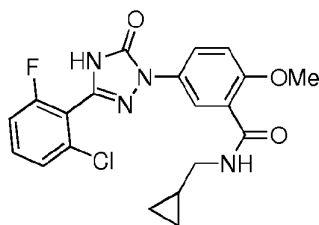


The mixture of methyl 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoate (step-2 of Intermediate-9, 0.100 g, 0.280 mmol), 1-[2-(trifluoromethyl)phenyl]methanamine (0.075 g, 0.430 mmol), 1,2,4-triazole (0.004 g, 0.05 mmol) and DBU (0.009 g, 0.005 mmol) was heated at 80-90°C in seal tube for 15 h. The reaction mass was quenched in water and extracted with ethyl acetate. The organic layer was washed with dilute HCl and water, dried over anhydrous sodium sulphate and

concentrated to afford 0.030 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 4.66-4.68 (d, J = 4.2 Hz, 2H), 7.45-7.75 (m, 7H), 8.05 (s, 4H), 9.16 (br s, 1H), 12.73 (br s, 1H); MS (m/z): 489.42 (M-H) $^-$.

Example-19

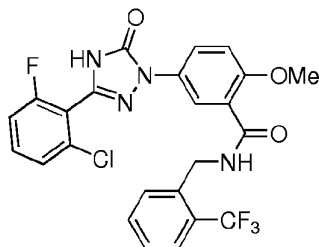
5-[3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-*N*-(cyclopropylmethyl)-2-methoxybenzamide



The title compound was prepared according to the procedure described in Example-17 by using 5-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-2-methoxybenzoic acid (Intermediate-11, 0.100 g, 0.270 mmol), THF (5 mL), di-isopropyl ethyl amine (1.0 mL), TBTU (0.177 g, 0.540 mmol) and 1-cyclopropylmethanamine (0.045 g, 0.410 mmol) to afford 0.035 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.23-0.24 (m, 2H), 0.41-0.44 (m, 2H), 1.03 (m, 1H), 3.17 (t, J = 7.2 Hz, 2H), 3.91 (s, 3H), 7.25 (d, J = 9.0 Hz, 1H), 7.49 (t, J = 8.7 Hz, 1H), 7.57 (d, J = 7.8 Hz, 1H), 7.65-7.70 (m, 1H), 7.98 (d, J = 9.0 Hz, 1H), 8.23 (s, 1H), 8.32 (m, 1H), 12.57 (s, 1H); MS (m/z): 415.23 (M-H) $^-$.

Example-20

5-[3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-2-methoxy-*N*-[2-(trifluoromethyl)benzyl]benzamide

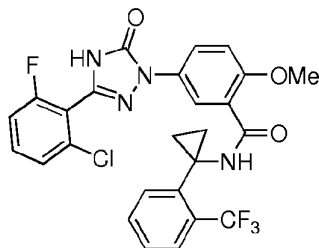


The title compound was prepared according to the procedure described in Example-17 by using 5-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-2-

methoxybenzoic acid (Intermediate-11, 0.100 g, 0.270 mmol), THF (5 mL), di-isopropyl ethyl amine (1.0 mL), TBTU (0.177 g, 0.540 mmol) and 1-[2-(trifluoromethyl)phenyl]methanamine (0.072 g, 0.410 mmol) to afford 0.035 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.95 (s, 3H), 4.62-4.71 (br s, 2H), 7.30 (d, J = 8.7 Hz, 1H), 7.46-7.52 (m, 2H), 7.56-7.59 (m, 2H), 7.65-7.75 (m, 3H), 8.02 (dd, J = 2.4 Hz, 1H), 8.28 (s, 1H), 8.90 (br s, 1H), 12.58 (s, 1H); MS (m/z): 519.30 (M-H) $^-$.

Example-21

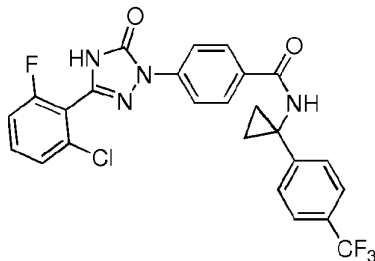
5-[3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-2-methoxy-*N*-{1-[2-(trifluoromethyl)phenyl]cyclopropyl}benzamide



The title compound was prepared according to the procedure described in Example-17 by using 5-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-2-methoxybenzoic acid (Intermediate-11, 0.100 g, 0.270 mmol), THF (5 mL), di-isopropyl ethyl amine (1.0 mL), TBTU (0.177 g, 0.540 mmol) and 1-[2-(trifluoromethyl)phenyl]cyclopropanamine (Intermediate-12, 0.066 g, 0.329 mmol) to afford 0.025 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.26-1.28 (m, 4H), 3.91 (s, 3H), 7.24 (d, J = 9.3 Hz, 1H), 7.11-7.48 (m, 6H), 7.98 (t, J = 7.5 Hz, 2H), 8.27 (s, 1H), 8.74 (s, 1H), 12.56 (br s, 1H); MS (m/z): 545.31 (M-H) $^-$.

Example-22

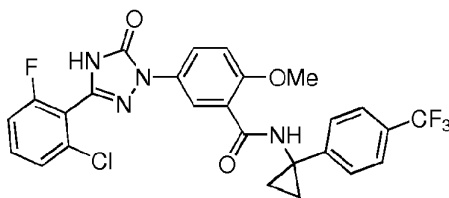
4-[3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-*N*-{1-[4-(trifluoromethyl)phenyl]cyclopropyl}benzamide



The title compound was prepared according to the procedure described in Example-17 by using 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoic acid (Intermediate-9, 0.100 g, 0.290 mmol), THF (5 mL), di-isopropyl ethyl amine (2.0 mL), TBTU (0.193 g, 0.590 mmol) and 1-[4-(trifluoromethyl)phenyl]cyclopropanamine (Intermediate-13, 0.073 g, 0.350 mmol) to afford 0.030 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.35 (s, 4H), 7.36 (d, *J* = 7.8Hz, 2H), 7.46-7.70 (m, 5H), 8.01 (s, 4H), 9.30 (s, 1H), 12-13 (br s, 1H); MS (*m/z*): 515.23 (M-H)⁻.

Example-23

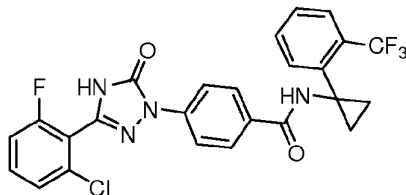
5-[3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-2-methoxy-*N*-{1-[4-(trifluoromethyl)phenyl]cyclopropyl}benzamide



The title compound was prepared according to the procedure described in Example-17 by using 5-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-2-methoxybenzoic acid (Intermediate-11, 0.100 g, 0.270 mmol), THF (10 mL), di-isopropyl ethyl amine (10.0 mL), TBTU (0.177 g, 0.540 mmol) and 1-[4-(trifluoromethyl)phenyl]cyclopropanamine (Intermediate-13, 0.066 g, 0.320 mmol) to afford 0.025 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.23 (s, 4H), 3.94 (s, 3H), 7.26 (d, *J* = 9.0Hz, 1H), 7.42 (d, *J* = 8.4Hz, 3H), 7.30-7.48 (m, 4H), 7.99 (d, *J* = 9.3Hz, 1H), 8.05 (s, 1H), 8.96 (s, 1H), 12.51 (br s, 1H); MS (*m/z*): 547.15 (M⁺).

Example-24

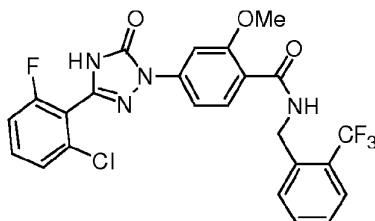
4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(1-(2-(trifluoromethyl)phenyl)cyclopropyl)benzamide



The title compound was prepared according to the procedure described in Example-17 by using 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoic acid (Intermediate-9, 0.100 g, 0.290 mmol), THF (10 mL), di-isopropyl ethyl amine (10.0 mL), TBTU (0.177 g, 0.540 mmol) and 1-[2-(trifluoromethyl)phenyl]cyclopropanamine (Intermediate-12, 0.035 g, 0.350 mmol) to afford 0.020 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.23 (s, 2H), 1.31 (s, 2H), 7.59 (d, *J* = 7.2 Hz, 2H), 7.65-7.68 (m, 4H), 7.82 (d, *J* = 12.0 Hz, 2H), 7.88-8.02 (m, 3H), 8.77 (s, 1H), 12.60 (br s, 1H); MS (*m/z*): 517.03 (*M*⁺).

Example-25

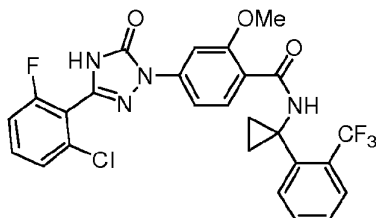
4-[3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]-2-methoxy-*N*-[2-(trifluoromethyl)benzyl]benzamide



The title compound was prepared according to the procedure described in Example-17 by using 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoic acid (Intermediate-15, 0.100 g, 0.270 mmol), THF (5 mL), di-isopropyl ethyl amine (1.0 mL), TBTU (0.177 g, 0.540 mmol) and 1-[2-(trifluoromethyl)phenyl]methanamine (0.072 g, 0.410 mmol) to afford 0.025 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.96 (s, 3H), 4.70 (d, *J* = 4.8 Hz, 2H), 7.45-7.50 (m, 3H), 7.57 (t, *J* = 8.4 Hz, 2H), 7.67 (t, *J* = 8.7 Hz, 2H), 7.71 (d, *J* = 9.3 Hz, 2H), 7.92 (d, *J* = 8.4 Hz, 1H), 8.80 (t, 1H), 12.77 (br s, 1H); MS (*m/z*): 519.43 (*M*-H)⁻.

Example-26

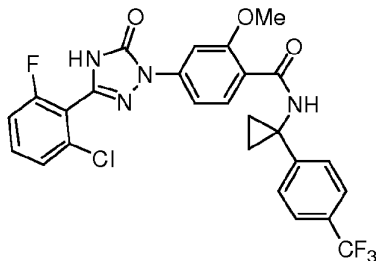
4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(1-(2-(trifluoromethyl)phenyl)cyclopropyl)benzamide



The title compound was prepared according to the procedure described in Example-17 by using 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoic acid (Intermediate-15, 0.100 g, 0.270 mmol), THF (5 mL), di-isopropyl ethyl amine (2.0 mL), TBTU (0.177 g, 0.540 mmol) and 1-[2-(trifluoromethyl)phenyl] cyclopropanamine (Intermediate-12, 0.082 g, 0.410 mmol) to afford 0.020 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.27-1.31 (m, 4H), 3.90 (s, 3H), 7.47 (t, *J* = 8.1 Hz, 2H), 7.54-7.69 (m, 6H), 7.87 (t, *J* = 8.4 Hz, 1H), 7.99 (d, *J* = 7.8 Hz, 1H), 8.61 (s, 1H); MS (*m/z*): 547.15 (*M*⁺).

Example-27

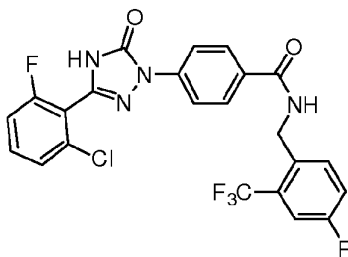
4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(1-(4-(trifluoromethyl)phenyl)cyclopropyl)benzamide



The title compound was prepared according to the procedure described in Example-17 by using 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoic acid (Intermediate-15, 0.100 g, 0.270 mmol), THF (5 mL), di-isopropyl ethyl amine (2.0 mL), TBTU (0.177 g, 0.540 mmol) and 1-[4-(trifluoromethyl)phenyl] cyclopropanamine (Intermediate-13, 0.066 g, 0.320 mmol) to afford 0.020 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.37 (s, 4H), 3.95 (s, 3H), 7.42 (d, *J* = 7.8 Hz, 2H), 7.50 (t, *J* = 8.7 Hz, 1H), 7.63 (t, *J* = 7.8 Hz, 4H), 7.73 (t, *J* = 12.3 Hz, 3H), 8.85 (s, 1H), 12.50 (br s, 1H); MS (*m/z*): 546.99 (*M*⁺).

Example-28

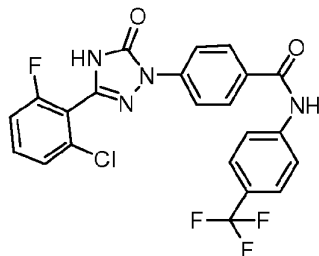
4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(4-fluoro-2-(trifluoromethyl)benzyl)benzamide



The title compound was prepared according to the procedure described in Example-17 by using 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoic acid (Intermediate-9, 0.100 g, 0.30 mmol), THF (5 mL), di-isopropyl ethyl amine (2.0 mL), TBTU (0.177 g, 0.540 mmol) and 4-fluoro-2-trifluoromethyl benzylamine (0.089 g, 0.600 mmol) to afford 0.025 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 4.63 (d, $J = 4.5$ Hz, 2H), 7.48-7.50 (m, 6H), 8.05 (s, 4H), 9.16 (t, 1H), 12.67 (br, 1H); MS (m/z): 509.05 ($\text{M}+\text{H}$) $^+$.

Example-29

4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(4-(trifluoromethyl)phenyl)benzamide

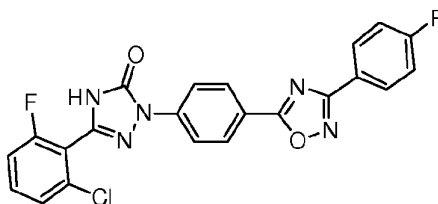


To a solution of 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzamide (Intermediate-17, 0.100 g, 0.301 mmol) in dry toluene (5 mL) was added 1-bromo-4-(trifluoromethyl)benzene (0.101 g, 0.451 mmol), sodium *tert*-butoxide (0.058 g, 0.602 mmol), tris(diphenylideneacetone)dipalladium(0) (0.005 g, 0.0006 mmol), 4,5-bis(diphenylphosphino) and 9,9-dimethylxanthene (0.005 g, 0.0009 mmol) under nitrogen atmosphere. The reaction mass was refluxed for 3-4 h. The reaction mass was quenched in water and extracted in ethyl acetate. The organic layer was concentrated to afford crude product which was purified by column chromatography to afford 0.015 g of

desired product. ^1H NMR (300 MHz, DMSO d_6): δ 7.40 (br s, 1H), 7.84 (t, $J = 9$ Hz, 1H), 7.55 (d, 1H), 7.72-7.75 (m, 3H), 8.02 (d, $J = 8.1$ Hz, 2H), 8.11 (br s, 3H) 10.61 (br s, 1H), 12.70 (br s, 1H). MS (m/z): 475.18 (M-H) $^-$.

Example-30

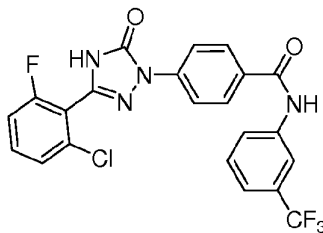
3-(2-Chloro-6-fluorophenyl)-1-(4-(3-(4-fluorophenyl)-1,2,4-oxadiazol-5-yl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



To a solution of 4-fluoro-*N*'-hydroxybenzimidamide (Intermediate-18, 0.073 g, 0.47 mmol) in dry DMF (3 mL) under nitrogen atmosphere was added 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzonitrile (step-4 of Intermediate-17, 0.100 g, 0.31 mmol) and ZnCl_2 (0.0087 g, 0.06 mmol). The reaction mass was heated at 100°C for 8-10 h. The reaction mass was quenched with water, extracted in DCM and column purified to afford 0.025 g of desired product. ^1H NMR (DMSO- d_6): δ 7.02 (br s, 1H), 7.33 (t, 1H), 7.45-7.54 (m, 2H), 7.61 (d, 1H), 7.70 (m, 1H), 7.83 (t, 1H), 8.09-8.17 (m, 2H), 8.26-8.30 (m, 2H), 12.81 (br s, 1H). MS (m/z): 452.05 [M+H] $^+$.

Example-31

4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(trifluoromethyl)phenyl)benzamide

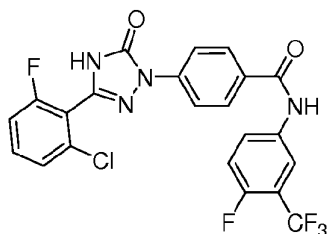


To a solution of methyl 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoate (step-2 of Intermediate-9, 0.100 g, 0.28 mmol) in dry toluene was added 3-trifluoromethyl aniline (0.070 g, 0.43 mmol) followed by addition of trimethyl

aluminium(2M solution in toluene) (0.5 mL). The reaction mixture was refluxed for 1 h. The reaction mass was quenched in water and acidified with dilute HCl and extracted in DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford crude product, which was column purified to afford 0.040 g of desired product. ^1H NMR (DMSO- d_6): δ 7.45-7.54 (m, 2H), 7.59 (m, 2H), 7.72 (q, J = 6.3 Hz, 1H), 8.05-8.12 (m, 5H), 8.25 (s, 1H), 10.58 (br s, 1H), 12.77 (br s, 1H); MS (m/z): 477.08 $[\text{M}+\text{H}]^+$.

Example-32

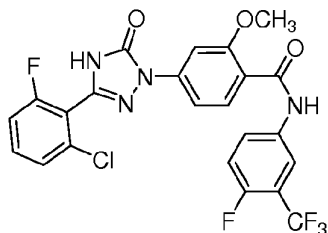
4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(4-fluoro-3-(trifluoromethyl)phenyl)benzamide



The title compound was prepared according to the procedure described in Example-31, by using methyl 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoate (step-2 of Intermediate-9, 0.100 g, 0.28 mmol), 4-fluoro-3-trifluoromethyl aniline (0.077 g, 0.43 mmol) and trimethyl aluminium (2M solution in toluene) (0.5 mL) to afford 0.043 g of desired product. ^1H NMR (DMSO- d_6): δ 7.46-7.59 (m, 3H), 7.69 (t, J = 6.3 Hz, 1H), 8.11 (br s, 5H), 8.26 (m, 1H), 10.59 (br s, 1H), 12.72 (br s, 1H); MS (m/z): 495.07 $[\text{M}+\text{H}]^+$.

Example-33

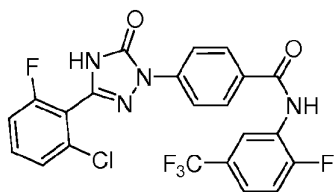
4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(4-fluoro-3-(trifluoromethyl)phenyl)-2-methoxybenzamide



The title compound was prepared according to the procedure described in Example-31, by using methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-2 of Intermediate-15, 0.100 g, 0.26 mmol), 4-floro-3-trifluoromethyl aniline (0.070 g, 1.5 mmol), trimethyl aluminium (2M solution in toluene) (0.5 mL) to afford 0.030 g of desired product. ¹HNMR (DMSO-*d*₆): δ 3.94 (s, 3H), 7.50-7.60 (m, 3H), 7.67-7.72 (m, 2H), 7.77-7.82 (m, 2H), 8.01 (m, 1H), 8.26 (m, 1H), 10.38 (br s, 1H), 12.80 (br s, 1H); MS (m/z): 525.17 [M+H]⁺.

Example-34

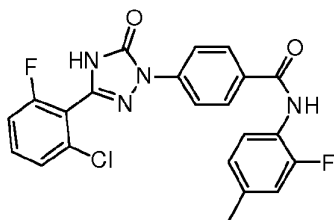
4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(2-fluoro-5-(trifluoromethyl)phenyl)benzamide



The title compound was prepared according to the procedure described in Example-31, by using methyl 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoate (step-2 of Intermediate-9, 0.100g, 0.28 mmol), 2-fluoro-5-trifluoromethyl aniline (0.077 g, 0.43 mmol) and trimethyl aluminium (2M solution in toluene) (0.5 mL) to afford 0.050 g of desired product. ¹HNMR (DMSO-*d*₆): δ 7.48-7.61 (m, 3H), 7.68-7.73 (m, 2H), 8.08-8.12 (m, 5H), 10.41 (br s, 1H), 12.78 (br s, 1H); MS (m/z): 493.09 [M-H]⁻.

Example-35

4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(2-fluoro-4-methylphenyl)benzamide

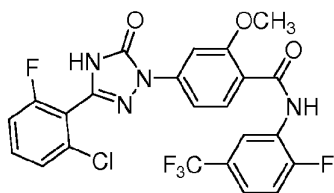


The title compound was prepared according to the procedure described in Example-31, by using methyl 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-

yl]benzoate (step-2 of Intermediate-9, 0.100 g, 0.28 mmol), 2-fluoro-4-methyl aniline (0.054 g, 0.43 mmol) and trimethyl aluminium (2M solution in toluene) (0.5 mL) to afford 0.055 g of desired product. ^1H NMR ($\text{DMSO}-d_6$): δ 2.30 (s, 3H), 7.00 (d, $J=7.8$ Hz, 1H), 7.11 (d, $J=11.7$ Hz, 1H), 7.39-7.48 (m, 2H), 7.51-7.58 (d, 1H), 7.65-7.72 (m, 1H), 8.07 (br s, 4H), 10.04 (br s, 1H), 12.73 (br s, 1H); MS (m/z): 439.13 $[\text{M}-\text{H}]^-$.

Example-36

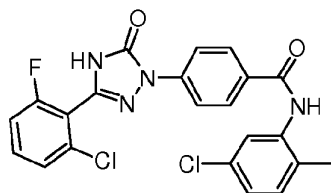
4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(2-fluoro-5-(trifluoromethyl)phenyl)-2-methoxybenzamide



The title compound was prepared according to the procedure described in Example-31, by using methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-2 of Intermediate-15, 0.100 g, 0.26 mmol), 2-fluoro-5-trifluoromethyl aniline (0.070 g, 0.39 mmol) and trimethyl aluminium (2M solution in toluene) (0.5 mL) to afford 0.049 g of desired product. ^1H NMR ($\text{DMSO}-d_6$): δ 4.05 (s, 3H), 7.50 (t, $J=8.4$ Hz, 1H), 7.57-7.70 (m, 3H), 7.72-7.77 (m, 2H), 7.85 (s, 1H), 8.08 (d, $J=8.4$ Hz, 1H), 8.67 (d, $J=6.3$ Hz, 1H), 10.38 (br s, 1H), 12.82 (br s, 1H); MS (m/z): 525.15 $[\text{M}+\text{H}]^+$.

Example-37

N-(5-Chloro-2-methylphenyl)-4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzamide

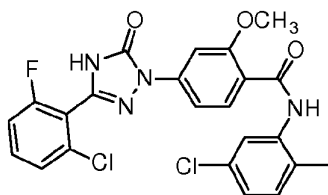


The title compound was prepared according to the procedure described in Example-31, by using methyl 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoate (step-2 of Intermediate-9, 0.100 g, 0.28 mmol), 5-chloro-2-methyl aniline

(0.062 g, 0.43 mmol) and trimethyl aluminium (2M solution in toluene) (0.5 mL) to afford 0.059 g of desired product. ^1H NMR (DMSO- d_6): δ 2.23 (s, 3H), 7.23 (d, J = 6.9 Hz, 1H), 7.30 (d, J = 8.4 Hz, 1H), 7.53 (m, 2H), 7.59 (m, 1H), 7.71 (m, 1H), 8.10 (br s, 4H), 9.98 (br s, 1H), 12.76 (br s, 1H); MS (m/z): 457.18 $[\text{M}]^+$.

Example-38

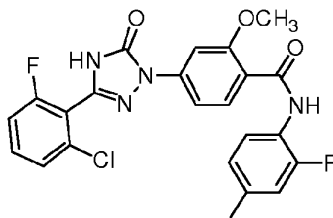
N-(5-Chloro-2-methylphenyl)-4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzamide



The title compound was prepared according to the procedure described in Example-31, by using methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-2 of Intermediate-15, 0.100 g, 0.26 mmol), 5-chloro-2-methyl aniline (0.057 g, 0.39 mmol) and trimethyl aluminium (2M solution in toluene) (0.5 mL) to afford 0.057 g of desired product. ^1H NMR (DMSO- d_6): δ 2.32 (s, 3H), 4.05 (s, 3H), 7.13 (d, J = 9.9 Hz, 1H), 7.30 (d, J = 8.1 Hz, 1H), 7.49 (t, 1H), 7.59 (d, 1H), 7.61-7.75 (m, 2H), 7.84 (s, 1H), 8.08 (d, J = 8.7 Hz, 1H), 8.19 (s, 1H), 9.94 (br s, 1H), 12.83 (br s, 1H); MS (m/z): 487.11 $[\text{M}]^+$.

Example-39

4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(2-fluoro-4-methylphenyl)-2-methoxybenzamide

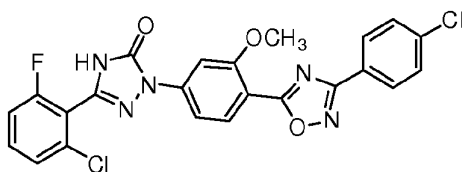


The title compound was prepared according to the procedure described in Example-31, by using methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-2 of Intermediate-15, 0.100 g, 0.26 mmol), 2-fluoro-4-methyl aniline (0.050 g, 0.39 mmol) and trimethyl aluminium (2M solution in toluene)

(0.5 mL) to afford 0.047 g of desired product. ^1H NMR ($\text{DMSO}-d_6$): δ 2.30 (s, 3H), 4.03 (s, 3H), 7.02 (d, $J = 8.1$ Hz, 1H), 7.14 (d, $J = 11.7$ Hz, 1H), 7.51 (t, 1H), 7.58 (d, 1H), 7.65-7.75 (m, 2H), 7.85 (s, 1H), 8.04-8.10 (m, 2H), 10.07 (br s, 1H), 12.65 (br s, 1H); MS (m/z): 469.16 [$\text{M}-\text{H}$].

Example-40

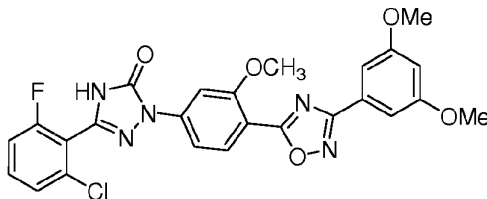
3-(2-Chloro-6-fluorophenyl)-1-(4-(3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl)-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



To a solution of 4-chloro-*N*-hydroxybenzimidamide (Intermediate-19, 0.069 g, 0.39 mmol) in dry toluene was added sodium hydride (0.016 g, 0.39 mmol). The reaction mixture was refluxed for 30 minutes followed by addition of methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-2 of Intermediate-15, 0.100 g, 0.26 mmol). The reaction mixture was refluxed for 5-6 h. The reaction mixture was quenched in water and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford crude product which was further purified by column chromatography to afford 0.015 g of desired product. ^1H NMR ($\text{DMSO}-d_6$): δ 3.98 (s, 3H), 7.45-7.48 (m, 1H), 7.54 (d, $J = 8.1$ Hz, 1H), 7.67 (d, $J = 8.4$ Hz, 3H), 7.83 (d, $J = 8.1$ Hz, 1H), 7.95 (s, 1H), 8.10 (d, $J = 8.4$ Hz, 2H), 8.18 (d, $J = 8.7$ Hz, 1H), 12.90 (br s, 1H). MS [$\text{M}+\text{H}$] $^+$:499.89.

Example-41

3-(2-Chloro-6-fluorophenyl)-1-(4-(3-(3,5-dimethoxyphenyl)-1,2,4-oxadiazol-5-yl)-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one

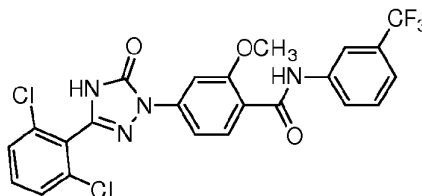


The title compound was prepared by following the procedure as described for Example-40 by using methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-

yl)-2-methoxybenzoate (step-2 of Intermediate -15, 0.100 g, 0.26 mmol), *N'*-hydroxy-3,5-dimethoxybenzimidamide (Intermediate-20, 0.077 g, 0.529 mmol), sodium hydride (0.006 g, 0.39 mmol) and dry THF (5.0 mL) at RT to afford 0.015 g of desired product. ¹HNMR (DMSO-*d*₆): δ 3.84 (s, 6H), 3.99 (s, 3H), 6.73 (s, 1H), 7.19 (s, 2H), 7.47 (t, *J* = 8.1 Hz, 1H), 7.56 (d, *J* = 8.4 Hz, 1H), 7.67 (t, *J* = 8.7 Hz, 1H), 7.81 (d, *J* = 8.1 Hz, 1H), 7.93 (s, 1H), 8.19 (d, *J* = 8.7 Hz, 1H), 12.61 (br s, 1H); MS (*m/z*): 523.89 [M+]⁺.

Example-42

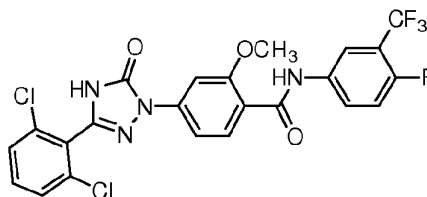
4-(3-(2,6-Dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide



The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-21, 0.100 g, 0.25 mmol), 3-(trifluoromethyl)aniline (0.062 g, 0.38 mmol), trimethyl aluminium (2M solution in toluene) (0.5 mL) and dry toluene (5.0 mL) to afford 0.035 g of desired product. ¹HNMR (DMSO-*d*₆): δ 3.95 (s, 3H), 7.45 (d, *J* = 7.5 Hz, 1H), 7.59 (t, *J* = 7.8 Hz, 1H), 7.76-7.68 (m, 5H), 7.82 (t, *J* = 8.7 Hz, 1H), 7.98 (d, *J* = 8.4 Hz, 1H), 8.25 (s, 1H), 10.37 (s, 1H), 12.75 (br s, 1H); MS (*m/z*): 524.96[M+H]⁺.

Example-43

4-(3-(2,6-Dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(4-fluoro-3-(trifluoromethyl)phenyl)-2-methoxybenzamide

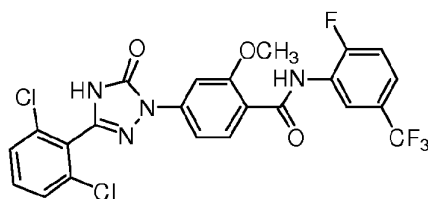


The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-

2-methoxybenzoate (Intermediate-21, 0.100 g, 0.25 mmol), 4-fluoro-3-(trifluoromethyl)aniline (0.069 g, 0.38 mmol), trimethyl aluminium (2M solution in toluene) (0.5 mL) and dry toluene (5.0 mL) to afford 0.050 g of desired product. ¹HNMR (DMSO-*d*₆): δ 4.09 (s, 3H), 7.52 (t, *J* = 9.6 Hz, 1H), 7.88-7.68 (m, 6H), 8.25 (s, 1H), 8.25 (d, *J* = 6.9 Hz, 1H), 10.37 (s, 1H), 12.74 (s, 1H); MS (*m/z*): 540.91[M]⁺.

Example-44

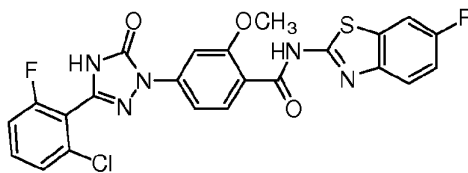
4-(3-(2,6-Dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(2-fluoro-5-(trifluoromethyl)phenyl)-2-methoxybenzamide



The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-21, 0.100 g, 0.25 mmol), 2-fluoro-5-(trifluoromethyl)aniline (0.069 g, 0.38 mmol), trimethyl aluminium (2M solution in toluene) (0.5 mL) and dry toluene (5.0 mL) to afford 0.035 g of desired product. ¹HNMR (DMSO-*d*₆): δ 4.06 (s, 3H), 7.59 (d, *J* = 7.8 Hz, 1H), 7.77-7.68 (m, 5H), 7.84 (s, 1H), 8.09 (d, *J* = 8.7 Hz, 1H), 8.67 (d, *J* = 6.9 Hz, 1H), 10.38 (s, 1H), 12.79 (br s, 1H); MS (*m/z*): 540.92 [M+H]⁺.

Example-45

4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(6-fluorobenzo[d]thiazol-2-yl)-2-methoxybenzamide

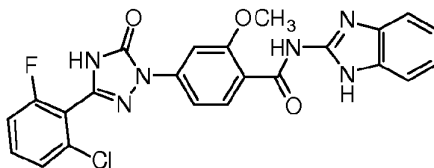


The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-2 of Intermediate-15, 0.100 g, 0.26 mmol), 6-fluorobenzo[d]thiazol-2-amine (0.067 g, 0.397 mmol), trimethyl aluminium (2M solution

in toluene) (0.5 mL) and dry toluene (5.0 mL) to afford 0.050 g of desired product. ¹HNMR (DMSO-*d*₆): δ 3.99 (s, 3H), 7.32 (t, *J* = 7.2 Hz, 1H), 7.51 (t, *J* = 9.3 Hz, 1H), 7.59 (d, *J* = 8.4 Hz, 1H), 7.68-7.80 (m, 4H), 7.94 (t, *J* = 6.3 Hz, 2H), 11.98 (s, 1H), 12.83 (br s, 1H); MS (*m/z*): 512.97 [M-H]⁻.

Example-46

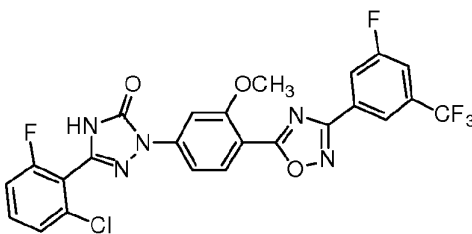
N-(1*H*-Benzo[d]imidazol-2-yl)-4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzamide



The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-2 of Intermediate-15, 0.100 g, 0.26 mmol), 1*H*-benzo[d]imidazol-2-amine (0.046 g, 0.34 mmol), trimethyl aluminium (2M solution in toluene) (0.5 mL) and dry toluene (5.0 mL) to afford 0.040 g of desired product. ¹HNMR (DMSO-*d*₆): δ 4.01 (s, 3H), 7.11 (m, 2H), 7.48-7.76 (m, 6H), 7.81 (s, 1H), 7.96 (d, *J* = 9.0 Hz, 1H), 11.16 (br s, 1H), 12.28 (br s, 1H), 12.76 (br s, 1H); MS (*m/z*): 478.90 [M]⁺

Example-47

3-(2-Chloro-6-fluorophenyl)-1-(4-(3-(3-fluoro-5-(trifluoromethyl)phenyl)-1,2,4-oxadiazol-5-yl)-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one

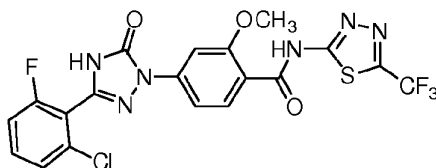


The title compound was prepared by following the procedure as described for Example-40 by using methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-2 of Intermediate-15, 0.100 g, 0.26 mmol), 3-fluoro-*N'*-hydroxy-5-(trifluoromethyl)benzimidamide (Intermediate-22, 0.088 g, 0.39 mmol), sodium hydride (0.017 g, 0.42 mmol) and dry THF (5.0 mL) at RT to afford 0.010 g of

desired product. ^1H NMR (DMSO- d_6): δ 4.01 (s, 3H), 7.50 (m, 1H), 7.60 (m, 1H), 7.70 (m, 1H), 7.84 (m, 1H), 7.92 (br s, 1H), 8.02 (d, 1H), 8.18 (m, 2H), 8.25 (d, 1H), 12.95 (br s, 1H); MS (m/z): 550.07 [M] $^+$.

Example-48

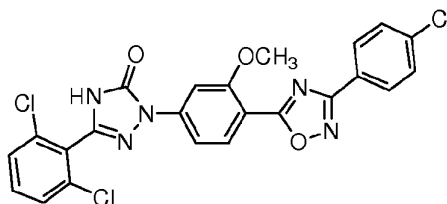
4-(3-(2-Chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(5-(trifluoromethyl)-1,3,4-thiadiazol-2-yl)benzamide



The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-2 of Intermediate -15, 0.100 g, 0.26 mmol), 5-(trifluoromethyl)-1,3,4-thiadiazol-2-amine (0.059 g, 0.34 mmol), trimethyl aluminium (2M solution in toluene) (0.5 mL) and dry toluene (5.0 mL) to afford 0.025 g of desired product. ^1H NMR (DMSO- d_6): δ 3.95 (s, 3H), 7.51 (t, J = 9.0 Hz, 1H), 7.87-7.76 (m, 3H), 7.78 (s, 1H), 7.85 (d, J = 8.7 Hz, 1H), 12.86 (br s, 2H); MS (m/z): 514.90 [M] $^+$.

Example-49

1-(4-(3-(4-Chlorophenyl)-1,2,4-oxadiazol-5-yl)-3-methoxyphenyl)-3-(2,6-dichlorophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one

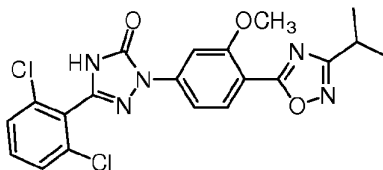


The title compound was prepared by following the procedure as described for Example-40 by using methyl 4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-21, 0.100 g, 0.25 mmol), 4-chloro-*N'*-hydroxybenzimidamide (Intermediate-19, 0.064 g, 0.38 mmol), sodium hydride (0.020 g, 0.50 mmol) and dry THF (5.0 mL) at RT to afford 0.015 g of desired product. ^1H NMR (DMSO- d_6): δ 4.00 (s, 3H), 7.66-7.72 (m, 5H), 7.81 (d, J = 8.7 Hz, 1H), 7.89 (s, 1H),

8.10 (d, $J = 8.4$ Hz, 2H), 8.22 (d, $J = 9.0$ Hz, 1H), 12.82 (br s, 1H); MS (m/z): 515.84 $[M+H]^+$.

Example-50

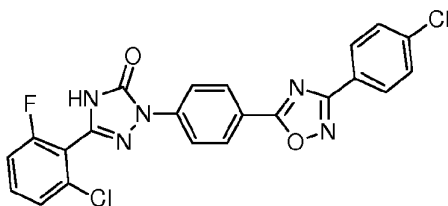
3-(2,6-Dichlorophenyl)-1-(4-(3-isopropyl-1,2,4-oxadiazol-5-yl)-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



The title compound was prepared by following the procedure as described for Example-40 by using methyl 4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-21, 0.100 g, 0.25 mmol), *N*'-hydroxyisobutyrimidamide (0.077 g, 0.529 mmol), sodium hydride (0.020 g, 0.50 mmol) and dry THF (5.0 mL) at RT to afford 0.019 g of desired product. ^1H NMR ($\text{DMSO}-d_6$): δ 1.31 (d, $J = 6.9$ Hz, 6H), 3.12 (m, 1H), 3.95 (s, 3H), 7.65-7.78 (m, 4H), 7.56 (s, 1H), 8.09 (d, $J = 8.7$ Hz, 1H), 12.83 (br s, 1H); MS (m/z): 447.96 $[M+H]^+$.

Example-51

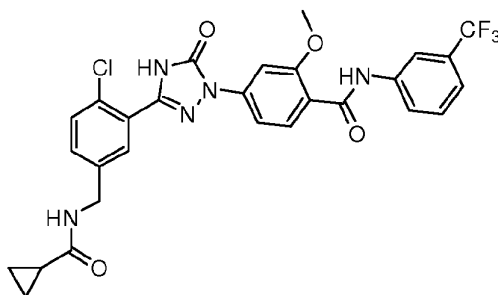
3-(2-Chloro-6-fluorophenyl)-1-(4-(3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



The title compound was prepared by following the procedure as described for Example-40 by using methyl 4-[3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl]benzoate (step-2 of Intermediate-9, 0.100 g, 0.28 mmol), 4-chloro-*N*'-hydroxybenzimidamide (Intermediate-19, 0.073 g, 0.43 mmol), sodium hydride (0.018 g, 0.43 mmol) and dry THF (5.0 mL) to afford 0.025 g of desired product. ^1H NMR ($\text{DMSO}-d_6$): δ 7.51 (t, $J = 8.7$ Hz, 1H), 7.59 (d, $J = 8.4$ Hz, 1H), 7.67-7.73 (m, 3H), 8.11 (d, $J = 8.1$ Hz, 2H), 8.24- 8.29 (m, 4H), 12.85 (br s, 1H); MS (m/z): 467.87 $[M+H]^+$.

Example-52

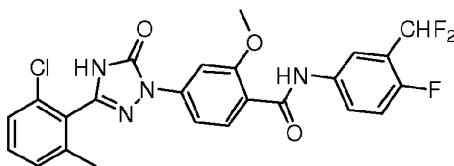
4-(3-(2-Chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide



The title compound was prepared by following the procedure as described for step- 2 of Intermediate-30 by using 4-(3-(5-(aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide (Intermediate-26 , 0.060 g, 0.11 mmol), cyclopropylcarbonyl chloride (0.019 g, 0.17 mmol), DIPEA (1 mL) and THF (5 mL) to afford 0.020 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.06 (m, 2H), 1.22 (m, 2H), 3.17 (m, 1H), 3.95 (s, 3H), 4.34 (d, *J* = 6.0 Hz, 2H), 7.44-7.84 (m, 8H), 7.95 (d, *J* = 9.0 Hz, 1H), 8.25 (s, 1H), 8.70 (m, 1H), 10.43 (m, 1H), 12.05 (m, 1H); MS (*m/z*): 584.35 (M+H⁺).

Example-53

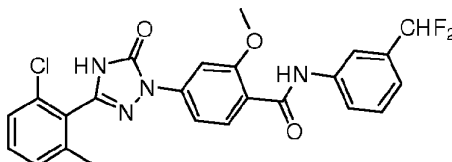
4-(3-(2-Chloro-6-methylphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(difluoromethyl)-4-fluorophenyl)-2-methoxybenzamide



The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2-chloro-6-methylphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-27, 0.100 g, 0.268 mmol), 3-(difluoromethyl)-4-fluoroaniline (Intermediate-28, 0.065 g, 0.403 mmol), trimethyl aluminium (2M solution in toluene) (0.5 mL) and dry toluene (5.0 mL) to afford 0.025 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 2.32 (s, 3H), 3.94 (s, 3H), 7.05-7.41 (m, 3H), 7.50 (s, 2H), 7.68-7.88 (m, 4H), 8.11 (m, 1H), 10.27 (s, 1H), 12.53 (s, 1H); MS (*m/z*): 503.10 (M+H⁺).

Example-54

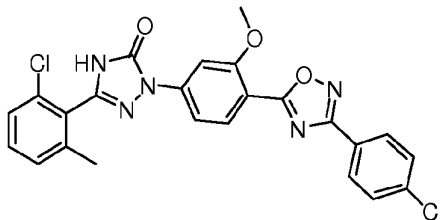
4-(3-(2-Chloro-6-methylphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(difluoromethyl)phenyl)-2-methoxybenzamide



The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2-chloro-6-methylphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-27, 0.100 g, 0.268 mmol), 3-(difluoromethyl)aniline (Intermediate-29, 0.058 g, 0.40 mmol), trimethyl aluminium (2M solution in toluene) (0.5 mL) and dry toluene (5.0 mL) to afford 0.035 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 2.33 (s, 3H), 3.95 (s, 3H), 7.04 (m, 1H), 7.35 (d, 1H), 7.41 (m, 1H), 7.50 (m, 3H), 7.69 (d, *J* = 8.7 Hz, 1H), 7.77-7.82 (m, 3H), 8.07 (s, 1H), 10.25 (s, 1H), 12.52 (br s, 1H); MS (*m/z*): 485.06 (*M*+*H*⁺).

Example-55

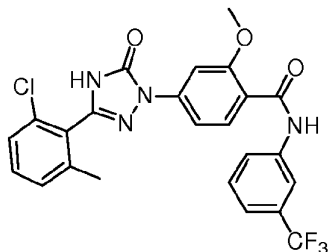
3-(2-Chloro-6-methylphenyl)-1-(4-(3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl)-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



The title compound was prepared by following the procedure as described for Example-40 by using methyl 4-(3-(2-chloro-6-methylphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-27, 0.100 g, 0.26 mmol), 4-chloro-*N'*-hydroxybenzimidamide (Intermediate-19, 0.069 g, 0.40 mmol), sodium hydride (0.022 g, 0.53 mmol) and dry THF (5.0 mL) to afford 0.020 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 2.33 (s, 3H), 4.00 (s, 3H), 7.41 (m, 1H), 7.51 (s, 2H), 7.67 (d, *J* = 8.7 Hz, 2H), 7.83 (d, *J* = 8.7 Hz, 1H), 7.90 (s, 1H), 8.10 (d, *J* = 8.4 Hz, 2H), 8.22 (d, *J* = 8.7 Hz, 1H), 12.63 (s, 1H); MS (*m/z*): 495.98 (*M*+*H*⁺).

Example-56

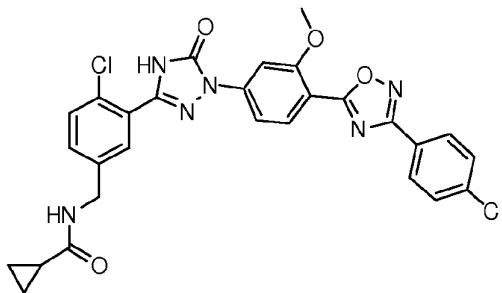
4-(3-(2-Chloro-6-methylphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide



The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2-chloro-6-methylphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-27, 0.100 g, 0.268 mmol), 3-(trifluoromethyl)aniline (0.040 g, 0.403 mmol), trimethyl aluminium (2M solution in toluene) (0.5 mL) and dry toluene (5.0 mL) to afford 0.030 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 2.33 (s, 3H), 3.95 (s, 3H), 7.41-7.59 (m, 4H), 7.68 (t, *J* = 6.9 Hz, 1H), 7.75 (d, 1H), 7.77-7.82 (m, 2H), 7.95 (d, *J* = 8.4 Hz, 1H), 8.25 (s, 1H), 10.36 (s, 1H), 12.53 (br s, 1H); MS (*m/z*): 503.17 (*M*⁺).

Example-57

N-(4-Chloro-3-(1-(4-(3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl)-3-methoxyphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)cyclopropanecarboxamide

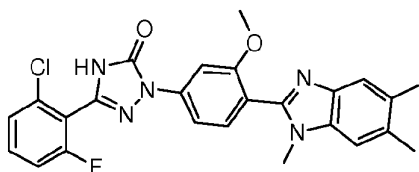


The title compound was prepared by following the procedure as described for Example-40 by using methyl 4-(3-(2-chloro-6-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-30, 0.100 g, 0.21 mmol), 4-chloro-*N*-hydroxybenzimidamide (Intermediate-19, 0.056 g, 0.32 mmol), sodium hydride (0.018 g, 0.43 mmol) and dry THF (5.0 mL) to afford 0.020 g of desired

product. ^1H NMR (300 MHz, DMSO d_6): δ 1.21 (m, 2H), 1.59 (m, 2H), 3.04 (m, 1H), 3.98 (s, 3H), 4.31 (m, 2H), 7.53 (m, 2H), 7.64 (m, 3H), 7.82 (m, 1H), 7.92 (m, 1H), 8.08 (m, 2H), 8.20 (m, 1H), 8.67 (s, 1H), 12.77 (br s, 1H); MS (m/z): 577.07 ($\text{M}+\text{H}^+$).

Example-58

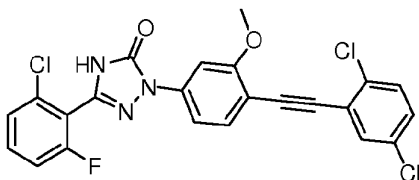
3-(2-Chloro-6-fluorophenyl)-1-(3-methoxy-4-(1,5,6-trimethyl-1*H*-benzo[d]imidazol-2-yl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (step-2 of Intermediate-15, 0.100 g, 0.26 mmol), *N*¹,4,5-trimethylbenzene-1,2-diamine (Intermediate-31, 0.048 g, 0.32 mmol), trimethyl aluminium (2M solution in toluene) (0.5 mL), dry toluene (5.0 mL) to afford 0.030 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 2.33 (s, 3H), 2.36 (s, 3H), 3.56 (s, 3H), 3.85 (s, 3H), 7.33 (s, 1H), 7.41 (s, 1H), 7.47-7.59 (m, 3H), 7.66-7.78 (m, 3H), 12.5 (br s, 1H); MS (m/z): 478.29 ($\text{M}+\text{H}^+$).

Example-59

3-(2-Chloro-6-fluorophenyl)-1-(4-((2,5-dichlorophenyl)ethynyl)-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one

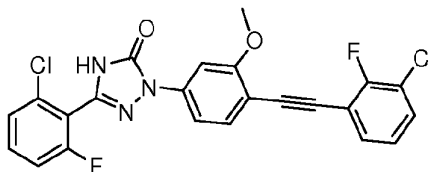


The title compound was prepared according to the procedure described in Example-3 using 3-(2-chloro-6-fluorophenyl)-1-(4-iodo-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-32, 0.050 g, 0.11 mmol), 1,4-dichloro-2-ethynylbenzene (Intermediate-23, 0.029 g, 0.16 mmol), TBAF (0.101 g, 0.33 mmol), bis(triphenylphosphine)palladium(II) chloride (0.005 g, 0.04 mmol) and DMSO (3.0 mL) at 80°C to afford 0.010 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ

3.89 (s, 3H), 7.49-7.56 (m, 3H), 7.59-7.73 (m, 6H), 12.90 (br s, 1H); MS (m/z): 487.95 (M+H⁺).

Example-60

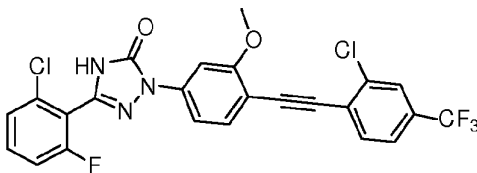
1-(4-((3-Chloro-2-fluorophenyl)ethynyl)-3-methoxyphenyl)-3-(2-chloro-6-fluorophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



The title compound was prepared according to the procedure described in Example-3 using 3-(2-chloro-6-fluorophenyl)-1-(4-iodo-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-32, 0.050 g, 0.11 mmol), 1-chloro-3-ethynyl-2-fluorobenzene (Intermediate-24, 0.033 g, 0.22 mmol), TBAF (0.080 g, 0.30 mmol), bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.022 mmol) and DMSO (3.0 ml) at 80°C to afford 0.015 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.89 (s, 3H), 7.28 (t, *J* = 7.8 Hz, 1H), 7.48 (t, *J* = 9.0 Hz, 1H), 7.56-7.70 (m, 7H), 12.79 (m, 1H); MS (m/z): 470.00 (M-H).

Example-61

1-(4-((2-Chloro-4-(trifluoromethyl)phenyl)ethynyl)-3-methoxyphenyl)-3-(2-chloro-6-fluorophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one

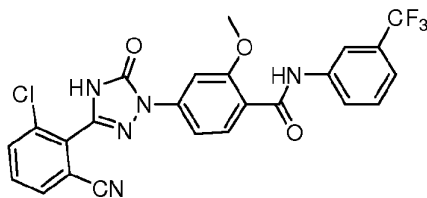


The title compound was prepared according to the procedure described in Example-3 using 3-(2-chloro-6-fluorophenyl)-1-(4-iodo-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-32, 0.050 g, 0.11 mmol), 2-chloro-1-ethynyl-4-(trifluoromethyl)benzene (Intermediate-25, 0.044 g, 0.22 mmol), TBAF (0.080 g, 0.30 mmol), bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.022 mmol) and DMSO (3.0 mL) at 80°C to afford 0.012 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ

3.89 (s, 3H), 7.52 (t, $J = 9.0$ Hz, 1H), 7.56-7.75 (m, 6H), 7.83 (d, $J = 8.1$ Hz, 1H), 8.02 (s, 1H), 12.78 (s, 1H); MS (m/z): 522.00 (M+H)⁺.

Example-62

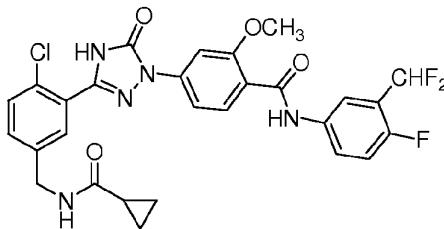
4-(3-(2-Chloro-6-cyanophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide



To a solution of 4-(3-(2-chloro-6-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide (Intermediate-34, 0.130 g, 0.211 mmol) in DMF (5 mL) was added CuCN (0.020 g, 0.232 mmol) and reaction mixture was heated at 80°C for 4-5 h. The reaction mass was quenched with aq. KMnO₄ solution and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 0.015 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.95 (s, 3H), 7.44 (d, $J = 7.8$ Hz, 1H), 7.59 (t, $J = 8.4$ Hz, 1H), 7.68 (d, $J = 8.1$ Hz, 1H), 7.80-7.86 (m, 3H), 7.95 (d, $J = 8.7$ Hz, 1H), 8.06-8.12 (m, 2H), 8.25 (s, 1H), 10.38 (s, 1H), 13.00 (br s, 1H); MS (m/z): 512.14 (M-H)⁻.

Example-63

4-(3-(2-Chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(difluoromethyl)-4-fluorophenyl)-2-methoxybenzamide

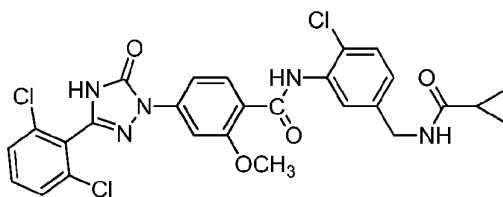


The title compound was prepared by following the procedure as described for step-2 of Intermediate-30, by using 4-(3-(5-(aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(difluoromethyl)-4-fluorophenyl)-2-methoxybenzamide (Intermediate-35, 0.050 g, 0.09 mmol), cyclopropylcarbonyl chloride (0.016 g, 0.14

mmol), DIPEA (2.0 mL) and THF (5 mL) to afford 0.015 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.67 (s, 4H), 1.59 (m, 1H), 3.92 (s, 3H), 3.32 (d, J = 4.8 Hz, 2H), 7.03-7.33 (m, 1H), 7.35-7.47 (m, 3H), 7.62-7.68 (m, 2H), 7.76-7.84 (m, 2H), 8.09 (m, 2H), 8.70 (m, 1H), 10.28 (s, 1H), 12.69 (brs, 1H); MS (m/z): 586.12 (M+H) $^+$.

Example-64

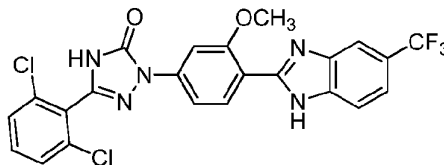
N-(2-Chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzamide



The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-21, 0.100 g, 0.25 mmol), *N*-(3-amino-4-chlorobenzyl)cyclopropanecarboxamide (Intermediate-36, 0.085 g, 0.38 mmol), trimethyl aluminium (2M solution in toluene) (1 mL) and dry toluene (5.0 mL) to afford 0.020 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.68 (d, J = 8.1 Hz, 4H), 1.61 (m, 1H), 4.11 (s, 3H), 4.28 (d, J = 5.4 Hz, 2H), 7.04 (d, J = 6.9 Hz, 1H), 7.52 (d, J = 8.7 Hz, 1H), 7.71-7.79 (m, 4H), 7.88 (s, 1H), 8.19 (d, J = 9.0 Hz, 1H), 8.44 (s, 1H), 8.67 (m, 1H), 10.53 (s, 1H), 12.82 (br s, 1H); MS (m/z): 587.96 (M+H) $^+$.

Example-65

3-(2,6-Dichlorophenyl)-1-(3-methoxy-4-(5-(trifluoromethyl)-1*H*-benzo[d]imidazol-2-yl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one

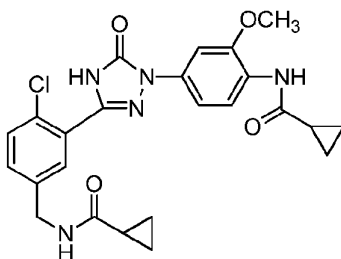


The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-21, 0.150 g, 0.30 mmol), 4-(trifluoromethyl)benzene-1,2-diamine (0.081 g, 0.45 mmol), trimethyl aluminium (2M solution in toluene) (2 mL),

dry toluene (5.0 mL) to afford 0.015 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 4.09 (s, 3H), 7.52 (s, 1H), 7.73-7.85 (m, 5H), 8.02-8.15 (m, 2H), 8.48 (m, 1H), 12.79 (br s, 1H), 13.33 (br s, 1H); MS (m/z): 520.14 (M+H) $^+$.

Example-66

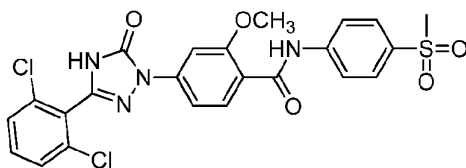
N-(4-Chloro-3-(1-(4-(cyclopropanecarboxamido)-3-methoxyphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)cyclopropanecarboxamide



The title compound was prepared by following the procedure as described for step-2 of Intermediate-30 by using *N*-(3-(1-(4-amino-3-methoxyphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-4-chlorobenzyl)cyclopropanecarboxamide (Intermediate-37, 0.050 g, 0.120 mmol), cyclopropylcarbonyl chloride (0.018 g, 0.180 mmol), DIPEA (2.0 mL) and THF (5 mL) to afford 0.015 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.69 (s, 4H), 0.77 (s, 4H), 1.60 (m, 1H), 2.08 (m, 1H), 3.88 (s, 3H), 4.33 (d, J = 5.4 Hz, 2H), 7.45 (m, 2H), 7.62 (s, 2H), 7.96 (d, J = 8.4 Hz, 1H), 8.69 (m, 1H), 9.49 (s, 1H), 12.54 (s, 1H); MS (m/z): 482.10 (M+H) $^+$.

Example-67

4-(3-(2,6-Dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(4-(methylsulfonyl)phenyl)benzamide

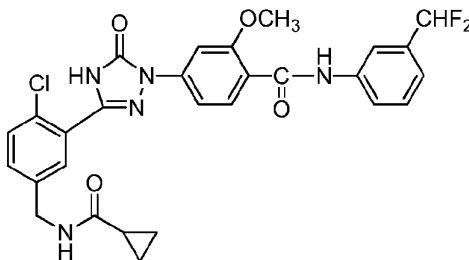


To a solution of 4-(3-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(4-(methylthio)phenyl)benzamide (Intermediate-38, 0.070 g, 0.13 mmol) in mixture of acetonitrile: water (3 mL:3 mL) was added oxone (0.053 g, 0.34 mmol). The reaction mass was stirred at RT for 24 h. The reaction mass was diluted with water and

extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 0.020 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.18 (s, 3H), 3.94 (s, 3H), 7.68-7.76 (m, 7H), 7.80 (d, J = 8.7 Hz, 1H), 7.90 (d, J = 8.7 Hz, 1H), 7.99 (d, J = 8.4 Hz, 1H), 10.48 (s, 1H), 12.76 (br s, 1H); MS (m/z): 531.14 (M+H) $^+$.

Example-68

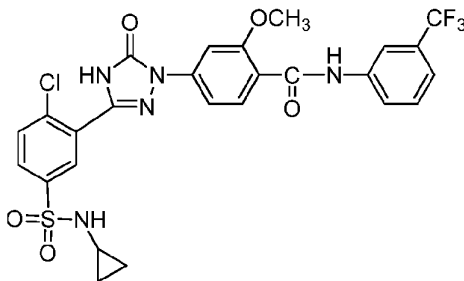
4-(3-(2-Chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-N-(3-(difluoromethyl)phenyl)-2-methoxybenzamide



The title compound was prepared by following the procedure as described for step-2 of Intermediate-30 by using 4-(3-(5-(aminomethyl)-2-chlorophenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-N-(3-(difluoromethyl)phenyl)-2-methoxybenzamide (Intermediate-39, 0.050 g, 0.10 mmol), cyclopropane carbonyl chloride (0.016 g, 0.15 mmol), DIPEA (2.0 mL) and THF (2 mL) to afford 0.015 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.68 (d, J = 8.7 Hz, 4H), 1.60 (m, 1H), 3.95 (s, 3H), 4.34 (d, J = 5.4 Hz, 2H), 7.05 (m, 1H), 7.29 (d, J = 7.9 Hz, 1H), 7.41-7.55 (m, 4H), 7.65 (d, J = 9.0 Hz, 1H), 7.74 (s, 1H), 7.82 (d, J = 8.7 Hz, 2H), 8.07 (s, 1H), 8.72 (m, 1H), 10.33 (s, 1H); MS (m/z): 566.29 (M-H) $^-$.

Example-69

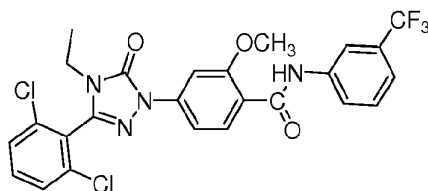
4-(3-(2-Chloro-5-(N-cyclopropylsulfamoyl)phenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-2-methoxy-N-(3-(trifluoromethyl)phenyl)benzamide



The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2-chloro-5-(*N*-cyclopropylsulfamoyl) phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-40, 0.040 g, 0.08 mmol), 3-(trifluoromethyl)aniline (0.020 g, 0.12 mmol), trimethyl aluminium (2M solution in toluene) (1 ml) and dry toluene (5.0 mL) to afford 0.015 g of crude product, which was purified by washing with methanol: DCM: DEE to afford 0.015 g of pure product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.42 (s, 2H), 0.54 (d, *J* = 5.4 Hz, 2H), 2.20 (m, 1H), 3.96 (s, 3H), 7.45 (d, *J* = 7.5 Hz, 1H), 7.61 (t, 1H), 7.70 (d, *J* = 7.8 Hz, 1H), 7.80-7.84 (m, 2H), 7.96 (s, 3H), 8.18 (m, 2H), 8.26 (s, 1H), 10.40 (s, 1H), 12.88 (br s, 1H); MS (*m/z*): 606.13(M-H)⁻.

Example-70

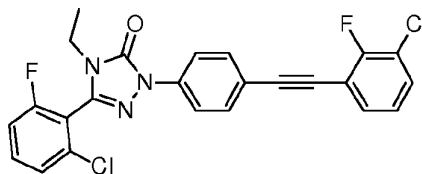
4-(3-(2,6-Dichlorophenyl)-4-ethyl-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide



The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2,6-dichlorophenyl)-4-ethyl-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxybenzoate (Intermediate-41, 0.040 g, 0.09 mmol), 3-(trifluoromethyl)aniline (0.010 g, 0.6 mmol), trimethyl aluminium (2M solution in toluene) (1 mL) and dry toluene (5.0 mL) to afford 0.015 g of crude product which was purified by column chromatography in basic alumina eluting with 0.2 % methanol: DCM to afford 0.015 g of pure product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.14 (t, 3H), 3.55-3.57 (m, 2H), 3.95 (s, 3H), 7.46 (m, 1H), 7.59 (t, 1H), 7.69-7.81 (m, 6H), 7.94 (m, 1H), 8.25 (s, 1H), 10.38 (s, 1H); MS (*m/z*): 551.11 (M+H)⁺.

Example-71

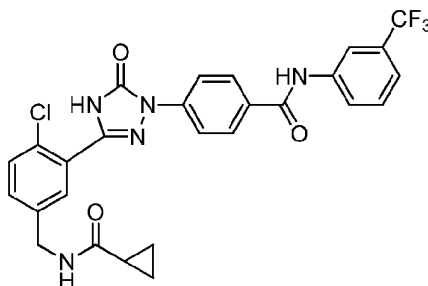
1-(4-((3-Chloro-2-fluorophenyl)ethynyl)phenyl)-3-(2-chloro-6-fluorophenyl)-4-ethyl-1*H*-1,2,4-triazol-5(4*H*)-one



The title compound was prepared by following the procedure as described for Intermediate-41 by using 1-(4-((3-chloro-2-fluorophenyl)ethynyl)phenyl)-3-(2-chloro-6-fluorophenyl)-1H-1,2,4-triazol-5-one (Example-12, 0.070 g, 0.15 mmol), dry DMF (3 mL), NaH (0.010 g, 0.23 mmol) and ethyl bromide (0.026 g, 0.23 mmol) to afford 0.030 g of the crude product which was purified by column chromatography in basic alumina eluting with 0.2 % methanol: DCM to afford 0.030 g of pure product. ^1H NMR (300 MHz, DMSO d_6): δ 1.09 (t, 3H), 3.56 (m, 2H), 7.29 (t, 1H), 7.52-7.84 (m, 7H), 8.04 (d, J = 8.1 Hz, 2H); MS (m/z): 470.06 ($M+H$) $^+$.

Example-72

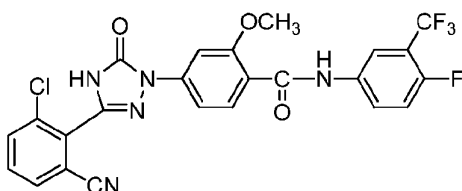
4-(3-(2-Chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)-N-(3-(trifluoromethyl)phenyl)benzamide



The title compound was prepared by following the procedure as described for Example-31 by using methyl 4-(3-(2-chloro-5-(cyclopropane carboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1H-1,2,4-triazol-1-yl)benzoate (Intermediate-42, 0.100 g, 0.24 mmol), 3-(trifluoromethyl)aniline (0.050 g, 0.36 mmol), trimethyl aluminium (2M solution in toluene) (1 mL) and dry toluene (8.0 mL). The reaction mass was quenched in water, extracted with DCM and the organic layer was concentrated to afford 0.010 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.70 (s, 4H), 1.62 (m, 1H), 4.33 (d, J = 6.3 Hz, 2H), 7.44-7.67 (m, 4H), 8.10-8.26 (m, 6H), 8.68 (m, 1H), 10.53 (s, 1H), 12.68 (br s, 1H); MS (m/z): 554.30 ($M-H$) $^-$.

Example-73

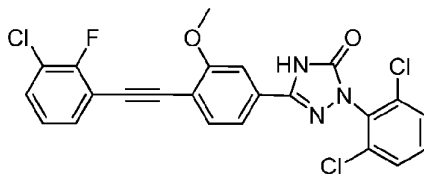
4-(3-(2-Chloro-6-cyanophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(4-fluoro-3-(trifluoromethyl)phenyl)-2-methoxybenzamide



The title compound was prepared by following the procedure as described for Example-62 by using 4-(3-(2-chloro-6-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(4-fluoro-3-(trifluoromethyl)phenyl)-2-methoxybenzamide (Intermediate-43, 0.130 g, 0.21 mmol), DMF (4 mL) and CuCN (0.020 g, 0.23 mmol) to afford 0.030 g of crude product which was further purified by column chromatography in basic alumina eluting with 50 % methanol: DCM and few drops of 10 % ammonia to afford 0.030 g of pure product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.94 (s, 3H), 7.52 (t, 1H), 7.68 (d, *J* = 7.8 Hz, 1H), 7.81 (m, 3H), 8.05-8.12 (m, 3H), 8.26 (s, 1H), 10.40 (s, 1H), 13.01 (br s, 1H); MS (*m/z*): 532.12 (*M*+*H*)⁺.

Example-74

3-(4-((3-Chloro-2-fluorophenyl)ethynyl)-3-methoxyphenyl)-1-(2,6-dichlorophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one

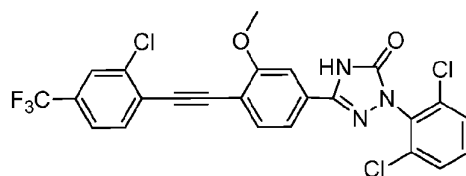


To a solution of 1-(2,6-dichlorophenyl)-3-(4-iodo-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-45, 0.070 g, 0.15 mmol) in DMSO (3.0 mL) was added 1-chloro-3-ethynyl-2-fluorobenzene (Intermediate-24, 0.035 g, 0.22 mmol), TBAF (0.080 g, 0.30 mmol) and bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.022 mmol). The reaction mass was stirred at 80°C for 5-6 h. The reaction mass was quenched in water and extracted with DCM and concentrated. The obtained product was purified with column chromatography on silica gel eluting with 2.0 % EA: DCM to afford 0.020 g of

the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.92 (s, 3H), 7.29 (t, J = 8.4 Hz, 1H), 7.48-7.72 (m, 8H), 12- 13 (br s, 1H); MS (m/z): 488.00 (M^+).

Example-75

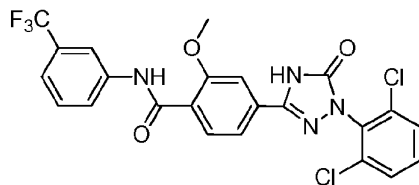
3-(4-((2-Chloro-4-(trifluoromethyl)phenyl)ethynyl)-3-methoxyphenyl)-1-(2,6-dichlorophenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



The title compound was prepared according to the procedure described in Example-3 using 1-(2,6-dichlorophenyl)-3-(4-iodo-3-methoxyphenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-45, 0.070 g, 0.15 mmol), 2-chloro-1-ethynyl-4-(trifluoromethyl)benzene (Intermediate-25, 0.046 g, 0.22 mmol), TBAF (0.080 g, 0.30 mmol), bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.022 mmol) and DMSO (3.0 mL) at 80°C to afford 0.020 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.95 (s, 3H), 7.50-7.53 (m, 2H), 7.56-7.79 (m, 5H), 7.88 (d, J = 8.4 Hz, 1H), 8.05 (s, 1H), 12.04 (s, 1H); MS (m/z): 538.07 (M^+).

Example-76

4-(1-(2,6-Dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide

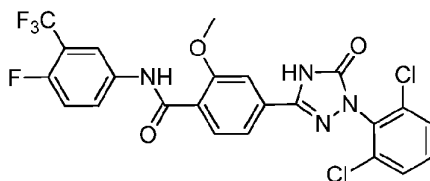


To a solution of methyl 4-(1-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-methoxybenzoate (Intermediate-46, 0.070g, 0.17 mmol) in dry toluene was added 3-(trifluoromethyl)aniline (0.043 g, 0.17mmol) followed by addition of trimethyl aluminum (2M solution in toluene) (0.5 mL). The reaction mixture was refluxed for 1 hr under inert atmosphere. The reaction mixture was brought to RT and quenched with

water. Few drops of dilute HCl were added and the reaction mixture was extracted with DCM and concentrated. The obtained product was purified with column chromatography to afford 0.050 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.95 (s, 3H), 7.46 (d, J = 6.9 Hz, 1H), 7.54-7.65 (m, 4H), 7.72-7.76 (m, 3H), 7.94 (d, J = 8.4 Hz, 1H), 8.24 (s, 1H), 10.51 (s, 1H), 12.85 (s, 1H); MS (m/z): 523.04 (M^+).

Example-77

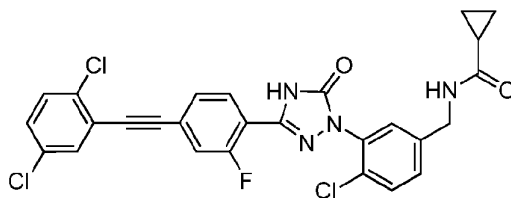
4-(1-(2,6-Dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-*N*-(4-fluoro-3-(trifluoromethyl)phenyl)-2-methoxybenzamide



The title compound was prepared by following the procedure as described for step-6 of Intermediate-26 by using methyl 4-(1-(2,6-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-methoxybenzoate (Intermediate-46, 0.070g, 0.17 mmol), 4-fluoro-3-(trifluoromethyl)aniline (0.048 g, 0.26 mmol), trimethyl aluminum (2M solution in toluene) (0.5 mL) and dry toluene (5.0 mL) to afford 0.040 g of desired product. ^1H NMR (300 MHz, DMSO d_6): δ 3.95 (s, 3H), 7.49-7.64 (m, 4H), 7.72-7.76 (m, 3H), 7.99 (m, 1H), 8.25 (m, 1H), 10.52 (s, 1H), 12.86 (s, 1H); MS (m/z): 541.05 (M^+).

Example-78

N-(4-Chloro-3-(3-(4-((2,5-dichlorophenyl)ethynyl)-2-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide

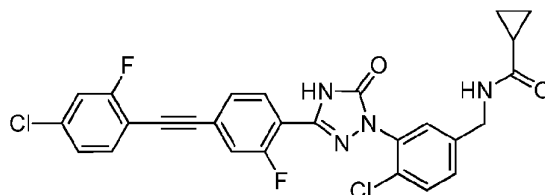


The title compound was prepared according to the procedure described in Example-3 by using *N*-(4-chloro-3-(3-(2-fluoro-4-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide (Intermediate-48, 0.060 g, 0.10 mmol), 1,4-dichloro-2-ethynylbenzene (Intermediate-23, 0.026 g, 0.15 mmol), TBAF (0.064 g, 0.20

mmol), bis(triphenylphosphine)palladium(II) chloride (0.064 g, 0.20 mmol) and DMSO (3.0 mL) at 80°C to afford 0.020 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.68 (m, 4H), 1.60 (m, 1H), 4.33 (d, *J* = 5.7 Hz, 2H), 7.38 (d, *J* = 7.8 Hz, 1H), 7.47 (s, 1H), 7.57-7.69 (m, 5H), 7.86-7.94 (m, 2H), 8.68 (m, 1H), 12.45 (br s, 1H); MS (*m/z*): 557.05 (*M*+H⁺).

Example-79

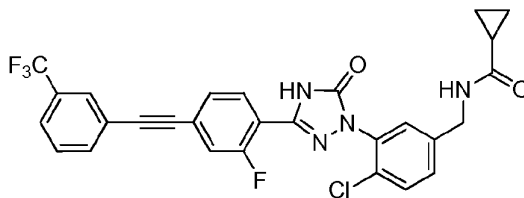
N-(4-Chloro-3-(3-(4-((4-chloro-2-fluorophenyl)ethynyl)-2-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide



The title compound was prepared according to the procedure described in Example-3 using *N*-(4-chloro-3-(3-(2-fluoro-4-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl) cyclopropanecarboxamide (Intermediate-48, 0.050 g, 0.097 mmol), 4-chloro-1-ethynyl-2-fluorobenzene (Intermediate-49, 0.022 g, 0.14 mmol), TBAF (0.061 g, 0.19 mmol), bis(triphenylphosphine)palladium(II) chloride (catalytic) and DMSO (1.0 mL) at 80°C to afford 0.025 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.65 - 0.68 (m, 4H), 1.60 (m, 1H), 4.32 (d, *J* = 6.0 Hz, 2H), 7.33-7.52 (m, 3H), 7.55-7.74 (m, 5H), 7.90 (m, 1H), 8.68 (m, 1H), 12.0 (br s, 1H); MS (*m/z*): 539.0 (*M*⁺).

Example-80

N-(4-Chloro-3-(3-(2-fluoro-4-((3-(trifluoromethyl)phenyl)ethynyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide

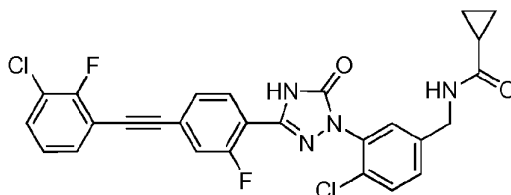


The title compound was prepared according to the procedure described in Example-3 by using *N*-(4-chloro-3-(3-(2-fluoro-4-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl) cyclopropanecarboxamide (Intermediate-48, 0.050 g, 0.097 mmol), 1-ethynyl-

3-(trifluoromethyl)benzene (Intermediate-50, 0.025 g, 0.14 mmol), TBAF (0.061 g, 0.19 mmol), bis(triphenylphosphine)palladium(II) chloride (catalytic) and DMSO (1.0 mL) at 80°C to afford 0.020 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.65-0.70 (m, 4H), 1.60 (m, 1H), 4.31 (d, *J* = 6.0 Hz, 2H), 7.28-7.41 (m, 2H), 7.51-7.61 (m, 4H), 7.67-7.97 (m, 4H), 8.68 (m, 1H), 12.0 (br s, 1H).; MS (*m/z*): 555.05 (M+H⁺).

Example-81

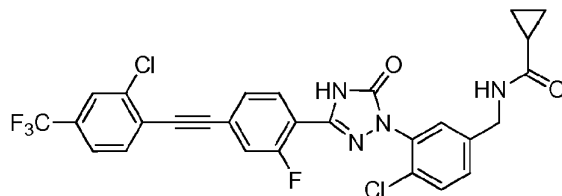
N-(4-Chloro-3-(3-(4-((3-chloro-2-fluorophenyl)ethynyl)-2-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide



The title compound was prepared according to the procedure described in Example-3 using *N*-(4-chloro-3-(3-(2-fluoro-4-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl) cyclopropanecarboxamide (Intermediate-48, 0.050 g, 0.097 mmol), 1-chloro-3-ethynyl-2-fluorobenzene (Intermediate-24, 0.022 g, 0.14 mmol), TBAF (0.061 g, 0.19 mmol), bis(triphenylphosphine)palladium(II) chloride (catalytic) and DMSO (1.0 mL) at 80°C to afford 0.030 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.69 (m, 4H), 1.60 (m, 1H), 4.33 (d, *J* = 5.7 Hz, 2H), 7.30-7.41 (m, 3H), 7.48-7.74 (m, 5H), 7.87-7.93 (m, 1H), 8.69 (m, 1H), 12.40- 12.58 (br s, 1H); MS (*m/z*): 539.36 (M⁺).

Example-82

N-(4-Chloro-3-(3-(4-((2-chloro-4-(trifluoromethyl)phenyl)ethynyl)-2-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide

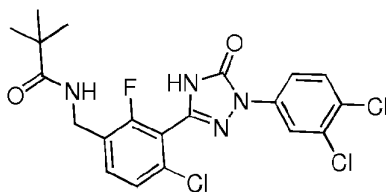


The title compound was prepared according to the procedure described in Example-3 by using *N*-(4-chloro-3-(3-(2-fluoro-4-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl) cyclopropanecarboxamide (Intermediate-48, 0.060 g, 0.117 mmol), 2-chloro-

1-ethynyl-4-(trifluoromethyl)benzene (Intermediate-25, 0.037 g, 0.175 mmol), TBAF (0.073 g, 0.234 mmol), bis(triphenylphosphine)palladium(II) chloride (catalytic) and DMSO (1.0 mL) at 80°C to afford 0.030 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.66- 0.69 (m, 4H), 1.60 (m, 1H), 4.33 (d, *J* = 6.0 Hz, 2H), 7.40 (d, *J* = 8.4 Hz, 1H), 7.48 (s, 1H), 7.62-7.64 (m, 2H), 7.73 (d, *J* = 11.4 Hz, 1H), 7.82 (d, *J* = 7.8 Hz, 1H), 7.92-7.97 (m, 2H), 8.09 (s, 1H), 8.69 (m, 1H), 12.00 (br s, 1H); MS (*m/z*): 589.14 (*M*⁺).

Example-83

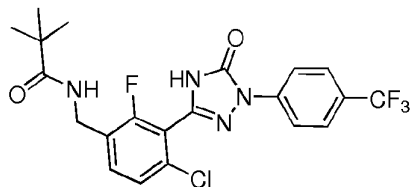
N-(4-Chloro-3-(1-(3,4-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide



To a solution of *tert*-butyl 2-(3,4-dichlorophenyl)hydrazinecarboxylate (Intermediate-52, 0.060 g, 0.21 mmol) in DCM (10 mL) was added solution of 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoyl isocyanate (Intermediate-51, 0.101 g, 0.32 mmol) in DCM and the reaction mixture was stirred for 20 h at room temperature, followed by addition of TFA (3 mL) and further stirred for 20 h at room temperature. The reaction mass was quenched in water, extracted with DCM and concentrated to afford crude product which was purified by column chromatography eluting with MeOH: DCM to afford 0.007 g of pure product. ¹H NMR (300 MHz, CDCl₃): δ 1.20 (s, 9H), 4.44 (d, *J* = 5.4 Hz, 2H), 6.20 (m, 1H), 7.29 (m, 1H), 7.41-7.49 (m, 2H), 7.91 (d, *J* = 8.1 Hz, 1H), 8.15 (s, 1H); MS (*m/z*): 471.26 (*M*⁺).

Example-84

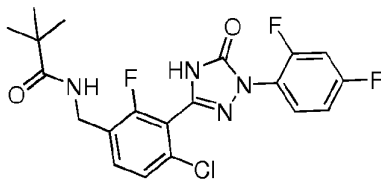
N-(4-Chloro-2-fluoro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(4-(trifluoromethyl)phenyl)hydrazinecarboxylate (Intermediate-53, 0.060 g, 0.21 mmol), 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoyl isocyanate (Intermediate-51, 0.101 g, 0.32 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.025 g of pure product. ^1H NMR (300 MHz, CDCl_3): δ 1.19 (s, 9H), 4.40 (m, 2H), 6.22 (m, 1H), 7.26 (m, 1H), 7.35 (m, 1H), 7.62 (d, $J = 8.1$ Hz, 2H), 8.12 (d, $J = 7.8$ Hz, 2H); MS (m/z): 471.18 ($\text{M}+\text{H}^+$).

Example-85

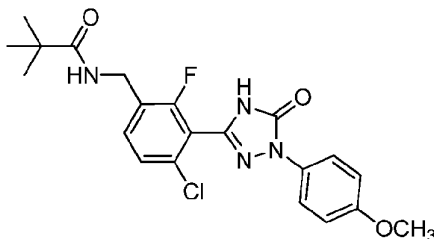
N-(4-Chloro-3-(1-(2,4-difluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(2,4-difluorophenyl)hydrazinecarboxylate (Intermediate-54, 0.060 g, 0.21 mmol), 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoyl isocyanate (Intermediate-51, 0.101 g, 0.32 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.018 g of pure product. ^1H NMR (300 MHz, CDCl_3): δ 1.16 (s, 9H), 4.38 (d, $J = 5.4$ Hz, 2H), 6.25 (m, 1H), 6.79- 6.82 (m, 1H), 6.90- 6.96 (m, 2H), 7.20-7.32 (m, 2H), 7.50-7.53 (m, 1H); MS (m/z): 437.26 ($\text{M}-\text{H}$).

Example-86

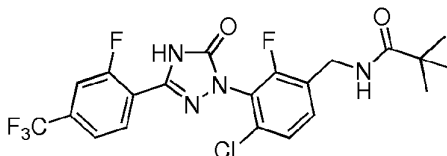
N-(4-Chloro-2-fluoro-3-(1-(4-methoxyphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(4-methoxyphenyl)hydrazinecarboxylate (Intermediate-55, 0.060 g, 0.21 mmol), 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoyl isocyanate (Intermediate-51, 0.101 g, 0.32 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.018 g of pure product. ^1H NMR (400 MHz, CDCl_3): δ 1.20 (s, 9H), 3.82 (s, 3H), 4.46 (d, J = 5.8 Hz, 2H), 6.12 (m, 1H), 6.95 (d, J = 8.9 Hz, 2H), 7.26-7.30 (m, 1H), 7.43 (t, J = 7.9 Hz, 1H), 7.82 (d, J = 8.9 Hz, 2H), 11.01 (br s, 1H); MS (m/z): 433.20 ($\text{M}+\text{H}^+$).

Example-87

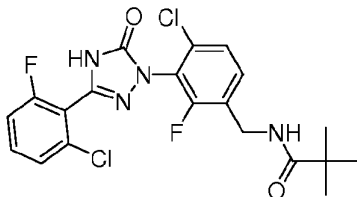
N-(4-Chloro-2-fluoro-3-(3-(2-fluoro-4-(trifluoromethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)pivalamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(6-chloro-2-fluoro-3-(pivalamidomethyl)phenyl)hydrazinecarboxylate (Intermediate-57, 0.050 g, 0.13 mmol), 2-fluoro-4-(trifluoromethyl)benzoyl isocyanate (Intermediate-56, 0.10 g, 0.26 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.030 g of pure product. ^1H NMR (400 MHz, CDCl_3): δ 1.20 (s, 9H), 4.48 (d, J = 5.7 Hz, 2H), 6.13 (m, 1H), 7.31 (d, J = 8.5 Hz, 1H), 7.41-7.51 (m, 3H), 8.12 (t, J = 7.9 Hz, 1H); MS (m/z): 489.20 ($\text{M}+\text{H}^+$).

Example-88

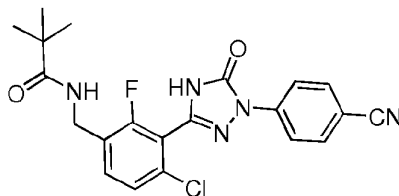
N-(4-Chloro-3-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-fluorobenzyl)pivalamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(6-chloro-2-fluoro-3-(pivalamidomethyl)phenyl)hydrazinecarboxylate (Intermediate-57, 0.050 g, 0.13 mmol), 2-chloro-6-fluorobenzoyl isocyanate (Intermediate-8, 0.060 g, 0.26 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.025 g of pure product. ^1H NMR (400 MHz, CDCl_3): δ 1.19 (s, 9H), 4.48 (m, 2H), 6.09 (m, 1H), 7.15 (t, $J = 8.4$ Hz, 1H), 7.32 (t, $J = 8.2$ Hz, 2H), 7.40-7.45 (m, 2H), 10.66 (br s, 1H); MS (m/z): 455.21 (M^+).

Example-89

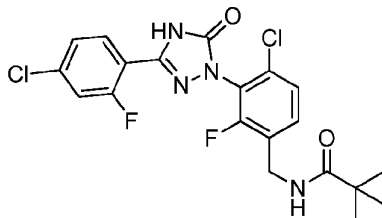
N-(4-chloro-3-(1-(4-cyanophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(4-cyanophenyl)hydrazinecarboxylate (step-2 of Intermediate17, 0.060 g, 0.25 mmol), 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoyl isocyanate (Intermediate-51, 0.160 g, 0.51 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.020 g of pure product. ^1H NMR (400 MHz, CDCl_3): δ 1.20 (s, 9H), 4.45 (d, $J = 5.9$ Hz, 2H), 6.19 (m, 1H), 7.32 (d, $J = 8.4$ Hz, 1H), 7.43 (t, $J = 7.9$ Hz, 1H), 7.73 (d, $J = 8.8$ Hz, 2H), 8.19 (d, $J = 8.4$ Hz, 2H), 10.72 (br s, 1H); MS (m/z): 426.27 (M-H^-).

Example-90

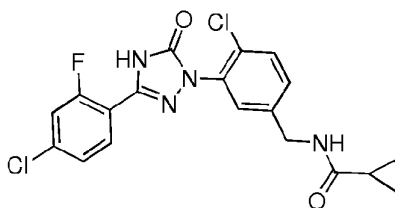
N-(4-Chloro-3-(3-(4-chloro-2-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-fluorobenzyl)pivalamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(6-chloro-2-fluoro-3-(pivalamidomethyl)phenyl)hydrazinecarboxylate (Intermediate-57, 0.050 g, 0.13 mmol), 4-chloro-2-fluorobenzoyl isocyanate (Intermediate-58, 0.060 g, 0.26 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.025 g of desired product. ^1H NMR (400 MHz, CDCl_3): δ 1.21 (s, 9H), 4.50 (t, $J = 7.2$ Hz, 2H), 6.07 (m, 1H), 7.22- 7.26 (m, 2H), 7.32 (d, $J = 8.4$ Hz, 1H), 7.44 (t, $J = 8.1$ Hz, 1H), 7.93 (t, $J = 8.6$ Hz, 1H); MS (m/z): 455.08 (M^+).

Example-91

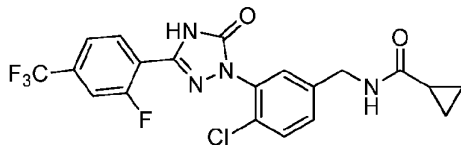
N-(4-Chloro-3-(3-(4-chloro-2-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl)hydrazinecarboxylate (step-7 of Intermediate-48, 0.060 g, 0.17 mmol), 4-chloro-2-fluorobenzoyl isocyanate (Intermediate-58, 0.035 g, 0.71 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.040 g of desired product. ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ 0.67-0.69 (m, 4H), 1.59 – 1.60 (m, 1H), 4.31 (d, $J = 5.9$ Hz, 2H), 7.36 (dd, $J = 6.24$ Hz, 1H), 7.45 (dd, $J = 8.5$ Hz, 2H), 7.59 (d, $J = 8.3$ Hz, 1H), 7.65 (d, $J = 8.8$ Hz, 1H), 7.83 (t, $J = 8.3$ Hz, 1H), 7.64 (t, $J = 6$ Hz, 1H), 12.51 (br s, 1H); MS (m/z): 421.08 (M^+).

Example-92

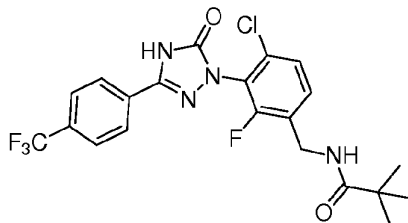
N-(4-Chloro-3-(3-(2-fluoro-4-(trifluoromethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(2-chloro-5-(cyclopropanecarboxamidomethyl)phenyl)hydrazinecarboxylate (step-7 of Intermediate-48, 0.060 g, 0.17 mmol), 2-fluoro-4-(trifluoromethyl)benzoyl isocyanate (Intermediate-56, 0.035 g, 0.82 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.045 g of desired product. ^1H NMR (400 MHz, DMSO d_6): δ 0.67 (m, 4H), 1.60 (m, 1H), 4.3 (d, J = 5.9 Hz, 2H), 7.37 (d, J = 6.4 Hz, 1H), 7.46 (s, 1H), 7.60 (d, J = 8.3 Hz, 1H), 7.74 (d, J = 8.2 Hz, 1H), 7.89 (d, J = 10.56 Hz, 1H), 8.05 (t, J = 7.7 Hz, 1H), 8.65 (t, J = 5.8 Hz, 1H), 12.6 (br s, 1H); MS (m/z): 454.98 (M^+).

Example-93

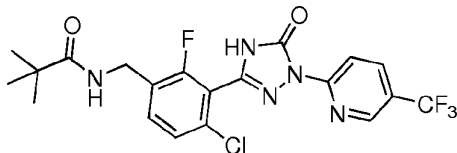
N-(4-Chloro-2-fluoro-3-(5-oxo-3-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)pivalamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(6-chloro-2-fluoro-3-(pivalamidomethyl)phenyl)hydrazinecarboxylate (Intermediate-57, 0.050 g, 0.13 mmol), 4-(trifluoromethyl)benzoyl isocyanate (Intermediate-59, 0.060 g, 0.26 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.045 g of desired product. ^1H NMR (400 MHz, DMSO d_6): δ 1.12 (s, 9H), 4.30 (d, J = 4.4 Hz, 2H), 7.41 (t, J = 8 Hz, 1H), 7.53 (d, J = 8.5 Hz, 1H), 7.91 (d, J = 8.4 Hz, 2H), 8.0 (d, J = 8.2 Hz, 2H), 8.17 (t, J = 5.8 Hz, 1H), 12.9 (br s, 1H); MS (m/z): 471.10 ($M+H^+$).

Example-94

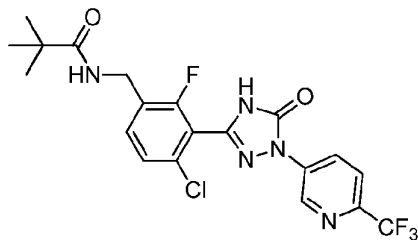
N-(4-Chloro-2-fluoro-3-(5-oxo-1-(5-(trifluoromethyl)pyridin-2-yl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(5-(trifluoromethyl)pyridin-2-yl)hydrazinecarboxylate (Intermediate-60, 0.050 g, 0.18 mmol), 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoyl isocyanate (Intermediate-51, 0.120 g, 0.36 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.012 g of desired product. ^1H NMR (400 MHz, DMSO d_6): δ 1.13 (s, 9H), 4.30 (d, J = 5.6 Hz, 2H), 7.11 (br s, 1H), 7.46 (t, J = 8 Hz, 1H), 7.52 (d, J = 8.8 Hz, 1H), 8.14 (m, 1H), 8.28 (d, J = 8.8 Hz, 1H), 8.35 (dd, J = 7.2 Hz, 1H), 8.87 (s, 1H); MS (m/z): 472.24 ($M+H^+$).

Example-95

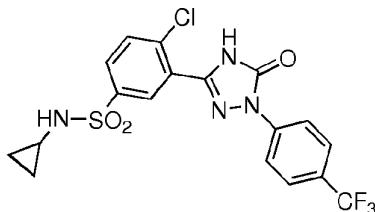
N-(4-Chloro-2-fluoro-3-(5-oxo-1-(6-(trifluoromethyl)pyridin-3-yl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(6-(trifluoromethyl)pyridin-3-yl) hydrazine carboxylate (Intermediate-61, 0.050 g, 0.18 mmol), 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoyl isocyanate (Intermediate-51, 0.120 g, 0.36 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.013 g of desired product. ^1H NMR (400 MHz, DMSO d_6): δ 1.13 (s, 9H), 4.30 (d, J = 5.6 Hz, 2H), 7.11 (br s, 1H), 7.46 (t, J = 8 Hz, 1H), 7.53 (d, J = 8.4 Hz, 1H), 8.02 (d, J = 8.4 Hz, 1H), 8.16 (m, 1H), 8.56 (dd, J = 6.4 Hz, 1H), 9.30 (m, 1H); MS (m/z): 472.26 ($M+H^+$).

Example-96

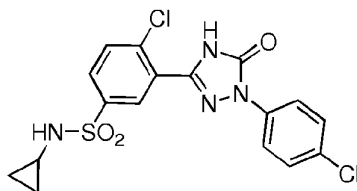
4-Chloro-*N*-cyclopropyl-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzenesulfonamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(4-(trifluoromethyl)phenyl)hydrazinecarboxylate (Intermediate-53, 0.050 g, 0.18 mmol), 2-chloro-5-(*N*-cyclopropylsulfamoyl)benzoyl isocyanate (step-4 of Intermediate-40, 0.108 g, 0.36 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.050 g of desired product. ^1H NMR (400 MHz, DMSO d_6): δ 0.40-0.42 (m, 2H), 0.50-0.54 (m, 2H), 2.17-2.21 (m, 1H), 7.85 (d, J = 8.8 Hz, 2H), 7.92-7.98 (m, 2H), 8.15-8.21 (m, 4H), 12.84 (s, 1H); MS (m/z): 457.15 ($M+H^+$).

Example-97

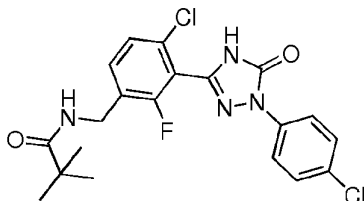
4-Chloro-3-(1-(4-chlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-*N*-cyclopropylbenzenesulfonamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(4-chlorophenyl)hydrazinecarboxylate (Intermediate-62, 0.050 g, 0.20 mmol), 2-chloro-5-(*N*-cyclopropylsulfamoyl)benzoyl isocyanate (step-4 of Intermediate-40, 0.123 g, 0.41 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.025 g of pure product. ^1H NMR (400 MHz, DMSO d_6): δ 0.39-0.41 (m, 2H), 0.49-0.52 (m, 2H), 2.17-2.20 (m, 1H), 7.52-7.55 (d, J = 8.0 Hz, 2H), 7.91-8.00 (m, 5H), 8.14 (m, 1H), 12.60 (br s, 1H); MS (m/z): 425.10 (M^+).

Example-98

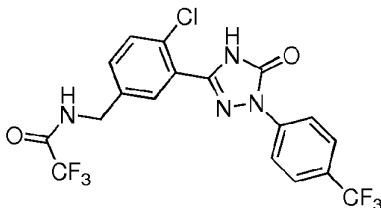
N-(4-Chloro-3-(1-(4-chlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide



The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(4-chlorophenyl)hydrazinecarboxylate (Intermediate-62, 0.050 g, 0.20 mmol), 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoyl isocyanate (Intermediate-51, 0.130 g, 0.41 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.040 g of pure product. ^1H NMR (400 MHz, DMSO d_6): δ 1.12 (s, 9H), 4.30 (d, J = 6.0 Hz, 2H), 7.24-7.47 (m, 2H), 7.52 (d, J = 7.2 Hz, 2H), 7.94 (d, J = 8.8 Hz, 2H), 8.16 (m, 1H), 12.10 (br s, 1H); MS (m/z): 437.16 (M^+).

Example-99

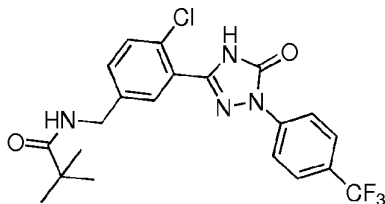
N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



The title compound was prepared according to the procedure described for Example-83 by using *tert*-butyl 2-(4-(trifluoromethyl)phenyl)hydrazinecarboxylate (Intermediate-53, 1.00 g, 3.63 mmol), 2-chloro-5-((2,2,2-trifluoroacetamido)methyl) benzoyl isocyanate (step-3 of Intermediate-26, 1.30 g, 4.03 mmol), DCM (20 mL) and TFA (5.0 mL) to afford 0.700 g of the desired product. ^1H NMR (400 MHz, DMSO d_6): δ 4.44-4.46 (d, J = 6 Hz, 2H), 7.48-7.50 (d, J = 8.4 Hz, 1H), 7.65-7.68 (m, 2H), 7.84-7.86 (d, J = 8.8 Hz, 2H), 8.17-8.19 (d, J = 8.4 Hz, 2H), 10.05 (br s, 1H), 12.69 (br s, 1H); MS (m/z): 463.17 ($M-H$).

Example-100

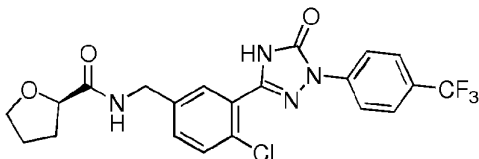
N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



To a solution of 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-63, 0.070 g, 0.189 mmol) in dry THF (5 mL) was added DIPEA (3 mL) and stirred for 20 minutes. The reaction mixture was cooled to 0°C and pivaloyl chloride (0.025 g, 0.283 mmol) was added and stirred for 3 h at RT. The reaction mass was quenched in water, extracted with DCM: MeOH and concentrated to afford of the crude product which was purified by column chromatography eluting with MeOH:DCM to afford 0.030 g of the desired title product. ¹H NMR (400 MHz, DMSO *d*₆): δ 1.12 (s, 9H), 4.28-4.30 (d, *J* = 6 Hz, 2 H), 7.39-7.42 (d, *J* = 8. Hz, 1H), 7.59-7.61 (d, *J* = 6.4 Hz, 2H), 7.84-7.86 (d, *J* = 8.8 Hz, 2H), 8.14-8.19 (m, 3H), 12.59 (br s, 1H); MS (*m/z*): 451.29 (M-H).

Example-101

(*R*)-*N*-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)tetrahydrofuran-2-carboxamide

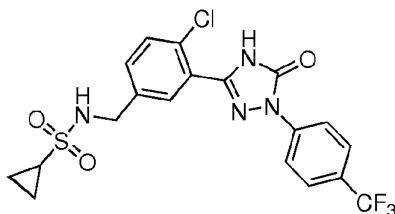


To a solution of 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-63, 0.070 g, 0.189 mmol) in dry THF: DMF (4: 1 mL) was added TBTU (0.182g, 0.567 mmol) and TEA (3 mL) and stirred for 1h under nitrogen atmosphere. To the reaction mixture (*R*)-(+)-tetrahydro-2-furoic acid (0.032 g, 0.283 mmol) was added and stirred for 18 h at room temperature. The reaction mass was quenched in water, extracted with ethyl acetate and concentrated to afford crude product which was purified by column chromatography eluting with MeOH: DCM to afford 0.025 g of the desired title product. ¹H NMR (400 MHz, DMSO *d*₆): δ 1.23 (s, 2H), 1.74-1.92 (m, 3H), 2.09-2.24, (m, 1H), 3.75-3.80 (m, 1H), 3.89-3.94 (m, 1H), 4.26-4.37 (m, 3H), 7.43-7.45 (d, *J* = 8.4 Hz, 1H), 7.61-7.63 (d, *J* = 8.4 Hz, 2H), 7.85-7.87 (d, *J* = 8.8 Hz,

2H), 8.19-8.21 (d, $J = 8.8\text{Hz}$, 2H), 8.50-8.53 (m, 1H), 12.71 (br s, 1H); MS (m/z): 465.55 ($M-H^+$).

Example-102

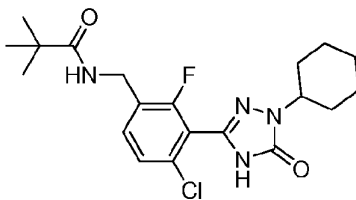
N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)cyclopropanesulfonamide



The title compound was prepared according to the procedure described in Example-100 by using 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-63, 0.070 g, 0.189 mmol), cyclopropanesulfonyl chloride (0.2 mL), DIPEA (3 mL), and dry THF (5 mL) to afford 0.029 g of pure product. ^1H NMR (400 MHz, DMSO d_6): δ 0.86-0.92 (m, 4H), 1.14-1.24 (m, 1H), 4.26-4.28 (d, $J = 6.4\text{ Hz}$, 2H), 7.57-7.60 (d, $J = 8.4\text{ Hz}$, 1H), 7.63-7.68 (d, $J = 8.4\text{Hz}$, 1H), 7.58-7.81 (m, 1H), 7.86-7.88 (d, $J = 8.8\text{ Hz}$, 2H), 8.19-8.21 (d, $J = 8.4\text{Hz}$, 2H), 12.89 (br s, 1H); MS (m/z): 471.30 ($M-H^+$).

Example-103

N-(4-Chloro-3-(1-cyclohexyl-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide

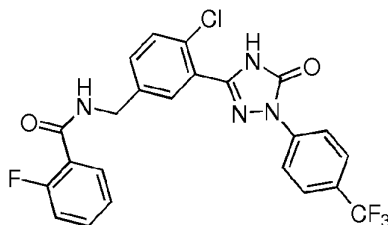


The title compound was prepared by following the procedure as described for Example-83 by using *tert*-butyl 2-(cyclohexyl)hydrazinecarboxylate (Intermediate-64, 0.050 g, 0.23 mmol), 6-chloro-2-fluoro-3-(pivalamidomethyl)benzoyl isocyanate (Intermediate-51, 0.225 g, 0.70 mmol), DCM (10 mL) and TFA (3 mL) to afford 0.015 g of pure product. ^1H NMR (400 MHz, DMSO d_6): δ 1.13 (s, 9H), 1.22-1.23 (m, 2H), 1.30-1.37 (m, 2H),

1.62-1.64 (m, 2H), 1.78-1.81 (m, 4H), 3.90-3.92 (m, 1H), 4.27-4.28 (d, $J = 4.8$ Hz, 2H), 7.37-7.41 (t, 1H), 7.46-7.48 (d, 1H), 8.17 (br s, 1H); MS (m/z): 409.36 ($M+H^+$).

Example-104

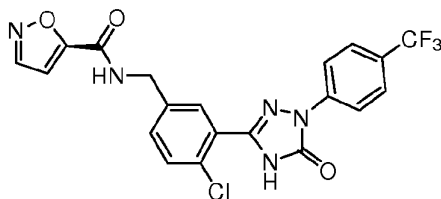
N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2-fluorobenzamide



To a solution of 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-63, 0.060 g, 0.162 mmol) in dry THF (5 mL) was added DIPEA (3 mL) and stirred for 20 minutes. The reaction mixture was cooled to 0°C and 2-fluorobenzoyl chloride (0.038 g, 0.243 mmol) was added and stirred for 3 h at room temperature. The reaction mass was quenched in water, extracted with DCM: MeOH and concentrated to afford crude product which was purified by column chromatography eluting with MeOH: DCM to afford 0.030 g of pure product. ¹H NMR (400 MHz, DMSO d_6): δ 4.50-4.52 (d, $J = 8$ Hz, 2H), 7.25-7.33 (m, 2H), 7.52-7.55 (m, 2H), 7.63-7.65 (m, 2H), 7.72-7.86 (d, $J = 2.4$ Hz, 1H), 7.83-7.86 (d, $J = 11.6$ Hz, 2H), 8.17-8.20 (d, $J = 11.6$ Hz, 2H), 8.95-8.99 (t, 1H), 12.69 (br s, 1H); MS (m/z): 491.17 ($M+H^+$).

Example-105

N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)isoxazole-5-carboxamide

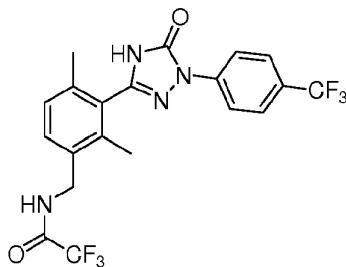


To a solution of 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-63, 0.060 g, 0.162 mmol) in dry THF (5 mL) was added DIPEA (3 mL) and stirred for 20 minutes. The reaction mixture was cooled to 0°C

and isoxazole-5-carbonyl chloride (0.032 g, 0.243 mmol) was added and stirred for 3 h at room temperature. The reaction mass was quenched in water, extracted with DCM: MeOH and concentrated to afford crude product which was purified by column chromatography eluting with MeOH: DCM to afford 0.030 g of pure product. ^1H NMR (400 MHz, DMSO d_6): δ 4.50-4.52 (d, J = 8 Hz, 2 H), 7.09-7.10 (d, J = 2 Hz, 1H), 7.43-7.45 (d, J = 8.4 Hz, 1H), 7.50-7.53 (d, J = 10.4 Hz, 1H), 7.59-7.64 (d, J = 18 Hz, 1H), 7.82-7.85 (d, J = 8.8 Hz, 2H), 8.17-8.19 (d, J = 8.4 Hz, 2H), 8.74-8.75 (d, J = 2 Hz, 1H), 9.58-9.61 (t, 1H), 12.61-12.74 (br s, 1H); MS (m/z): 464.11 (M+H $^+$).

Example-106

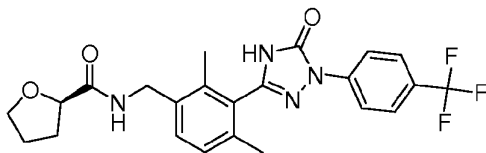
N-(3-(1-(4-(Trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)-2,2,2-trifluoroacetamide



The title compound was prepared according to the procedure described in Example-83 by using *tert*-butyl 2-(4-(trifluoromethyl)phenyl)hydrazinecarboxylate (Intermediate-53, 1.00 g, 3.63 mmol), 3-((2,2,2-trifluoroacetamido)methyl)-2,6-dimethylbenzoyl isocyanate (Intermediate-65, 1.00 g, 3.33 mmol), DCM (20 mL), trifluoro acetic acid (5.0 mL) to afford 0.700 g of the desired product. ^1H NMR (400 MHz, DMSO d_6): δ 2.20 (s, 3H), 2.22 (s, 3H), 4.40 (d, J = 4.2 Hz, 2H), 7.21 (d, J = 6.0 Hz, 1H), 7.29 (d, J = 6.0 Hz, 1H), 7.82 (d, J = 6.6 Hz, 1H), 8.17 (d, J = 6.6 Hz, 1H), 9.95 (t, J = 6.9 Hz, 1H), 12.32 (br s, 1H); MS (m/z): 457.26 (M-H $^-$).

Example-107

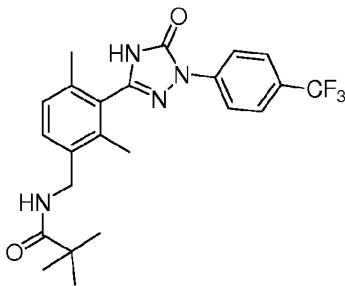
(*R*)-*N*-(2,4-Dimethyl-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)tetrahydrofuran-2-carboxamide



To a solution of 5-(3-(aminomethyl)-2,6-dimethylphenyl)-2-(4-(trifluoromethyl)phenyl)-2H-1,2,4-triazol-3(4H)-one (Intermediate-66, 0.050 g, 0.138 mmol) in dry THF:DMF (4 mL:1 mL), TBTU (0.132 g, 0.414 mmol) and TEA (1.0 mL) were added and the reaction mass was stirred for 1h under nitrogen atmosphere. To the reaction mass, (R)-(-)-tetrahydrofuran-2-carboxylic acid (0.024 g, 0.207 mmol) was added and stirred for 18 h at room temperature. After completion of the reaction, the reaction mass was quenched in water, extracted with ethyl acetate and concentrated to afford crude product which was purified by column chromatography by eluting with solution of 2% ammonia in 10% MeOH:DCM to afford 0.017 g of the title product. ^1H NMR (400 MHz, DMSO d_6): δ 1.79 (m, 3H), 1.87 (m, 3H), 2.20 (s, 6H), 3.74-3.79 (m, 1H), 3.88-3.92 (m, 1H), 4.25 (d, J = 4.5 Hz, 2H), 7.16 (d, J = 6.0 Hz, 1H), 7.25 (d, J = 6.0 Hz, 1H), 7.82 (d, J = 6.6 Hz, 1H), 8.17 (d, J = 6.3 Hz, 2H), 8.24 (t, J = 4.8 Hz, 2H), 12.30 (br s, 1H); MS (m/z): 461.14 ($M+H$) $^+$.

Example-108

N-(3-(1-(4-(Trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1H-1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)pivalamide

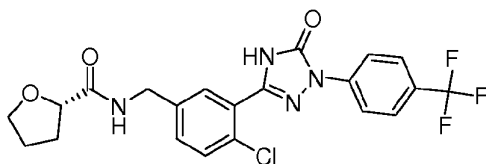


To a solution of 5-(3-(aminomethyl)-2,6-dimethylphenyl)-2-(4-(trifluoromethyl)phenyl)-2H-1,2,4-triazol-3(4H)-one (Intermediate-66, 0.050 g) in dry THF (5 mL), TEA (3 mL) was added and stirred the reaction mixture for 20 minutes. The reaction mixture was cooled to 0°C and pivaloyl chloride (0.05 g, 0.563 mmol) was added and stirred for 3 h at RT. After completion of the reaction, the reaction mass was quenched with water,

extracted with DCM:MeOH. The organic layer was separated, concentrated, and purified by column chromatography to afford 0.012 g of the desired title product. ^1H NMR (400 MHz, DMSO d_6): δ 1.15 (s, 9H), 2.18 (s, 3H), 2.22 (s, 3H), 4.24 (d, J = 12.6 Hz, 2H), 7.19 (d, 1H), 7.25 (d, 1H), 7.84 (d, J = 5.7 Hz, 2H), 8.0 (t, 1H), 8.20 (d, J = 6.4 Hz, 2H), 12.30 (br s, 1H); MS (m/z): 447.20 ($M+H$) $^+$.

Example-109

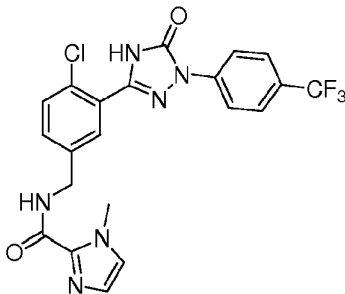
(S)-N-(4-Chloro-3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1H-1,2,4-triazol-3-yl)benzyl)-tetrahydrofuran-2-carboxamide



The title compound was prepared according to the procedure described in Example-107 by using 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1H-1,2,4-triazol-5(4H)-one (Intermediate-63, 0.070 g, 0.189 mmol), THF:DMF (5 mL:1 mL), (S)-(-) tetrahydrofuran-2-carboxylic acid (0.032 g, 0.283 mmol), TBTU (0.182 g, 0.567 mmol), and TEA (3.0 mL). The obtained crude was purified by column chromatography on basic alumina by eluting solution of 2% ammonia in 10.0 % MeOH:DCM to afford 0.010 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.80 (m, 3H), 2.15 (m, 1H), 3.75 (m, 1H), 3.94 (m, 1H), 4.27-4.32 (m, 3H), 7.42 (d, 1H), 7.60 (d, 2H), 7.85 (d, J = 5.4 Hz, 2H), 8.20 (d, J = 4.5 Hz, 2H), 8.52 (s, 1H), 12.50 (s, 1H); MS (m/z): 467.08 ($M+H$) $^+$.

Example-110

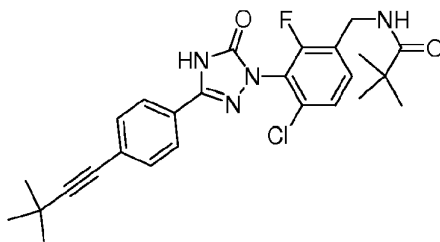
N-(4-Chloro-3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1H-1,2,4-triazol-3-yl)benzyl)-1-methyl-1H-imidazole-2-carboxamide



The title compound was prepared according to the procedure described in Example-108 by using 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-63, 0.070 g, 0.189 mmol), 1-methyl-1*H*-imidazole-2-carbonyl chloride (0.026 g, 0.283 mmol), dry THF (5 mL), DIPEA (2 mL). The obtained crude was purified by column chromatography using 2% ammonia in 20 % MeOH:DCM as mobile phase to afford 0.020 g of the desired title product. ¹H NMR (300 MHz, DMSO *d*₆): δ 3.50 (s, 3H), 4.41 (d, *J* = 18.0 Hz, 1H), 6.99 (s, 1H), 7.43 (br, 2H), 7.67 (d, 1H), 7.78 (s, 1H), 8.28 (d, *J* = 4.5 Hz, 2H), 8.30 (s, 1H), 9.08 (t, 1H); MS (*m/z*): 477.10 (M+H)⁺.

Example-111

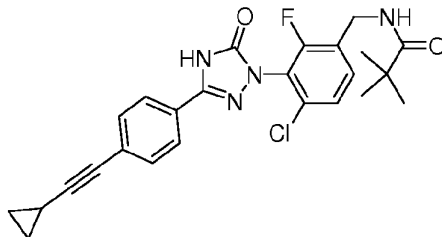
N-(4-Chloro-2-fluoro-3-(4,5-dihydro-3-(4-(3,3-dimethylbut-1-ynyl)phenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide



To a solution of *N*-(4-chloro-2-fluoro-3-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide (Intermediate-68, 0.050 g, 0.094 mmol) in DMSO (3.0 mL), 3,3-dimethylbut-1-yne (0.011 g, 0.142 mmol), TBAF (0.074 g, 0.283 mmol) and bis(triphenylphosphine)palladium(II) chloride (0.020 g, 0.022 mmol) were added and the reaction mass was stirred at 80⁰C for 5-6 h. After completion of the reaction, the reaction mass was quenched with water and extracted with DCM and concentrated. The obtained crude product was purified with column chromatography on silica gel column and 2.0 % EA: DCM as mobile phase to afford 0.022 g of the desired title product. ¹H NMR (400 MHz, DMSO *d*₆): δ 1.09 (s, 9H), 1.25 (s, 9H), 4.31 (br s, 2H), 7.35-7.51 (m, 4H), 7.80 (d, *J* = 8.3 Hz, 2H), 8.18-8.20 (m, 1H); MS (*m/z*): 481.14 (M-H)⁻.

Example-112

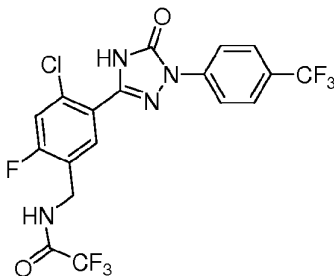
N-(4-Chloro-3-(3-(4-(2-cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)-2-fluorobenzyl)pivalamide



The title compound was prepared according to the procedure described in Example-111 by using *N*-(4-chloro-2-fluoro-3-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide (Intermediate-68, 0.050 g, 0.094 mmol), ethynylcyclopropane (0.009 g, 0.142 mmol), TBAF (0.074 g, 0.283 mmol), bis(triphenylphosphine)palladium(II) chloride (0.003 g, 0.003 mmol) and DMSO (3.0 mL) to afford 0.018 g of the desired title product. ^1H NMR (400 MHz, DMSO d_6): δ 0.75 (m, 2H), 0.77-0.91 (m, 2H), 1.36 (s, 9H), 1.53-1.59 (m, 2H), 7.38 (t, $J = 5.9$ Hz, 1H), 7.46-7.52 (m, 3H), 8.79 (d, $J = 4.4$ Hz, 2H), 8.19 (t, $J = 4.3$ Hz, 1H); MS (m/z): 467.15 ($M+H$) $^+$.

Example-113

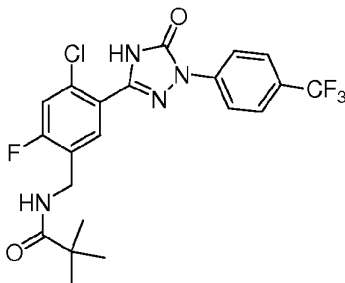
N-(4-Chloro-2-fluoro-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



The title compound was prepared according to the procedure described in Example-83 by using *tert*-butyl 2-(4-(trifluoromethyl)phenyl)hydrazinecarboxylate (Intermediate-53, 0.300 g), 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzoyl isocyanate (Intermediate-69, 0.300 g), DCM (20 mL), trifluoro acetic acid (5.0 mL) to afford 0.030 g of the desired product. ^1H NMR (400 MHz, DMSO d_6): δ 4.49 (m, 2H), 7.74-7.77 (m, 1H), 7.78-7.80 (m, 1H), 7.87 (d, $J = 5.1$ Hz, 2H), 8.19 (d, $J = 4.2$ Hz, 2H), 10.06 (m, 1H), 12.76 (s, 1H); MS (m/z): 467.15 ($M+H$) $^+$.

Example-114

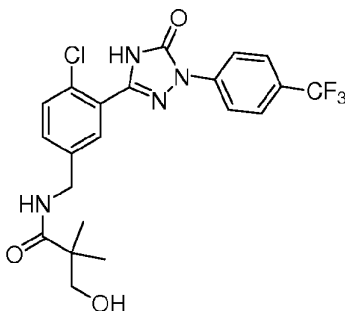
N-(4-Chloro-2-fluoro-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(4-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-70, 0.080 g, 0.206 mmol), pivaloyl chloride (0.027 g, 0.227 mmol), THF (5.0 mL) and TEA (0.2 mL) to afford 0.043 g of the desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 1.13 (s, 9H), 4.31-4.32 (m, 2H), 7.86 (d, *J* = 6.5 Hz, 2H), 8.13-8.18 (m, 3H), 12.70 (s, 1H); MS (*m/z*): 471.14 (M+H)⁺.

Example-115

N-(4-Chloro-3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-3-hydroxy-2,2-dimethylpropanamide

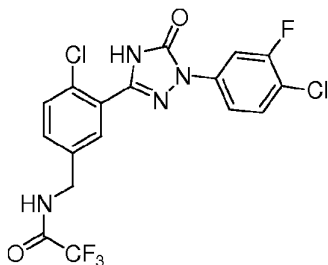


The title compound was prepared according to the procedure described in Example-107 by using 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-63, 0.100 g, 0.271 mmol), THF:DMF (5 mL:1 mL), 3-methoxy-2,2-dimethylpropanoic acid (0.053 g, 0.407 mmol), TBTU (0.261 g, 0.813 mmol), TEA (2.0 mL). The obtained crude was purified by column chromatography using basic alumina as stationary phase and eluting with the solution of 2% ammonia in

10.0% MeOH:DCM to afford 0.012 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.05 (s, 6H), 3.30 (s, 2H), 4.30 (d, $J = 4.5$ Hz, 1H), 4.36 (s, 1H), 7.53 (d, $J = 6.3$ Hz, 1H), 7.75 (s, 1H), 8.08 (t, $J = 4.5$ Hz, 1H), 8.21 (d, $J = 6.3$ Hz, 2H); MS (m/z): 469.04 ($M+H$) $^+$.

Example-116

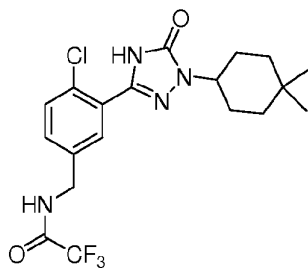
N-(4-Chloro-3-(1-(4-chloro-3-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



The title compound was prepared according to the procedure described in Example-83 by using *tert*-butyl 2-(4-chloro-3-fluorophenyl)hydrazinecarboxylate (Intermediate-71, 0.828 g, 0.003 mmol), 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate (step-3 of Intermediate-26, 1.3 g, 0.043 mmol), DCM (20 mL), trifluoro acetic acid (5.0 mL) to afford 0.500 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 4.45 (d, 2H), 7.48 (d, $J = 15.0$ Hz, 1H), 7.68-7.83 (m, 3H), 7.85 (d, 1H), 7.99 (d, $J = 17.1$ Hz, 1H), 10.08 (d, $J = 12.0$ Hz, 1H), 12.73 (s, 1H).

Example-117

N-(4-Chloro-3-(4,5-dihydro-1-(4,4-dimethylcyclohexyl)-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide

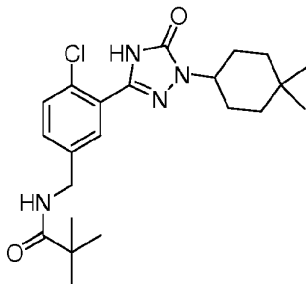


The title compound was prepared according to the procedure described in Example-83 by using 4,4-dimethyl *tert*-butyl 2-cyclohexyl hydrazinecarboxylate (Intermediate-72, 0.269

g, 1.1 mmol), 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate (step-2 of Intermediate-26, 0.400 g, 1.2 mmol), DCM (20 mL), trifluoro acetic acid (5.0 mL) to afford 0.200 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.33 (s, 6H), 1.43-1.91 (m, 8H), 3.86-3.94 (m, 1H), 4.41 (d, $J = 4.5$ Hz, 2H), 7.38 (d, $J = 7.5$ Hz, 1H), 7.53 (s, 1H), 7.58 (d, $J = 6.3$ Hz, 1H), 10.02-10.04 (m, 1H), 11.95 (s, 1H); MS (m/z): 431.17 (M+H) $^+$.

Example-118

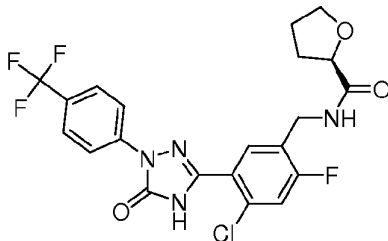
N-(4-Chloro-3-(4,5-dihydro-1-(4,4-dimethylcyclohexyl)-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



To a solution of 5-(5-(aminomethyl)-2-chlorophenyl)-2-(4,4-dimethylcyclohexyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-73, 0.070 g, 0.202 mmol) in dry THF (5.0 mL), DIPEA (2.0 mL) and pivaloyl chloride (0.30 g, 0.242 mmol) were added and the reaction mass was stirred at RT for 16 h. After completion of the reaction, the reaction mass was quenched with water and extracted with 10 % MeOH:DCM. The organic layer was washed with dilute HCl and concentrated. The obtained crude was purified with column chromatography on basic alumina by eluting with 2.0 % MeOH:DCM to afford 0.040 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.44 (s, 6H), 1.47 (s, 9H), 1.81-1.94 (m, 8H), 3.86-3.90 (m, 1H), 4.40 (s, 2H), 7.33 (d, $J = 1.5$ Hz, 1H), 7.45 (s, 1H), 7.50 (d, $J = 2.1$ Hz, 1H), 8.16 (t, 1H), 11.93 (s, 1H); MS (m/z): 419.21 (M+H) $^+$.

Example-119

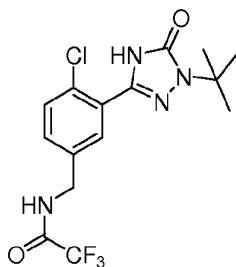
(*R*)-*N*-(4-chloro-2-fluoro-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)tetrahydrofuran-2-carboxamide



The title compound was prepared according to the procedure described in Example-107 by using 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(4-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-70, 0.090 g, 0.233 mmol), THF:DMF (5 mL:1 mL), (R)-(-)-tetrahydrofuran-2-carboxylic acid (0.040 g, 0.349 mmol), TBTU (0.224 g, 0.699 mmol), and TEA (0.5 mL). The obtained crude was purified with column chromatography using basic alumina as stationary phase and 2% ammonia in 10.0 % MeOH:DCM solution as mobile phase to afford 0.015 g of the desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 1.78-1.91 (m, 2H), 2.09-2.15 (m, 1H), 3.74 (m, 1H), 3.88-3.94 (m, 1H), 4.28 (m, 1H), 4.43 (m, 2H), 7.65 (m, 2H), 7.86 (d, *J* = 8.72 Hz, 2H), 8.18 (d, *J* = 8.52 Hz, 2H), 8.45 (d, *J* = 16.0 Hz, 1H), 12.71 (s, 1H).

Example-120

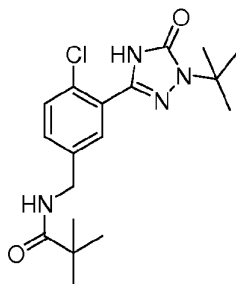
N-(3-(1-*tert*-Butyl-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-4-chlorobenzyl)-2,2,2-trifluoroacetamide



The title compound was prepared according to the procedure described in Example-83 by using *tert*-butyl 2-*tert*-butylhydrazinecarboxylate (Intermediate-74, 0.233 g, 1.2 mmol), 2-chloro-5-((2,2,2-trifluoroacetamido)methyl)benzoyl isocyanate (step-3 of Intermediate-26, 0.400 g, 1.2 mmol), DCM (10 mL), trifluoro acetic acid (5.0 mL) to afford 0.100 g of the desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 1.52 (s, 9H), 4.42 (s, 2H), 7.39 (d, 1H), 7.41 (s, 1H), 7.58 (d, *J* = 2.1 Hz, 2H), 10.0 (br s, 1H), 12.0 (br s, 1H); MS (*m/z*): 377.20 (M+H)⁺.

Example-121

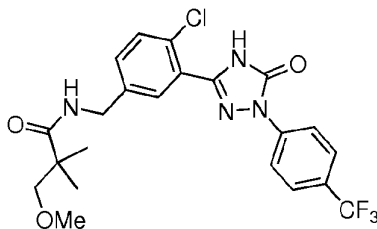
N-(3-(1-*tert*-Butyl-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-4-chlorobenzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 2-*tert*-butyl-5-(5-(aminomethyl)-2-chlorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-75, 0.250 g), pivaloyl chloride (1.0 mL), DIPEA (2.0 mL), and dry THF (5 mL) to afford 0.180 g of desired product. ^1H NMR (400 MHz, DMSO d_6): δ 1.08 (s, 9H), 1.49 (s, 9H), 4.24 (d, J = 4.5 Hz, 2H), 7.32 (dd, J = 4.5 Hz, 3.6Hz, 2H), 7.43 (s, 1H), 7.50 (s, 1H), 8.14 (d, J = 4.5 Hz, 1H); MS (m/z): 365.20 ($M+H$) $^+$.

Example-122

N-(4-Chloro-3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-3-methoxy-2,2-dimethylpropanamide

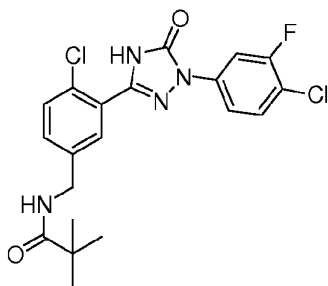


To a solution of 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-63, 0.250 g, 0.678 mmol) in THF:DMF (5 mL:1 mL), TBTU (0.653 g, 2.3 mmol), TEA (4.0 mL) were added and the reaction mass was stirred at RT for 1 h. To the reaction mixture, 3-methoxy-2,2-dimethylpropanoic acid (0.134 g, 1.01 mmol) was added and stirring was continued at RT for 2 days. After completion of the reaction, the reaction mass was quenched with water and extracted with ethyl acetate. The organic layer was separated, dried and concentrated to obtain crude product. The obtained crude was purified with column chromatography on basic alumina by eluting with 5.0 % MeOH:DCM to afford 0.110 g of the desired product. ^1H NMR

(400 MHz, DMSO d_6): δ 1.15 (s, 6H), 1.18 (s, 3H), 3.09 (t, 2H), 4.32 (d, $J = 11.4$ Hz, 2H), 7.43 (dd, $J = 4.5$ Hz, 3.6 Hz, 1H), 7.45 (d, 1H), 7.86 (d, $J = 5.4$ Hz, 2H), 8.15 (t, 1H), 8.20 (d, 2H), 12.50 (br s, 1H); MS (m/z): 483.20 (M+H)⁺.

Example-123

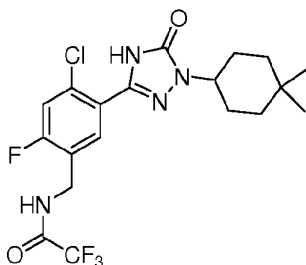
N-(4-Chloro-3-(1-(4-chloro-3-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chlorophenyl)-2-(4-chloro-3-fluorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-76, 0.250 g), pivaloyl chloride (1.0 mL), DIPEA (2.0 mL), dry THF (5 mL) to afford 0.110 g of the desired product. ¹H NMR (400 MHz, DMSO d_6): δ 1.24 (s, 9H), 4.30 (d, $J = 14.7$ Hz, 2H), 7.40 (dd, $J = 4.5$ Hz, 3.6 Hz, 1H), 7.43 (d, 1H), 7.60 (d, $J = 5.4$ Hz, 1H), 7.74 (d, 1H), 7.98 (d, 1H), 8.18 (t, $J = 15.0$ Hz, 1H), 12.70 (s, 1H); MS (m/z): 437.59 (M+H)⁺.

Example-124

N-(4-Chloro-2-fluoro-5-(4,5-dihydro-1-(4,4-dimethylcyclohexyl)-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide

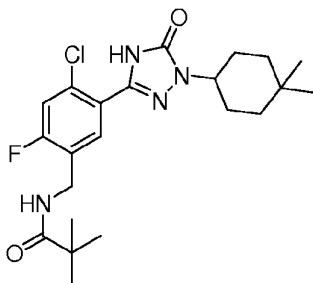


The title compound was prepared according to the procedure described in Example-83 by using 4,4-dimethyl *tert*-butyl 2-cyclohexyl hydrazinecarboxylate (Intermediate-72, 0.400g), 5-((2,2,2-trifluoroacetamido)methyl)-2-chloro-4-fluorobenzoyl isocyanate

(Intermediate-69, 0.400 g), DCM (10 mL), trifluoro acetic acid (5.0 mL) to afford 0.150 g of the desired product. ^1H NMR (400 MHz, DMSO d_6): δ 0.93 (s, 6H), 1.32-1.35 (m, 2H), 1.36-1.46 (m, 2H), 1.59-1.62 (m, 2H), 1.83-1.87 (m, 2H), 3.89 (m, 1H), 4.42 (m, 2H), 7.72 (m, 2H), 10.0 (s, 1H), 12.01 (s, 1H); MS (m/z): 449.48 (M+H) $^+$.

Example-125

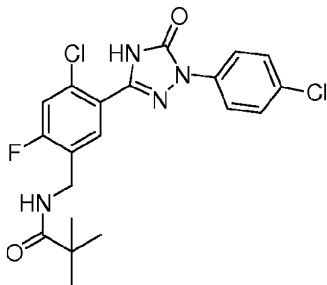
N-(4-Chloro-2-fluoro-5-(4,5-dihydro-1-(4,4-dimethylcyclohexyl)-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(4,4-dimethylcyclohexyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-77, 0.200 g), pivaloyl chloride (0.5 mL), TEA (2.0 mL), dry THF (5 mL) to afford 0.033 g of the desired product. ^1H NMR (300 MHz, CDCl_3): δ 1.36 (s, 6H), 1.37 (s, 9H), 1.62 (m, 2H), 1.79 (m, 1H), 1.82 (m, 2H), 1.89 (m, 2H), 3.46-3.94 (m, 1H), 4.28 (s, 2H), 7.47 (d, $J = 7.9$ Hz, 1H), 7.59 (d, $J = 9.72$ Hz, 1H), 8.15 (m, 1H), 11.95 (s, 1H); MS (m/z): 436.55 (M+H) $^+$.

Example-126

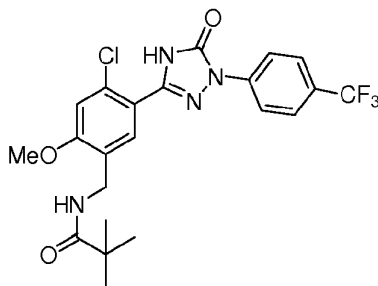
N-(4-Chloro-5-(1-(4-chlorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(4-chlorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-78, 0.120 g), pivaloyl chloride (0.5 mL), TEA (2.0 mL), dry THF (5 mL) to afford 0.033 g of the desired product. ¹H NMR (300 MHz, DMSO): δ 1.12 (s, 9H), 4.30 (br s, 2H), 7.53-7.63 (m, 4H), 7.95 (d, *J* = 12.8 Hz, 2H), 8.14 (s, 1H), 12.61 (s, 1H); MS (*m/z*): 437.47 (*M*+H)⁺.

Example-127

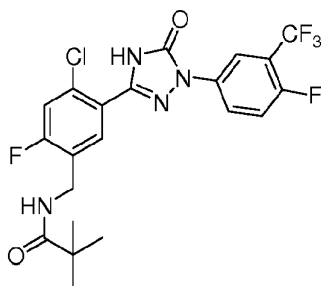
N-(4-Chloro-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-methoxybenzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chloro-4-methoxyphenyl)-2-(4-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-79, 0.200 g), pivaloyl chloride (0.2 mL), TEA (2.0mL), dry THF (5 mL) to afford 0.015 g of the desired product. ¹H NMR (300 MHz, CDCl₃): δ 1.15 (s, 9H), 3.91 (s, 3H), 4.22 (m, 2H), 7.26 (s, 1H), 7.42 (s, 1H), 7.87 (m, 2H), 7.99 (m, 1H), 8.15 (m, 2H), 12.57 (s, 1H); MS (*m/z*): 483.38 (*M*+H)⁺.

Example-128

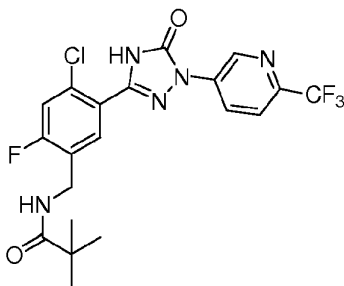
N-(4-Chloro-2-fluoro-5-(1-(4-fluoro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(4-fluoro-3-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-81, 0.300 g), pivaloyl chloride (0.3 mL), TEA (2.0 mL), dry THF (5 mL) to afford 0.065 g of the desired product. ¹H NMR (300 MHz, CDCl₃): δ 1.12 (s, 9H), 4.30 (m, 2H), 7.61-7.69 (m, 3H), 8.16 (br s, 1H), 8.25 (br s, 2H), 12.73 (s, 1H); MS (m/z): 489.47 (M+H)⁺.

Example-129

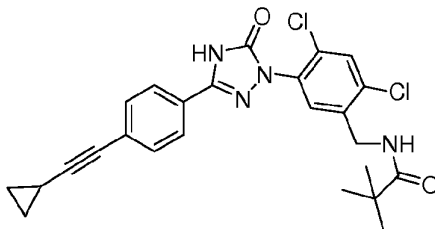
N-(4-Chloro-2-fluoro-5-(1-(6-(trifluoromethyl)pyridin-3-yl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(6-(trifluoromethyl)pyridin-3-yl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-82, 0.300 g), pivaloyl chloride (0.3 mL), TEA (2.0 mL), dry THF (10 mL) to afford 0.050 g of the desired product. ¹H NMR (300 MHz, CDCl₃): δ 1.12 (s, 9H), 4.31 (d, *J* = 6.8 Hz, 2H), 7.63-7.71 (m, 2H), 8.05-8.07 (m, 1H), 8.16 (m, 1H), 8.55 (d, *J* = 13.2 Hz, 1H), 9.29 (s, 1H), 12.85 (s, 1H); MS (m/z): 472.41 (M+H)⁺.

Example-130

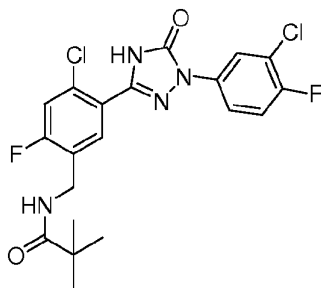
N-(2,4-Dichloro-5-(3-(4-(2-cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-111 using *N*-(2,4-dichloro-5-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl) pivalamide (Intermediate-84, 0.150 g, 0.028 mmol), ethynylcyclopropane (0.038 g, 0.56 mmol), TBAF (0.268 g, 0.85 mmol), bis(triphenylphosphine)palladium(II) chloride (catalytic) and DMSO (3.0 mL) to afford 0.050 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.77 (m, 2H), 0.93 (m, 2H), 1.21 (s, 9H), 1.58 (m, 1H), 4.32 (d, *J* = 6.3 Hz, 2H), 7.50-7.53 (m, 3H), 7.82-7.91 (m, 2H), 7.95 (s, 1H), 8.21 (m, 1H)

Example-131

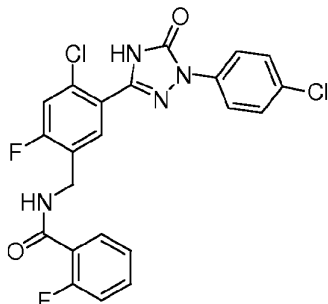
N-(4-Chloro-5-(1-(3-chloro-4-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(3-chloro-4-fluorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-86, 0.300 g), pivaloyl chloride (0.3 mL), TEA (2.0 mL), dry THF (10 mL) to afford 0.083 g of the desired product. ¹H NMR (300 MHz, DMSO): δ 1.13 (s, 9H), 4.30 (d, *J* = 6.4 Hz, 2H), 7.53-7.66 (s, 3H), 7.91-8.15 (s, 3H), 12.67 (s, 1H); MS (*m/z*): 455.19 (*M*+*H*)⁺.

Example-132

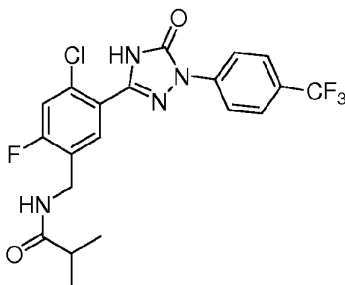
N-(4-Chloro-5-(1-(4-chlorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)-2-fluorobenzamide



A mixture of 2-fluoro benzoic acid (0.400 g), thionyl chloride (20 mL) and DMF (cat. amt.) was refluxed for 5 h and excess of solvent was removed under reduced pressure to obtain crude product. The obtained crude product was added to the solution of 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(4-chlorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-78, 0.250 g) and DIPEA (1.0 mL) in THF (20 mL) and the reaction mass was stirred for 3 h. After completion of reaction, reaction mass was quenched with water and extracted with ethyl acetate. The organic layer was separated, concentrated to afford 0.032 g of the desired product. ¹H NMR (300 MHz, DMSO): δ 4.53 (br s, 2H), 7.29 (m, 2H), 7.70 (m, 3H), 7.78 (m, 3H), 7.95 (br s, 2H), 8.94 (br s, 1H), 12.63 (br s, 1H); MS (m/z): 475.39 (M+H)⁺.

Example-133

N-(4-Chloro-2-fluoro-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)isobutyramide

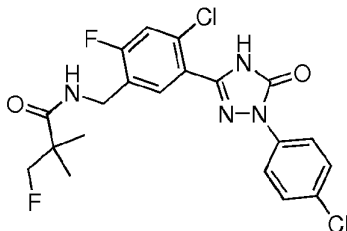


The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(4-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-70, 0.100 g), 2-methylpropanoyl chloride (0.1 mL), TEA (0.1 mL), dry THF (5 mL) to afford 0.028 g of the desired product. ¹H NMR (300 MHz, DMSO): δ 1.01 (s, 6H), 4.32 (d, *J* = 7.6 Hz, 2H), 7.67-7.71 (m, 2H), 7.85 (d, *J*

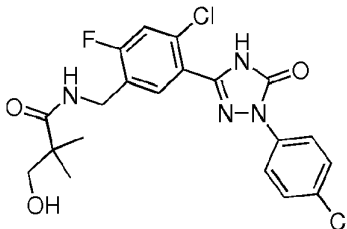
= 11.2 Hz, 2H), 8.17 (d, J = 11.6 Hz, 2H), 8.36 (m, 1H), 12.73 (s, 1H); MS (m/z): 457.37 ($M+H$)⁺.

Example-134

N-(4-Chloro-5-(1-(4-chlorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)-3-fluoro-2,2-dimethylpropanamide



Step-1:- Preparation of *N*-(4-chloro-5-(1-(4-chlorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)-3-hydroxy-2,2-dimethylpropanamide



The title compound was prepared according to the procedure described in Example-107 by using 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(4-chlorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-78, 0.250 g, 0.708 mmol), THF:DMF (5 mL: 1 mL), 3-hydroxy-2,2-dimethylpropanoic acid (0.091 g, 0.407 mmol), TBTU (0.340 g, 1.062 mmol), TEA (2.0 mL) to afford 0.150 g of the desired product. ¹H NMR (300 MHz, CDCl₃): δ 1.05 (s, 6H), 2.68 (s, 2H), 4.32 (d, J = 8.0 Hz, 2H), 7.53 (d, J = 12.0 Hz, 2H), 7.64-7.67 (m, 2H), 7.95-7.98 (m, 2H), 8.08 (br s, 1H); MS (m/z): 453.27 (M^+H)⁺.

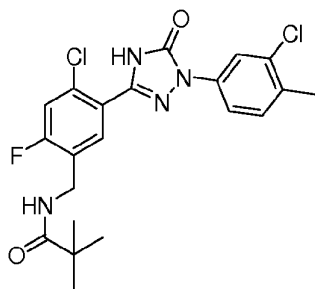
Step-2:- Preparation *N*-(4-chloro-5-(1-(4-chlorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)-3-fluoro-2,2-dimethylpropanamide

To a solution of *N*-(4-chloro-5-(1-(4-chlorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)-3-hydroxy-2,2-dimethylpropanamide (0.150 g, 0.330 mmol) in THF (5.0 mL), DAST (0.079 g, 0.495 mmol) was added at 0-5°C and stirred at RT for 5-6 h. After completion of the reaction, the reaction mass was quenched with water and extracted with ethyl acetate. The organic layer was separated, concentrated and the

obtained crude was purified with column chromatography using basic alumina by eluting with solution of 2% ammonia in 10.0 % MeOH: DCM to afford 0.013 g of the desired product. ^1H NMR (300 MHz, CDCl_3): δ 1.13 (s, 6H), 4.32 (m, 3H), 4.46 (s, 1H), 7.55 (m, 2H), 7.69 (m, 2H), 7.94-7.97 (m, 2H), 8.31 (s, 1H), 12.61 (s, 1H); MS (m/z): 454.25 (M+H) $^+$.

Example-135

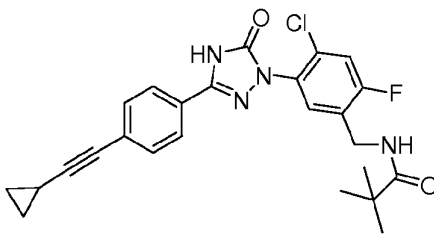
N-(4-Chloro-5-(1-(3-chloro-4-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chloro-4-fluorophenyl)-2-(3-chloro-4-methylphenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-88, 0.200 g), pivaloyl chloride (0.5 mL), TEA (2.0 mL), dry THF (5 mL) to afford 0.074 g of the desired product. ^1H NMR (300 MHz, $\text{DMSO } d_6$): δ 1.11 (s, 9H), 2.32 (s, 3H), 4.29 (d, $J = 8.8\text{Hz}$, 2H), 7.43 (d, $J = 11.6\text{Hz}$, 1H), 7.60-7.67 (m, 1H), 7.79 (d, $J = 10.4\text{ Hz}$, 1H), 7.96 (s, 1H), 8.15 (m, 1H), 12.60 (s, 1H); MS (m/z): 451.52 (M+H) $^+$.

Example-136

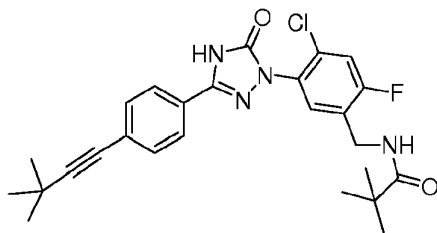
N-(4-Chloro-5-(3-(4-(2-cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)-2-fluorobenzyl)pivalamide



The title compound was prepared according to the procedure described in Example-111 using *N*-(4-chloro-2-fluoro-5-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide (Intermediate-90, 0.200 g, 0.370 mmol), ethynylcyclopropane (0.050 g, 0.75 mmol), TBAF (0.360 g, 1.13 mmol), bis(triphenylphosphine)palladium(II) chloride (catalytic) and DMSO (3.0 mL) at 80°C to afford 0.080 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.77 (br s, 9H), 0.92 (br s, 2H), 1.10 (s, 3H), 1.57 (m, 1H), 4.30 (d, *J* = 5.4Hz, 2H), 7.43-7.50 (m, 3H), 7.65 (d, *J* = 5.4Hz, 2H), 7.77 (d, *J* = 5.1Hz, 2H), 12.57 (br s, 1H); MS (*m/z*): 467.36 (M+H)⁺.

Example-137

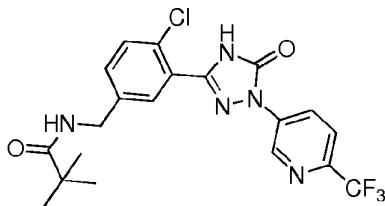
N-(4-Chloro-2-fluoro-5-(4,5-dihydro-3-(4-(3,3-dimethylbut-1-ynyl)phenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-111 using *N*-(4-chloro-2-fluoro-5-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide (Intermediate-90, 0.100 g, 0.189 mmol), 3,3-dimethylbut-1-yne (0.031 g, 0.378 mmol), TBAF (0.178g, 0.567 mmol), bis(triphenylphosphine)palladium(II) chloride (catalytic) and DMSO (3.0 mL) to afford 0.050 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.10 (s, 9H), 1.30 (s, 9H), 4.30 (d, *J* = 9.0Hz, 2H), 7.43-7.50 (m, 3H), 7.65 (d, *J* = 9.3Hz, 1H), 7.78 (d, *J* = 8.1Hz, 2H), 8.14 (br s, 1H), 12.60 (br s, 1H); MS (*m/z*): 483.29 (M+H)⁺.

Example-138

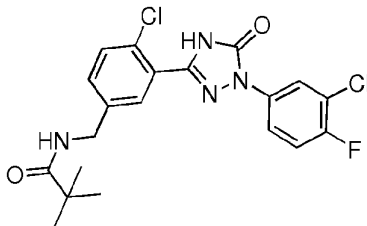
N-(4-Chloro-3-(1-(6-(trifluoromethyl)pyridin-3-yl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chlorophenyl)-2-(6-(trifluoromethyl)pyridin-3-yl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-91, 0.350 g, 0.945 mmol), pivaloyl chloride (0.120 g, 0.945 mmol), DIPEA (2.0 mL), dry THF (10 mL) to afford 0.080 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.11 (s, 9H), 4.30 (d, *J* = 6.3 Hz, 2H), 7.43 (d, *J* = 8.4 Hz, 1H), 7.61-7.64 (m, 2H), 8.06 (d, *J* = 9.0 Hz, 1H), 8.19 (m, 1H), 8.58 (d, *J* = 8.4 Hz, 1H), 9.32 (s, 1H), 12.82 (s, 1H); MS (*m/z*): 454.50 (*M*+H)⁺.

Example-139

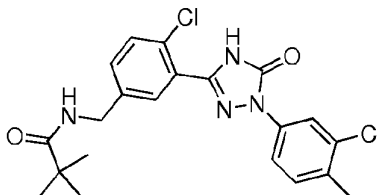
N-(4-Chloro-3-(1-(3-chloro-4-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chlorophenyl)-2-(3-chloro-4-fluorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-92, 0.300 g, 0.852 mmol), pivaloyl chloride (0.108 g, 0.852 mmol), DIPEA (2.0 mL), dry THF (10 mL) to afford 0.050 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.13 (s, 9H), 4.29 (d, *J* = 5.4 Hz, 2H), 7.40 (d, *J* = 7.8 Hz, 1H), 7.52-7.59 (m, 3H), 7.91-8.18 (m, 3H), 12.65 (s, 1H); MS (*m/z*): 437.23 (*M*+H)⁺.

Example-140

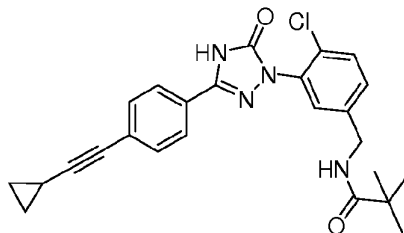
N-(4-Chloro-3-(1-(3-chloro-4-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chlorophenyl)-2-(3-chloro-4-methylphenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-93, 0.350 g, 1.00 mmol), pivaloyl chloride (0.128 g, 1.0 mmol), DIPEA (2.0 mL), dry THF (10 mL) to afford 0.200 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.13 (s, 9H), 2.34 (s, 3H), 4.29 (d, J = 6.3 Hz, 2H), 7.39-7.47 (m, 2H), 7.58-7.61 (m, 2H), 7.82 (d, J = 8.7 Hz, 1H), 8.01 (s, 1H), 8.18 (t, 1H), 12.58 (s, 1H); MS (m/z): 433.19 ($M+H$) $^+$.

Example-141

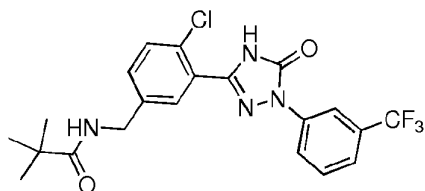
N-(4-Chloro-3-(3-(4-(2-cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide



To a solution of *N*-(4-chloro-3-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide (Intermediate-95, 0.100 g, 0.196 mmol) in DMSO (3.0 mL), ethynylcyclopropane (0.035 g, 0.530 mmol), TBAF (0.150 g, 0.574) and bis(triphenylphosphine)palladium(II) chloride (catalytic) were added and the reaction mass was stirred at 80 $^{\circ}$ C for 5-6 h. After completion of the reaction, the reaction mass was quenched with water and extracted with DCM and concentrated to afford 0.050 g of the desired title product. ^1H NMR (300 MHz, DMSO d_6): δ 0.76-0.91 (m, 2H), 0.91-0.92 (m, 2H), 1.11 (s, 9H), 1.57 (m, 1H), 4.28 (br d, 2H), 7.31-7.34 (br d, 1H), 7.42 (br s, 1H), 7.47-7.50 (br m, 2H), 7.60 (d, J = 8.4 Hz, 1H), 7.78 (d, J = 8.4 Hz, 2H), 8.18 (s, 1H), 12.58 (s, 1H); MS (m/z): 449.52 ($M+H$) $^+$.

Example-142

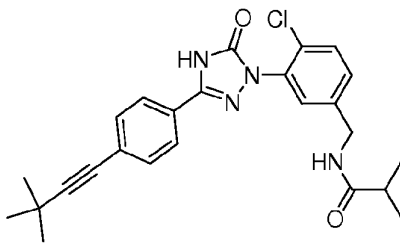
N-(4-Chloro-3-(1-(3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chlorophenyl)-2-(3-(trifluoromethyl)phenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-96, 0.250 g, 0.678 mmol), pivaloyl chloride (0.3 mL), TEA (2.0 mL), and dry THF (5 mL) to afford 0.150 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.13 (s, 9H), 4.30 (d, *J* = 6.0 Hz, 2H), 7.41-7.43 (m, 1H), 7.59-7.62 (m, 3H), 7.71-7.76 (m, 1H), 8.19-8.29 (m, 3H), 12.68 (s, 1H); MS (*m/z*): 453.38 (*M*+*H*)⁺.

Example-143

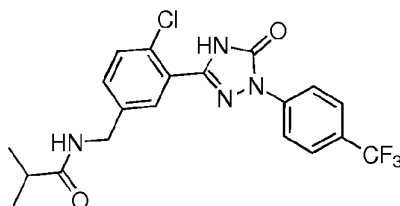
N-(4-Chloro-3-(4,5-dihydro-3-(4-(3,3-dimethylbut-1-ynyl)phenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide



The title compound was prepared according to the procedure described in Example-111 by using *N*-(4-chloro-3-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide (Intermediate-98, 0.060 g, 0.120 mmol), 3,3-dimethylbut-1-yne (0.014 g, 0.181 mmol), TBAF (0.094 g, 0.362 mmol), bis(triphenylphosphine)palladium (II)chloride (0.003 g, 0.003 mmol) and DMSO (3.0 mL) to afford 0.015 g of the desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 1.03 (s, 6H), 1.30 (s, 9H), 2.50 (m, 1H), 4.27 (d, *J* = 5.4 Hz, 2H), 7.35 (d, *J* = 8.7 Hz, 1H), 7.44-7.50 (m, 3H), 7.61 (d, *J* = 8.1 Hz, 1H), 7.80 (d, *J* = 8.1 Hz, 2H), 8.38 (m, 1H); MS (*m/z*): 449.9 (*M*-*H*)⁻.

Example-144

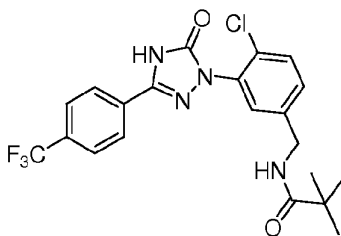
N-(4-Chloro-3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)isobutyramide



The title compound was prepared according to the procedure described in Example-108 by using 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-63, 0.250 g, 0.678 mmol), isobutyryl chloride (0.108 g, 1.01 mmol), TEA (2.0 mL), and dry THF (5 mL) to afford 0.350 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.14 (d, *J* = 6.6 Hz, 6H), 2.41-2.43 (m, 1H), 4.30 (d, *J* = 6.3 Hz, 2H), 7.43 (d, *J* = 8.1 Hz, 1H), 7.61-7.64 (m, 2H), 7.86 (d, *J* = 9.3 Hz, 2H), 8.20 (d, *J* = 8.4 Hz, 2H), 8.38 (m, 1H), 12.69 (s, 1H); MS (*m/z*): 439.24 (M+H)⁺.

Example-145

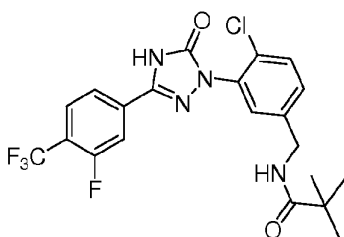
N-(4-Chloro-3-(3-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-83 by using *tert*-butyl 2-(2-chloro-5-((2,2-dimethylpropanoyl)amino)methyl)phenylhydrazinecarboxylate (Intermediate-94, 0.150 g, 0.420 mmol), 4-(trifluoromethyl)benzoyl isocyanate (Intermediate-59, 0.181 g, 0.84 mmol), DCM (15 mL), and trifluoro acetic acid (3.0 mL) to afford 0.070 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.13 (s, 9H), 4.29 (d, *J* = 5.4 Hz, 2H), 7.36 (d, *J* = 8.7 Hz, 1H), 7.45 (s, 1H), 7.62 (d, *J* = 8.4 Hz, 1H), 7.92 (d, *J* = 8.4 Hz, 2H), 8.06 (d, *J* = 8.4 Hz, 2H), 8.19 (m, 1H), 12.79 (s, 1H); MS (*m/z*): 456.59 (M+H)⁺.

Example-146

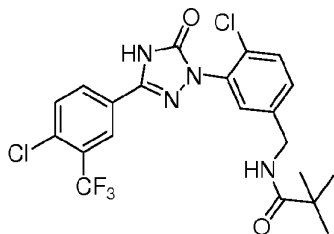
N-(4-Chloro-3-(3-(3-fluoro-4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-83 by using 3-fluoro-4-(trifluoromethyl)benzoyl isocyanate (Intermediate-99, 2.0 g), *tert*-butyl 2-(2-chloro-5-{[(2,2-dimethylpropanoyl)amino]methyl}phenyl)hydrazinecarboxylate (Intermediate-94, 1.0 g), DCM (15 mL), and trifluoro acetic acid (3.0 mL) to afford 0.900 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.12 (s, 9H), 4.29 (d, J = 5.4 Hz, 2H), 7.35 (d, J = 8.4 Hz, 1H), 7.44 (s, 1H), 7.62 (d, J = 8.7 Hz, 1H), 7.86-8.10 (m, 3H), 8.19 (m, 1H), 12.86 (s, 1H); MS (m/z): 471.35 ($\text{M}+\text{H}$) $^+$.

Example-147

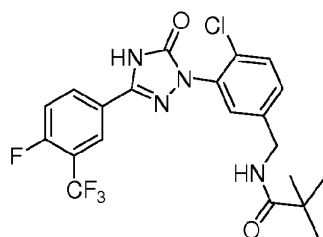
N-(4-Chloro-3-(3-(4-chloro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide



The title compound was prepared according to the procedure described in Example-83 by using 4-chloro-3-(trifluoromethyl)benzoyl isocyanate (Intermediate-100, 2.0 g), *tert*-butyl 2-(2-chloro-5-{[(2,2-dimethylpropanoyl)amino]methyl}phenyl)hydrazinecarboxylate (Intermediate-94, 1.0 g), DCM (15 mL), and trifluoro acetic acid (3.0 mL) to afford 0.900 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.12 (s, 9H), 4.29 (d, J = 5.1 Hz, 2H), 7.36 (d, J = 9.3 Hz, 1H), 7.43 (s, 1H), 7.61 (d, J = 8.4 Hz, 1H), 7.93 (d, J = 9.0 Hz, 1H), 8.10-8.18 (m, 2H), 8.27 (s, 1H), 12.82 (s, 1H); MS (m/z): 487.29 ($\text{M}+\text{H}$) $^+$.

Example-148

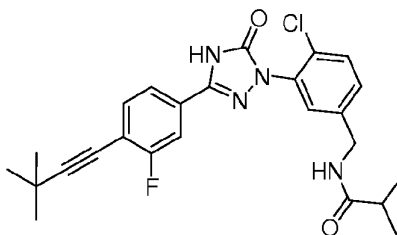
N-(4-Chloro-3-(3-(4-fluoro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide



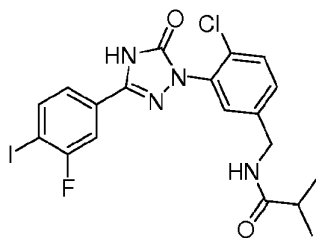
The title compound was prepared according to the procedure described in Example-83 by using 4-fluoro-3-(trifluoromethyl)benzoyl isocyanate (Intermediate-101, 0.196 g, 0.84 mmol), *tert*-butyl 2-(2-chloro-5-[(2,2-dimethylpropanoyl)amino]methyl]phenyl)hydrazinecarboxylate (Intermediate-94, 0.150 g, 0.420 mmol), DCM (15 mL), and trifluoro acetic acid (3.0 mL) to afford 0.090 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.12 (s, 9H), 4.29 (d, J = 5.7 Hz, 2H), 7.35 (d, J = 8.4 Hz, 1H), 7.43 (s, 1H), 7.60 (d, J = 8.7 Hz, 1H), 7.70-7.76 (m, 1H), 8.19-8.23 (m, 3H), 12.76 (s, 1H); MS (m/z) : 471.32 ($M+H$) $^+$.

Example-149

N-(4-Chloro-3-(3-(3-fluoro-4-(3,3-dimethylbut-1-ynyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide



Step1:- Preparation of *N*-(4-chloro-3-(3-(3-fluoro-4-iodophenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide



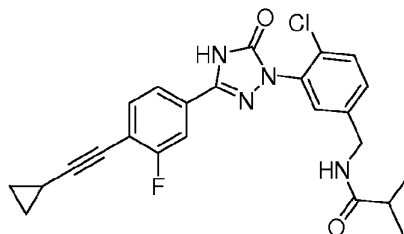
The title compound was prepared according to the procedure described in Example-83 by using 3-fluoro-4-iodobenzoyl isocyanate (Intermediate-102, 1.0 g), *tert*-butyl 2-(2-chloro-5-[(2-methylpropanoyl)amino]methyl}phenyl)hydrazinecarboxylate (Intermediate-97, 0.800 g), DCM (15 mL), and trifluoro acetic acid (3.0 mL) to afford 0.900 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.30 (s, 6H), 2.43-2.45 (m, 1H), 4.26 (m, 2H), 7.01 (m, 1H), 7.46-7.49 (m, 1H), 7.63-7.66 (m, 1H), 7.84-7.91 (m, 1H), 8.01-8.06 (m, 1H), 8.22 (s, 1H), 8.36 (br s, 1H), 12.66 (m, 1H); MS (m/z): 515.21 ($M+H$) $^+$.

Step-2:- Preparation of *N*-(4-chloro-3-(3-(3-fluoro-4-(3,3-dimethylbut-1-ynyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide

The title compound was prepared according to the procedure described in Example-111 by using *N*-(4-chloro-3-(3-(3-fluoro-4-iodophenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide (0.080 g, 0.155 mmol), 3,3-dimethylbut-1-yne (0.019 g, 0.233 mmol), TBAF (0.121 g, 0.466 mmol), bis(triphenylphosphine)palladium(II)chloride (0.004 g, 0.006 mmol) and DMSO (3.0 mL) to afford 0.009 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.02 (d, J = 6.9 Hz, 6H), 1.31 (s, 9H), 2.39-2.42 (m, 1H), 4.29 (d, J = 5.7 Hz, 2H), 7.35 (d, J = 7.5 Hz, 1H), 7.43 (s, 1H), 7.57-7.67 (m, 4H), 8.37 (m, 1H), 13.0 (br s, 1H); MS (m/z): 469.31 ($M+H$) $^+$.

Example-150

N-(4-Chloro-3-(3-(4-(2-cyclopropylethynyl)-3-fluorophenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide

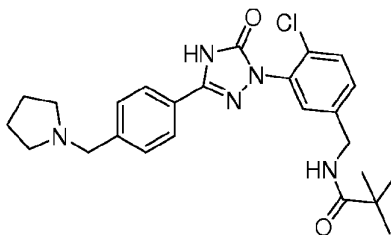


The title compound was prepared according to the procedure described in Example-111 by using *N*-(4-chloro-3-(3-(3-fluoro-4-iodophenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide (step 1 of Example-149, 0.080 g, 0.155 mmol), ethynylcyclopropane (0.019 g, 0.233 mmol), TBAF (0.121 g, 0.466 mmol), bis(triphenylphosphine)palladium(II)chloride (0.004 g, 0.006 mmol) and DMSO (3.0

mL) to afford 0.009 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.79-0.92 (m, 2H), 1.011 (m, 2H), 1.30 (s, 6H), 1.63 (m, 1H), 2.39 (m, 1H), 4.28 (d, $J = 5.7$ Hz, 2H), 7.35 (m, 1H), 7.43 (s, 1H), 7.57-7.67 (m, 4H), 8.37 (m, 1H); MS (m/z): 453.24 ($\text{M}+\text{H}$) $^+$.

Example-151

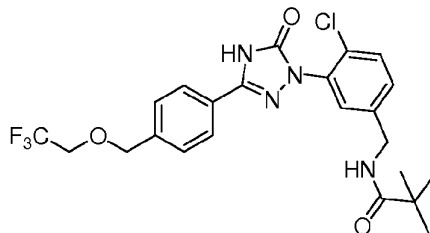
N-(4-Chloro-3-(4,5-dihydro-5-oxo-3-(4-((pyrrolidin-1-yl)methyl)phenyl)-1,2,4-triazol-1-yl)benzyl)pivalamide



To a cold solution of pyrrolidine (0.060 g, 0.830 mmol) in DMF was added NaH (0.034 g, 0.830 mmol) at 0°C and stirred the reaction mass for 1 h. Then solution of *N*-(3-(3-(4-(bromomethyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)-4-chlorobenzyl)pivalamide (Intermediate-103, 0.200 g, 0.419) in DMF was added at 0°C and continued stirring at 5-10°C for 2-3 h. The reaction mass was quenched in ice and pH adjusted to 6-7 and extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated. The obtained crude product was purified by column chromatography on basic alumina, eluting with 5 % MeOH:DCM to afford 0.020 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.23 (s, 9H), 1.79 (br s, 4H), 2.53 (br s, 4H), 3.66 (s, 2H), 4.48 (d, $J = 5.7$ Hz, 2H), 6.08 (m, 1H), 7.26-7.54 (m, 5H), 7.79 (d, $J = 7.8$ Hz, 2H); MS (m/z): 468.51 ($\text{M}+\text{H}$) $^+$.

Example-152

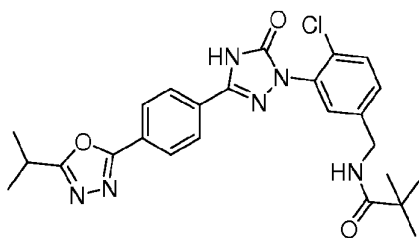
N-(3-(3-(4-((2,2,2-Trifluoroethoxy)methyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)-4-chlorobenzyl)pivalamide



To a solution of 2,2,2-trifluoro ethanol (0.042 g, 0.410 mmol) in THF was added NaH (0.017 g, 0.041 mmol) at 10°C and continued stirring at RT for 1 h. Then solution of *N*-(3-(3-(4-(bromomethyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)-4-chlorobenzyl)pivalamide (Intermediate-103, 0.100 g, 0.041 mmol) in THF and TBAI (0.004 g, 0.0012 mmol) were added to the reaction mixture. The reaction mass was refluxed for 2-3 h. The reaction mass was quenched in ice and pH adjusted to 6-7 and extracted with DCM. The organic layer was dried over anhydrous sodium sulphate and concentrated to afford 0.060 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.12 (s, 9H), 4.11-4.17 (m, 2H), 4.28 (d, *J* = 5.4 Hz, 2H), 4.71 (s, 2H), 7.28 (d, *J* = 9.0 Hz, 1H), 7.41 (s, 1H), 7.46 (d, *J* = 7.8 Hz, 2H), 7.57 (d, *J* = 8.1 Hz, 1H), 7.85 (d, *J* = 8.1 Hz, 2H), 8.18 (t, 1H); MS (*m/z*): 497.31 (M+H)⁺.

Example-153

N-(4-Chloro-3-(4,5-dihydro-3-(4-(5-isopropyl-1,3,4-oxadiazol-2-yl)phenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide

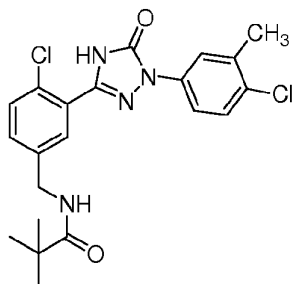


The title compound was prepared according to the procedure described in Example-83 by using 4-(5-isopropyl-1,3,4-oxadiazol-2-yl)benzoyl isocyanate (Intermediate-104, 0.196 g, 0.84 mmol), *tert*-butyl 2-(2-chloro-5-[(2,2-dimethylpropanoyl)amino]methyl]phenyl)hydrazinecarboxylate (Intermediate-94, 0.150 g, 0.420 mmol), DCM (15 mL), and trifluoro acetic acid (3.0 mL) to afford 0.090 g of the

desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.06 (s, 9H), 1.35 (s, 6H), 3.25 (m, 1H), 4.27 (d, J = 5.4 Hz, 2H), 7.34 (d, J = 8.1 Hz, 1H), 7.44 (s, 1H), 7.59 (d, J = 8.4 Hz, 1H), 8.01-8.04 (m, 2H), 8.10-8.17 (m, 3H), 12.73 (s, 1H); MS (m/z) : 495.39 (M+H) $^+$.

Example-154

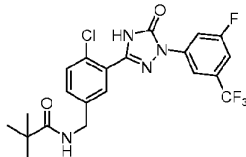
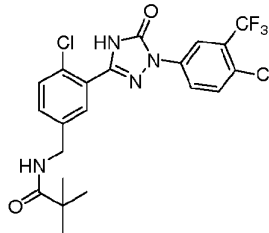
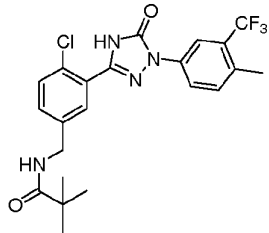
N-(4-Chloro-3-(1-(4-chloro-3-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide

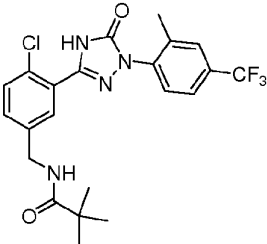
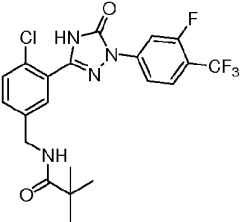
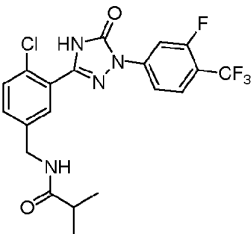


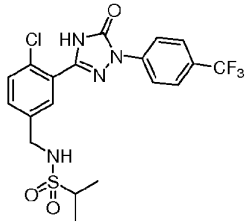
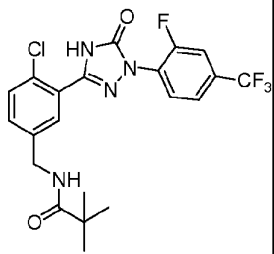
The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chlorophenyl)-2-(4-chloro-3-methylphenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (Intermediate-106, 0.300 g, 0.859 mmol), pivaloyl chloride (0.130 g, 1.03 mmol), TEA (2.0 mL), dry THF (5 mL) to afford 0.140 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 1.13 (s, 9H), 2.37 (s, 3H), 4.29 (d, J = 5.7 Hz, 2H), 7.41 (d, J = 8.7 Hz, 1H), 7.51 (d, J = 8.7 Hz, 1H), 7.58 (s, 2H), 7.70 (br d, 1H), 7.91 (s, 1H), 8.20 (d, J = 6.0 Hz, 1H), 12.56 (s, 1H); MS (m/z) : 433.38 (M+H) $^+$.

Example-155 to Example-163 were prepared by following the procedure described in Example-108 by using corresponding intermediates mentioned in table below, TEA and THF.

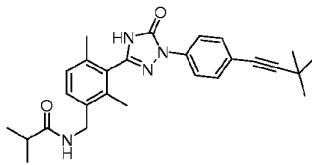
Intermediates used	Example No. and Structure	Example chemical name and characterization data
Intermediate-106 + Isobutyryl chloride	<u>Example-155</u> 	<i>N</i> -(4-Chloro-3-(1-(4-chloro-3-methylphenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)isobutyramide. ^1H NMR (300 MHz, DMSO d_6): δ 1.02 (s, 3H), 1.05 (s, 3H), 2.38 (s, 3H), 4.29 (d, J = 6.0Hz, 2H), 7.42 (d, J = 9.0

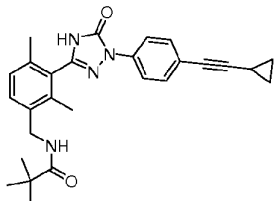
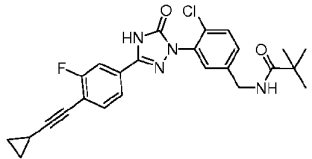
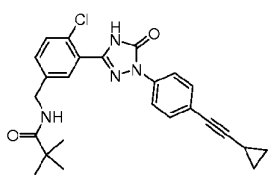
Intermediates used	Example No. and Structure	Example chemical name and characterization data
		Hz, 1H), 7.52 (d, $J = 8.7$ Hz, 1H), 7.60 (s, 2H), 7.82 (br d, 1H), 7.91 (s, 1H), 8.38 (t, 1H), 12.57 (s, 1H); MS (m/z) : 433.30 (M+H) ⁺ .
Intermediate-130 + Pivaloyl chloride	<u>Example-156</u> 	<i>N</i> -(4-Chloro-3-(1-(3-fluoro-5-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.13 (s, 9H), 4.30 (d, $J = 6.3$ Hz, 2H), 7.42 (d, $J = 8.40$ Hz, 1H), 7.61 (m, 3H), 8.09 (d, 1H), 8.14 (s, 1H), 8.20 (t, 1H), 12.80 (s, 1H); MS (m/z) : 471.39 (M+H) ⁺ .
Intermediate-131 + Pivaloyl chloride	<u>Example-157</u> 	<i>N</i> -(4-Chloro-3-(1-(4-chloro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.13 (s, 9H), 4.30 (d, $J = 6.0$ Hz, 2H), 7.42 (d, $J = 7.20$ Hz, 1H), 7.62 (d, $J = 8.10$ Hz, 2H), 7.86 (d, $J = 8.7$ Hz, 1H), 8.19-8.26 (m, 2H), 8.42 (s, 1H), 12.76 (s, 1H); MS (m/z) : 487.43 (M+H) ⁺ .
Intermediate-133 + Pivaloyl chloride	<u>Example-158</u> 	<i>N</i> -(4-Chloro-3-(1-(3-(trifluoromethyl)-4-methylphenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.13 (s, 9H), 2.46 (s, 3H), 4.31 (d, $J = 5.7$ Hz, 2H), 7.39 (d, $J = 7.8$ Hz, 2H), 7.57 (d, 2H), 7.68 (d, 2H), 7.80 (s, 1H), 12.47 (s, 1H); MS (m/z) : 467.22 (M+H) ⁺ .

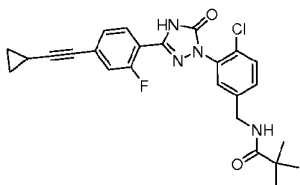
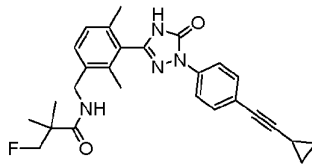
Intermediates used	Example No. and Structure	Example chemical name and characterization data
Intermediate-134 + Pivaloyl chloride	<u>Example-159</u> 	<i>N</i> -(4-Chloro-3-(1-(4-(trifluoromethyl)-2-methylphenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.12 (s, 9H), 2.39 (s, 3H), 4.28 (d, 2H), 7.43 (d, 1H), 7.55-7.60 (m, 3H), 8.12 (d, 1H), 8.20 (m, 1H), 8.29 (s, 1H), 12.65 (s, 1H); MS (m/z) : 467.15 (M+H) ⁺ .
Intermediate-135 + Pivaloyl chloride	<u>Example-160</u> 	<i>N</i> -(4-Chloro-3-(1-(3-fluoro-4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.13 (s, 9H), 4.29 (d, <i>J</i> = 4.8 Hz, 2H), 7.43 (m, 1H), 7.60-7.61 (m, 2H), 7.91-7.94 (m, 1H), 7.98-8.06 (m, 2H), 8.19 (t, 1H); MS (m/z): 471.18 (M+H) ⁺ .
Intermediate-135 + Isobutyryl chloride	<u>Example-161</u> 	<i>N</i> -(4-Chloro-3-(1-(3-fluoro-4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)isobutyramide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.02 (s, 3H), 1.05 (s, 3H), 2.43 (m, 1H), 4.29 (d, <i>J</i> = 6.6 Hz, 2H), 7.42 (d, <i>J</i> = 7.5 Hz, 1H), 7.60 (s, 1H), 7.63 (br s, 1H), 7.91 (t, <i>J</i> = 6.9 Hz, 1H), 7.99-8.08 (m, 2H), 8.40 (t, 1H); MS (m/z) : 455.01 (M+H) ⁺ .
Intermediate-63 + Isopropyl sulfonyl	<u>Example-162</u>	<i>N</i> -(4-Chloro-3-{5-oxo-1-[4-(trifluoromethyl)phenyl]-4,5-dihydro-1 <i>H</i> -1,2,4-triazol-3-yl}benzyl)propane-2-sulfonamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.22 (s, 3H), 1.24 (s, 3H), 3.17 (m, 1H), 4.23 (d, <i>J</i> = 6.0 Hz,

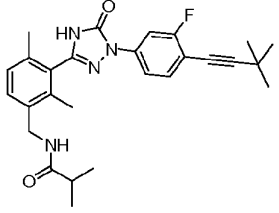
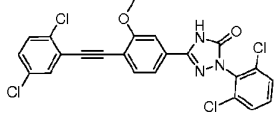
Intermediates used	Example No. and Structure	Example chemical name and characterization data
chloride		2H), 7.56 (d, $J = 7.8$ Hz, 1H), 7.65-7.72 (m, 3H), 7.87 (d, $J = 8.4$ Hz, 2H), 8.19 (d, $J = 8.7$ Hz 2H), 12.72 (s, 1H); MS (m/z) : 475.15 (M+H) ⁺ .
Intermediate-136 + Pivaloyl chloride	<u>Example-163</u> 	<i>N</i> -(4-Chloro-3-(1-(2-fluoro-4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.12 (s, 9H), 7.28 (d, $J = 5.7$ Hz, 2H), 7.40 (d, $J = 7.8$ Hz, 1H), 7.57-7.62 (m, 2H), 7.75 (d, $J = 7.8$ Hz, 1H), 7.90-7.98 (m, 2H), 8.18 (t, 1H), 12.59 (s, 1H); MS (m/z) : 471.17 (M+H) ⁺ .

Example-164 to Example-171 were prepared by following the procedure described in Example-111 by using corresponding intermediates mentioned in table below, bis(triphenylphosphine)palladium(II) chloride, TBAF and DMSO.

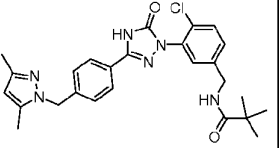
Intermediates used	Example No. and Structure	Example chemical name and Characterization data
Intermediate-140 + 3,3-dimethylbut-1-yne	<u>Example-164</u> 	<i>N</i> -(3-(4,5-Dihydro-1-(4-(3,3-dimethylbut-1-ynyl)phenyl)-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)isobutyramide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.01 (s, 3H), 1.04 (s, 3H), 1.26 (s, 9H), 2.16 (s, 3H), 2.20 (s, 3H), 2.42 (m, 1H), 4.24 (d, $J = 6.3$ Hz, 2H), 7.17 (d, $J = 7.8$ Hz, 1H), 7.27 (d, $J = 8.4$ Hz, 1H), 7.43 (d, $J = 8.7$ Hz, 2H), 7.92 (d, $J = 9.0$ Hz, 2H), 8.19 (t,

Intermediates used	Example No. and Structure	Example chemical name and Characterization data
		1H), 12.23 (s, 1H); MS (m/z) : 445.26 (M+H) ⁺ .
Intermediate-138 + Ethyne-cyclopropane	<u>Example-165</u> 	<i>N</i> -(3-(1-(4-(2-Cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 0.73 (m, 2H), 0.88 (m, 2H), 1.14 (s, 9H), 1.54 (s, 3H), 2.16 (s, 3H), 2.19 (s, 3H), 4.23 (d, <i>J</i> = 6.0 Hz, 2H), 7.16 (d, <i>J</i> = 8.4 Hz, 1H), 7.23 (d, <i>J</i> = 7.8 Hz, 1H), 7.43 (d, <i>J</i> = 8.4 Hz, 2H), 7.91 (d, <i>J</i> = 9.0 Hz, 2H), 8.00 (t, 1H), 12.22 (s, 1H) ; MS (m/z) : 443.51 (M+H) ⁺ .
Intermediate-119 + Ethyne-cyclopropane	<u>Example-166</u> 	<i>N</i> -(4-Chloro-3-(3-(4-(2-cyclopropylethynyl)-3-fluorophenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 0.80 (m, 2H), 0.93-0.95 (m, 2H), 1.12 (s, 9H), 1.63 (m, 1H), 4.28 (d, <i>J</i> = 6.3 Hz, 2H), 7.35 (d, <i>J</i> = 9.0 Hz, 1H), 7.43 (s, 1H), 7.59-7.67 (m, 4H), 8.19 (t, 1H), 12.66 (s, 1H) ; MS (m/z) : 467.24 (M+H) ⁺ .
Intermediate-139 + Ethyne-cyclopropane	<u>Example-167</u> 	<i>N</i> -(4-Chloro-3-(1-(4-(2-cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 0.73 (m, 2H), 0.89 (m, 2H), 1.13 (s, 9H), 1.54 (m, 1H), 4.30 (d, 2H),

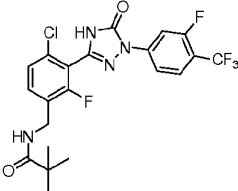
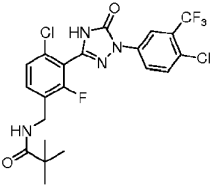
Intermediates used	Example No. and Structure	Example chemical name and Characterization data
		7.41-7.47 (m, 2H), 7.58 (s, 2H), 7.81 (m, 1H), 7.91 (d, $J = 6.3$ Hz, 2H), 8.18 (br s, 1H), 12.56 (s, 1H); MS (m/z) : 449.30 (M+H) ⁺ .
Intermediate-124 + Ethynylcyclopropane	<u>Example-168</u> 	<i>N</i> -(4-Chloro-3-(3-(4-(2-cyclopropylethynyl)-2-fluorophenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 0.78 (m, 2H), 0.93 (m, 2H), 1.12 (s, 9H), 1.59 (m, 1H), 4.28 (d, $J = 6.6$ Hz, 2H), 7.34 (d, $J = 8.4$ Hz, 2H), 7.41 (m, 2H), 7.62 (d, $J = 7.8$ Hz, 1H), 7.75 (d, $J = 7.8$ Hz, 1H), 8.19 (t, 1H), 12.40 (s, 1H); MS (m/z) : 467.20 (M) ⁺ .
Intermediate-143 + Ethynylcyclopropane	<u>Example-169</u> 	<i>N</i> -(3-(1-(4-(2-Cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)-3-fluoro-2,2-dimethylpropanamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 0.74 (m, 2H), 0.85 (m, 2H), 1.15 (s, 6H), 1.54 (m, 1H), 2.15 (s, 3H), 2.19 (s, 3H), 4.27 (d, 2H), 4.43 (s, 1H), 4.49 (s, 1H), 7.17-7.24 (m, 2H), 7.43 (d, $J = 8.4$ Hz, 2H), 7.90 (d, $J = 8.7$ Hz, 2H), 8.16 (s, 1H), 12.22 (s, 1H); MS (m/z): 461.21 (M+H) ⁺ .
Intermediate-141 + 3,3-	<u>Example-170</u>	<i>N</i> -(3-(1-(3-Fluoro-4-(3,3-dimethylbut-1-ynyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)isobutyramide. ¹ H NMR

Intermediates used	Example No. and Structure	Example chemical name and Characterization data
dimethylbut-1-yne		(300 MHz, DMSO d_6): δ 1.01 (s, 3H), 1.04 (s, 3H), 1.23 (s, 9H), 2.15 (s, 3H), 2.20 (s, 3H), 4.25 (d, 2H), 7.18 (m, 1H), 7.26 (m, 1H), 7.51 (m, 1H), 7.84 (m, 2H), 8.19 (t, 1H), 12.35 (s, 1H); MS (m/z): 463.151 (M+H) ⁺ .
Intermediate-45 + 1,4-dichloro-2-ethynylbenzene	<u>Example-171</u> 	2-(2,6-Dichlorophenyl)-5-(4-(2-(2,5-dichlorophenyl)ethynyl)-3-methoxyphenyl)-2H-1,2,4-triazol-3(4H)-one. ¹ H NMR (300 MHz, DMSO d_6): δ 3.92 (s, 3H), 7.54 (m, 4H), 7.58-7.69 (m, 4H), 7.75 (s, 1H), 12-13 (br s, 1H); MS (m/z): 506.01 (M+H) ⁺ .

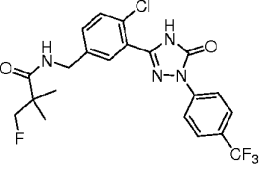
Example-172 was prepared by following the procedure described in Intermediate-41 by using corresponding intermediates mentioned in table below, NaH and DMF.

Intermediate used	Example No. and Structure	Example chemical name and characterization data
Intermediate-103 + 3,5-dimethyl-1H-pyrazole	<u>Example-172</u> 	N-(4-Chloro-3-(4,5-dihydro-3-(4-((3,5-dimethyl-1H-pyrazol-1-yl)methyl)phenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, DMSO d_6): δ 1.11 (s, 9H), 2.10 (s, 3H), 2.15 (s, 3H), 4.28 (d, J = 5.1 Hz, 2H), 5.24 (s, 2H), 5.87 (s, 1H), 7.19 (d, J = 7.8 Hz, 2H), 7.32 (d, J = 8.7 Hz, 1H), 7.22 (s, 1H), 7.59 (d, J = 8.1 Hz, 1H), 7.79 (d, J = 7.5 Hz, 1H), 8.18 (t, 1H); MS (m/z) : 493.36 (M) ⁺ .

Example-173 and Example-174 were prepared by following the procedure as described in Example-83 by using corresponding intermediates mentioned in table below, TFA and DCM.

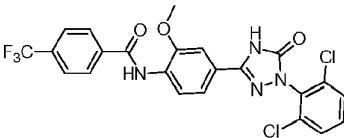
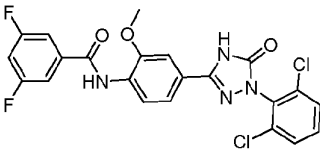
Intermediates used	Example No. and Structure	Example chemical name and characterization data
Intermediate-109 + Intermediate-51	Example-173 	<i>N</i> -(4-Chloro-2-fluoro-3-(1-(3-fluoro-4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)pivalamide. ¹ H NMR (300 MHz, CDCl ₃): δ 1.21 (s, 9H), 4.46 (d, <i>J</i> = 5.7 Hz, 2H), 6.21 (t, 1H), 6.42 (d, 1H), 7.32 (s, 1H), 7.42 (t, 1H), 7.65 (t, 1H), 7.96 (m, 2H); MS (m/z) : 489.56 (M) ⁺ .
Intermediate-111 + Intermediate-51	Example-174 	<i>N</i> -(4-Chloro-3-(1-(4-chloro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.13 (s, 9H), 4.30 (d, <i>J</i> = 6.0 Hz, 2H), 7.46 (d, <i>J</i> = 7.8 Hz, 1H), 7.55 (d, <i>J</i> = 8.1 Hz, 1H), 7.85 (d, <i>J</i> = 8.1 Hz, 1H), 8.20 (t, 2H), 8.41 (s, 1H), 12.90 (m, 1H); MS (m/z): 505.27 (M+H) ⁺ .

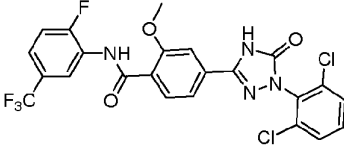
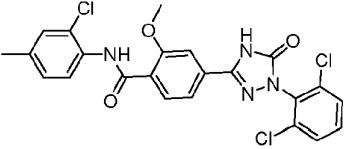
Example-175 was prepared by following the procedure as described in described for step-2 of Example-134 by using Example-115, DAST and THF.

Intermediates used	Example No. and Structure	Example chemical name and characterization data
Example-115	Example-175 	<i>N</i> -(4-Chloro-3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)benzyl)-3-fluoro-2,2-dimethylpropanamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 1.15 (s, 6H), 4.33 (m, 4H), 7.41 (d, <i>J</i> = 7.2 Hz, 1H), 7.61 (d,

Intermediates used	Example No. and Structure	Example chemical name and characterization data
		$J = 7.8$ Hz, 2H), 7.85 (d, $J = 9.3$ Hz, 2H), 8.20 (d, $J = 8.4$ Hz, 2H), 8.35 (t, 1H), 12.71 (m, 1H); MS (m/z) : 471.28 (M+H) ⁺ .

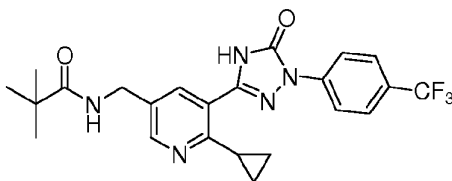
Example-176 to Example-179 were prepared by following the procedure as described for step-6 of Intermediate-26 by using corresponding intermediates used mentioned in table below, trimethyl aluminium (2M solution in toluene) and dry toluene.

Intermediates used	Example No. and Structures	Example chemical name and characterization data
Step-5 of Intermediate-45 + methyl 4-(trifluoromethyl)benzoate	<u>Example-176</u> 	<i>N</i> -(4-(1-(2,6-Dichlorophenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2-methoxyphenyl)-4-(trifluoromethyl)benzamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 3.91 (s, 3H), 7.49-7.64 (m, 3H), 7.72-7.74 (m, 2H), 7.91 (d, $J = 8.4$ Hz, 2H), 7.98 (d, $J = 8.4$ Hz, 1H), 8.15 (d, $J = 8.1$ Hz, 2H), 9.90 (s, 1H), 12.70 (s, 1H); MS (m/z): 523.10 (M+H) ⁺ .
Step-5 of Intermediate-45 + methyl 3,5-difluorobenzoate	<u>Example-177</u> 	<i>N</i> -(4-(1-(2,6-Dichlorophenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2-methoxyphenyl)-3,5-difluorobenzamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 3.90 (s, 3H), 7.47-7.61 (m, 4H), 7.64-7.72 (m, 4H), 7.88 (d, $J = 7.8$ Hz, 1H), 9.86 (s, 1H), 12.69 (s, 1H); MS (m/z): 491.03 (M+H) ⁺ .
Intermediate-	<u>Example-178</u>	4-(1-(2,6-Dichlorophenyl)-4,5-dihydro-

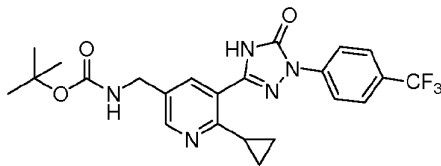
Intermediates used	Example No. and Structures	Example chemical name and characterization data
46 + 2-fluoro-5-(trifluoromethyl)aniline		5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)- <i>N</i> -(2-fluoro-5-(trifluoromethyl)phenyl)-2-methoxybenzamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 4.06 (s, 3H), 7.60-7.65 (m, 5H), 7.55-7.67 (m, 2H), 8.03 (d, <i>J</i> = 7.8 Hz, 1H), 8.60 (s, 1H), 10.44 (s, 1H), 12.90 (s, 1H); MS (<i>m/z</i>): 541.05 (<i>M</i> + <i>H</i>) ⁺ .
Intermediate-46 + 2-chloro-4-methyl aniline	<u>Example-179</u> 	<i>N</i> -(2-Chloro-4-methylphenyl)-4-(1-(2,6-dichlorophenyl)-4,5-dihydro-5-oxo-1 <i>H</i> -1,2,4-triazol-3-yl)-2-methoxybenzamide. ¹ H NMR (300 MHz, DMSO <i>d</i> ₆): δ 2.30 (s, 3H), 4.13 (s, 3H), 7.20 (d, <i>J</i> = 8.7 Hz, 1H), 7.41 (s, 1H), 7.41-7.75 (m, 5H), 8.15 (d, <i>J</i> = 7.8 Hz, 1H), 8.30 (d, <i>J</i> = 8.4 Hz, 1H), 10.47 (s, 1H), 12.91 (s, 1H); MS (<i>m/z</i>): 503.00 (<i>M</i> + <i>H</i>) ⁺ .

Example-180

N-((6-Cyclopropyl-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)pivalamide



Step-1:- Preparation of *tert*-butyl (6-cyclopropyl-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methylcarbamate



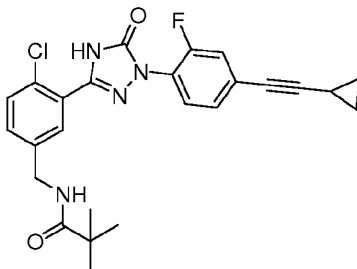
A solution of 6-cyclopropyl-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)pyridine-3-carbonitrile (0.110 g) in ethanol (10 mL) and BOC anhydride (0.150 g), TEA (1.0 mL), Pd/C (catalytic amount) was stirred under hydrogen atmosphere under 35-40 psi pressure in Parr apparatus for 4-5 h. The reaction mass was filtered and the obtained filtrate was concentrated to afford 0.100 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.93-0.98 (m, 4H), 1.36 (s, 9H), 2.09 (m, 1H), 4.12 (d, *J* = 5.7 Hz, 2H), 7.51 (m, 1H), 7.69 (s, 1H), 7.88 (d, *J* = 8.4 Hz, 2H), 8.43 (d, *J* = 7.8 Hz, 2H); MS (*m/z*) : 476.13 (M+H)⁺.

Step-2:- Preparation *N*-((6-cyclopropyl-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)pivalamide

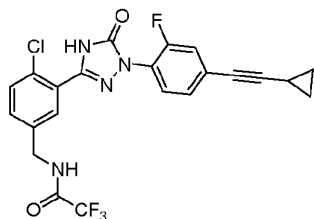
Stirred a solution of *tert*-butyl (6-cyclopropyl-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methylcarbamate (0.100 g) in EtOC:HCl (5.0 mL) for 6 h at RT. Excess of solvent was removed and added DCM (5.0 mL), TEA (0.5 mL) and pivaloyl chloride (0.045 g) under nitrogen atmosphere o the reaction mixture. The reaction mass was stirred at RT for 4 h. Excess of solvent was removed under vacuum and the residue was diluted with water, extracted with EtOAC and concentrated to afford 0.020 g of the product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.98 (m, 4H), 1.12 (s, 9H), 2.73 (m, 1H), 4.27 (d, *J* = 6.0 Hz, 2H), 7.78 (s, 1H), 7.85 (d, *J* = 8.7 Hz, 2H), 8.13 (t, 1H), 8.19 (d, *J* = 9.3 Hz, 2H), 8.40 (s, 1H), 12.75 (s, 1H); MS (*m/z*) : 460.30 (M+H)⁺

Example-181

N-(4-Chloro-3-(1-(4-(2-cyclopropylethynyl)-2-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide

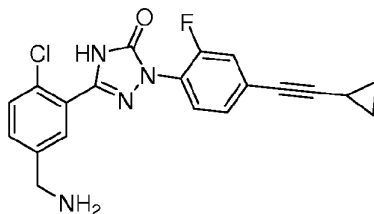


Step 1:- Preparation of *N*-(4-chloro-3-(1-(4-(2-cyclopropylethynyl)-2-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide



The title compound was prepared according to the procedure described in Example-111 by using *N*-(4-chloro-3-(1-(2-fluoro-4-iodophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (Intermediate-126, 1.100 g, 3.88 mmol), ethynylcyclopropane (0.127 g, 1.93 mmol), TBAF (1.10 g, 3.88 mmol), bis(triphenylphosphine)palladium(II) chloride (0.036 g, 0.051 mmol) and DMSO (3.0 mL) to afford 0.400 g of the desired title product. ^1H NMR (400 MHz, DMSO d_6): δ 0.95 (m, 4H), 1.56 (m, 1H), 4.44 (br s, 2H), 7.32 (d, $J = 7.8$ Hz, 1H), 7.42-7.46 (m, 2H), 7.58 (t, $J = 7.8$ Hz, 1H), 7.65 (m, 3H), 10.12 (br s, 1H); MS (m/z): 479.11 ($M+H$) $^+$.

Step 2:- Preparation of 5-(5-(aminomethyl)-2-chlorophenyl)-2-(4-(2-cyclopropylethynyl)-2-fluorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one



The title compound was prepared by following the procedure as described in step-2 of Intermediate-106 by using *N*-(4-chloro-3-(1-(4-(2-cyclopropylethynyl)-2-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide (0.300 g),

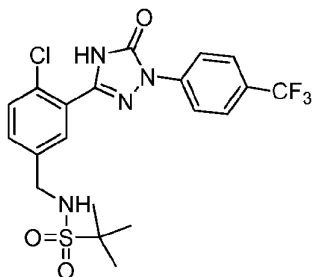
KOH (0.300 g), water (2.0 mL), THF (10.0 mL) to afford 0.200 g of desired product. MS (m/z): 383.16 (M+H)⁺.

Step-3:-Preparation of *N*-(4-chloro-3-(1-(4-(2-cyclopropylethynyl)-2-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide

The title compound was prepared according to the procedure described in Example-108 by using 5-(5-(aminomethyl)-2-chlorophenyl)-2-(4-(2-cyclopropylethynyl)-2-fluorophenyl)-2*H*-1,2,4-triazol-3(4*H*)-one (0.100 g, 0.266 mmol), pivaloyl chloride (0.040 g, 0.319 mmol), THF (10.0 mL) and TEA (2 mL) to afford 0.040 g of the desired product. ¹H NMR (400 MHz, DMSO *d*₆): δ 0.78 (m, 2H), 0.919 (m, 2H), 1.11 (br s, 9H), 1.56 (m, 1H), 4.28 (br d, 2H), 7.32-7.41 (m, 3H), 7.56 (br s, 3H), 8.16 (s, 1H), 12.46 (s, 1H); MS (m/z): 467.22 (M+H)⁺.

Example-182

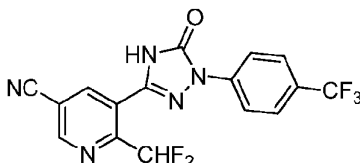
N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2-methylpropane-2-sulfonamide



To a solution of *N*-(4-chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2-methylpropane-2-sulfonamide (Intermediate-144, 0.075 g, 0.158 mmol) in mixture of DCM : acetonitrile : water (0.2 ml : 0.2 mL : 0.3 mL), sodium periodate (0.050 g, 0.238 mmol) and ruthenium chloride (0.001 g, 0.003 mmol) were added and the reaction mass was stirred at RT for 4 h. After completion of the reaction the reaction mass was filtered through celite bed and extracted with DCM. The organic layer was separated, dried over anhydrous sodium sulphate and concentrated. The obtained product was purified with column chromatography on neutral alumina eluting with 10 % MeOH: DCM to afford 0.020 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.30 (s, 9H), 4.31 (d, *J* = 6.0 Hz, 2H), 7.55-7.71 (m, 4H), 7.86 (d, *J* = 8.7 Hz, 2H), 8.19 (d, *J* = 9.0 Hz, 2H), 12.71 (s, 1H). MS (m/z): 489.04 (M+H)⁺.

Example-183

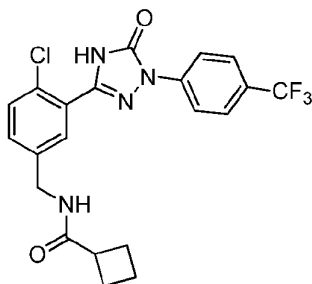
6-(Difluoromethyl)-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)nicotinonitrile



The title compound was prepared by following the procedure as described in Example-83 by using 5-cyano-2-(difluoromethyl)nicotinoyl isocyanate (Intermediate-145, 0.900 g, 4.03 mmol), *tert*-butyl 2-(4-(trifluoromethyl)phenyl)hydrazinecarboxylate (Intermediate-53, 1.0 g, 3.63 mmol), TFA (10 mL), DCM (40 mL) to afford 0.350 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 7.45-7.56 (t, *J* = 53.4 Hz, 1H), 7.90 (d, *J* = 9.3 Hz, 2H), 8.20 (d, *J* = 8.7 Hz, 2H), 8.77 (s, 1H), 9.32 (s, 1H), 13.01 (s, 1H); MS (*m/z*): 382.15 (M+H)⁺.

Example-184

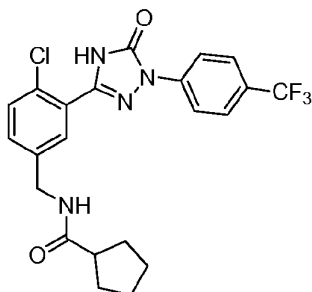
N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)cyclobutanecarboxamide



The title compound was prepared according to the procedure described in Example-108 by using 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-63, 0.100 g, 0.271 mmol), TEA (1.0 mL), DCM (10 mL), cyclobutanecarbonyl chloride (0.041 g, 0.352 mmol) to afford 0.020 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 2.05 (m, 6H), 3.08 (m, 1H), 4.29 (m, 2H), 7.44 (m, 1H), 7.60 (m, 2H), 7.86 (m, 2H), 8.17 (m, 3H), 12.69 (s, 1H); MS (*m/z*): 451.07 (M+H)⁺.

Example-185

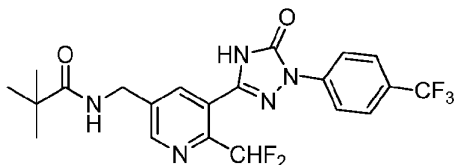
N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)cyclopentanecarboxamide



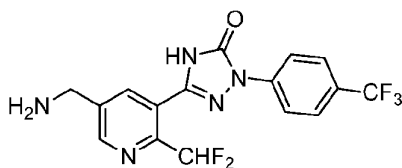
The title compound was prepared according to the procedure described in Example-17 by using 3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (Intermediate-63, 0.100 g, 0.271 mmol), TEA (1.0 mL), TBTU (0.261 g, 0.813 mmol), THF : DMF (10 mL), cyclopentanecarboxylic acid (0.061 g, 0.542 mmol) to afford 0.025 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.50-1.76 (m, 8H), 2.62 (s, 1H), 4.30 (d, 2H), 7.45 (m, 1H), 7.61 (m, 2H), 7.85 (d, *J* = 9.6 Hz, 2H), 8.19 (d, *J* = 8.1 Hz, 2H), 8.40 (m, 1H), 12.69 (s, 1H); MS (*m/z*): 465.08 (M+H)⁺.

Example-186

N-((6-(Difluoromethyl)-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)pivalamide



Step-1:- Preparation of 3-(5-(aminomethyl)-2-(difluoromethyl)pyridin-3-yl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



The title compound was prepared according to the procedure described in Intermediate-16 by using 6-(difluoromethyl)-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)nicotinonitrile (Example-183, 0.050 g, 0.13 mmol), Raney Ni (catalytic

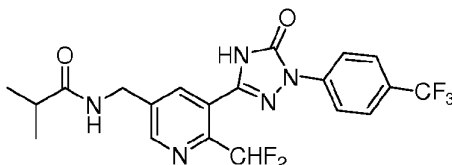
amount), TEA (0.040 g, 0.39 mmol) in ethanol (20 mL) to afford 0.50 g of the desired product.

Step-2:- Preparation of *N*-((6-(difluoromethyl)-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)pivalamide

The title compound was prepared by following the procedure as described in Example-108 by using 3-(5-(aminomethyl)-2-(difluoromethyl)pyridin-3-yl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (0.50 g, 0.12 mmol), pivaloyl chloride (0.018 g, 0.14 mmol), TEA (0.038 g, 0.37 mmol) in DCM (10 mL) to afford 0.010 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.18 (s, 9H), 4.38 (s, 2H), 7.24-7.63 (m, 1H), 7.64 (d, *J* = 8.4 Hz, 2H), 7.99 (s, 1H), 8.10 (d, *J* = 8.4 Hz, 2H), 8.61 (s, 1H); MS (*m/z*): 470.25 (M+H)⁺.

Example-187

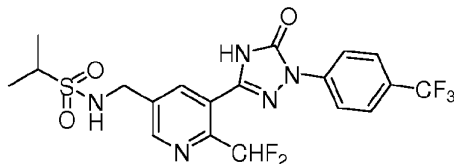
N-((6-(Difluoromethyl)-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)isobutyramide



The title compound was prepared by following the procedure as described in Example-108 by using 3-(5-(aminomethyl)-2-(difluoromethyl)pyridin-3-yl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (step-1 of Example-186, 0.50 g, 0.12 mmol), isobutyryl chloride (0.017 g, 0.15 mmol), TEA (0.042 g, 0.41 mmol) and DCM (10 mL) to afford 0.015 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.14-1.21 (m, 6H), 2.47 (m, 1H), 4.47 (s, 2H), 7.32-7.54 (m, 1H), 7.72 (d, *J* = 8.7 Hz, 2H), 8.09 (br s, 1H), 8.18 (d, *J* = 7.8 Hz, 2H), 8.70 (s, 1H); MS (*m/z*): 456.14 (M+H)⁺.

Example-188

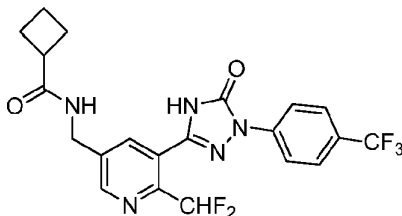
N-((6-(Difluoromethyl)-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)propane-2-sulfonamide



The title compound was prepared by following the procedure as described in Example-108 by using 3-(5-(aminomethyl)-2-(difluoromethyl)pyridin-3-yl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (step-1 of Example-186, 0.75 g, 0.19 mmol), isopropyl sulphonyl chloride (0.042 g, 0.29 mmol), TEA (0.059 g, 0.58 mmol), and DCM (10 mL) to afford 0.012 g of desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.32 (s, 3H), 1.35 (s, 3H), 3.15 (m, 1H), 4.36 (s, 2H), 7.26-7.66 (m, 3H), 8.08-8.11 (m, 3H), 8.65 (s, 1H); MS (m/z): 492.08 (M+H)⁺.

Example-189

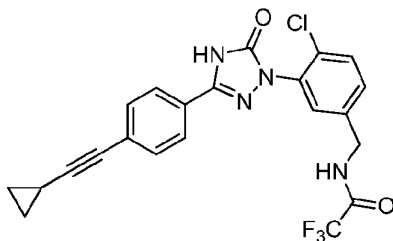
N-((6-(Difluoromethyl)-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)cyclobutanecarboxamide



The title compound was prepared by following the procedure as described in Example-108 by using 3-(5-(aminomethyl)-2-(difluoromethyl)pyridin-3-yl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (step-1 of Example-186, 0.75 g, 0.19 mmol), cyclobutane carbonyl chloride (0.035 g, 0.29 mmol), TEA (0.059 g, 0.58 mmol) and DCM (10 mL) to afford 0.020 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 1.89-2.29 (m, 7H), 4.45 (s, 2H), 7.31-7.52 (m, 1H), 7.72 (d, *J* = 8.4 Hz, 2H), 8.05 (s, 1H), 8.18 (d, *J* = 8.4 Hz, 2H), 8.66 (s, 1H); MS (m/z): 468.26 (M+H)⁺.

Example-190

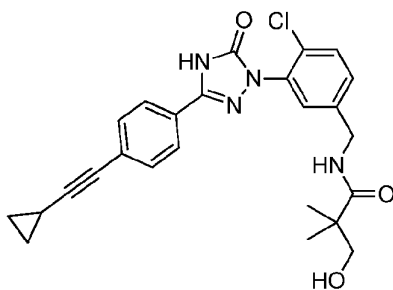
N-(4-Chloro-3-(3-(4-(cyclopropylethynyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)-2,2,2-trifluoroacetamide



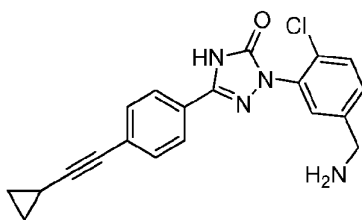
The title compound was prepared according to the procedure described in Example 111 using *N*-(4-chloro-3-(3-(4-iodophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)-2,2,2-trifluoroacetamide (Intermediate-146, 0.300 g, 0.568 mmol), ethynylcyclopropane (0.056 g, 0.852 mmol), TBAF (0.444 g, 1.72 mmol), bis(triphenylphosphine)palladium(II) chloride (0.016 g, 0.022 mmol) and DMSO (3.0 mL) to afford 0.020 g of the desired product. ^1H NMR (300 MHz, DMSO d_6): δ 0.75 (m, 2H), 0.90 (m, 2H), 1.55 (m, 1H), 4.43 (d, $J = 6.0$ Hz, 2H), 7.38 (d, $J = 6.6$ Hz, 1H), 7.46–7.51 (m, 3H), 7.64 (d, $J = 8.1$ Hz, 1H), 7.76 (d, $J = 8.4$ Hz, 2H), 10.06 (m, 1H), 12–13 (br s, 1H). MS (m/z): 461.12 ($M+H$) $^+$.

Example-191

N-(4-Chloro-3-(3-(4-(cyclopropylethynyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)-3-hydroxy-2,2-dimethylpropanamide



Step-1:- Preparation of 1-(5-(aminomethyl)-2-chlorophenyl)-3-(4-(cyclopropylethynyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one



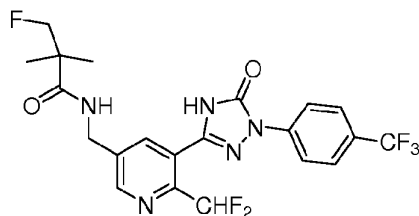
The title compound was prepared by following the procedure as described in step-2 of Intermediate-106 by using *N*-(4-chloro-3-(3-(4-(cyclopropylethynyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)-2,2,2-trifluoroacetamide (Example-190, 0.100 g), LiOH (0.100 g), water (2.0 mL), THF (10.0 mL) to afford 0.050 g of the desired product. MS (m/z): 365.83 (M+H)⁺.

Step-2:- Preparation of *N*-(4-chloro-3-(3-(4-(cyclopropylethynyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)-3-hydroxy-2,2-dimethylpropanamide

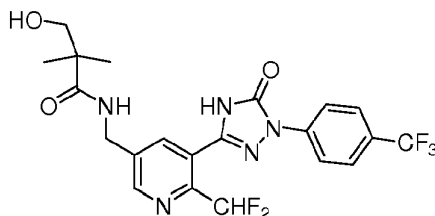
The title compound was prepared according to the procedure described in Example-107 by using 1-(5-(aminomethyl)-2-chlorophenyl)-3-(4-(cyclopropylethynyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (0.080 g, 0.228 mmol), DMF (5 mL), 3-methoxy-2,2-dimethylpropanoic acid (0.040 g, 0.342 mmol), BOP (0.151 g, 0.342 mmol), TEA (2.0 mL) to afford 0.010 g of the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.75 (m, 2H), 0.83 (m, 2H), 1.10 (s, 6H), 1.51 (m, 1H), 3.34 (m 2H), 4.31 (br s, 2H), 4.85 (m, 1H), 7.37-7.50 (m, 3H), 7.58 (d, *J* = 7.8 Hz, 1H), 7.77-7.83 (m, 3H), 8.11 (br s, 1H), 12.56 (s, 1H); MS (m/z): 465.07 (M+H)⁺.

Example-192

3-Fluoro-*N*-((6-(difluoromethyl)-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)-2,2-dimethylpropanamide



Step-1:- Preparation of *N*-((6-(difluoromethyl)-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)-3-hydroxy-2,2-dimethylpropanamide



The title compound was prepared by following the procedure as described in Example-107 by using 3-(5-(aminomethyl)-2-(difluoromethyl)pyridin-3-yl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one (step-1 of Example-186, 0.130 g, 0.33 mmol), DMF (5 mL), 3-methoxy-2,2-dimethylpropanoic acid (0.060 g, 0.50 mmol), BOP (0.224 g, 0.50 mmol), TEA (2.0 mL) to afford 0.070 g of the desired product. MS (m/z): 486.22 (M+H)⁺.

Step-2:- Preparation of 3-fluoro-*N*-((6-(difluoromethyl)-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)-2,2-dimethylpropanamide

The title compound was prepared by following the procedure as described in step-2 of Example-134 by using *N*-((6-(difluoromethyl)-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)-3-hydroxy-2,2-dimethylpropanamide (0.060 g, 0.12 mmol), DAST (0.060 g, 0.37 mmol), DCM (10 mL) to afford 0.005 g the desired product. ¹H NMR (300 MHz, DMSO *d*₆): δ 0.80 (br s, 6H), 4.22 (br s, 1H), 4.38-4.42 (m, 3H), 7.41-7.66 (m, 3H), 7.95 (br s, 1H), 8.08-8.11 (m, 2H), 8.61 (s, 1H); MS (m/z): 488.23 (M+H)⁺.

Pharmacological activity

In-vitro Protocol for screening of mPGES-1 inhibitors:

mPGES-1 (Microsomal prostaglandin E synthase-1) is a microsomal enzyme that converts endoperoxide substrate PGH₂ (prostaglandin H₂) to product PGE₂ (prostaglandin E₂) by isomerization in the presence of reduced glutathione (GSH). mPGES-1 inhibitors were screened by assessing their ability to inhibit formation of PGE₂ from PGH₂ in the presence of mPGES-1 using an anti-PGE₂ antibody based detection method. Recombinant human mPGES-1 was generated in-house by expression in CHO cells (Ouellet M *et al.* (2002), Protein Expression and Purification 26: 489 – 495). The assay was set up using crude microsomal fractions at a protein concentration of 40-60 µg/mL. Test compounds were prepared in 100 % dimethyl sulfoxide (DMSO) to obtain 20 mM stock solution and then diluted using assay buffer comprising 0.1 M Potassium phosphate buffer with 2 mM EDTA. The final concentration of DMSO in the reaction was 0.5 % (v/v). Negative controls were comprised of all assay reagents except the enzyme. Positive controls were comprised of the enzyme reaction in the absence of any inhibitor. Test

compounds were incubated for 10 minutes in assay buffer containing 2.5 mM GSH and mPGES-1 enzyme followed by addition of PGH₂ at a concentration of 15 μ M for 1 minute. The reaction was stopped by addition of Stannous chloride (11mg/ml) and PGE₂ levels were measured (Masse F *et al.* (2005), Journal of Biomolecular Screening 10(6) 599 – 605; Goedken RE *et al.* (2008), Journal of Biomolecular Screening 13 (7): 619 – 625) by HTRF kit (CisBio)).

Inhibition of mPGES-1 enzyme activity was measured using the percent of reaction occurring in the positive control. Concentration response curves were plotted using percent inhibition of maximum enzyme reaction. The IC₅₀ value was calculated from the concentration response curve by nonlinear regression analysis using GraphPad PRISM software.

The compounds prepared were tested using the above assay procedure and the results obtained are given in Table 1. Percentage inhibition at concentrations of 1.0 μ M, 3.0 μ M or 10.0 μ M are given in the table along with IC₅₀ (nM) details for selected examples. The compounds prepared were tested using the above assay procedure and were found to have IC₅₀ less than 200nM, preferably less than 100nM, more preferably less than 50nM or most preferably less than 20nM.

The IC₅₀ (nM) values of some of the compounds are set forth in Table 1 wherein “A” refers to an IC₅₀ value of less than 50 nM, “B” refers to IC₅₀ value in range of 50.01 to 100.0 nM and “C” refers to IC₅₀ values more than 100 nM.

Table 1:

Sr. No.	Example No.	Percentage inhibition at			IC ₅₀ (nM)
		1 μ M	3 μ M	10 μ M	
1.	Example-1	9.32	-	68.18	-
2.	Example-2	17.97	-	18.97	-
3.	Example-3	88.90	96.79	-	A
4.	Example-4	46.43	-	69.70	-
5.	Example-5	91.83	-	86.21	A
6.	Example-6	83.89	-	87.67	A
7.	Example-7	89.12	-	95.25	A

Sr. No.	Example No.	Percentage inhibition at			IC ₅₀ (nM)
		1 μ M	3 μ M	10 μ M	
8.	Example-8	84.86	-	87.24	B
9.	Example-9	1.53	-	23.19	-
10.	Example-10	84.88	-	91.21	A
11.	Example-11	2.34	-	71.72	-
12.	Example-12	84.76	-	87.54	A
13.	Example-13	86.27	-	90.42	C
14.	Example-14	95.99	-	95.82	B
15.	Example-15	64.15	-	83.79	-
16.	Example-16	40.88	-	92.28	-
17.	Example-17	2.52	-	44.08	-
18.	Example-18	56.23	-	98.10	-
19.	Example-19	5.11	-	29.66	-
20.	Example-20	22.37	-	87.36	-
21.	Example-21	76.06	-	91.78	-
22.	Example-22	39.26	-	99.73	-
23.	Example-23	13.25	-	52.94	-
24.	Example-24	70.96	-	83.09	-
25.	Example-25	74.44	-	88.92	C
26.	Example-26	80.56	-	88.14	C
27.	Example-27	72.84	-	97.54	-
28.	Example-28	72.22	-	95.51	-
29.	Example-29	79.21	-	85.78	C
30.	Example-30	89.60	-	94.52	A
31.	Example-31	96.88	-	98.63	B
32.	Example-32	88.85	-	93.97	B
33.	Example-33	89.42	-	96.78	A
34.	Example-34	70.73	-	92.28	-

Sr. No.	Example No.	Percentage inhibition at			IC ₅₀ (nM)
		1 μ M	3 μ M	10 μ M	
35.	Example-35	37.43	-	59.15	-
36.	Example-36	97.34	-	98.90	A
37.	Example-37	50.96	-	94.31	-
38.	Example-38	87.36	-	84.91	B
39.	Example-39	90.06	-	96.37	B
40.	Example-40	89.05	-	90.68	A
41.	Example-41	58.81	-	87.08	-
42.	Example-42	96.55	-	99.65	A
43.	Example-43	98.58	-	99.05	A
44.	Example-44	97.37	-	99.03	A
45.	Example-45	81.65	-	90.11	A
46.	Example-46	59.77	-	72.09	-
47.	Example-47	89.46	-	96.51	A
48.	Example-48	98.33	-	100.00	A
49.	Example-49	89.15	-	96.34	A
50.	Example-50	95.32	-	99.85	B
51.	Example-51	88.29	-	85.96	A
52.	Example-52	96.44	-	98.34	A
53.	Example-53	82.70	-	97.89	A
54.	Example-54	89.36	-	97.50	B
55.	Example-55	80.95	91.85	-	A
56.	Example-56	98.08	-	98.54	A
57.	Example-57	98.73	98.55	-	A
58.	Example-58	95.20	-	97.32	B
59.	Example-59	93.91	-	96.80	A
60.	Example-60	95.39	-	93.07	A
61.	Example-61	92.44	-	96.85	A

Sr. No.	Example No.	Percentage inhibition at			IC ₅₀ (nM)
		1 μ M	3 μ M	10 μ M	
62.	Example-62	97.77	-	98.29	A
63.	Example-63	92.41	-	94.79	A
64.	Example-64	28.94	-	63.78	-
65.	Example-65	75.45	-	94.48	-
66.	Example-66	87.85	-	98.83	C
67.	Example-67	72.71	-	96.32	-
68.	Example-68	97.39	-	96.41	A
69.	Example-69	86.44	-	89.11	C
70.	Example-70	0.55	-	6.63	-
71.	Example-71	12.95	-	17.88	-
72.	Example-72	99.24	99.05	-	A
73.	Example-73	92.38	-	92.43	A
74.	Example-74	93.25	-	98.71	B
75.	Example-75	99.07	-	98.31	A
76.	Example-76	95.62	-	95.38	A
77.	Example-77	93.21	-	98.06	B
78.	Example-78	99.94	-	99.88	A
79.	Example-79	99.14	-	97.14	A
80.	Example-80	96.99	-	97.10	A
81.	Example-81	99.76	-	98.05	A
82.	Example-82	98.26	-	98.96	A
83.	Example-83	97.49	-	99.86	A
84.	Example-84	98.81	-	99.51	A
85.	Example-85	22.58	-	69.28	-
86.	Example-86	79.87	-	96.68	C
87.	Example-87	93.36	-	99.19	B
88.	Example-88	3.34	-	39.18	-

Sr. No.	Example No.	Percentage inhibition at			IC ₅₀ (nM)
		1 μ M	3 μ M	10 μ M	
89.	Example-89	95.30	-	99.06	A
90.	Example-90	81.74	-	97.20	C
91.	Example-91	66.15	-	94.31	-
92.	Example-92	76.02	-	99.37	C
93.	Example-93	96.83	-	99.60	A
94.	Example-94	87.02	-	100.00	C
95.	Example-95	98.71	-	99.04	A
96.	Example-96	52.25	-	68.70	-
97.	Example-97	20.19	-	6.88	-
98.	Example-98	97.52	-	99.45	A
99.	Example-99	95.30	-	98.55	A
100.	Example-100	99.97	-	98.56	A
101.	Example-101	94.97	-	98.18	A
102.	Example-102	96.52	-	98.03	A
103.	Example-103	19.92	-	51.12	-
104.	Example-104	96.91	-	99.50	A
105.	Example-105	73.88	-	95.96	C
106.	Example-106	92.03	-	96.77	C
107.	Example-107	56.07	-	90.50	-
108.	Example-108	89.82	-	99.72	C
109.	Example-109	97.10	-	97.15	B
110.	Example-110	47.19	-	95.07	-
111.	Example-111	87.20	-	96.15	A
112.	Example-112	99.53	-	99.83	A
113.	Example-113	99.30	-	99.06	A
114.	Example-114	99.06	-	98.39	A
115.	Example-115	96.49	-	98.22	A

Sr. No.	Example No.	Percentage inhibition at			IC ₅₀ (nM)
		1 μ M	3 μ M	10 μ M	
116.	Example-116	97.97	-	95.50	A
117.	Example-117	67.93	-	95.02	-
118.	Example-118	76.25	-	87.82	C
119.	Example-119	88.70	-	94.30	A
120.	Example-120	11.45	-	46.01	-
121.	Example-121	16.28	-	72.53	-
122.	Example-122	92.22	-	93.49	A
123.	Example-123	93.78	-	100.00	A
124.	Example-124	27.96	-	88.96	-
125.	Example-125	79.87	-	94.99	-
126.	Example-126	100.00	-	99.29	A
127.	Example-127	32.43	-	58.48	-
128.	Example-128	91.83	-	89.35	A
129.	Example-129	93.50	-	92.14	A
130.	Example-130	96.56	-	97.79	B
131.	Example-131	100.00	-	99.23	A
132.	Example-132	91.76	-	89.83	A
133.	Example-133	98.00	-	97.89	A
134.	Example-134	91.45	-	93.88	A
135.	Example-135	92.26	-	98.89	A
136.	Example-136	95.06	-	94.79	A
137.	Example-137	93.36	-	93.06	A
138.	Example-138	93.30	-	92.91	A
139.	Example-139	93.94	-	99.06	A
140.	Example-140	97.36	-	98.11	A
141.	Example-141	97.57	-	97.48	A
142.	Example-142	97.25	-	93.98	A

Sr. No.	Example No.	Percentage inhibition at			IC ₅₀ (nM)
		1 μ M	3 μ M	10 μ M	
143.	Example-143	99.21	-	90.70	A
144.	Example-144	97.49	-	92.73	A
145.	Example-145	94.24	-	98.86	A
146.	Example-146	100.00	-	99.73	A
147.	Example-147	98.64	-	97.61	A
148.	Example-148	98.91	-	97.78	A
149.	Example-149	98.34	-	94.61	A
150.	Example-150	98.38	-	99.52	A
151.	Example-151	64.96	-	95.89	-
152.	Example-152	96.01	-	100.00	A
153.	Example-153	93.94	-	99.65	B
154.	Example-154	95.18	-	100.00	A
155.	Example-155	99.74	-	97.08	A
156.	Example-156	98.50	-	99.25	A
157.	Example-157	98.54	-	96.46	A
158.	Example-158	99.05	-	99.70	A
159.	Example-159	75.40	-	96.76	C
160.	Example-160	96.62	-	97.33	A
161.	Example-161	98.04	-	99.36	A
162.	Example-162	99.35	-	99.33	A
163.	Example-163	91.82	-	98.66	A
164.	Example-164	98.30	-	97.19	A
165.	Example-165	97.91	-	99.51	A
166.	Example-166	99.68	-	100.00	A
167.	Example-167	99.03	-	98.43	A
168.	Example-168	100.00	-	98.89	A
169.	Example-169	95.54	-	99.17	C

Sr. No.	Example No.	Percentage inhibition at			IC ₅₀ (nM)
		1 μ M	3 μ M	10 μ M	
170.	Example-170	99.83	-	98.87	A
171.	Example-171	95.22	-	99.60	A
172.	Example-172	91.86	-	99.04	C
173.	Example-173	96.77	-	98.95	A
174.	Example-174	99.90	-	100.00	A
175.	Example-175	99.44	-	99.44	A
176.	Example-176	45.84	-	89.54	-
177.	Example-177	29.89	-	71.07	-
178.	Example-178	42.72	-	91.55	-
179.	Example-179	50.59	-	95.17	-
180.	Example-180	72.34	-	92.51	C
181.	Example-181	88.24	-	94.28	A
182.	Example-182	92.94	-	94.05	A
183.	Example-183	41.55	-	85.5	-
184.	Example-184	98.59	-	94.26	A
185.	Example-185	96.06	-	98.3	A
186.	Example-186	96.84	-	98.02	A
187.	Example-187	98.63	-	97.51	A
188.	Example-188	89.49	-	99.32	B
189.	Example-189	96.17	-	98.48	A
190.	Example-190	100	-	100	A
191.	Example-191	93.5	-	100	C
192.	Example-192	99.3	-	99.31	A

Screening for mPGES-1 inhibitors using the A549 cell based assay

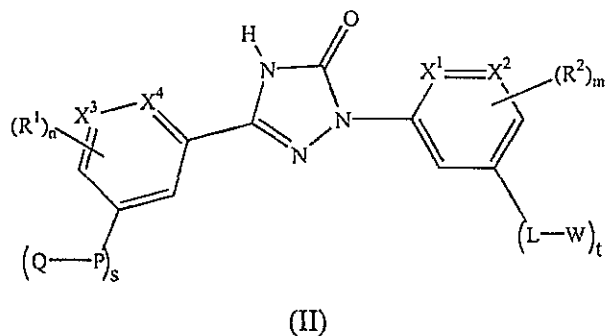
The inhibition of mPGES-1 enzyme in the A549 cell line was monitored as inhibition of IL-1 β induced PGE₂ release. A549 cells were maintained in DMEM medium with 10% FBS and 1% Penicillin-Streptomycin Solution in 5% CO₂ at 37°C.

Cells were seeded 24 h prior to the assay in 96 well plates in DMEM containing 1% Penicillin-Streptomycin and 2% FBS so as to get ~ 40,000 cells per well on the day of experiment. The assay was carried out in a total volume of 200 μ L. Test compounds were dissolved in dimethyl sulfoxide (DMSO) to prepare 2 mM stock solution and then diluted using plain DMEM. The final concentration of DMSO in the reaction was 0.55% (v/v). Cells were treated with test compounds for 30 minutes followed by addition of IL-1 β at a final concentration of 10 ng/mL for 16-20 h. Plates were then centrifuged at 1000 rpm for 10 min at 4⁰C. Supernatants were collected and analyzed by the addition of PGE₂-D2 and anti-PGE₂ cryptate conjugate supplied by the CisBio HTRF kit in a 96 well half area Blackwell EIA/RIA plate. The assay plate was incubated overnight at 4-5⁰ C before being read in an Artemis (K-101) (Japan) HTRF plate reader and levels of PGE₂ calculated by extrapolation from the standard curve.

The concentration response curves were plotted as a percentage of maximal response obtained in the absence of test antagonist. The IC₅₀ value was calculated from the concentration response curve by nonlinear regression analysis using GraphPad PRISM software.

WHAT IS CLAIMED IS:

1. A compound of formula (II)



or a pharmaceutically acceptable salt thereof,

wherein,

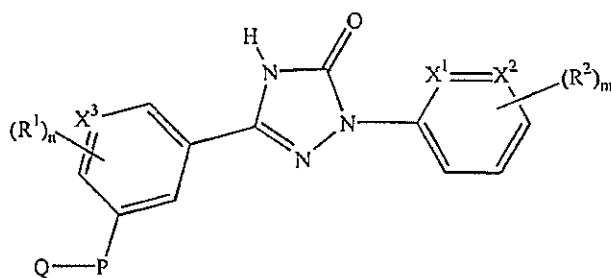
- X^1 , X^2 , X^3 and X^4 are each independently selected from CH and N;
- each occurrence of L is independently selected from $-CH_2NHC(O)-$ and $-CH_2NHS(O)_2-$;
- each occurrence of P is independently selected from $-CH_2NHC(O)-$ and $-CH_2NHS(O)_2-$;
- each occurrence of Q is independently selected from isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-methoxy-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl, cyclopropyl, cyclobutyl, cyclopentyl, 2-fluorophenyl, tetrahydrofuranyl, tetrahydrofuran-2-yl, (S)-tetrahydrofuran-2-yl, (R)-tetrahydrofuran-2-yl, isoxazolyl and 1-methyl-1*H*-imidazole-2-yl;
- each occurrence of W is independently selected from isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl and cyclopropyl;
- each occurrence of R^1 is independently selected from cyano, Cl, F, CHF_2 , CF_3 , OCH_3 , CH_3 , (2,2,2-trifluoroethoxy)methyl, cyclopropyl, (pyrrolidin-1-yl)methyl, 5-isopropyl-1,3,4-oxadiazol-2-yl, (3,5-dimethyl-1*H*-pyrazol-1-yl)methyl, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, (2,5-dichlorophenyl)ethynyl, (4-chloro-2-fluorophenyl)ethynyl, (3-chloro-2-fluorophenyl)ethynyl, (3-(trifluoromethyl)phenyl)ethynyl and (2-chloro-4-(trifluoromethyl)phenyl)ethynyl;
- each occurrence of R^2 is independently selected from cyano, Cl, F, CH_3 , CF_3 , OCH_3 , 3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl, cyclopropanecarboxamido, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, $-CONH-[3-(trifluoromethyl)phenyl]$, $-CONH-[3-(difluoromethyl)-4-fluorophenyl]$ and $-CONH-[3-(difluoromethyl)phenyl]$;
- 'm' is an integer ranging from 0 to 3, both inclusive;
- 'n' is an integer ranging from 0 to 3, both inclusive;

's' is an integer ranging from 0 to 1, both inclusive; and

't' is an integer ranging from 0 to 1, both inclusive;

with the provisos that (i) 's' and 't' are not '0' simultaneously, and (ii) 'm' and 'n' are not '0' simultaneously.

2. The compound according to claim 1, wherein
'n' is 1 or 2.
3. The compound according to claim 1 or 2, wherein
'm' is 1 or 2.
4. The compound according to any one of claims 1 to 3, wherein
's' is 1; and 't' is 0.
5. The compound according to any one of claims 1 to 4, wherein 't' is 1; and 's' is 0.
6. A compound of the formula (III)



(III)

or a pharmaceutically acceptable salt thereof,

wherein,

X¹, X² and X³ are each independently selected from CH and N;

P is selected from -CH₂NHC(O)- and -CH₂NHS(O)₂-;

Q is selected from isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-methoxy-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl, cyclopropyl, cyclobutyl, cyclopentyl, 2-fluorophenyl, tetrahydrofuranyl, tetrahydrofuran-2-yl, (S)-tetrahydrofuran-2-yl, (R)-tetrahydrofuran-2-yl, isoxazolyl and 1-methyl-1*H*-imidazole-2-yl;

each occurrence of R¹ is independently selected from cyano, Cl, F, CHF₂, CF₃, OCH₃, CH₃ and cyclopropyl;

each occurrence of R² is independently selected from independently cyano, Cl, F, CH₃, CF₃, OCH₃, 3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl, cyclopropanecarboxamido, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, -CONH-[3-(trifluoromethyl)phenyl], -CONH-[3-(difluoromethyl)-4-fluorophenyl] and -CONH-[3-(difluoromethyl)phenyl];

'm' is an integer ranging from 0 to 3, both inclusive; and

'n' is an integer ranging from 0 to 3, both inclusive; with the proviso that 'm' and 'n' are not '0' simultaneously.

7. The compound according to claim 6, wherein X^1 , X^2 and X^3 are CH.

8. The compound according to claim 6 or 7, wherein

each R^1 is, independently, cyano, Cl, F, CHF_2 , CF_3 , OCH_3 , CH_3 or cyclopropyl;

each R^2 is, independently, cyano, Cl, F, CH_3 , CF_3 or OCH_3 ;

P is $-CH_2NHC(O)-$;

'm' is 1 or 2;

'n' is 0, 1 or 2; and

Q is isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-methoxy-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl, cyclopropyl, cyclobutyl, cyclopentyl, 2-fluorophenyl, tetrahydrofuranyl, tetrahydrofuran-2-yl, (S)-tetrahydrofuran-2-yl, (R)-tetrahydrofuran-2-yl, isoxazolyl or 1-methyl-1*H*-imidazole-2-yl.

9. A compound selected from

4-(3-(2-Chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-methoxy-*N*-(3-(trifluoromethyl)phenyl)benzamide;

N-(4-Chloro-3-(1-(4-(3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl)-3-methoxyphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)cyclopropanecarboxamide;

4-(3-(2-Chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(difluoromethyl)-4-fluorophenyl)-2-methoxybenzamide;

N-(4-Chloro-3-(1-(4-(cyclopropanecarboxamido)-3-methoxyphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)cyclopropanecarboxamide;

4-(3-(2-Chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(difluoromethyl)phenyl)-2-methoxybenzamide;

4-(3-(2-Chloro-5-(cyclopropanecarboxamidomethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-*N*-(3-(trifluoromethyl)phenyl)benzamide;

N-(4-Chloro-3-(3-(4-((2,5-dichlorophenyl)ethynyl)-2-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide;

N-(4-Chloro-3-(3-(4-((4-chloro-2-fluorophenyl)ethynyl)-2-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide;

N-(4-Chloro-3-(3-(2-fluoro-4-((3-(trifluoromethyl)phenyl)ethynyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide;

N-(4-Chloro-3-(3-(4-((3-chloro-2-fluorophenyl)ethynyl)-2-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide;

N-(4-Chloro-3-(3-(4-((2-chloro-4-(trifluoromethyl)phenyl)ethynyl)-2-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide;

N-(4-Chloro-3-(1-(3,4-dichlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide;

N-(4-Chloro-2-fluoro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(2,4-difluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide;

N-(4-Chloro-2-fluoro-3-(1-(4-methoxyphenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-2-fluoro-3-(3-(2-fluoro-4-(trifluoromethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-3-(3-(2-chloro-6-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-fluorobenzyl)pivalamide;

N-(4-Chloro-3-(1-(4-cyanophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide;

N-(4-Chloro-3-(3-(4-chloro-2-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)-2-fluorobenzyl)pivalamide;

N-(4-Chloro-3-(3-(4-chloro-2-fluorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide;

N-(4-Chloro-3-(3-(2-fluoro-4-(trifluoromethyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)cyclopropanecarboxamide;

N-(4-Chloro-2-fluoro-3-(5-oxo-3-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-2-fluoro-3-(5-oxo-1-(5-(trifluoromethyl)pyridin-2-yl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-2-fluoro-3-(5-oxo-1-(6-(trifluoromethyl)pyridin-3-yl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(4-chlorophenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide;

N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide;

N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

(*R*)-*N*-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)tetrahydrofuran-2-carboxamide;

N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)cyclopropanesulfonamide;

N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2-fluorobenzamide;

N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)isoxazole-5-carboxamide;

N-(3-(1-(4-(Trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)-2,2,2-trifluoroacetamide;

(*R*)-*N*-(2,4-Dimethyl-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)tetrahydrofuran-2-carboxamide;

N-(3-(1-(4-(Trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)pivalamide;

(*S*)-*N*-(4-Chloro-3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-tetrahydrofuran-2-carboxamide;

N-(4-Chloro-3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-1-methyl-1*H*-imidazole-2-carboxamide;

N-(4-Chloro-2-fluoro-3-(4,5-dihydro-3-(4-(3,3-dimethylbut-1-ynyl)phenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-3-(3-(4-(2-cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)-2-fluorobenzyl)pivalamide;

N-(4-Chloro-2-fluoro-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide;

N-(4-Chloro-2-fluoro-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-3-hydroxy-2,2-dimethylpropanamide;

N-(4-Chloro-3-(1-(4-chloro-3-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide;

(*R*)-*N*-(4-chloro-2-fluoro-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)tetrahydrofuran-2-carboxamide;

N-(4-Chloro-3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-3-methoxy-2,2-dimethylpropanamide;

N-(4-Chloro-3-(1-(4-chloro-3-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-5-(1-(4-chlorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide;

N-(4-Chloro-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-methoxybenzyl)pivalamide;

N-(4-Chloro-2-fluoro-5-(1-(4-fluoro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-2-fluoro-5-(1-(6-(trifluoromethyl)pyridin-3-yl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(2,4-Dichloro-5-(3-(4-(2-cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-5-(1-(3-chloro-4-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide;

N-(4-Chloro-5-(1-(4-chlorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)-2-fluorobenzamide;

N-(4-Chloro-2-fluoro-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)isobutyramide;

N-(4-Chloro-5-(1-(4-chlorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)-3-fluoro-2,2-dimethylpropanamide;

N-(4-Chloro-5-(1-(3-chloro-4-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide;

N-(4-Chloro-5-(3-(4-(2-cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)-2-fluorobenzyl)pivalamide;

N-(4-Chloro-2-fluoro-5-(4,5-dihydro-3-(4-(3,3-dimethylbut-1-ynyl)phenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(6-(trifluoromethyl)pyridin-3-yl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(3-chloro-4-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(3-chloro-4-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(3-(4-(2-cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(4,5-dihydro-3-(4-(3,3-dimethylbut-1-ynyl)phenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide;

N-(4-Chloro-3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)isobutyramide;

N-(4-Chloro-3-(3-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-3-(3-(3-fluoro-4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-3-(3-(4-chloro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-3-(3-(4-fluoro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-3-(3-(3-fluoro-4-(3,3-dimethylbut-1-ynyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide;

N-(4-Chloro-3-(3-(4-(2-cyclopropylethynyl)-3-fluorophenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)isobutyramide;

N-(4-Chloro-3-(4,5-dihydro-5-oxo-3-(4-((pyrrolidin-1-yl)methyl)phenyl)-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(3-(3-(4-((2,2,2-Trifluoroethoxy)methyl)phenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)-4-chlorobenzyl)pivalamide;

N-(4-Chloro-3-(4,5-dihydro-3-(4-(5-isopropyl-1,3,4-oxadiazol-2-yl)phenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(4-chloro-3-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(4-chloro-3-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)isobutyramide;

N-(4-Chloro-3-(1-(3-fluoro-5-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(4-chloro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(3-(trifluoromethyl)-4-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(4-(trifluoromethyl)-2-methylphenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(3-fluoro-4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(3-fluoro-4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)isobutyramide;

N-(4-Chloro-3-{5-oxo-1-[4-(trifluoromethyl)phenyl]-4,5-dihydro-1*H*-1,2,4-triazol-3-yl}benzyl)propane-2-sulfonamide;

N-(4-Chloro-3-(1-(2-fluoro-4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(3-(4,5-Dihydro-1-(4-(3,3-dimethylbut-1-ynyl)phenyl)-5-oxo-1*H*-1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)isobutyramide;

N-(3-(1-(4-(2-Cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)pivalamide;

N-(4-Chloro-3-(3-(4-(2-cyclopropylethynyl)-3-fluorophenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(4-(2-cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(3-(4-(2-cyclopropylethynyl)-2-fluorophenyl)-4,5-dihydro-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(3-(1-(4-(2-Cyclopropylethynyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)-3-fluoro-2,2-dimethylpropanamide;

N-(3-(1-(3-Fluoro-4-(3,3-dimethylbut-1-ynyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2,4-dimethylbenzyl)isobutyramide;

N-(4-Chloro-3-(4,5-dihydro-3-(4-((3,5-dimethyl-1*H*-pyrazol-1-yl)methyl)phenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

N-(4-Chloro-2-fluoro-3-(1-(3-fluoro-4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(1-(4-chloro-3-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)-2-fluorobenzyl)pivalamide;

N-(4-Chloro-3-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)-3-fluoro-2,2-dimethylpropanamide;

N-((6-Cyclopropyl-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)pivalamide;

N-(4-Chloro-3-(1-(4-(2-cyclopropylethynyl)-2-fluorophenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)benzyl)pivalamide;

N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2-methylpropane-2-sulfonamide;

N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)cyclobutanecarboxamide;

N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)cyclopentanecarboxamide;

N-((6-(Difluoromethyl)-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)pivalamide;

N-((6-(Difluoromethyl)-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)isobutyramide;

N-((6-(Difluoromethyl)-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)propane-2-sulfonamide;

N-((6-(Difluoromethyl)-5-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)cyclobutanecarboxamide;

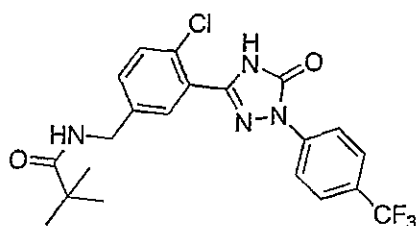
N-(4-Chloro-3-(3-(4-(cyclopropylethynyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)-2,2,2-trifluoroacetamide;

N-(4-Chloro-3-(3-(4-(cyclopropylethynyl)phenyl)-5-oxo-4,5-dihydro-1*H*-1,2,4-triazol-1-yl)benzyl)-3-hydroxy-2,2-dimethylpropanamide;

3-Fluoro-*N*-((6-(difluoromethyl)-5-(1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-5-oxo-1*H*-1,2,4-triazol-3-yl)pyridin-3-yl)methyl)-2,2-dimethylpropanamide;

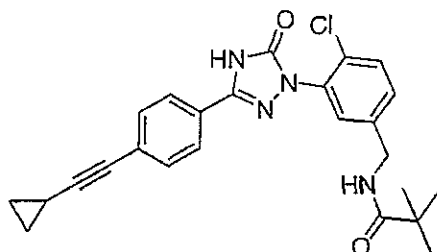
and pharmaceutically acceptable salts thereof.

10. A compound of the formula



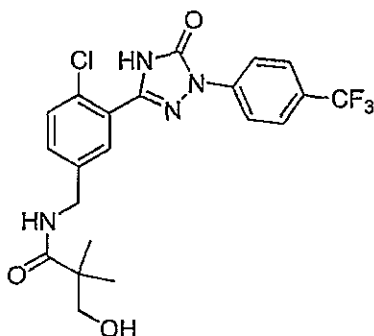
or a pharmaceutically acceptable salt thereof.

11. A compound of the formula



or a pharmaceutically acceptable salt thereof.

12. A compound of formula



or a pharmaceutically acceptable salt thereof.

13. A compound selected from

3-(5-(aminomethyl)-2-chlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1*H*-1,2,4-triazol-5(4*H*)-one;

N-(4-Chloro-3-(5-oxo-1-(4-(trifluoromethyl)phenyl)-4,5-dihydro-1*H*-1,2,4-triazol-3-yl)benzyl)-2,2,2-trifluoroacetamide;

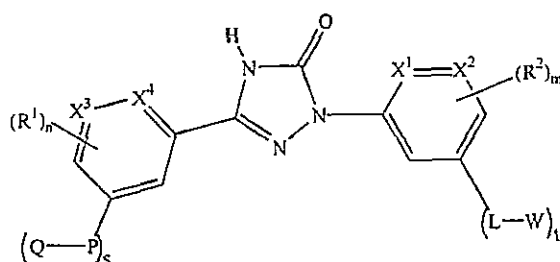
N-(4-Chloro-3-(4,5-dihydro-3-(4-iodophenyl)-5-oxo-1,2,4-triazol-1-yl)benzyl)pivalamide;

and pharmaceutically acceptable salts thereof.

14. A pharmaceutical composition comprising a compound according to any one of claims 1 to 12 and a pharmaceutically acceptable excipient.

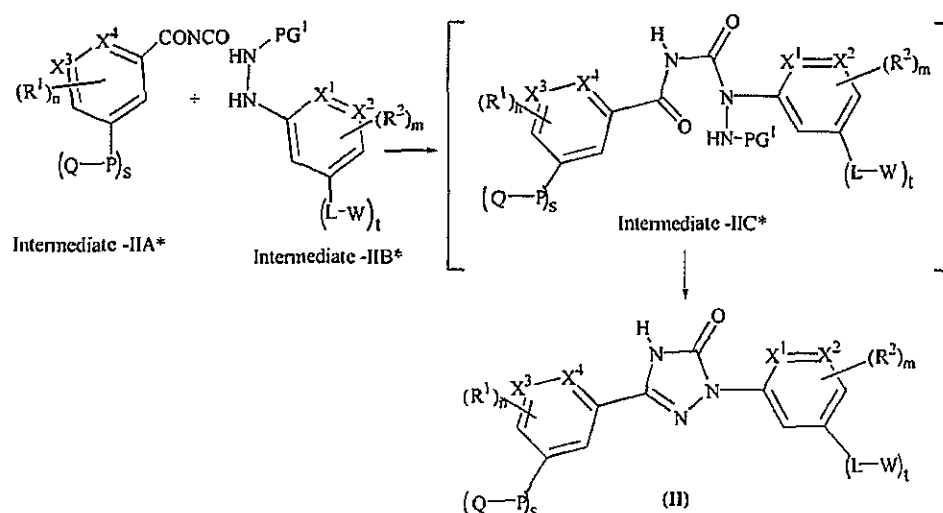
15. The pharmaceutical composition according to claim 14, wherein the pharmaceutically acceptable excipient is a carrier or diluent.

16. A process for preparing a compound of formula (II)



(II)

or a pharmaceutically acceptable salt thereof, which comprises: reacting a compound of formula (Intermediate-IIA*) with a compound of formula (Intermediate-IIB*) to form a compound of formula (Intermediate-IIC*); and converting the compound of formula (Intermediate-IIC*) to a compound of formula (II):



wherein,

X^1 , X^2 , X^3 and X^4 are each independently selected from CH and N;

each occurrence of L is independently selected from $-CH_2NHC(O)-$ and $-CH_2NHS(O)_2-$;

each occurrence of P is independently selected from $-CH_2NHC(O)-$ and $-CH_2NHS(O)_2-$;

each occurrence of Q is independently selected from isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-methoxy-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl, cyclopropyl, cyclobutyl, cyclopentyl, 2-fluorophenyl, tetrahydrofuranyl, tetrahydrofuran-2-yl, (S)-tetrahydrofuran-2-yl, (R)-tetrahydrofuran-2-yl, isoxazolyl and 1-methyl-1*H*-imidazole-2-yl;

each occurrence of W is independently selected from isopropyl, *tert*-butyl, trifluoromethyl, 1-fluoro-2-methylpropan-2-yl, 1-hydroxy-2-methylpropan-2-yl and cyclopropyl;

each occurrence of R^1 is independently selected from cyano, Cl, F, CHF_2 , CF_3 , OCH_3 , CH_3 , (2,2,2-trifluoroethoxy)methyl, cyclopropyl, (pyrrolidin-1-yl)methyl, 5-isopropyl-1,3,4-oxadiazol-2-yl, (3,5-dimethyl-1*H*-pyrazol-1-yl)methyl, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, (2,5-dichlorophenyl)ethynyl, (4-chloro-2-fluorophenyl)ethynyl, (3-chloro-2-fluorophenyl)ethynyl, (3-(trifluoromethyl)phenyl)ethynyl and (2-chloro-4-(trifluoromethyl)phenyl)ethynyl;

each occurrence of R^2 is independently selected from cyano, Cl, F, CH_3 , CF_3 , OCH_3 , 3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl, cyclopropanecarboxamido, 3,3-dimethylbut-1-ynyl, 2-cyclopropylethynyl, $-CONH-[3-(trifluoromethyl)phenyl]$, $-CONH-[3-(difluoromethyl)-4-fluorophenyl]$ and $-CONH-[3-(difluoromethyl)phenyl]$;

PG^1 represents an amine protecting group;

'm' is an integer ranging from 0 to 3, both inclusive;

'n' is an integer ranging from 0 to 3, both inclusive;

's' is an integer ranging from 0 to 1, both inclusive; and

't' is an integer ranging from 0 to 1, both inclusive;

with the provisos that (i) 's' and 't' are not '0' simultaneously; and (ii) that 'm' and 'n' are not '0' simultaneously.

17. The process according to claim 16, wherein the isocyanate of formula (Intermediate-IIA*) is obtained from the corresponding acid, amide or acid halide derivative.

18. The process according to claim 16, wherein a compound of formula (Intermediate-IIA*) is reacted with a compound of formula (Intermediate-IIB*) in DCM, toluene or EDC.

19. The process according to claim 16, wherein Intermediate-IIC* is not isolated.
20. The process according to claim 16, wherein PG¹ is *tert*-butoxycarbonyl.
21. The process according to claim 16, wherein a compound of formula (Intermediate-IIC*) is converted to compound of formula (II) using an organic acid.