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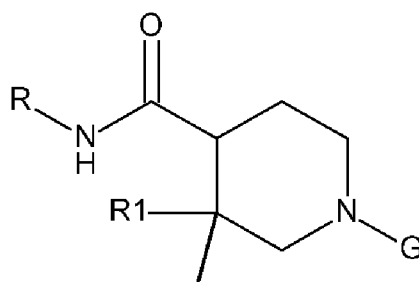
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(54) Title: NOVEL METHYL-PIPERIDINE COMPOUNDS USEFUL FOR INHIBITING MICROSOMAL PROSTAGLANDIN E₂ SYNTHASE-1

(1)

(57) Abstract: The present invention provides compounds of Formula (1), or a pharmaceutically acceptable salt thereof, formula (1), where R, R₁, and G are as described herein; methods of preparing the compounds; and use of the compounds to treat pain and/or inflammation associated with arthritis or osteoarthritis.

Novel Methyl-Piperidine Compounds Useful for Inhibiting Microsomal Prostaglandin E₂ Synthase-1

The present invention relates to novel compounds; to pharmaceutical compositions
5 comprising the compounds; methods of using the compounds to treat pain and/or inflammation
associated with arthritis; and intermediates and processes useful in the synthesis of the
compounds.

Arthritis involves inflammation of the joints and is often accompanied by pain and
stiffness. Osteoarthritis, the most common form of arthritis, is a complex degenerative disease of
10 the joints characterized by progressive destruction of articular cartilage; peri-articular structures
including bones, synovium, and associated fibrous joint tissues; and varying degrees of
inflammation. Existing drug therapies using non-steroidal, anti-inflammatory drugs (NSAIDs)
and cyclooxygenase-2 inhibitors (COX-2 inhibitors) can reduce pain associated with
osteoarthritis, but may be only moderately effective over time and each has variable risk/benefit
15 considerations.

The NSAIDs and COX-2 inhibitors reduce inflammation and pain through inhibition of
the COX-2 enzymes. In response to pro-inflammatory stimuli, the COX-2 enzymes metabolize
arachidonic acid to prostaglandin H₂ (PGH₂). PGH₂ is further metabolized by a variety of
enzymes to other eicosanoids including prostaglandin E₂ (PGE₂), prostaglandin I₂ (PGI₂),
20 prostaglandin F_{2α} (PGF_{2α}), prostaglandin D₂ (PGD₂), and thromboxane A₂ (TXA₂). These
metabolites are known to induce physiological and pathophysiological effects. It is thought that
a drug-mediated imbalance of PGI₂ and TXA₂ may explain why NSAIDs and COX-2 inhibitors
produce deleterious gastrointestinal and cardiovascular side-effects. Consequently, these classes
of drugs may be contraindicated for many patients due to pre-existing or emergent
25 cardiovascular and/or gastrointestinal conditions. Additionally, patients can become refractory
over time to specific drug treatments.

Of the arachidonic acid metabolites, PGE₂ has been identified as an important mediator of
conditions associated with osteoarthritis; for example, fever, pain, and inflammation.
Prostaglandin E₂ is specifically produced through the metabolism of PGH₂ by microsomal

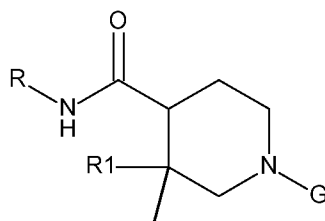
prostaglandin E₂ synthase-1 (mPGES-1). It is thought that selectively inhibiting mPGES-1 may provide a new treatment option for patients suffering from arthritis.

Publication WO 2013/146970 discloses tri-substituted quinoline compounds and suggests that the disclosed compounds may be useful for treating inflammatory diseases *inter alia*.

5 However, that publication does not disclose compounds as claimed in this application.

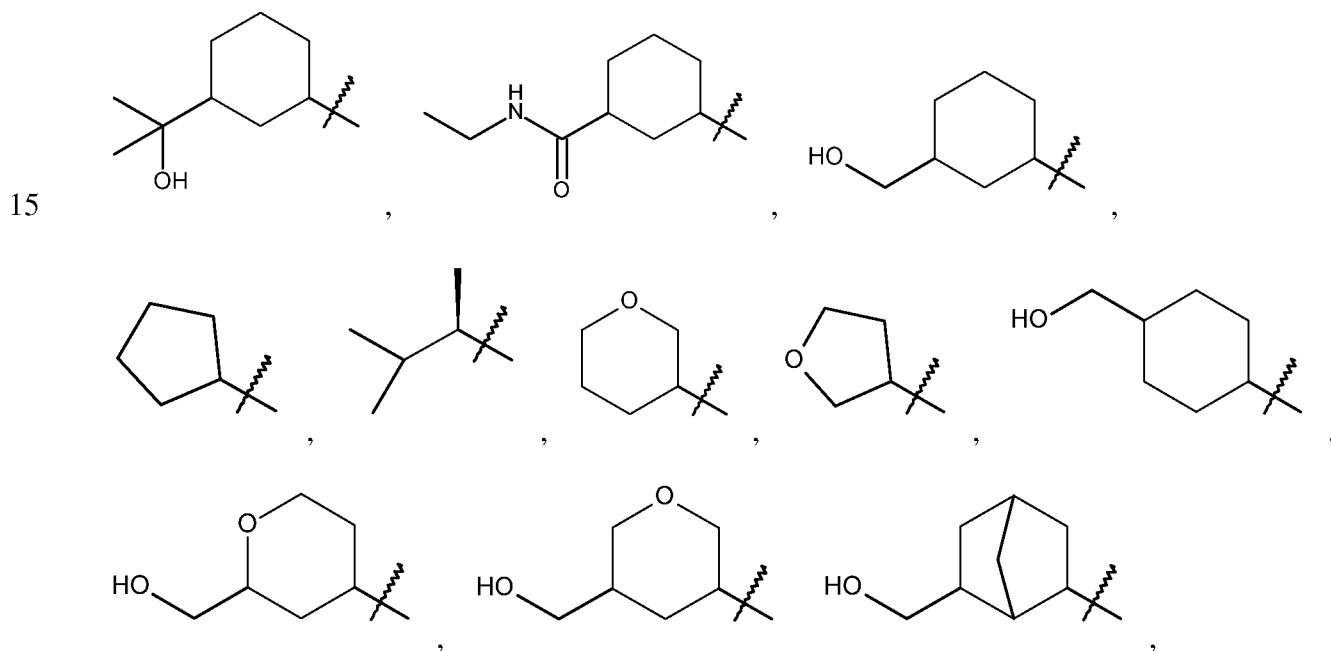
There remains a need for additional options to treat the inflammation and alleviate the pain associate with arthritis. The present invention provides novel compounds that inhibit mPGES-1 and that may be beneficial for treating patients suffering from arthritis and osteoarthritis.

10 The present invention provides compounds according to Formula 1, or pharmaceutically acceptable salts thereof,

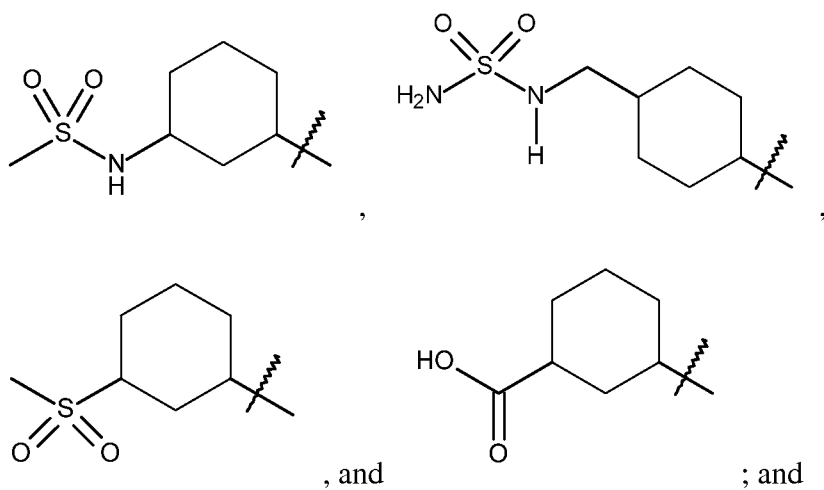


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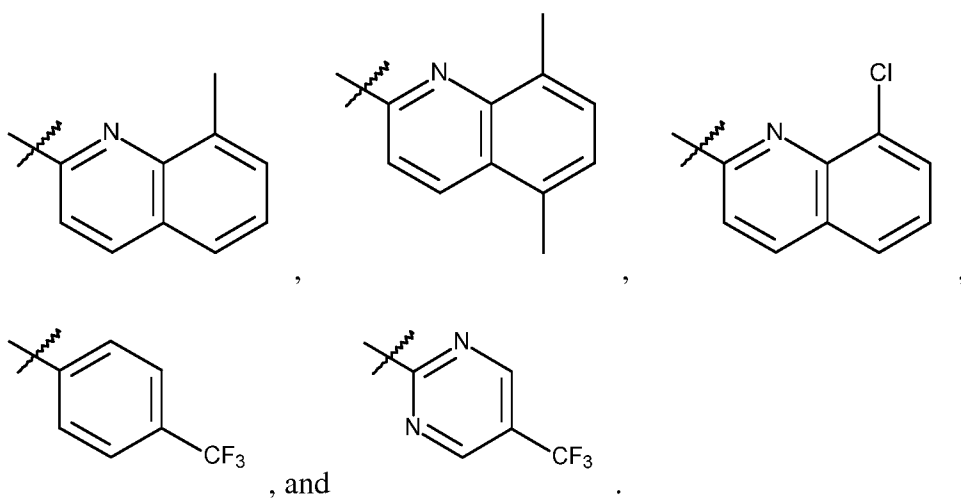
where R1 is H or -CH₃; R is selected from:



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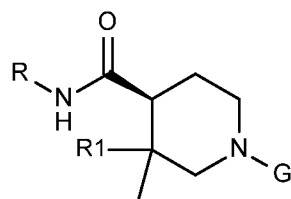


G is selected from:



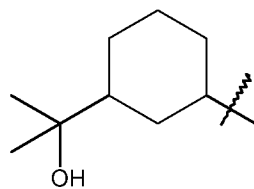
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The present invention provides compounds according to Formula 2, or pharmaceutically acceptable salts thereof:

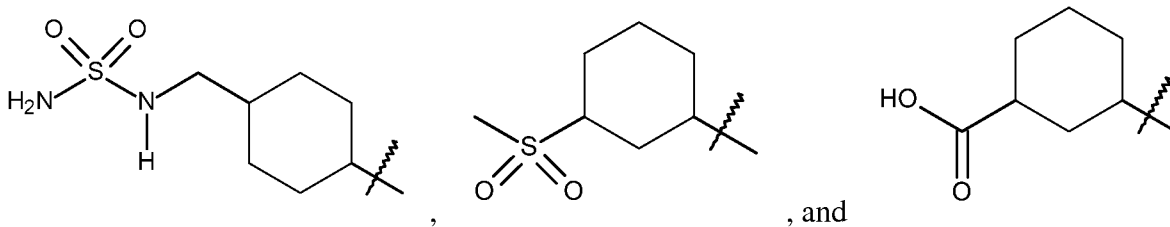
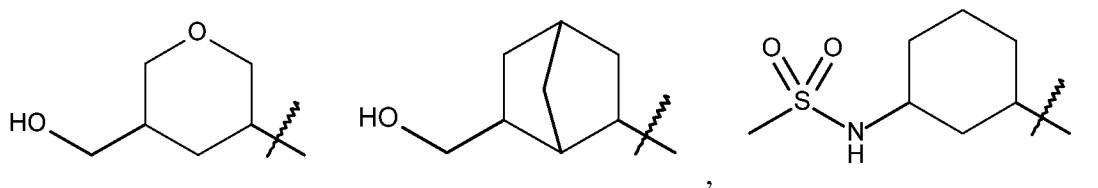
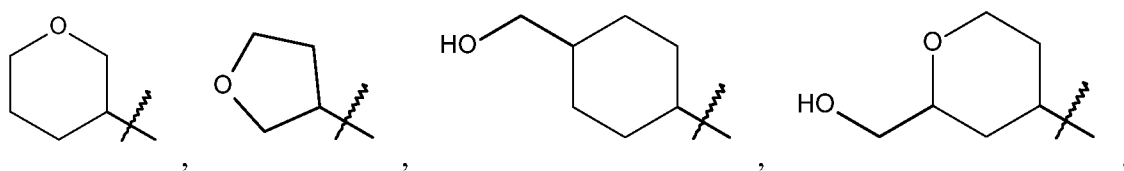
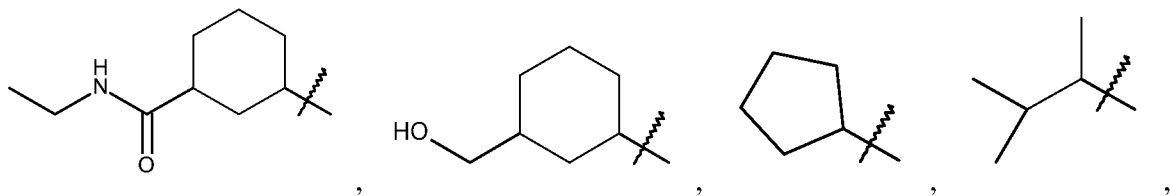


2,

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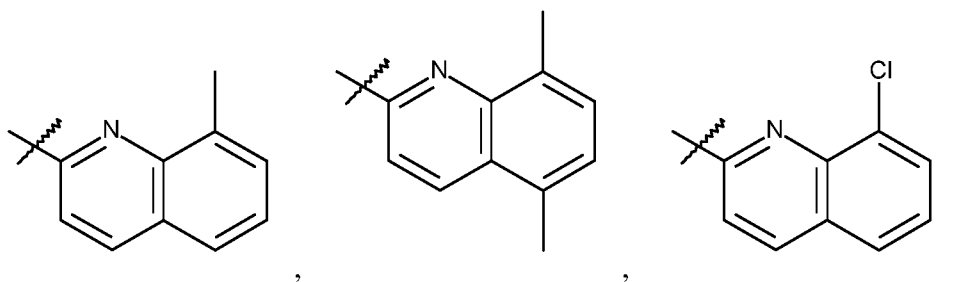
where: R₁ is H or -CH₃; R is selected from:

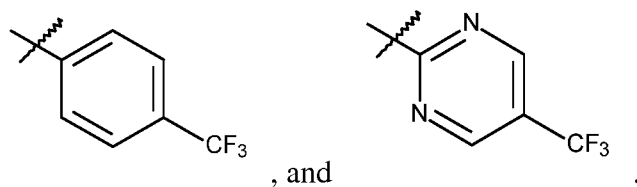


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and

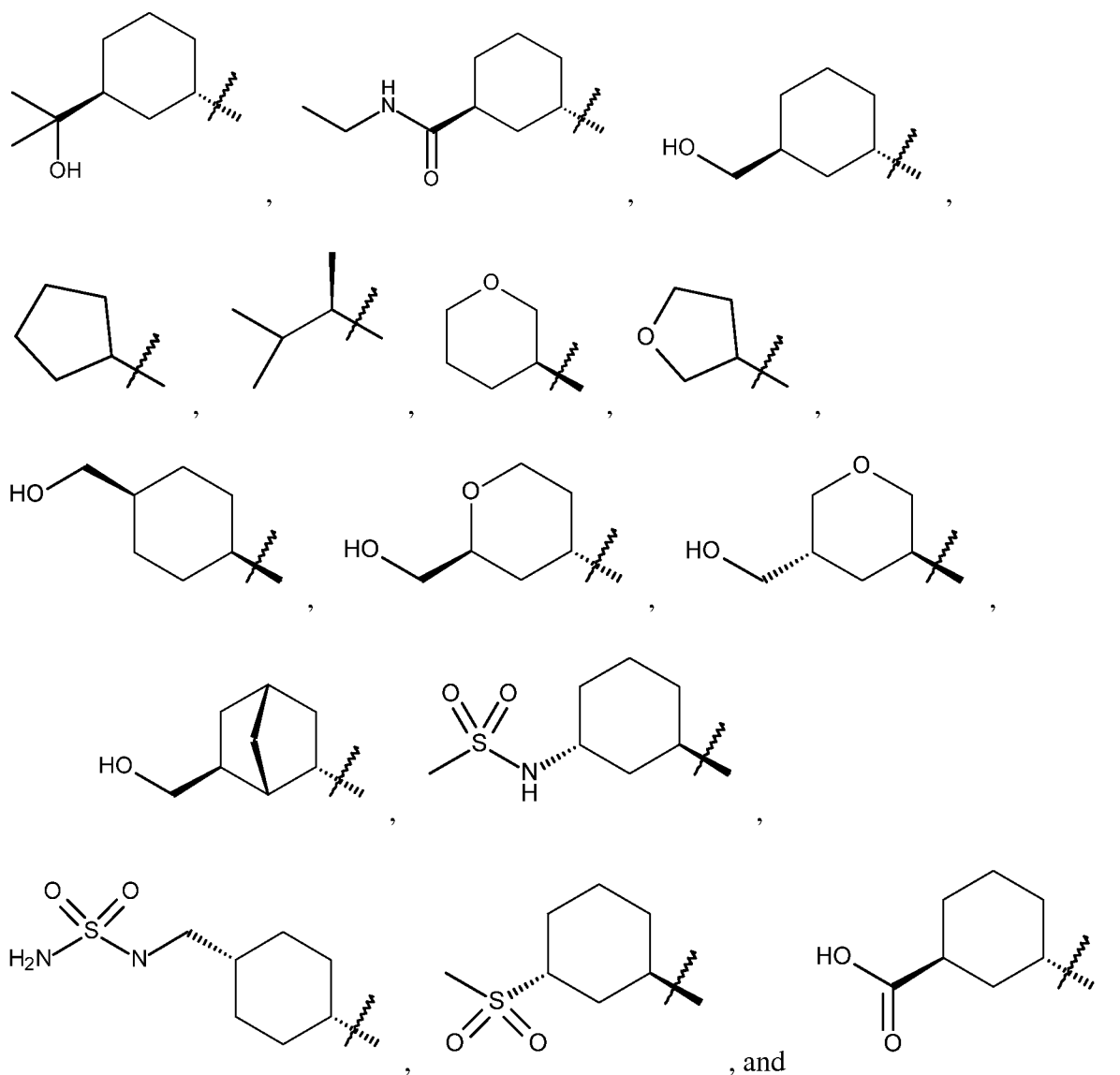
G is selected from:



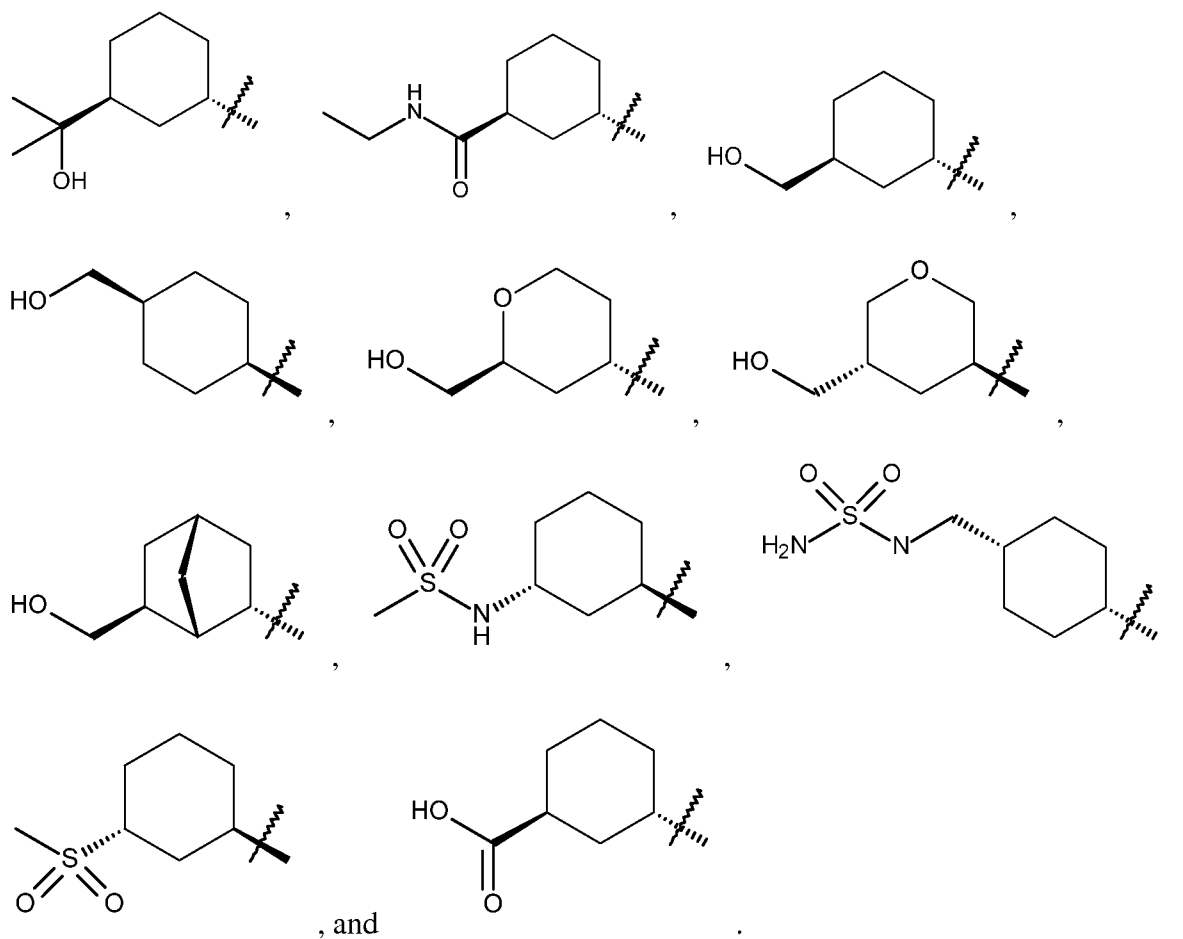


The present invention also provides compounds according to Formulae 1 or 2, or pharmaceutically acceptable salts thereof, where R1 is -CH₃.

5 For preferred compounds of Formulae 1 or 2, or pharmaceutically acceptable salts thereof, R is selected from:

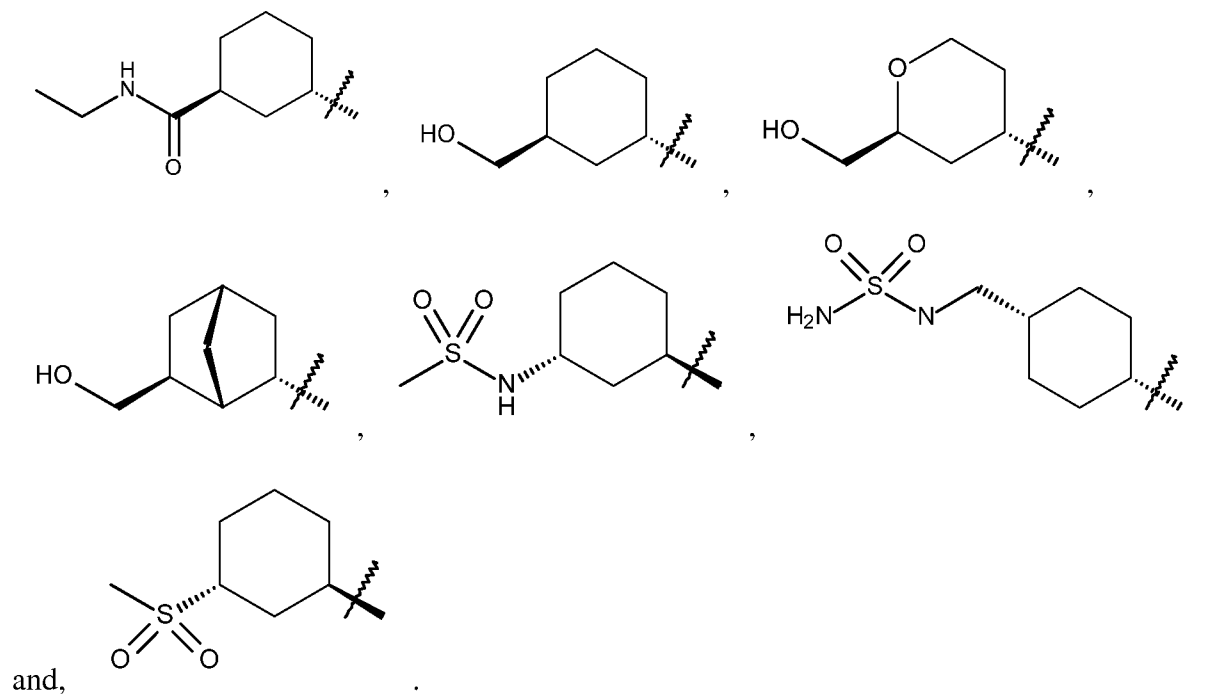


For more preferred compounds of Formulae 1 or 2, or pharmaceutically acceptable salts thereof, R is selected from:

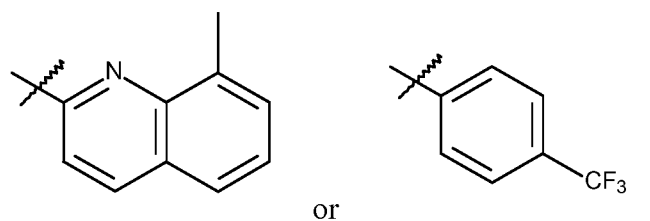


For still more preferred compounds of Formulae 1 or 2, or pharmaceutically acceptable salts thereof, R is selected from:

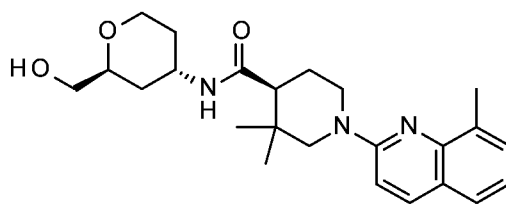
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5 For preferred compounds of Formulae 1 or 2, or pharmaceutical salts thereof, G is:



The present invention also provides a compound according to Formula 3, or a pharmaceutically acceptable salt thereof.



3

In one embodiment, compounds of Formulae 1, 2, or 3 are provided as a neutral species. In another embodiment, the compounds are provided as pharmaceutically acceptable salts. In one preferred salt form, the compound of Formula 3 is provided as hydrochloride salt.

The present invention also provides a pharmaceutically acceptable composition that
5 includes a compound according to Formulae 1, 2, or 3, or a pharmaceutically acceptable salt thereof, and at least one of a pharmaceutically acceptable carrier, diluent, or excipient.

The present invention also provides a method of treating a patient in need of treatment for pain associated with arthritis or osteoarthritis. The method comprises administering to the patient an effective amount of a pharmaceutically acceptable composition that includes a
10 compound according to Formulae 1, 2, or 3, or a pharmaceutically acceptable salt thereof, and at least one of a pharmaceutically acceptable carrier, diluent, or excipient.

The present invention also provides a method of treating a patient in need of treatment for arthritis or osteoarthritis. In one form, the present invention provides a method of treating a patient for the signs and/or symptoms of osteoarthritis. The method comprises administering to
15 the patient an effective amount of a compound according to Formulae 1, 2, or 3, or a pharmaceutically acceptable salt thereof.

The present invention also provides a method of treating a patient in need of treatment for inflammation associated with arthritis or osteoarthritis. The method comprises administering to the patient an effective amount of a pharmaceutically acceptable composition that includes a
20 compound according to Formulae 1, 2, or 3, or a pharmaceutically acceptable salt thereof, and at least one of a pharmaceutically acceptable carrier, diluent, or excipient.

The present invention also provides a method of treating a patient in need of treatment for pain associated with arthritis. The method comprises administering to the patient an effective amount of a compound according to Formulae 1, 2, or 3, or a pharmaceutically acceptable salt
25 thereof. The present invention further provides a method of treating a patient for pain associated with osteoarthritis. The method comprising administering to the patient an effective amount of a compound according to Formulae 1, 2, or 3, or a pharmaceutically acceptable salt thereof.

The present invention provides a compound according to Formulae 1, 2, or 3 for use as a
30 medicament. The medicament or compound can be used for therapy. The therapy can include

the treatment of the patient for arthritis or osteoarthritis. In one embodiment the therapy can be the treatment of pain or inflammation associated with arthritis or osteoarthritis.

The present invention also provides the use of a compound for the manufacture of a medicament to treat arthritis or osteoarthritis. In one form, the medicament is used to treat pain
5 or inflammation associated with arthritis or osteoarthritis.

As used herein, the terms “treating” or “to treat” includes stopping or reducing the severity of an existing symptom or disorder, in particular the pain and/inflammation, associated with arthritis or osteoarthritis.

As used herein, the term "patient" refers to a mammals, such as a guinea pigs, rats, dogs,
10 cats, cows, horses, sheep, goats or humans; or fowl such as a chickens or ducks. The preferred patient is a mammal, more preferably a human.

The exemplified compounds of the present invention can be formulated into pharmaceutical compositions in accordance within accepted practices. Examples of pharmaceutically acceptable carriers, excipients, and diluents can be found in Remington's
15 Pharmaceutical Sciences, Gennaro, Ed., Mack Publishing Co. Easton Pa. 1990. Non-limiting examples include the following: starch, sugars, mannitol, and silica derivatives; binding agents such as carboxymethyl cellulose and other cellulose derivatives, glycerol monostearate; adsorptive carriers such as kaolin and bentonite; and lubricants such as talc, calcium, and magnesium stearate, and solid polyethyl glycols. In one form, the pharmaceutical formulation
20 includes 20% Captisol in 25 mM phosphate buffer pH 2.

Preferred pharmaceutical compositions can be formulated as a tablet or capsule for oral administration or as an injectable solution. The tablet, capsule, or solution will include a compound of the present invention in an amount effective for treating a patient in need of treatment.

25 As used herein, the term “effective amount” refers to the amount or dose of a compound of the invention, or a pharmaceutically acceptable salt thereof, which, upon a single or multiple dose administration to the patient, provides the desired effect, such as, the reduction or elimination of pain and/or inflammation the patient under diagnosis or treatment.

An effective amount can be readily determined by the attending diagnostician by using
30 known techniques and by observing results obtained under analogous circumstances. In

determining the effective amount for a patient, a number of factors can be considered by the attending diagnostician, including, but not limited to: the species of mammal, its size, age, and general health; the severity of the symptoms; the response of the individual patient; the mode of administration; the bioavailability characteristics of the compound of Formula 1, 2, or 3 or its pharmaceutically acceptable salt form, as a formulated drug product in the dose regimen selected; the use of concomitant medication; and other relevant circumstances.

In one embodiment, the effective amount can be from about 0.0005 mg/kg of body weight to about 100 mg/kg. More preferably, the effective amount can be from about 0.001 mg/kg to about 50 mg/kg. Still more preferably, the effective amount can be from about 0.001 mg/kg to about 20 mg/kg.

A compound of the present invention can be combined with other treatment methods and/or additional therapeutic agents. Preferably a compound of the present invention can be combined with other agents that also are effective for the treatment of arthritis or osteoarthritis. Examples of these additional therapeutic agents include: NSAIDs or COX-2 inhibitors such as ibuprofen, aspirin, acetaminophen, celecoxib, naproxen, and ketoprofen; opioids such as oxycodone, and fentanyl; and corticosteroids such as hydrocortisone, prednisolone, and prednisone.

The compound of the invention and the additional therapeutic agent(s) can be administered either together through the same delivery route and device such as a single pill, capsule tablet, or solution; or separately administered either at the same time in separate delivery devices or administered sequentially.

The compound of the present invention can be provided as a pharmaceutically acceptable salt. "Pharmaceutically-acceptable salt" refers to salts of the compound of the invention considered to be acceptable for clinical and/or veterinary use. Pharmaceutically acceptable salts and common methodology for preparing them are well known in the art. See, *e.g.*, P. Stahl, *et al.*, Handbook of Pharmaceutical Salts: Properties, Selection and Use, (VCHA/Wiley-VCH, 2002); S.M. Berge, *et al.*, "Pharmaceutical Salts," *Journal of Pharmaceutical Sciences*, Vol. 66, No. 1, January 1977.

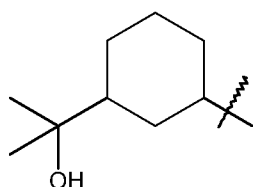
The compound of the present invention, or a salt thereof, may be prepared by a variety of procedures known in the art, some of which are illustrated in the Schemes, Preparations, and

Examples below. One of ordinary skill in the art recognizes that the specific synthetic steps for each of the routes described may be combined in different ways, or in conjunction with steps from different schemes, to prepare compounds of the invention, or salts thereof. The products of each step in the schemes below can be recovered or purified by conventional methods, including
5 extraction, evaporation, precipitation, chromatography, supercritical fluid chromatography, filtration, trituration, and crystallization.

Individual isomers, enantiomers, or diastereomers may be separated or resolved by one of ordinary skill in the art at any convenient point in the synthesis of compounds of Formula 1 by methods such as selective crystallization techniques or chiral chromatography (See for example,
10 J. Jacques, et al., "*Enantiomers, Racemates, and Resolutions*", John Wiley and Sons, Inc., 1981, and E.L. Eliel and S.H. Wilen, "*Stereochemistry of Organic Compounds*", Wiley-Interscience, 1994). Additionally, the intermediates described in the following preparations contain nitrogen and oxygen protecting groups. The protection and deprotection conditions are well known to the skilled artisan and are described in the literature (See for example "*Greene's Protective Groups in Organic Synthesis*", Fourth Edition, by Peter G.M. Wuts and Theodora W. Greene, John
15 Wiley and Sons, Inc. 2007).

The reagents and starting materials are generally readily available to one of ordinary skill in the art. Others may be made by standard techniques of organic and heterocyclic chemistry and the procedures described in the Preparations and the Examples below.

20 The depiction of a bond with a line through it as illustrated below indicates the point of attachment of the substituent to the rest of the molecule.



The abbreviations used herein are defined according to *Aldrichimica Acta*, Vol. 17, No. 1, 1984. Other abbreviations are defined as follows: "δ" refers to part per million down-field from
25 tetramethylsilane; "ATCC" refers to American type culture collection; "BOP" refers to (benzotriazol-1-yloxy)tris(dimethylamino)phosphonium hexafluorophosphate; "BSA" refers to Bovine Serum Albumin; "CDI" refers 1,1'-carbonyldiimidazole; "DCC" refers to 1,3-

dicyclohexylcarbodiimide; “DCM” refers to dichloromethane; “DIC” refers to 1,3-diisopropylcarbodiimide; “DMF” refers to dimethylformamide; “DMSO” refers to dimethylsulfoxide; “EDCI” refers to 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride; “EDTA” refers to ethylenediaminetetraacetic acid; “ee” refers to enantiomer excess; “EIA” refers to enzyme immunoassay; “EIMS” refers to electron ionized mass spectrometry; “ESMS” refers to electrospray mass spectrometry; “EtOAc” refers to ethyl acetate; “EtOH” refers to ethanol or ethyl alcohol; “Ex. No.” refers to Example Number; “HATU” refers to (1-[bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxid hexafluorophosphate); “HBTU” refers to (1H-benzotriazol-1-yloxy)(dimethylamino)-N,N-dimethylmethaniminium hexafluorophosphate; “HOAT” refers to 1-hydroxy-7-azobenzotriazole; “HOBt” refers to 1-hydroxylbenzotriazole hydrate; “IC₅₀” refers to the concentration of an agent that produces 50% of the maximal inhibitory response possible for that agent; “*i*-PrOH” refers to isopropanol or isopropyl alcohol; “LPS” refers to lipopolysaccharide; “MeOH” refers to methanol; “min” refers to minutes; NSAIDs” refers to nonsteroidal anti-inflammatory drugs; “PBS” refers to Phosphate Buffered Saline; “PGE₂” refers to prostaglandin E₂; “PGH₂” refers to prostaglandin H₂; “PGI₂” refers to prostaglandin I₂; “PyBOP” refers to (benzotriazol-1-yl-oxytripyrrolidinophosphonium hexafluorophosphate); “PyBrOP” refers to bromo(tri-pyrrolidinyl)phosphoniumhexafluorophosphate; “rhIL-1 β ” refers to recombinant human interleukin 1 β ; “SCF” refers to supercritical fluid; “SFC” refers to supercritical fluid chromatography; “TBME” refers to *t*-butyl ether methyl ether; “THF” refers to tetrahydrofuran; and “t_R” refers to retention time.

General LCMS Methods. All analyses are performed using an Agilent 1200 Infinity Series Liquid Chromatography (LC) system, consisting of a 1260 HiP degasser (G4225A), 1260 Binary Pump (G1312B), 1290 auto-sampler (G4226A), 1290 thermo-stated column compartment (G1316C) and a 1260 Diode Array Detector (G4212B) coupled to an Agilent 6150 single quadrupole mass spectrometry (MS) detector. The MS is operated with an electro-spray ionization source (ESI) in both positive & negative ion mode. The nebulizer pressure is set to 50 psi, the drying gas temperature and flow to 350 °C and 12.0 L/min respectively. The capillary voltages used are 4000 V in positive mode and 3500 V in negative mode. Data acquisition is performed with Agilent Chemstation software.

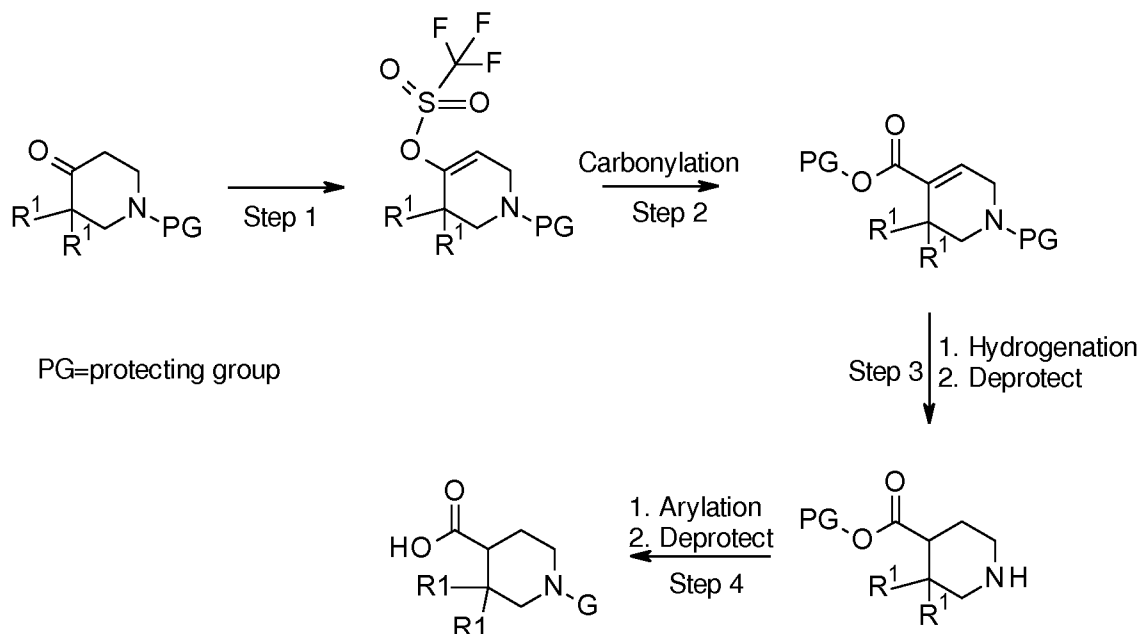
HPLC Method 1. Analyses are carried out on a Daicel ChiralPak AD-3R column (100 mm length, 4.6 mm internal diameter, 3 μ m particle size). The mobile phase used is: A2=Water with 10 mM ammonium bicarbonate, adjusted to pH 9 with ammonium hydroxide; and B2 = acetonitrile. The run is performed at a temperature of 25 °C and a flow rate of 1.5 mL/min with a gradient elution from 50 % to 95 % (B2) over 3.0 min followed by a 3.0 min hold at 95 % (B2). The UV (DAD) acquisition is performed at 40 Hz, with a scan range of 190–400 nm (by 5 nm step). A 1:1 flow split is used before the MS detector. The MS acquisition range is set to 100–800 m/z with a step size of 0.2 m/z in both polarity modes. Fragmentor voltage is set to 70 (ESI⁺) or 120 (ESI⁻), Gain to 0.40 (ESI⁺) or 1.00 (ESI⁻) and the ion count threshold to 4000 (ESI⁺) or 1000 (ESI⁻). The overall MS scan cycle time is 0.15 s/cycle.

SFC Method 1. Analyses are carried out on a Daicel ChiralPak OJ-H column (100 mm length, 4.6 mm internal diameter, 5 μ m particle size). The mobile phase is: 8 % (20 mM NH₃ in *i*-PrOH) and 92 % CO_{2(scf)} at a pressure of 100 bar. The run is performed at a temperature of 35 °C and a flow rate of 3 mL/minute. The UV (DAD) acquisition is performed at a wavelength of 220 nm.

SFC Method 2. Analyses are carried out on a Daicel ChiralPak AS-H column (100 mm length, 4.6 mm internal diameter, 5 μ m particle size). The mobile phase is: 20 % (20 mM NH₃ in *i*-PrOH) and 80 % CO_{2(scf)} at a pressure of 100 bar. The run is performed at a temperature of 35 °C and a flow rate of 5 mL/minute. The UV (DAD) acquisition is performed at a wavelength of 220 nm.

The following Schemes further illustrate the invention.

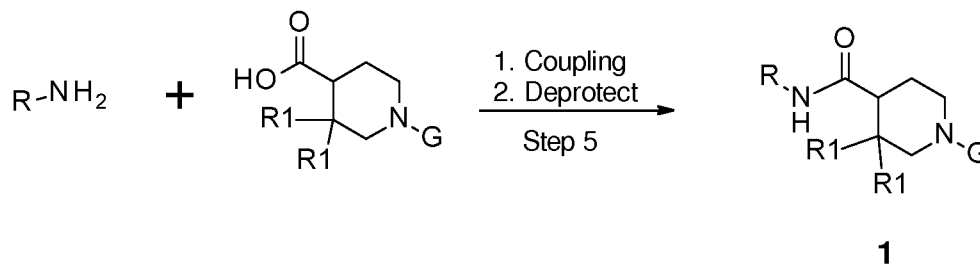
Scheme 1



In Scheme 1, Step 1, a trifluoromethyl sulfonyl group is installed on the piperidine to act
 5 as a leaving group in the subsequent reaction of Step 2. “PG” is a protecting group developed
 for the amino group, such as carbamates and amides. The product of Step 1 is then subjected to
 carbonylation using a palladium catalyst to give the ester product of Step 2. The double bond in
 the tetrahydropyridine can be reduced under hydrogenation conditions. In Step 3, substep 2, the
 piperidine amine can be deprotected under acidic conditions. The piperidine product of Step 3
 10 can then be reacted with a halogen substituted G group under conditions suitable for nucleophilic
 aromatic substitution give the product of Step 4. For example, an inorganic base such as K_2CO_3
 or an organic base such as N,N-diisopropylethylamine, pyridine, or triethylamine can be used to
 give the product of Step 4, substep 1. Alternatively, the piperidine product of step 3 can be
 reacted with a quinolone N-oxide using an activating agent, such as PyProp, to give the product
 15 of Step 4. Deprotection of the piperidine carboxy group can be accomplished under standard
 conditions with an inorganic base such as aqueous sodium hydroxide or lithium hydroxide to
 give the product of Step 4, substep 2 or under acidic conditions using 4 M HCl or aqueous
 sulfuric acid.

15

Scheme 2



In Scheme 2, the piperidine carboxylic acid product of Step 4 can be coupled with an appropriate amine, R-NH₂, under amidation conditions using an organic base such as triethylamine or diisopropylethylamine, a coupling reagent such as EDCI, and a coupling additive such as HOBT to give the compounds of Step 5, substep 1. One skilled in the art will recognize that there are a number of methods and reagents for amide formation resulting from the reaction of carboxylic acids and amines. For example, the reaction of the amine compound with an appropriate carboxylic acid in the presence of a coupling reagent with or without an organic base can provide a compound of Formula 1. Coupling reagents include carbodiimides, such as DCC, DIC, EDCI or a carbonyldiimidazole such as CDI. Amide coupling additives, such as HOBT and HOAt can also be used to enhance the reaction. Additionally, uronium or phosphonium salts of non-nucleophilic anions, such as HBTU, HATU, BOP, PyBOP, and PyBrOP could be used in place of the more traditional coupling reagents. An additive such as DMAP may be used to enhance the reaction. The product of Step 5, substep 1 can then be deprotected, if necessary, under standard conditions with an inorganic base such as aqueous sodium hydroxide or lithium hydroxide in a solvent such as MeOH and THF to give compounds of Formula 1.

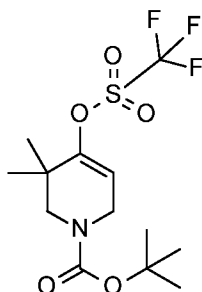
A pharmaceutically acceptable salt of the compounds of the invention, such as a hydrochloride salt, can be formed, for example by reaction of an appropriate free base of Formula 1 and an appropriate pharmaceutically acceptable acid, such as hydrochloric acid, in a suitable solvent such as diethyl ether under standard conditions. Additionally, the formation of such salts can occur simultaneously upon deprotection of a nitrogen protecting group. The formation of such salts is well known and appreciated in the art. See, for example, Gould, P.L., "Salt selection for basic drugs," *International Journal of Pharmaceutics*, **33**: 201-217 (1986); Bastin, R.J., *et al.* "Salt Selection and Optimization Procedures for Pharmaceutical New

Chemical Entities,” *Organic Process Research and Development*, 4: 427-435 (2000); and Berge, S.M., *et al.*, “Pharmaceutical Salts,” *Journal of Pharmaceutical Sciences*, 66: 1-19, (1977).

The following Preparations and Examples further illustrate the invention.

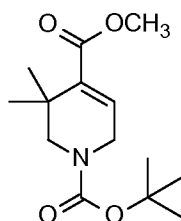
Preparation 1

tert-Butyl 3,3-dimethyl-4-(((trifluoromethyl)sulfonyl)oxy)-3,6-dihydropyridine-1(2H)-carboxylate



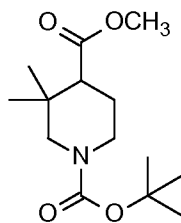
Under a nitrogen atmosphere, cool a solution of diisopropylamine (260 mL, 1.85 mol) in THF (1.0 L) to -20°C then add a solution of *n*-butyllithium (2.50 M in hexanes, 650 mL, 1.60 mol) drop-wise over 30 minutes. Allow the mixture to warm to -10°C and stir for 1 hour. Cool the mixture to -74°C and add a solution of *tert*-butyl 3,3-dimethyl-4-oxopiperidine-1-carboxylate (260 g, 1.14 mol) in THF (1.0 L mL) drop-wise over 60 minutes. Stir the mixture at -74°C for 2 hours, and then add a solution of *N*-phenylbis(trifluoromethanesulfonylimide) (430 g, 1.20 mol) in THF (1.0 L). Warm the mixture to 0°C and stir for 2 hours. Allow the mixture to warm to room temperature and stir overnight. Quench the reaction with saturated aqueous NH_4Cl (1.0 L); dilute with water (2.0 L); separate the layers; and extract the aqueous layer with EtOAc (2×2 L). Combine the organic extracts; dry over Na_2SO_4 ; filter; collect the filtrate; and concentrate the filtrate under reduced pressure. Subject the resulting crude material to silica gel flash chromatography, eluting with a gradient of 0 % to 15 % TBME in hexanes, to provide the title compound as an orange oil in about 75% purity by mass as estimated by ^1H NMR (430 g, 78%). ^1H NMR (400 MHz, $\text{DMSO}-d_6$) δ 1.05 (s, 6H), 1.40 (s, 9H), 3.36 (s, 2H), 4.02 (d, $J = 3.4$ Hz, 2H), 5.82 (br s, 1H).

Preparation 2

O1-tert-Butyl-O4-methyl 3,3-dimethyl-2,6-dihydropyridine-1,4-dicarboxylate

Combine palladium(II) acetate (4.40 g, 20.0 mmol), 1,1'-bis(diphenylphosphino)ferrocene (13.3 g, 22.8 mmol), *tert*-butyl 3,3-dimethyl-4-(((trifluoromethyl)sulfonyl)oxy)-3,6-dihydropyridine-1(2H)-carboxylate (72.0 g, 150 mmol), anhydrous acetonitrile (850 mL), anhydrous MeOH (570 mL), and triethylamine (36.0 mL, 245 mmol) in a 2-L PARR™ autoclave fitted with a mechanical stirrer. Seal the autoclave. Purge and then pressurize the autoclave with carbon monoxide to 689 kPa. Heat the mixture to 65 °C for 2.25 hours; cool the mixture to room temperature; and carefully vent the autoclave. (*Caution! Poison gas!*). Concentrate under reduced pressure to give crude material. Combine this material with five other batches of material prepared by an analogous procedure on similar scales. Subject the combined material to silica gel flash chromatography, eluting with a gradient of 0 % to 20 % TBME in hexanes, to give the title compound as a yellow oil in about 83% purity by mass (260 g, 88%). MS (*m/z*): 214 (*M* – *t*-Bu + 2H)⁺.

Preparation 3

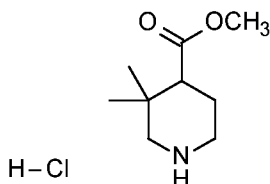
(±)-O1-tert-Butyl-O4-methyl 3,3-dimethylpiperidine-1,4-dicarboxylate

Suspend palladium (10 wt% on carbon, 5.4 g, 5.1 mmol) in MeOH (700 mL) then add a solution of *O1-tert-butyl-O4-methyl 3,3-dimethyl-2,6-dihydropyridine-1,4-dicarboxylate* (130 g, 396 mmol) dissolved in MeOH (700 mL) in a 2.25 L PARR™ reactor. Seal the reactor and purge it first with nitrogen gas then with hydrogen gas. Pressurize the reactor to 414 kPa with hydrogen and stir the mixture at room temperature for 1.5 hours. Release the pressure and filter

the mixture to remove the catalyst. Combine the filtrate with that obtained from another essentially identical reaction and concentrate under reduced pressure to give the title compound as a yellow oil in about 85% purity by mass (240 g, 95%). MS (m/z): 216 ($M - t\text{-Bu} + 2H$)⁺.

Preparation 4

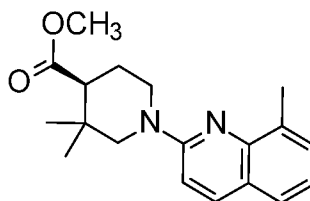
5 (±)-Methyl 3,3-dimethylpiperidine-4-carboxylate hydrochloride



Add HCl (4.0 M solution in 1,4-dioxane, 2.0 L, 8.0 mol) to a solution of *O1-tert-butyl-O4-methyl 3,3-dimethylpiperidine-1,4-dicarboxylate* (240 g, 752 mmol) in 1,4-dioxane (500 mL). Stir the resulting mixture at room temperature overnight and concentrate under reduced pressure. Dilute the residue with TBME (500 mL) and collect the resulting solids by filtration. Rinse the filter cake with TBME (2 × 400 mL) and dry the solid in a vacuum oven at 35 °C overnight to give the title compound as a white solid (144 g, 92%). MS (m/z): 172 ($M + H$)⁺.

Preparation 5

(-)-Methyl (4*S*)-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxylate

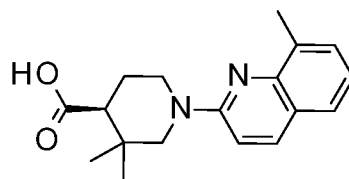


15 Add K₂CO₃ (210 g, 1.52 mol) to a mixture of methyl 3,3-dimethylpiperidine-4-carboxylate hydrochloride (144 g, 693 mmol) and 2-chloro-8-methylquinoline (125 g, 704 mmol) in DMSO (1.4 L). Stir the resulting mixture over night at 131 ± 1 °C. Cool the mixture to room temperature; filter to remove the solids; collect the filtrate; dilute the filtrate with water (2 L); then extract with EtOAc (2 × 3 L). Wash the combined organic extracts with water (3 × 1.5 L); dry over Na₂SO₄; filter; collect and concentrate the filtrate under reduced pressure. Subject the resulting crude material to silica gel flash chromatography, eluting with a gradient of 25 % to 30 % (10 % TBME in DCM) in hexanes, to provide the racemate of the title compound. Dissolve this material in MeOH (7.5 L) and filter. Subject the material to chiral SFC (Chiralpak

OJ-H, 50 mm × 250 mm × 5 μm) using 15 % (0.2 % dimethylethylamine in *i*-PrOH) in CO_{2(scf)} as the mobile phase at a flow rate of 400 g/min, by injecting 5 mL of solution every 95 seconds until all of the material has been subjected. For each injection, collect the first fraction to elute (t_R = 2.57 min by SFC Method 1). Combine the collected fractions with those from a previous reaction prepared similarly to provide 98 g of crude methyl 3,3-dimethylpiperidine-4-carboxylate hydrochloride. Concentrate mixture under reduced pressure and recrystallize the material from hot EtOH (1.38 L). Collect the crystals and dry the crystalline material in a vacuum oven at 40 °C overnight to give the title compound as a white crystalline solid (156 g, 43 % yield on a batch-proportional basis). MS (m/z): 313 ($M + H$)⁺, $[\alpha]^{20}_D -45^\circ$ (c 0.21, DCM). ee = >99 % as determined by SFC Method 1.

Preparation 6

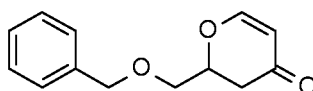
(-)-(4*S*)-3,3-Dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxylic acid



Treat methyl (4*S*)-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxylate (154 g, 493 mmol) with sulfuric acid (10% v/v in water, 2.31 L, 2.75 mol) and reflux the mixture overnight. Cool the mixture to room temperature and add NaOH (50 wt% in water) until the pH reaches 13. Add TBME (500 mL) to provide a triphasic mixture. Label the layers from top to bottom as: "Layer A," "Layer B," and "Layer C." Remove Layer C (the bottom layer). Add water (600 mL) to Layers A and B and separate the aqueous layer from the organic layers. Set aside the organic layers (A and B). Combine the aqueous layer with Layer C. Extract the combined aqueous layers with TBME (500 mL); separate the layers; and set aside the organic extracts. Add HCl (5.0 M) to the aqueous layer until the pH is 6.5. Extract the resulting aqueous mixture with TBME (2 × 400 mL). Combine all of the organic extracts (including Layers A and B); dry over MgSO₄; filter; collect and concentrate the filtrate under reduced pressure to give the title compound as a white solid in 96 % purity (145 g, 95%). MS (m/z): 299 ($M + H$)⁺. $[\alpha]^{20}_D -59.6^\circ$ (c 3.12, CH₃OH).

Preparation 7

(±)-2-((Benzyloxy)methyl)-2,3-dihydro-4H-pyran-4-one

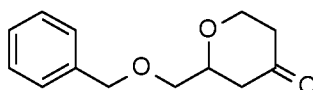


Add a solution of ZnCl_2 in THF (0.5 M, 1.21 L, 607 mmol) to a cold (0 °C) solution of
5 (E)-1-methoxy-3-(trimethylsilyl)oxy-1,3-butadiene (95% pure, 100 g, 551 mmol) and
(benzyloxy)acetaldehyde (97 % pure, 80 mL, 551.370 mmol) in toluene (551 mL) over 90
minutes while maintaining the internal temperature of the reaction mixture below 10 °C. Allow
the mixture to warm to room temperature and stir overnight. Divide the reaction mixture into
two equal portions; perform the following procedures on each portion. Add trifluoroacetic acid
10 (35 mL, 457 mmol) in four portions. After 20 minutes, concentrate the resulting mixture under
reduced pressure; dilute with EtOAc; add excess saturated NaHCO_3 ; filter to remove the solids;
collect the filtrate; separate and collect the organic layer. Wash the organic solution with
saturated aqueous NaCl ; isolate the organic extracts; dry over MgSO_4 ; filter; collect the filtrate;
and concentrate under reduced pressure. Combine the resulting material from each of the
15 previously separated portions. Subject the resulting material to silica gel flash chromatography,
eluting with a gradient of 20% to 50% EtOAc in hexanes, to give the title compound as an
orange oil in 92% purity (96 g, 73%). ^1H NMR (400 MHz, CDCl_3) δ 7.40–7.28 (m, 6H), 5.42 (d,
 $J = 6.0$ Hz, 1H), 4.65–4.56 (m, 3H), 3.74–3.67 (m, 2H), 2.75 (dd, $J = 16.8, 14.3$ Hz, 1H), 2.42
(dd, $J = 16.9, 3.2$ Hz, 1H).

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Preparation 8

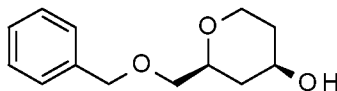
(±)-2-(Benzyloxymethyl)tetrahydropyran-4-one



Stir a mixture of EtOAc (880 mL), (±)-2-((benzyloxy)methyl)-2,3-dihydro-4H-pyran-4-
one (96 g, 0.440 mol), triethylamine (123 mL, 0.882 mol), and palladium (10% on carbon, 4.68
25 g, 4.40 mmol) under an atmosphere of hydrogen at room temperature for 73 hours. Filter the
mixture through diatomaceous earth; rinse the filter cake with EtOAc (250 mL); and sequentially
wash the filtrate with aqueous HCl (0.1 M), saturated aqueous NaHCO_3 , and saturated aqueous

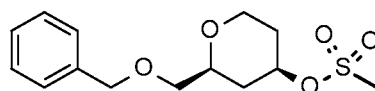
NaCl. Dry the organic layer over MgSO_4 ; filter; collect the filtrate; and concentrate under reduced pressure to give the title compound as a light yellow material (81.4 g, 84%). LC/MS (ESI^+): 221 $[\text{M}+\text{H}]^+$, 238 $[\text{M}+\text{NH}_4]^+$, 243 $[\text{M}+\text{Na}]^+$.

Preparation 9

5 (2*S*,4*R*)-2-(Benzyloxymethyl)tetrahydropyran-4-ol

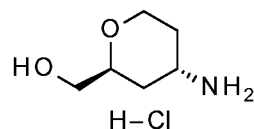
Cool a solution of ($\pm$)-2-(benzyloxymethyl)tetrahydropyran-4-one (40.6 g, 184 mmol) in THF (1.08 L) to -78°C . Add a solution of LiAlH_4 (1.0 M in THF, 240 mL, 240 mmol) drop-wise over 15 minutes. Allow the mixture to warm to 0°C after the addition is complete and
10 slowly add water (8.4 mL) drop-wise. After stirring the mixture for 5 minutes, add a solution of NaOH (15 mass % in water, 8.4 mL) and stir for an additional 5 minutes. Then add water (3×8.4 mL) and allow the mixture to warm to room temperature. After 30 minutes, filter the suspension to remove the solids and concentrate the filtrate under reduced pressure to provide the title compound as a colorless oil. Rinse the solids once with THF; filter; and concentrate the
15 filtrate under reduced pressure. Combine this material with those obtained from previously reactions performed essentially according to the same procedure starting with 40 g, 22 g, and 45 g of ($\pm$)-2-(benzyloxymethyl)tetrahydropyran-4-one. Dissolve the combined material in *i*-PrOH (637 mL). Subject the material to chiral SFC (Chiralpak AS-H, 50 mm $\times$ 150 mm $\times$ 5 μm) using 25% *i*-PrOH in $\text{CO}_{2(\text{scf})}$ as the mobile phase at a flow rate of 300 g/min, by injecting 1.35 g of
20 solution every 114 seconds until all of the material has been injected. For each injection, collect the first fraction to elute (major isomer). Combine all of the collected fractions to give the title compound in $>99\%$ ee as determined by SFC Method 2 (62.8 g, 42 % yield on a batch proportional basis). LC/MS (ESI): 223 $[\text{M}+\text{H}]^+$, 240 $[\text{M}+\text{NH}_4]^+$, 245 $[\text{M}+\text{Na}]^+$, 467 $[2\text{M}+\text{Na}]^+$.

Preparation 10

((2*S*,4*R*)-2-(Benzyloxymethyl)tetrahydropyran-4-yl)methanesulfonate

Drop-wise add a solution of methanesulfonyl chloride (11.7 mL, 150.6 mmol) in DCM
 5 (70 mL) to a mixture of (2*S*,4*R*)-2-(benzyloxymethyl)tetrahydropyran-4-ol (31 g, 139.5 mmol),
 DCM (1.40 L), and *N,N*-diisopropylethylamine (73.0 mL, 418.4 mmol) at 0 °C. Allow the
 mixture to warm to room temperature and stir it overnight. Thereafter add a solution of
 methanesulfonyl chloride (0.537 mL, 6.98 mmol) in DCM (5 mL) and stir the mixture for 5
 minutes at room temperature. Cool the mixture to 0 °C, pour it into water; separate; and collect
 10 the organic layer. Wash the organic layer with water (500 mL); combine the organic layer with
 organic layers/extracts obtained from other, essentially identical reactions. Dry the combined
 organic solutions over MgSO₄; filter; collect the filtrate; and concentrate the filtrate under
 reduced pressure to give the title compound as a light orange oil (84.7 g, 97 % yield on a batch
 proportional basis). ¹H NMR (400 MHz, *d*₆-DMSO) δ 7.36–7.24 (m, 5H), 4.86–4.77 (m, 1H),
 15 4.47 (s, 2H), 3.96 (dd, *J* = 7.6, 4.4 Hz, 1H), 3.60–3.54 (m, 1H), 3.48–3.36 (m, 3H), 3.18 (s, 3H)
 2.05 (d, *J* = 7.8 Hz, 1H), 1.98 (d, *J* = 7.8 Hz, 1H), 1.58 (app qd, *J* = 12.0, 4.8 Hz, 1H), 1.40 (app
 q, *J* = 11.8 Hz, 1H).

Preparation 11

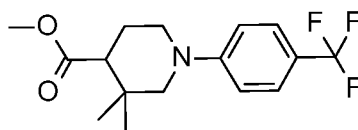
((2*S*,4*S*)-4-Aminotetrahydro-2H-pyran-2-yl)methanol hydrochloride

20 Add sodium azide (12.85 g, 191.7 mmol) to a solution of ((2*S*,4*R*)-2-
 (benzyloxymethyl)tetrahydropyran-4-yl)methanesulfonate (32.0 g, 106.5 mmol) in DMF (304
 mL). Heat the resulting mixture to 100 °C and stir it for 4 hours. Cool the reaction mixture to
 room temperature. Pour the mixture into water (400 mL) and then extract with EtOAc (2 × 400
 25 mL). Combine organic extracts and wash with saturated aqueous NaCl (3 × 200 mL). Dry the
 organic extracts over MgSO₄; filter; and collect the filtrate. Concentrate filtrate under reduced
 pressure to about 100 mL total volume. Dissolve resulting mixture in EtOAc (700 mL) and add

it to a PARR™ vessel containing PtO₂ (2.7 g, 12 mmol) and EtOAc (700 mL). Purge the vessel with nitrogen; then pressurize it with hydrogen to 414 kPa. Agitate the mixture for 4 hours at room temperature. Filter the mixture and concentrate the colorless filtrate under reduced pressure to provide 23.2 g of a colorless oil. Combine with 36.2 g of material from a previous reaction prepared by essentially the same procedure. Dissolve the combined material in EtOH (600 mL) and add HCl (37 wt% in H₂O, 50 mL). Add the resulting mixture to a PARR™ vessel which contains Pd (10% on carbon, 10.0 g, 9.40 mmol) and EtOH (600 mL). Purge the reactor with nitrogen and pressurize it with hydrogen to 414 kPa. Agitate the mixture at room temperature for 1 hour. Filter the mixture; collect the filtrate; and concentrate the colorless filtrate under reduced pressure to give the title compound as a dark, thick oil in about 72 % purity (54.2 g, 85 %). LC/MS (ESI): 132 [M+H]⁺.

Preparation 12

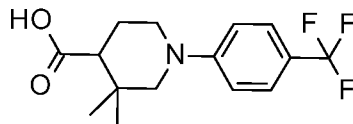
(±)-(Methyl 3,3-dimethyl-1-[4-(trifluoromethyl)phenyl]piperidine-4-carboxylate



Add K₂CO₃ (460 mg, 3.33 mmol) to a mixture of methyl 3,3-dimethylpiperidine-4-carboxylate hydrochloride (300 mg, 1.44 mmol), 1-fluoro-4-(trifluoromethyl)benzene (475 mg, 2.89 mmol) and DMSO (2 mL). Stir the resulting mixture at 130 °C for 2 days. Cool the mixture to room temperature and stir for 3 days. Subject the resulting material to reverse-phase flash chromatography on C₁₈ silica gel, eluting with a gradient of 10 % to 100 % acetonitrile (0.1 % formic acid) in water (0.1 % formic acid), to give the title compound as a yellow oil (113 mg, 25 %). MS (*m/z*): 316 (M + H)⁺.

Preparation 13

(±)-3,3-Dimethyl-1-[4-(trifluoromethyl)phenyl]piperidine-4-carboxylic acid.



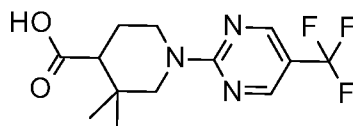
Add 2M aqueous NaOH (1 mL, 2 mmol) to a mixture of methyl (±)-3,3-dimethyl-1-[4-(trifluoromethyl)phenyl]piperidine-4-carboxylate (113 mg, 0.36 mmol), MeOH (1 mL), and THF

(5 mL). Stir the resulting mixture at 50 °C overnight. Cool the mixture to room temperature, and add HCl (33 wt % in water) until the pH of the mixture is 3. Concentrate the mixture under reduced pressure to give the title compound as a yellow solid in 88 % purity (123 mg, 99 % yield). MS (m/z): 302 ($M + H$)⁺.

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Preparation 14

(±)-3,3-Dimethyl-1-[5-(trifluoromethyl)pyrimidin-2-yl]piperidine-4-carboxylic acid

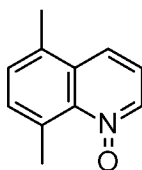


Add triethylamine (7.65 mL, 54.9 mmol) to a mixture of (±)-methyl 3,3-dimethylpiperidine-4-carboxylate hydrochloride (8.55 g, 41.2 mmol), 2-chloro-5-(trifluoromethyl)pyrimidine (5.0 g, 27.4 mmol), and acetonitrile (67 mL). Heat the mixture in a microwave to 180 °C for 1 hour; then cool the mixture to room temperature; and concentrate under reduced pressure. Add MeOH (40 mL), THF (80 mL), and 2 M aqueous NaOH (40 mL, 80 mmol). Stir the resulting mixture for 2 days at 50 °C. Add 2 M aqueous NaOH (45 mL, 90 mmol) and stir the resulting mixture for 4 hours at 50 °C. Cool the mixture to room temperature, and then add HCl (33 wt % in water) until the pH reaches 3. Concentrate under reduced pressure. Dilute with 1 N aqueous HCl (100 mL) and extract with EtOAc (2 × 100 mL). Wash the combined organic extracts with brine (100 mL); dry over Na₂SO₄; filter; collect the filtrate; and concentrate under reduced pressure to give the title compound (7.60 g, 61 %). MS (m/z): 302 ($M + H$)⁺.

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Preparation 15

5,8-Dimethylquinoline-1-oxide



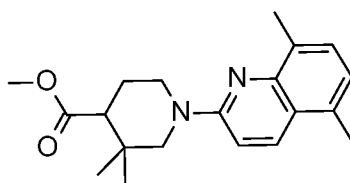
25

Add 3-chloroperoxybenzoic acid (5.96 g, 24.1 mmol) to a mixture of 5,8-dimethylquinoline (2 g, 12.1 mmol) and DCM (80 mL) maintained at 0 °C. After 1 hour, add Na₂SO₄ (5 g) and filter the mixture. Collect the filtrate. Stir the resulting mixture at

room temperature overnight. Dilute with DCM (100 mL); wash with 1 N aqueous NaOH (3 × 50 mL); dry over Na₂SO₄; filter; collect the filtrate; and concentrate to dryness under reduced pressure. Subject the resulting crude material to reverse-phase flash chromatography on C₁₈ silica gel, eluting with a gradient of 15 % to 70 % acetonitrile (10 mM ammonium bicarbonate) in water (10mM ammonium bicarbonate) to give the title compound (372 mg, 18 %). MS (*m/z*): 327 (M + H)⁺.

Preparation 16

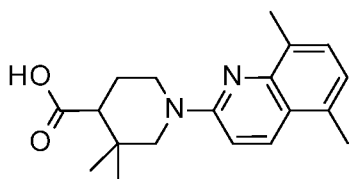
(±)-Methyl 1-(5,8-dimethyl-2-quinolyl)-3,3-dimethyl-piperidine-4-carboxylate.



Add N,N-diisopropylethylamine (2.02 mL, 11.6 mmol) and PyBroP (1.75 g, 3.75 mmol) to a mixture of (±)-methyl 3,3-dimethylpiperidine-4-carboxylate hydrochloride (600 mg, 2.89 mmol), 5,8-dimethylquinoline-1-oxide (678 mg, 3.91 mmol), and DCM (15 mL). Stir the resulting mixture at room temperature for 3 days. Subject the crude reaction to reverse-phase flash chromatography on C₁₈ silica gel, eluting with a gradient of 10% to 100% acetonitrile (0.1% formic acid) in water (0.1% formic acid). Combine the fractions containing the desired product and concentrate under vacuum to approximately 30 mL volume. Adjust the pH to 6 with addition of a 1N aqueous NaOH solution. Extract the aqueous solution with EtOAc (2 x 30 mL), wash with a pH 6 aqueous buffer solution (4 x 30 mL), dry over MgSO₄; filter; and concentrate under reduced pressure to give the title compound (166 mg, 18%). MS (*m/z*): 327 (M + H)⁺.

Preparation 17

(±)-1-(5,8-Dimethyl-2-quinolyl)-3,3-dimethyl-piperidine-4-carboxylic acid.

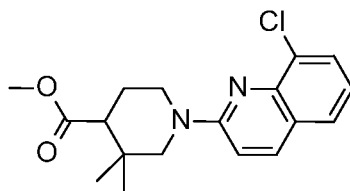


Combine a mixture of methyl (±)-1-(5,8-dimethyl-2-quinolyl)-3,3-dimethyl-piperidine-4-carboxylate (166 mg, 0.51 mmol), MeOH (0.1 mL), THF (0.5 mL), and 1 M aqueous NaOH (2.5

mL, 2.5 mmol) in a microwave vessel. Heat the resulting mixture at 120 °C for 2 hours in a microwave reactor. Cool the mixture to room temperature and add 1 N aqueous HCl until the pH is 6. Extract with CHCl₃ (2 × 25 mL); dry over Na₂SO₄; filter; collect the filtrate; and concentrate under reduced pressure to give the title compound (135 mg, 85 %). MS (*m/z*): 312 (M + H)⁺.

Preparation 18

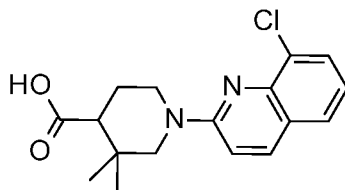
(±)-Methyl 1-(8-chloro-2-quinolyl)-3,3-dimethyl-piperidine-4-carboxylate.



Add K₂CO₃ (1.07 g, 7.75 mmol) to a mixture of methyl (±)-3,3-dimethylpiperidine-4-carboxylate hydrochloride (700 mg, 3.37 mmol), 2,8-dichloroquinoline (876 mg, 4.38 mmol), and DMSO (4.5 mL). Stir the resulting mixture at 130 °C overnight. Cool the mixture to room temperature; filter to remove the solids; dilute with water (5 mL); and extract with EtOAc (4 × 20 mL). Wash the combined organic extracts with water (20 mL) then brine (20 mL). Dry the organic extracts over MgSO₄; filter; collect the filtrate; and concentrate under reduced pressure. Subject the resulting crude material to flash chromatography on silica gel, eluting with a gradient of 0 % to 40 % EtOAc in hexanes, to give the title compound (1.1 g, 98 %). MS (*m/z*): 332 (M + H)⁺.

Preparation 19

(±)-1-(8-Chloro-2-quinolyl)-3,3-dimethyl-piperidine-4-carboxylic acid

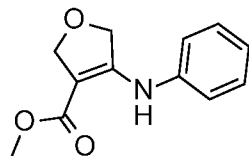


Combine a mixture of (±)-methyl 1-(8-chloro-2-quinolyl)-3,3-dimethyl-piperidine-4-carboxylate (1.1 g, 3.3 mmol), MeOH (0.8 mL), THF (3.3 mL), and 1 M aqueous NaOH (3.3 mL, 17 mmol) in a microwave vessel. Heat the resulting mixture at 120 °C for 2 hours in a microwave. Cool the mixture to room temperature and then add 5 N aqueous HCl until the pH is

6. Extract the mixture with EtOAc (3×10 mL). Combine the organic extracts; dry over Na_2SO_4 ; filter; collect the filtrate; and concentrate under reduced pressure to give the title compound (815 mg, 77 %). MS (m/z): 318 ($M + H$)⁺.

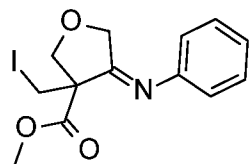
Preparation 20

5 Methyl 4-anilino-2,5-dihydrofuran-3-carboxylate



Add *p*-toluenesulfonic acid (500 mg, 3.16 mmol) to a mixture of ($\pm$)-methyl 4-oxotetrahydrofuran-3-carboxylate (6.6 g, 45.8 mmol), aniline (4.3 mL, 47.2 mmol) and toluene (70 mL). Fit the reaction vessel with a Dean-Stark trap and heat to 140 °C. Stir the resulting mixture at room temperature overnight. Heat the reaction to 140 °C and stir for 2 hours. Cool the mixture and add diethyl ether (100 mL) and water (100 mL). Extract the mixture with diethyl ether (3×50 mL); dry over MgSO_4 ; filter; collect the filtrate; and concentrate under reduced pressure. Subject the resulting crude material to flash chromatography on silica gel, eluting with a gradient of 0 % to 100 % EtOAc in heptane, to give the title compound (6.7 g, 67 %). MS (m/z): 220 ($M + H$)⁺.

Preparation 21

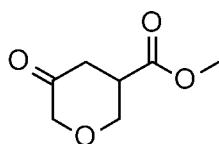
 $(\pm)$ -Methyl-3-(iodomethyl)-4-phenylimino-tetrahydrofuran-3-carboxylate

Add a solution of ($\pm$)-methyl 4-anilino-2,5-dihydrofuran-3-carboxylate (3.6 g, 16.4 mmol) and 18-crown-6 (4.8 g, 18 mmol) in toluene (15 mL) to a mixture of potassium *tert*-butoxide (2.0 g, 17.8 mmol) and toluene (15 mL). Stir the resulting mixture at room temperature for 30 minutes. Add diiodomethane (4 mL) and stir the reaction at room temperature overnight. Add water (50 mL) and extract with diethyl ether (2×50 mL). Collect the organic extracts and wash with brine (50 mL). Dry the organic extracts over MgSO_4 ; filter; collect the filtrate; and concentrate under reduced pressure. Subject the resulting crude material to flash

chromatography on silica gel, eluting with a gradient of 0 % to 60 % EtOAc in heptane, to give the title compound (1.0 g, 17 %). MS (m/z): 359 ($M + H$)⁺.

Preparation 22

(±)-Methyl 5-oxotetrahydropyran-3-carboxylate



5

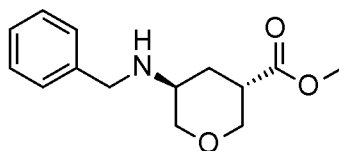
Heat a mixture of tri-*n*-butyltin hydride (1.12 mL, 4.18 mmol), 2,2'-azobis(2-methylpropionitrile) (35 mg, 0.21 mmol) and toluene (10 mL) to reflux. Add a solution of (±)-methyl-3-(iodomethyl)-4-phenylimino-tetrahydrofuran-3-carboxylate (1.0 g, 2.78 mmol) in toluene (50 mL) drop-wise over three hours. Stir the resulting mixture at reflux for 3 hours.

10 Concentrate the reaction under reduced pressure. Subject the resulting crude material to flash chromatography on silica gel, eluting with a gradient of 0 % to 80 % EtOAc in heptane, to give the title compound (0.37 g, 84 %). ¹H NMR (400 MHz, CDCl₃) δ 4.09-3.99 (m, 4H), 3.72 (s, 3H), 3.16 (m, 1H), 2.82 (dd, J= 7.2, 16.9 Hz, 1H), 2.66 (dd, J= 6.3, 16.9 Hz, 1H).

Preparation 23

15

(±)-Methyl *trans*-5-(benzylamino)tetrahydropyran-3-carboxylate

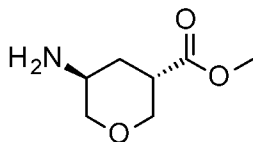


Add sodium triacetoxyborohydride (0.96 g, 4.53 mmol) to a mixture of (±)-methyl 5-oxotetrahydropyran-3-carboxylate (0.33 g, 2.08 mmol), benzylamine (0.31 mL, 2.84 mmol) and 1,2-dichloroethane (5 mL). Stir the resulting mixture at room temperature for 2.5 hours. Add a
20 saturated aqueous solution of sodium bicarbonate (10 mL); extract with DCM (3 × 10 mL); separate organic layer; wash with brine (10 mL); dry over MgSO₄; filter; collect the filtrate; and concentrate under reduced pressure. Subject the resulting crude material to flash chromatography on silica gel, eluting with a gradient of 0 % to 60 % EtOAc in heptane, to give the title compound (0.085 g, 16 % yield). MS (m/z): 250 ($M + H$)⁺.

25

Preparation 24

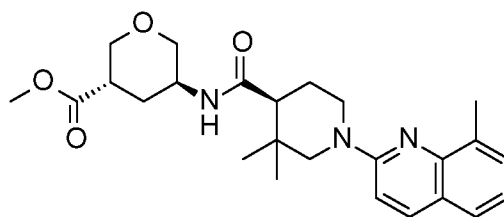
(±)-Methyl *trans*-5-aminotetrahydropyran-3-carboxylate



Add 10 % palladium on carbon (0.07 g, 0.66 mmol) to a mixture of (±)-methyl *trans*-5-(benzylamino)tetrahydropyran-3-carboxylate (0.085 g, 0.34 mmol) and ethanol (8 mL). Purge the mixture with hydrogen and stir the resulting mixture at room temperature under hydrogen (1 atm) overnight. Filter the mixture and concentrate under reduced pressure to give the title compound (0.045 g, 83 % yield). ¹H NMR (400 MHz, CDCl₃) δ 3.82-3.77 (m, 2H), 3.68 (m, 4H), 3.38-3.31 (m, 1H), 3.11-3.08 (m, 1H), 2.85-2.80 (m, 1H), 2.17-2.11 (m, 1H), 1.78-1.71 (m, 3H).

Preparation 25

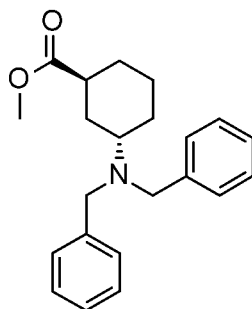
(±)-Methyl *trans*-5-[[[(4*S*)-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carbonyl]amino]tetrahydropyran-3-carboxylate



Add EDCI (0.087 g, 0.45 mmol) to a mixture of (–)-(4*S*)-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxylic acid (0.087 g, 0.29 mmol), (±)-methyl-*trans*-5-aminotetrahydropyran-3-carboxylate (0.045 g, 0.28 mmol), HOBt (9 mg, 0.06 mmol), triethylamine (0.15 mL, 1.08 mmol), in DCM (5 mL). Stir the resulting mixture at room temperature for 3 hours. Subject the resulting mixture to flash chromatography on silica gel, eluting with a gradient of 5 % to 100 % EtOAc in heptane, to give the title compound (0.085 g, 66 %). MS (*m/z*): 439 (M + H)⁺.

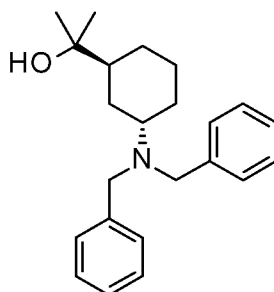
30

Preparation 26

(±)-Methyl *trans*-3-(dibenzylamino)cyclohexanecarboxylate

Add potassium carbonate (3.53 g, 25.6 mmol) and benzyl bromide (1.9 mL, 15.9 mmol) to a mixture of (±)-methyl-*trans*-3-aminocyclohexanecarboxylate (1.07 g, 6.8 mmol) in acetonitrile (40 mL). Heat the resulting mixture to 80 °C and stir for 5 hours. Cool the mixture and add acetonitrile (40 mL). Filter through diatomaceous earth and concentrate the filtrate under reduced pressure. Subject the resulting crude material to flash chromatography on silica gel, eluting with a gradient of 0 % to 20 % EtOAc in hexanes, to give the title compound (1.6 g, 70 %). MS (*m/z*): 338 (*M* + *H*)⁺.

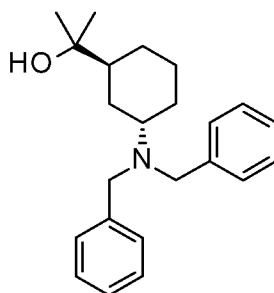
Preparation 27

(±)-2-[*trans*-3-(Dibenzylamino)cyclohexyl]propan-2-ol

Add methylmagnesium bromide (3.0 M solution in diethyl ether, 16 mL, 48 mmol) to a mixture of (±)-methyl-*trans*-3-(dibenzylamino)cyclohexanecarboxylate (1.6 g, 4.7 mmol) and THF (50 mL), maintained at 0 °C. Allow the reaction to warm to room temperature and stir overnight. Add water (25 mL) and filter through diatomaceous earth. Collect the aqueous filtrate. Extract the resulting aqueous material with EtOAc (50 mL). Collect the organic extracts; dry over Na₂SO₄; filter; collect the filtrate; and concentrate under reduced pressure. Add EtOAc (50 mL); wash with saturated aqueous NH₄Cl (2 × 25 mL); dry over Na₂SO₄; filter;

collect the filtrate; and concentrate under reduced pressure to give the title compound (1.1 g, 69 %). MS (m/z): 338 ($M + H$)⁺.

Preparation 28

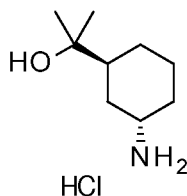
2-[(1*S*,3*S*)-3-(Dibenzylamino)cyclohexyl]propan-2-ol

5

Dissolve (±)-2-[*trans*-3-(dibenzylamino)cyclohexyl]propan-2-ol (1.1 g, 3.26 mmol) in ethanol (13 mg/mL). Subject the material to chiral SFC (Chiralpak IA, 2 cm × 15 cm) using 12 % (0.1 % dimethylethylamine in *i*-PrOH) in CO_{2(scf)} as the mobile phase at 220 nm and a flow rate of 70 mL/minute. Collect and combine the first fraction to elute and concentrate under reduced pressure, to provide the title compound (0.54 g, 49 %), ee = >99 %. MS (m/z): 338 ($M + H$)⁺. SFC Analytical method (Chiralpak IA, (15 cm × 0.46 cm) using 10 % (0.1 % dimethylethylamine in *i*-PrOH) in CO_{2(scf)} as the mobile phase at 220 nm and a flow rate of 3 mL/minutes).

10

Preparation 29

2-[(1*S*,3*S*)-3-Aminocyclohexyl]propan-2-ol hydrochloride

15

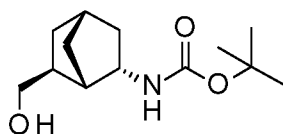
Add palladium hydroxide (20% on carbon, 0.51 g, 3.7 mmol) to a mixture of 2-[(1*S*,3*S*)-3-(dibenzylamino)cyclohexyl]propan-2-ol (0.54 g, 1.6 mmol) in MeOH (20 mL) in a ParrTM shaker vessel. Purge and pressurize the vessel with hydrogen to 414 kPa. Shake mixture for 4 days at room temperature. Filter the mixture through diatomaceous earth and concentrate the filtrate under reduced pressure. Dilute the mixture in diethyl ether (25 mL) and DCM (25 mL).

20

Add HCl (4.0 M solution in 1,4-dioxane, 0.8 mL) and concentrate under reduced pressure to give the title compound (0.24 g, 77 %). MS (m/z): 158 ($M + H$)⁺.

Preparation 30

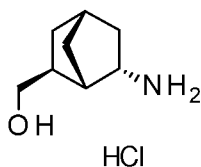
tert-Butyl N-[(1RS,2RS,4SR,6RS)-6-(hydroxymethyl)norbornan-2-yl]carbamate



Combine (acetylacetonato)dicarbonylrhodium (10 mg, 0.03 mmol), 1,1'-bis(diphenylphosphino)ferrocene (27 mg, 0.05 mmol), methyl bicyclo[2.2.1]hept-2-ene-5-carboxylate (2.5 g, 11.9 mmol) and *tert*-butanol (25 mL) to a PARRTM autoclave with mechanical stirrer. Purge and pressurize the vessel with Syngas (1:1 mix of carbon monoxide and hydrogen, 310 kPa). Stir overnight. Concentrate the mixture under reduced pressure. Dilute in DCM (29 mL) and MeOH (2.9 mL). Add sodium borohydride (0.26 g, 6.9 mmol) and stir for 20 min at room temperature. Add DCM (60 mL) and wash with saturated aqueous sodium carbonate (2 x 25 mL) and brine (25 mL). Dry the organic layer over Na₂SO₄; filter; collect the filtrate; and concentrate under reduced pressure. Subject the resulting crude material to flash chromatography on silica gel eluting with a solution of 30 % DCM, 30 % *tert*-butyl methyl ether and 40 % hexanes, to give the title compound as a mixture of diastereomers (0.58 g, 21 %). ¹H NMR (400 MHz, CDCl₃) δ 4.71-4.59 (m, 1H), 3.92-3.81 (m, 1H), 3.46-3.45 (m, 2H), 2.36 (d, J= 3.0 Hz, 1H), 2.21-2.05 (m, 3H), 1.55 (br s, 1H), 1.49-1.25(m, 12H), 1.16-1.09 (m, 1H), 0.68 (ddd, J= 12.9, 4.5, 2.4 Hz, 1H).

Preparation 31

[(1RS,2RS,4SR,6RS)-6-Aminonorbornan-2-yl]methanol hydrochloride



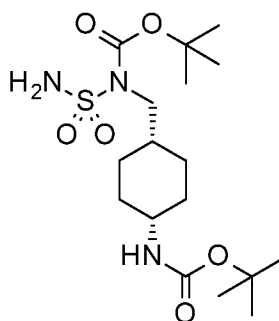
Add HCl (4.0 M solution in 1,4-dioxane, 2.1 mL, 8.3 mmol) to a mixture of *tert*-butyl N-[(1RS,2RS,4SR,6RS)-6-(hydroxymethyl)norbornan-2-yl]carbamate (0.2 g, 0.83 mmol) in DCM (8 mL) maintained at 0 °C. Allow the resulting mixture to warm to room temperature and stir for

3 hours. Concentrate the mixture under reduced pressure to give the title compound (0.15 g, 100%). ^1H NMR (400 MHz, CDCl_3) δ 8.17-8.02 (m, 3H), 4.93-4.90 (br, 1H), 3.47-3.45 (m, 1H), 3.19-3.15 (m, 2H), 2.31 (m, 1H), 2.14-2.12 (m, 1H), 2.01-1.94 (m, 1H), 1.92-1.88 (m, 1H), 1.44-1.35 (m, 2H), 1.22 (m, 1H), 1.04-0.98 (m, 2H).

5

Preparation 32

tert-Butyl ((*cis*-4-((*tert*-butoxycarbonyl)amino)cyclohexyl)methyl)(sulfamoyl)carbamate

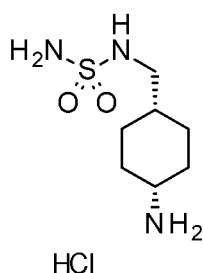


Add diisopropyl azodicarboxylate (1.6 mL, 7.8 mmol) to a mixture of *tert*-butyl *cis*-4-(hydroxymethyl)cyclohexylcarbamate (1.5g, 6.5 mmol), *tert*-butyl *N*-sulfamoylcarbamate (1.9 g, 9.8 mmol), triphenylphosphine (2.1 g, 7.8 mmol), and EtOAc (33 mL). Stir overnight at room temperature. Add water (50 mL); extract with EtOAc (2 $\times$ 50 mL); collect the organic extracts. Wash the organic extracts with brine (25 mL); dry over MgSO_4 ; filter; collect the filtrate; and concentrate under reduced pressure. Subject the resulting crude material to flash chromatography on silica gel, eluting with a gradient of 10 % to 80 % EtOAc in hexane, to give the title compound (1.78 g, 67 %). MS (m/z): 430 ($\text{M} + \text{Na}$) $^+$.

15

Preparation 33

cis-1-amino-4-[(sulfamoylamino)methyl]cyclohexane hydrochloride



Add HCl (4.0 M solution in 1,4-dioxane, 15 mL, 60 mmol) to *tert*-butyl ((*cis*-4-((*tert*-butoxycarbonyl)amino)cyclohexyl)methyl)(sulfamoyl)carbamate (1.78 g, 4.37 mmol). Stir the

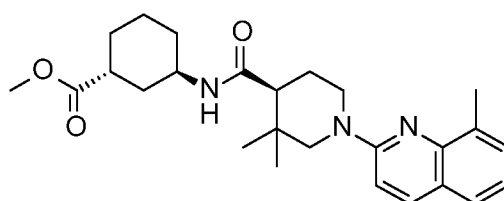
20

resulting mixture overnight at room temperature. Concentrate under reduced pressure. Dilute in MeOH (10 mL) and add drop-wise diethyl ether (150 mL) while vigorously stirring. Collect the resulting white precipitate by vacuum filtration to give the title compound (0.68 g, 56 %). MS (m/z): 208 ($M + H$)⁺.

5

Preparation 34

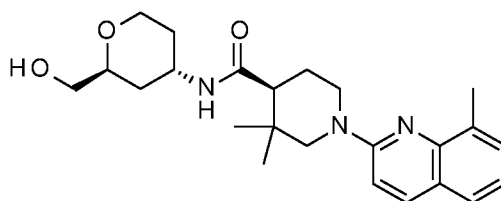
Methyl (1*RS*,3*RS*)-3-((*S*)-3,3-dimethyl-1-(8-methylquinolin-2-yl)piperidine-4-carboxamido)cyclohexane-1-carboxylate



Add triethylamine (0.9 mL, 7 mmol) to a mixture of (–)-(4*S*)-3,3-dimethyl-1-(8-methyl-
 10 2-quinolyl)piperidine-4-carboxylic acid (0.8 g, 3 mmol), (±)-methyl *trans*-3-aminocyclohexanecarboxylate hydrochloride (0.5 g, 3 mmol), BOP (2.0 g, 3 mmol) and DMF (5 mL). Stir the resulting mixture at room temperature overnight. Add a saturated aqueous solution of sodium bicarbonate (20 mL) and extract with EtOAc (2 × 25 mL). Collect the organic
 15 extracts; wash with brine (25 mL); dry over MgSO₄; filter; collect the filtrate; and concentrate under reduced pressure. Subject the resulting crude material to flash chromatography on silica gel, eluting with a gradient of 10 % to 90 % EtOAc in hexanes, to give the title compound (1.00 g, 80 %). MS (m/z): 438 ($M + H$)⁺.

Example 1

(*S*)-*N*-((2*S*,4*S*)-2-(Hydroxymethyl)tetrahydro-2H-pyran-4-yl)-3,3-dimethyl-1-(8-methylquinolin-
 20 2-yl)piperidine-4-carboxamide



Add benzotriazol-1-yl-oxytripyrrolidinophosphonium hexafluorophosphate (153 g, 293 mmol) as a slurry in DMF (152 mL) to a cold (10 °C) mixture of (–)-(4*S*)-3,3-dimethyl-1-(8-

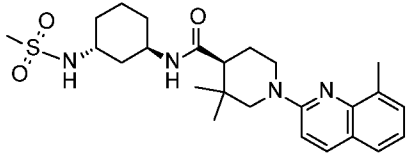
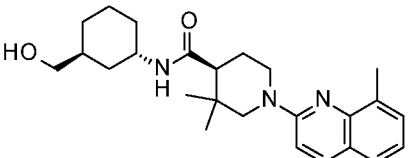
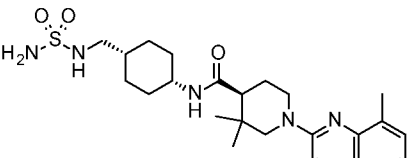
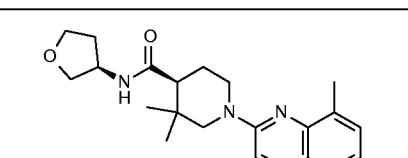
methyl-2-quinolyl)piperidine-4-carboxylic acid (76.0 g, 245 mmol), ((2*S*,4*S*)-4-aminotetrahydro-2H-pyran-2-yl)methanol hydrochloride (72 % pure, 57.37 g, 246 mmol), triethylamine (153 mL, 1.10 mol), and DMF (380 mL) while maintaining the internal reaction temperature below 20 °C. Thereafter allow the mixture to warm to room temperature and stir it for 30 minutes. Pour the mixture into ice water (1.5 L) while stirring it. Extract the mixture with DCM (2 × 600 mL) and wash the combined organic extracts with a half-saturated aqueous NaCl solution (1.0 L). Dry the organic solution over MgSO₄; filter; collect the filtrate; and concentrate the filtrate under reduced pressure to give a wet solid. Triturate the wet solid with water and isolate the solid by filtration. Triturate the solid again with water (1.5 L) and isolate the solid by filtration. Wash the solid with water (2 × 250 mL). Set this first batch of isolated solid aside. Collect the aqueous washings; extract with DCM (2 × 800 mL); combine the DCM extracts. Add the first batch of isolated solids to the combined DCM extracts and wash with aqueous HCl (2.5 M, 2 × 800 mL). Separate the aqueous layer and add 50% aqueous NaOH solution to the aqueous layer until the pH reaches 12 to induce precipitation. Filter the mixture to collect the resulting solid. Rinse the solid with water (2 × 300 mL). Dry the solid in a vacuum oven at 50 °C for 3 hours. Dissolve the solid in MeOH (1.2 L); add mercaptopropyl mesoporous silica scavenging resin (1.2 mmol/g, 715 m²/g, 54 μm average particle size); and stir at 50 °C overnight. Filter to remove the solids collecting the filtrate. Rinse the solid with 1:1 CH₂Cl₂:MeOH (2 × 500 mL). Combine the filtrates and concentrate under reduced pressure to give a white solid. Triturate the solid with TBME (1.5 L), and collect the solids. Crystallize the solid from hot EtOH (1.8 L). Dry the solid in a vacuum oven at 50 °C for 2.5 hours to give the title compound as a crystalline, white solid (76.7 g, 76 %). LC/MS (ESI): 412 [M+H]⁺.

The following Examples in Table 1 can be prepared essentially by the procedure of Preparation 34 using the appropriate starting amine in place of (±)-methyl-*trans*-3-aminocyclohexanecarboxylate hydrochloride and the appropriate starting carboxylic acid in place of (–)-(4*S*)-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxylic acid.

Table 1

Ex. No.	Structure	Chemical Name	MS (m/z) (M+H)	Purif.-Meth
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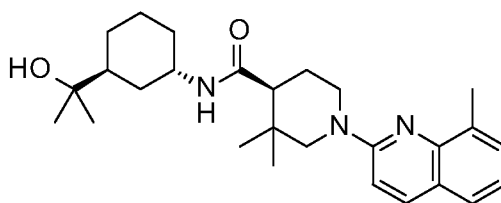
2		(4S)-N-[4-(Hydroxymethyl)- <i>cis</i> -cyclohexyl]-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxamide	410	A
3		(4S)-N-[6-(1S,2S,4R,6S) - (Hydroxymethyl)norbornan-2-yl]-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxamide	422	B
4		(4S)-1-(8-Chloro-2-quinolyl)-N-[4-(hydroxymethyl)- <i>cis</i> -cyclohexyl]-3,3-dimethylpiperidine-4-carboxamide	430	A
5		(4S)-N-[(1S,3S)-3-(Hydroxymethyl)cyclohexyl]-3,3-dimethyl-1-[4-(trifluoromethyl)phenyl]piperidine-4-carboxamide	413	C
6		(4S)-1-(5,8-Dimethyl-2-quinolyl)-N-[4-(hydroxymethyl)- <i>cis</i> -cyclohexyl]-3,3-dimethylpiperidine-4-carboxamide	424	B
7		(4S)-N-Cyclopentyl-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxamide	366	B
8		(4S)-N-[(1R)-1,2-Dimethylpropyl]-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxamide	368	B

9		(4S)-N-[3-(1S,3S)-(Methanesulfonamido)cyclohexyl]-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxamide	473	B
10		(4S)-N-[3-(1S,3S)-(Hydroxymethyl)cyclohexyl]-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxamide	410	A
11		(4S)-3,3-Dimethyl-1-(8-methyl-2-quinolyl)-N-[4-cis-[(sulfamoylamino)methyl]cyclohexyl]piperidine-4-carboxamide	488	A
12		(4S)-3,3-Dimethyl-1-(8-methyl-2-quinolyl)-N-[(3R)-tetrahydrofuran-3-yl]piperidine-4-carboxamide	368	A

Purification Methods (Purif Meth): A= Crude product is purified by silica gel normal phase column chromatography eluting with gradient of EtOAc in hexanes. B= Crude product is purified by C18 reverse phase column chromatography eluting with a gradient of acetonitrile and water. C= Crude product is purified by chiral column chromatography with Chiralpak AS-H, 21 × 150 mm) using 15% IPA in CO₂ as the mobile phase at a flow rate of 70 mL/min.

Example 13

(4S)-N-[(1S,3S)-3-(1-Hydroxy-1-methyl-ethyl)cyclohexyl]-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxamide



Add *N,N*-diisopropylethylamine (0.7 mL, 4 mmol) and HATU (0.31 g, 0.8 mmol) to a mixture of (–)-(4S)-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxylic acid (0.2 g, 0.67 mmol), 2-[(1S,3S)-3-aminocyclohexyl]propan-2-ol hydrochloride (0.13 g, 0.67 mmol) and

DMF (5 mL). Stir the resulting mixture at room temperature for 45 minutes. Subject the resulting reaction mixture to reverse phase flash chromatography on C18, eluting with a gradient of 5 % to 80 % ACN in water, to give the title compound (0.26 g, 88 %). MS (m/z): 438 ($M + H$)⁺.

- 5 Prepare the following Examples in Table 2 essentially by the procedure for Example 13 using the appropriate starting carboxylic acid in place of (–)-(4*S*)-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxylic acid.

Table 2

Ex. No.	Structure	Chemical Name	MS (m/z) ($M+H$)
14		(4 <i>S</i>)-N-[(1 <i>S</i> ,3 <i>S</i>)-3-(1-Hydroxy-1-methyl-ethyl)cyclohexyl]-3,3-dimethyl-1-[4-(trifluoromethyl)phenyl]piperidine-4-carboxamide	441
15		(4 <i>S</i>)-N-[(1 <i>S</i> ,3 <i>S</i>)-3-(1-Hydroxy-1-methyl-ethyl)cyclohexyl]-3,3-dimethyl-1-[5-(trifluoromethyl)pyrimidin-2-yl]piperidine-4-carboxamide	442

- 10 Prepare the following Examples in Table 3 essentially by the procedure for Preparation 25 using the appropriate starting amine in place of methyl (±)-*trans*-5-aminotetrahydropyran-3-carboxylate and the appropriate starting carboxylic acid in place of (–)-(4*S*)-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carboxylic acid.

Table 3

Ex. No.	Structure	Chemical Name	MS (m/z) ($M+H$)	Purification Method
---------	-----------	---------------	------------------------	---------------------

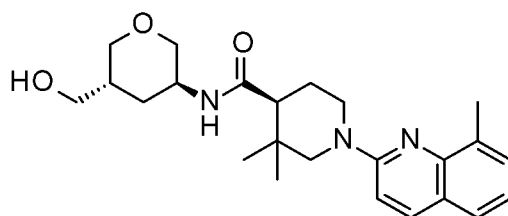
16		(4S)-1-(8-Methyl-2-quinolyl)-3,3-dimethyl-N-[(3S)-tetrahydropyran-3-yl]piperidine-4-carboxamide	382	A
17		(4S)-3,3-Dimethyl-1-(8-methyl-2-quinolyl)-N-[(1S,3S)-3-methylsulfonylcyclohexyl]piperidine-4-carboxamide	458	C

Purification Methods: A= Crude product is purified by silica gel normal phase column chromatography, eluting with a gradient of EtOAc in hexanes. C= Crude product is purified by chiral column chromatography using chiral SFC (Chiralcel OD-H, 21 × 250 mm) using 25%

5 EtOH(0.2% IPAm) in CO_{2(scf)} as the mobile phase at a flow rate of 70 mL/min.

Example 18

(S)-N-((3RS,5SR)-5-(Hydroxymethyl)tetrahydro-2H-pyran-3-yl)-3,3-dimethyl-1-(8-methylquinolin-2-yl)piperidine-4-carboxamide



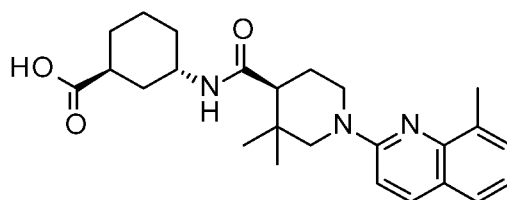
10 Add a 5 N aqueous NaOH solution (0.5 mL) to a mixture of methyl (3RS,5RS)-5-((4S)-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carbonyl]amino]tetrahydropyran-3-carboxylate (0.085 g, 0.19 mmol), MeOH (2 mL), and THF (2 mL). Stir the resulting mixture at room temperature for 1 hour. Add a 5 N aqueous HCl solution (0.5 mL) and concentrate. Dilute the crude mixture in THF (6 mL) and cool to 0 °C. Add a 2 M solution of borane-THF complex

15 in THF (0.15 mL, 0.3 mmol) drop-wise. Slowly warm the mixture to room temperature and stir at room temperature overnight. Add a saturated aqueous solution of sodium bicarbonate (10 mL) and extract the mixture with EtOAc (3 × 10 mL). Collect the organic extracts; wash with brine

(10 mL); dry over MgSO_4 ; filter; collect the filtrate; and concentrate under reduced pressure. Subject the resulting crude material to flash chromatography on silica gel eluting with 100 % EtOAc to give the title compound (0.046 g, 59 % yield over 2 steps). MS (m/z): 411 ($M + H$)⁺.

Example 19

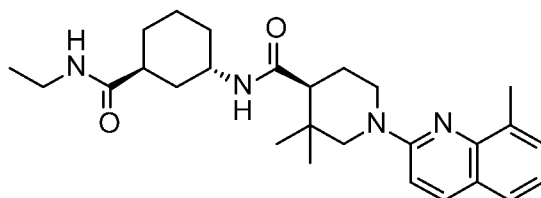
5 Methyl (1*RS*,3*RS*)-3-((*S*)-3,3-Dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carbonyl)amino]cyclohexanecarboxylic acid



10 Add a 5 N aqueous NaOH solution (2.2 mL) to a mixture of methyl ($\pm$)-*trans*-[[(*4S*)-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carbonyl]amino]cyclohexane-3-carboxylate (1.0 g, 2.2 mmol), MeOH (2 mL), and THF (8 mL). Stir the resulting mixture at room temperature overnight. Add a 5 N aqueous HCl solution (2.2 mL) to adjust the pH to 6.5. Extract with EtOAc (2 $\times$ 25 mL) and collect the organic extracts. Wash the organic extracts with brine (25 mL); dry over MgSO_4 ; filter; collect the filtrate; and concentrate under reduced pressure to give the title compound (0.935 g, 98 %). MS (m/z): 424 ($M + H$)⁺.

15 Example 20

(*S*)-*N*-((1*RS*,3*RS*)-3-(Ethylcarbamoyl)cyclohexyl)-3,3-dimethyl-1-(8-methylquinolin-2-yl)piperidine-4-carboxamide



20 Add triethylamine (0.7 mL, 5 mmol) to a mixture of ($\pm$)-*trans*-3-[[(*4S*)-3,3-dimethyl-1-(8-methyl-2-quinolyl)piperidine-4-carbonyl]amino]cyclohexanecarboxylic acid (0.9 g, 2 mmol), ethanamine (2 mL, 3 mmol), BOP (1.0 g, 3 mmol) and DMF (4 mL). Stir the resulting mixture at room temperature for 3 hours. Add a saturated aqueous solution of sodium bicarbonate (20 mL). Extract the resulting mixture with EtOAc (2 $\times$ 50 mL). Collect the organic extracts; wash

with brine (3 × 25 mL); dry over MgSO₄; filter; collect the filtrate; and concentrate under reduced pressure. Subject the resulting material to reverse-phase flash chromatography on C₁₈ silica gel, eluting with a gradient of 20 % to 80 % acetonitrile (0.1 % formic acid) in water (0.1 % formic acid), to give the title compound (0.25 g, 30 %). MS (*m/z*): 450 (M + H)⁺.

5 Biological Assays

Human mPGES-1 enzyme inhibition assay

Human mPGES-1 (Invitrogen™ (Cat# 97002RG, clone ID 6374722)) is subcloned into pcDNA3.1 and transiently expressed in 293E cells. Microsomes are prepared from cell pellets based on published methods (Oullet *et al.*, Purification and characterization of recombinant
10 microsomal prostaglandin E synthase-1, Protein Expression and Purification, 26 pp 489-495 (2002); and Thoren *et al.*, Human Microsomal Prostanglandin E Synthase-1, J. Biol Chem. 278(25) pp 22199-22209 (2003)). Briefly, pellets are brought up in homogenization buffer (15 mM Tris-HCl, pH 8.0; 0.25 M sucrose; 0.1 mM EDTA; 1 mM glutathione) and sonicated 5 × 30 seconds on ice. Homogenate is centrifuged at 5,000 × g for 10 minutes at 4 °C. The supernatant
15 fraction is decanted and loaded into Beckman Quick-Seal® tubes and centrifuged at 150,000 × g for 90 minutes at 4 °C. The supernatant fraction is discarded by decantation and the pellets are re-suspended in assay buffer (10 mM sodium phosphate, pH 7.0; 10% glycerol; 2.5 mM glutathione; Complete Protease Inhibitor Cocktail (Roche)). Protein concentration is determined using the Pierce Coomassie Plus™ reagent.

20 For the enzyme assay, the microsomes are diluted into assay buffer and 14 µL/well of the resulting solution is added to 384 well assay plates. Compound dilution plates (Nunc Cat#249944) are generated on a Tecan_MC384™ and 4 µL/well are added to the assay plates. Prostaglandin H₂ (PGH₂) is diluted into assay buffer immediately before use and 14 µL/well is added to the assay plates. Final concentrations are 6.55 µg/mL microsomes and 1.67 µM PGH₂.
25 After a 1.5 minute incubation at room temperature, 5 µL/well of 1 mg/mL SnCl₂ in 0.5 N HCl is added to stop the reaction. Five µL of the stopped reaction are transferred to a 384 well plate containing 45 µL of 0.1% formic acid, and the plates are stored at -80 °C. The plates are shipped to Agilent Technologies, formerly Biocius Lifesciences (Wakefield, MA 01880) for standard LC/MS analysis for PGE₂. The data are used to calculate the IC₅₀ (µM). The
30 compounds of the Examples inhibit human mPGES-1 with an IC₅₀ µM value of less than 100

nM. The exemplified compound of Example 1 inhibits human mPGES-1 with an IC_{50} μ M value of $0.00193, \pm 0.00064, n=17$. This result demonstrates that the exemplified compound of Example 1 is a potent inhibitor of the mPGES-1 enzyme in an isolated enzyme preparation.

5 Cell Based Assay for measuring Eicosanoid Selectivity

Human epithelial lung carcinoma cell line A549 is obtained from ATCC (CCL-185) and maintained in Kaighn's F12 cell culture medium + 10% fetal bovine serum (FBS) (plating medium) under standard 5% CO₂ humidified atmosphere at 37 °C. The cells are passaged at 1:3 twice per week.

10 For the assays, cells are released from flasks by washing once with PBS, then once with Trypsin/EDTA. After 3-5 minutes at 37 °C, the cells are suspended in plating medium and centrifuged at 2,000 rpm, 25 °C, for 5 minutes. The supernatant is aspirated and the cell pellet is resuspended in F12K plating medium. The cell number is determined by counting an aliquot of cells which has been diluted in PBS and Trypan blue on a hemocytometer. Cells are plated at
15 40,000/well in 96 well Falcon plates 24 hours prior to treatment. Compounds are diluted in DMSO to 100 $\times$ of the final concentration in Screen Mates tubes. The medium is removed from the cells and fresh medium (90 μ L/well) is added to the cells. The compounds are added at 1 μ L / well, $n=2$, to give seven concentrations each. Cells are pretreated with compounds for 30 minutes at 37 °C, 5% CO₂. Prostaglandin E₂ production is induced by the addition of
20 recombinant human interleukin (rhIL-1 β) diluted in plating medium to 10 $\times$ final. A 10 μ L/well aliquot is added to give a final rhIL-1 β concentration of 0.1 ng/mL. The treatment period is approximately 18 hours. Conditioned medium is removed to v-bottom polypropylene plates. The conditioned medium is assayed for levels of PGE₂ and prostaglandin I₂ (PGI₂) by specific enzyme immune-assay EIAs, according to the manufacturer's protocols (Cayman). Briefly,
25 conditioned medium (1 μ L) is added to each well of a 96 well plate coated with a capture antibody and containing EIA buffer (49 μ L) supplied by the manufacturer. The tracer is diluted with the EIA buffer (50 μ L). The detection antibody is diluted with the EIA buffer (50 μ L). The plate is covered with adhesive sealing film and is incubated for 1 hour at room temperature on an orbital shaker at 100 rpm. The wash buffer is diluted into MILLIPORE purified water, and the
30 plate is washed 5 $\times$ 350 μ L/well, using a plate washer. The substrate (Ellman's reagent) is

diluted with MILLIPORE purified water and then added to the plate at 200 μ L/well. After approximately 90-120 minutes at room temperature on an orbital shaker at 100 rpm, the plates are read at A412 on a plate reader. A standard curve of PGE₂ is used to calibrate the unknowns. The exemplified compound of Example 1 inhibits PGE₂ formation with an IC₅₀ of 0.00471 μ M $\pm$ 0.00301, n=2.

Human Whole Blood Assay

Blood is collected from normal volunteer donors into sodium heparin VACUTAINER tubes. Donors are selected, in part, on their confirmation that they have not taken NSAIDs, aspirin, Celebrex®, or glucocorticoids within two weeks of the donation. All tubes/donor are pooled into 250 mL Corning conical centrifuge tubes and 436.5 μ L/well is distributed into deep well polypropylene plates. Compounds are diluted in DMSO to 100 $\times$ final and 4.5 μ L/well in duplicate or triplicate is added to give 7 point curves. The blood is pretreated with compounds at 37 $^{\circ}$ C, 5% CO₂, in a humidified atmosphere, covered with a MicroClima Environmental Microplate lid, for 30 minutes after which 9 μ L/well of a solution of 5 mg/mL of LPS (Sigma, serotype 0111:B4) in 1 mg/mL BSA/PBS is added to give a final LPS concentration of 100 μ g/mL. The plates are incubated for 20-24 hours, at 37 $^{\circ}$ C, 5% CO₂, in a humidified atmosphere. The plates are tightly sealed with the aluminum foil lids and are chilled on ice for approximately 1 hour. Then the plates are centrifuged at 1,800 $\times$ g, 10 minutes, 4 $^{\circ}$ C, in an Eppendorf 5810R centrifuge. Plasma is removed from the cell layer using the Rainin L200 with sterile filtered tips and transferred to v-bottom polypropylene plates. One hundred microliters is quantitatively transferred to Costar cluster tubes blocks and 400 μ L/well of the MeOH stop reagent and internal standards, *d*₄-PGE₂, *d*₄-PGF_{2 α} , and *d*₄-TXA_{2 β} are added. Samples are vortexed for 5 minutes and are placed at -20 $^{\circ}$ C for at least one hour. Samples are centrifuged for 10 minutes at 4000 rpm in an Eppendorf 5810R.

Solid phase extraction is performed using Waters HLB 30 mg/bed 96 well plates on a vacuum manifold: Step 1, the matrix is washed with MeOH (1 mL), followed by 0.1% formic acid in water (1 mL); Step 2, 400 μ L sample is applied along with 0.1% formic acid in water (900 μ L) and allowed to bind for 5 minutes; Step 3, the matrix is washed with 0.1% formic acid in water (600 μ L), followed by 80/20 water/MeOH (600 μ L); Step 4, the products are eluted with

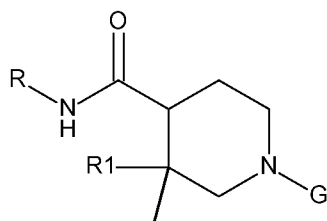
2-500 μ L volumes of EtOAc; Step 5 the samples are dried under nitrogen and reconstituted in 75/25 water/acetonitrile with 0.1% formic acid (50 μ L). The products are analyzed by LC/MS/MS. Example 1 inhibits The PGE₂ formation in this assay with an IC₅₀ of 0.00205 ± 0.00082 , n=11. This result supports that Example 1 inhibits PGE₂ synthesis in human whole blood.

5

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What is claimed is:

1. A compound of Formula 1:



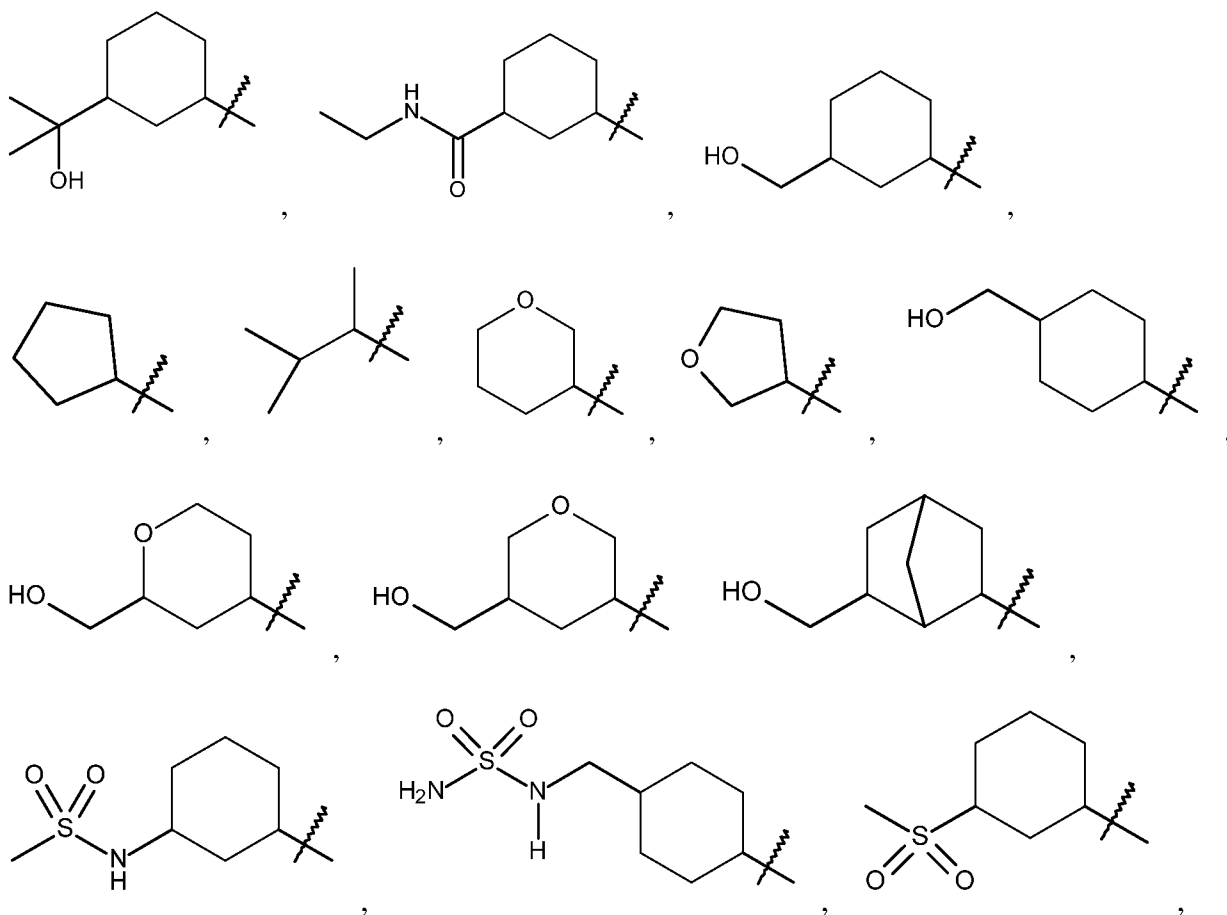
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1,

wherein:

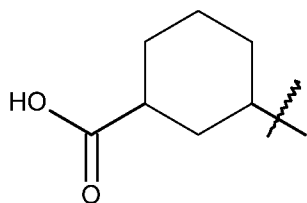
R1 is H or -CH₃;

R is selected from:



10

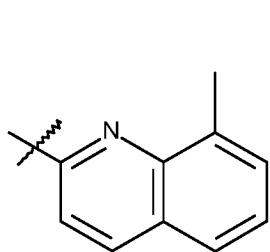
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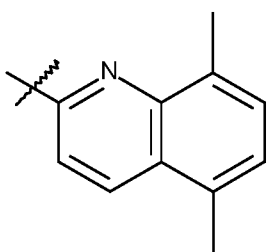
and

;

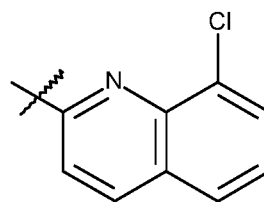
G is selected from:



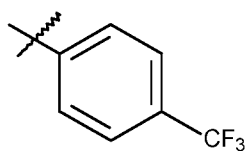
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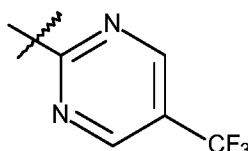
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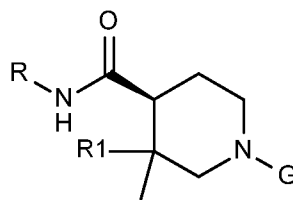
, and



;

5 or a pharmaceutically acceptable salt thereof.

2. A compound according to Formula 2:



2,

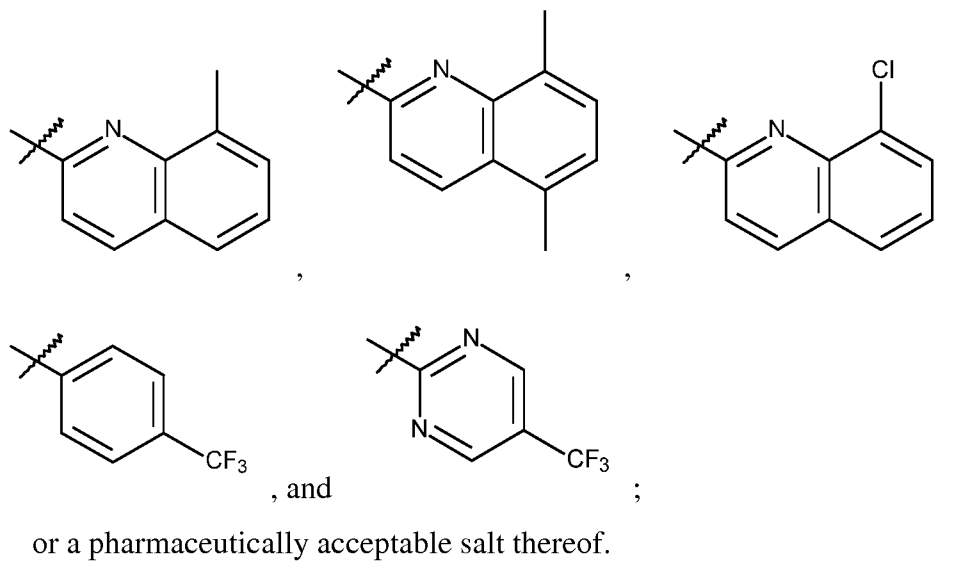
10 wherein:

R1 is H or -CH₃;

R is selected from:

5 and 0 ;
G is selected from:

48

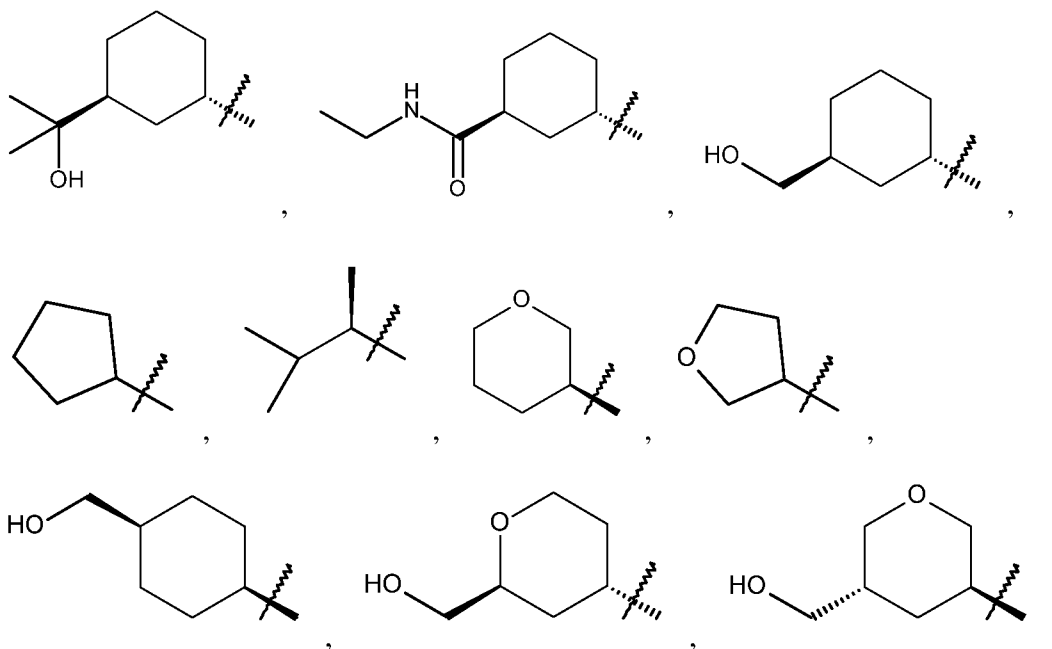


5 3. A compound according claim 1 or 2:

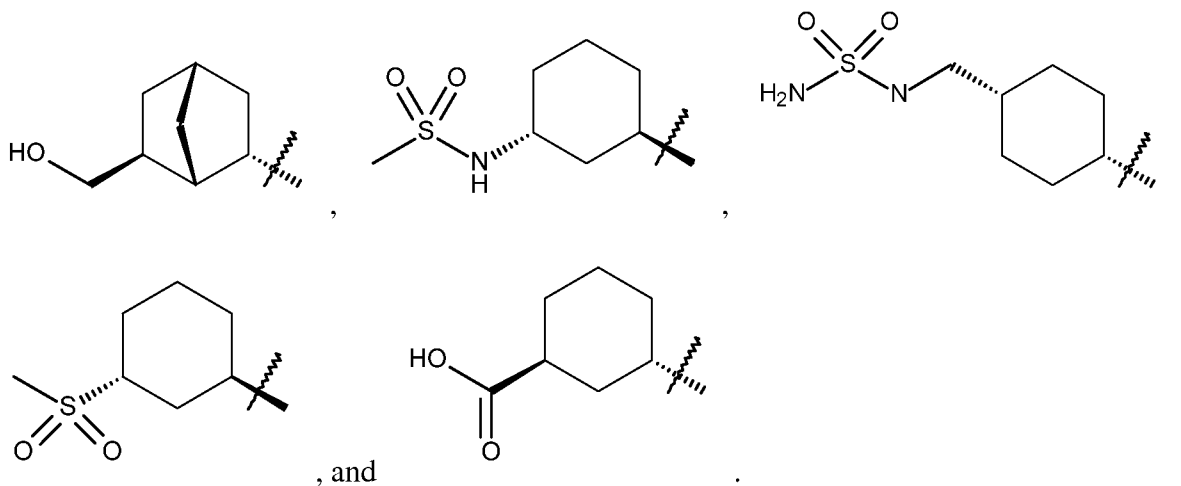
wherein:

R1 is H or -CH₃;

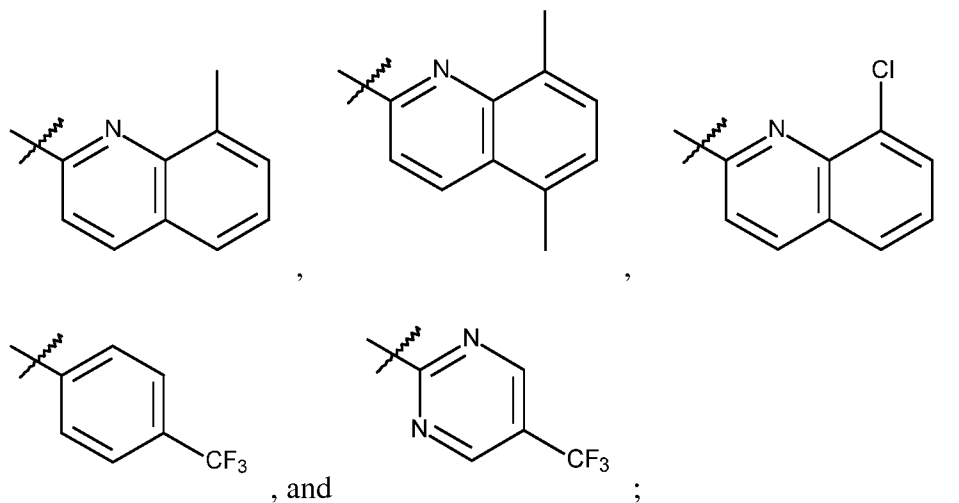
R is selected from:



49

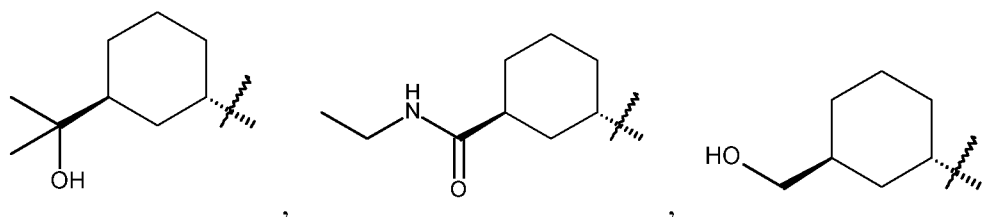


G is selected from:

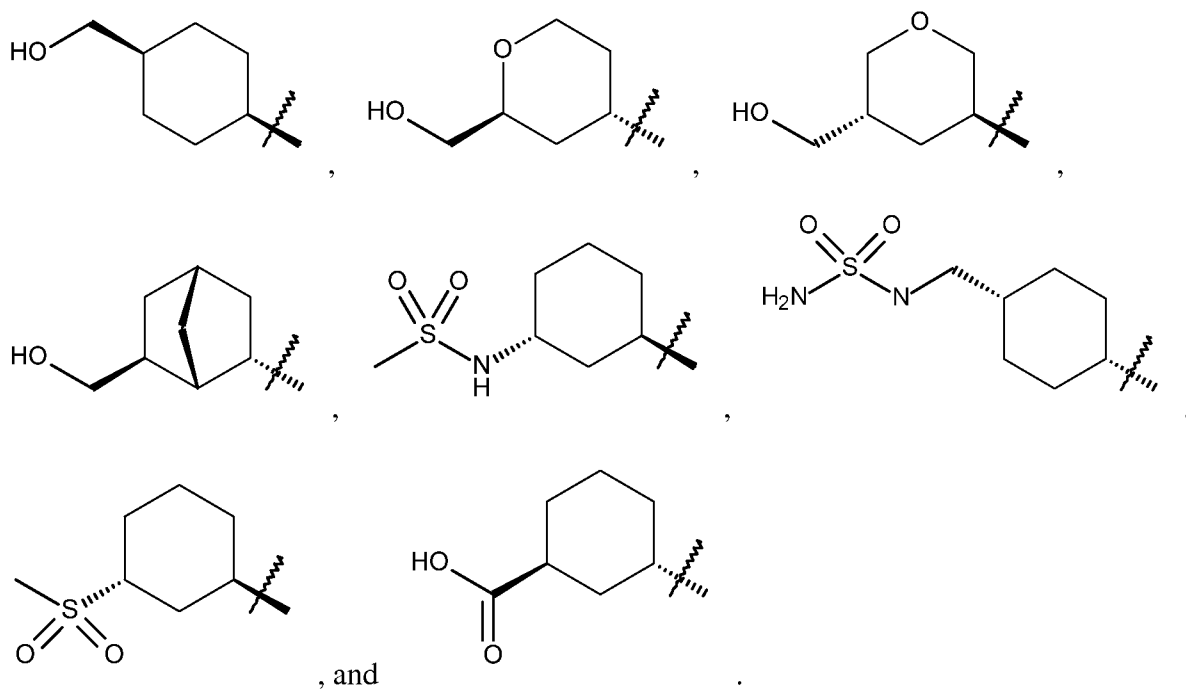


or a pharmaceutically acceptable salt thereof

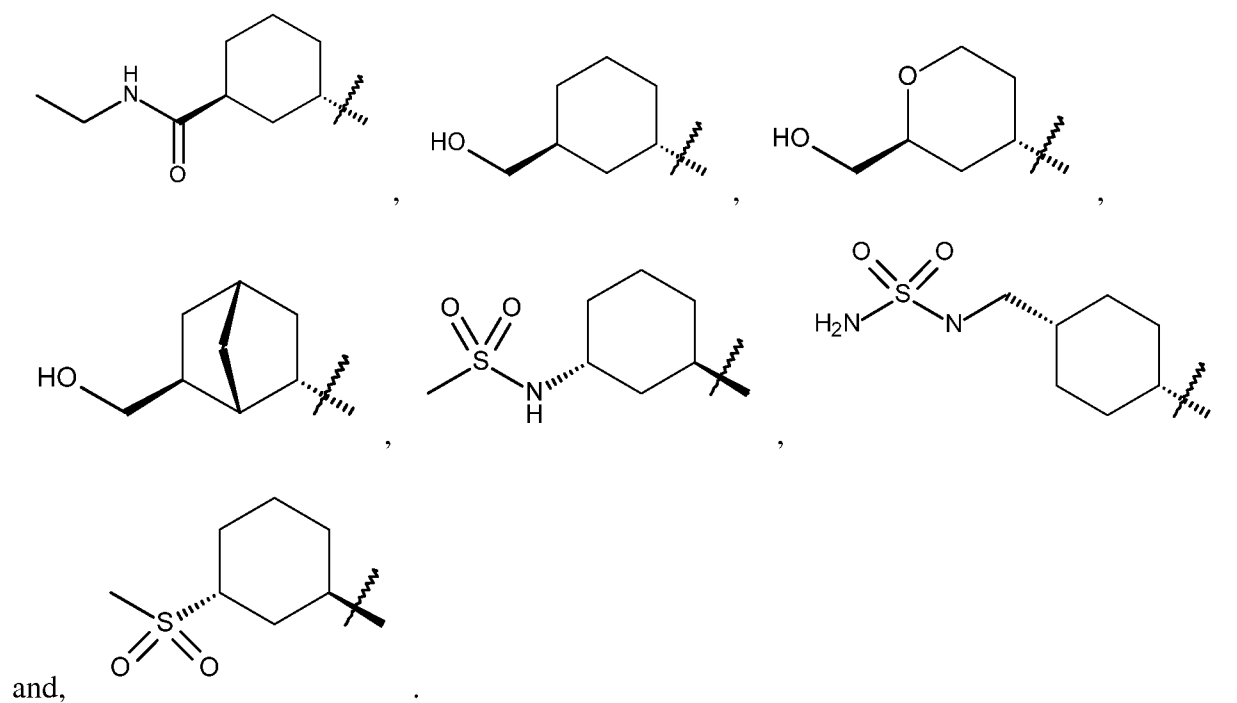
4. A compound according to any one of claims 1 to 3, or a pharmaceutically acceptable salt thereof, wherein R is selected from:



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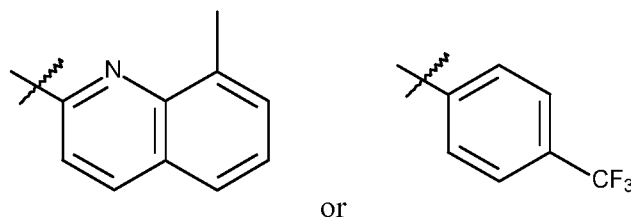


5. A compound according to any one of claims 1 to 4, or a pharmaceutically acceptable salt thereof, wherein R is selected from:



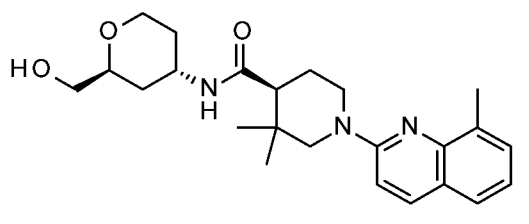
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6. A compound according any one of claims 1 to 5, or a pharmaceutically acceptable salt thereof, wherein G is:



5 7. A compound according to any one of claims 1 to 6, or a pharmaceutically acceptable salt thereof, wherein R1 is $-\text{CH}_3$.

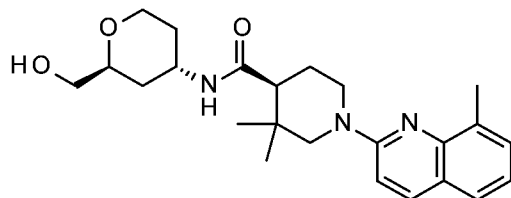
8. A compound which is:



10

or a pharmaceutically acceptable salt thereof.

9. A compound according to claim, which is:



15

10. A compound according to any one of claims 1 to 9 wherein the pharmaceutically acceptable salt is a hydrochloride salt.

11. A pharmaceutically acceptable composition comprising a compound according to any one of claims 1 to 10 and at least one of a pharmaceutically acceptable carrier, diluent, or excipient.

5 12. A method of treating a patient in need of treatment for pain associated with arthritis or osteoarthritis said method comprising administering to the patient an effective amount of a pharmaceutically acceptable composition according to claim 11.

10 13. A method of treating a patient in need of treatment for inflammation associated with arthritis or osteoarthritis, said method comprising administering to the patient an effective amount of a pharmaceutically acceptable composition according to claim 11.

15 14. A method of treating a patient in need of treatment for arthritis or osteoarthritis, said method comprising administering to the patient an effective amount of a compound according to any one of claims 1 to 10.

20 15. A method of treating a patient in need of treatment for pain associated with arthritis, said method comprising administering to the patient an effective amount of a compound according to any one of claims 1 to 10.

16. A method of treating a patient in need of treatment for pain associated with osteoarthritis, said method comprising administering to the patient an effective amount of a compound according to any one of claims 1 to 10.

25 17. A method of treating a patient in need of treatment for inflammation associated with arthritis, said method comprising administering to the patient an effective amount of a compound according to any one of claims 1 to 10.

18. A method of treating a patient in need of treatment for inflammation associated with osteoarthritis, said method comprising administering to the patient an effective amount of a compound according to any one of claims 1 to 10.

5 19. A compound according to any one of claims 1 to 10 for use as a medicament.

20. A compound according to any one of claims 1 to 10 for use in therapy.

10 21. A compound according to any one of claims 1 to 10 for use in the treatment of arthritis or osteoarthritis.

22. A compound according to any one of claims 1 to 10 for use in the treatment of pain or inflammation associated with arthritis or osteoarthritis.

15 23. Use of a compound according to any one of claims 1 to 10 for the manufacture of a medicament to treat arthritis or osteoarthritis.

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2015/056960

A. CLASSIFICATION OF SUBJECT MATTER INV. C07D405/14 C07D401/04 C07D211/62 A61K31/506 A61K31/4709 A61K31/451 A61P29/00 A61P19/02 ADD. According to International Patent Classification (IPC) or to both national classification and IPC											
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) C07D Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) EPO-Internal, WPI Data											
C. DOCUMENTS CONSIDERED TO BE RELEVANT <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 10%;">Category*</th> <th style="width: 70%;">Citation of document, with indication, where appropriate, of the relevant passages</th> <th style="width: 20%;">Relevant to claim No.</th> </tr> </thead> <tbody> <tr> <td style="text-align: center; vertical-align: top;">Y</td> <td style="vertical-align: top;"> WO 2013/146970 A1 (DAIICHI SANKYO CO LTD [JP]) 3 October 2013 (2013-10-03) cited in the application the whole document; in particular the claims; the abstract; specific compounds, e.g. 48, 87, 88, 92-94, 99-105, 115 & DATABASE WPI [Online] THOMSON SCIENTIFIC, LONDON, GB; 2013, Database accession no. 2013-Q08373 ----- </td> <td style="text-align: center; vertical-align: top;">1-23</td> </tr> <tr> <td style="text-align: center; vertical-align: top;">Y</td> <td style="vertical-align: top;"> EP 1 627 869 A1 (AJINOMOTO KK [JP]) 22 February 2006 (2006-02-22) the whole document; in particular the claims; paragraphs [0001], [0007], [0010] ----- </td> <td style="text-align: center; vertical-align: top;">1-23</td> </tr> </tbody> </table>			Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.	Y	WO 2013/146970 A1 (DAIICHI SANKYO CO LTD [JP]) 3 October 2013 (2013-10-03) cited in the application the whole document; in particular the claims; the abstract; specific compounds, e.g. 48, 87, 88, 92-94, 99-105, 115 & DATABASE WPI [Online] THOMSON SCIENTIFIC, LONDON, GB; 2013, Database accession no. 2013-Q08373 -----	1-23	Y	EP 1 627 869 A1 (AJINOMOTO KK [JP]) 22 February 2006 (2006-02-22) the whole document; in particular the claims; paragraphs [0001], [0007], [0010] -----	1-23
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.									
Y	WO 2013/146970 A1 (DAIICHI SANKYO CO LTD [JP]) 3 October 2013 (2013-10-03) cited in the application the whole document; in particular the claims; the abstract; specific compounds, e.g. 48, 87, 88, 92-94, 99-105, 115 & DATABASE WPI [Online] THOMSON SCIENTIFIC, LONDON, GB; 2013, Database accession no. 2013-Q08373 -----	1-23									
Y	EP 1 627 869 A1 (AJINOMOTO KK [JP]) 22 February 2006 (2006-02-22) the whole document; in particular the claims; paragraphs [0001], [0007], [0010] -----	1-23									
☐ Further documents are listed in the continuation of Box C. ☒ See patent family annex.											
* Special categories of cited documents : "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family											
Date of the actual completion of the international search 15 December 2015		Date of mailing of the international search report 23/12/2015									
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016		Authorized officer Hanisch, Inken									

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US2015/056960

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 2013146970	A1	03-10-2013	NONE

EP 1627869	A1	22-02-2006	AT 556067 T 15-05-2012
		CA 2526361 A1	02-12-2004
		EP 1627869 A1	22-02-2006
		EP 2281823 A2	09-02-2011
		JP 5279987 B2	04-09-2013
		US 2006128755 A1	15-06-2006
		US 2009258900 A1	15-10-2009
		WO 2004103954 A1	02-12-2004
