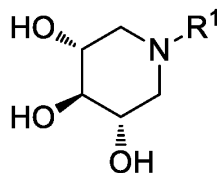


WHAT IS CLAIMED IS:

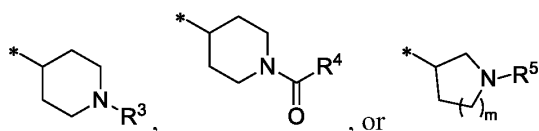
1. A compound of Formula (I) or a pharmaceutically acceptable salt thereof:



(I)

wherein

R^1 is $(CH_2)_nR^2$, wherein n is 1, 2, or 3, and R^2 is cyclohexyl, phenyl, thiophen-2-yl, thiophen-3-yl, pyridine-2-yl, adamantyl, 2,3-dihydro-1H-inden-2-yl, 1,2,3,4-tetrahydronaphthalen-2-yl, spiro[2.5]octan-6-yl, spiro[3.5]nonan-7-yl, spiro[4.5]decan-8-yl, (5S,8s)-3,3-dimethyl-2-oxaspiro[4.5]decan-8-yl, (benzo[d][1,3]dioxol-5-yl)methyl, (2,3-dihydrobenzo[b][1,4]dioxin-6-yl)methyl, 1-cyclohexylazetidin-3-yl,



each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C_{1-6} alkyl, C_{2-6} alkenyl, cyclopropyl, difluoromethyl, 1,1-difluoroethyl-1-yl, 2-fluoropropan-2-yl, methoxymethyl, OCF_3 , CF_3 , phenyl, pyrrolidin-1-yl, piperidin-1-yl, 4-morpholino, C_{1-6} alkoxy, cyclopropylmethoxy, (tetrahydrofuran-3-yl)oxy, (tetrahydro-2H-pyran-3-yl)oxy, (tetrahydro-2H-pyran-4-yl)oxy, phenoxy, (tetrahydrofuran-3-yl)methoxy, tetrahydro-2H-pyran-4-yl, 3,5-dimethylisoxazol-4-yl, and/or 3,5-dimethyl-1H-pyrazol-4-yl;

where R^3 is C_{1-6} alkyl, phenyl, pyridine-2-yl, pyridin-3-yl, benzo[d]thiazol-2-yl, cyclohexylcarbamothioyl, or cyclohexylcarbamoyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C_{1-6} alkyl, C_{2-6} alkenyl, cyclopropyl, C_{1-6} alkoxy, OCF_3 , and/or CF_3 ; and

where R^4 is C_{1-6} alkyl, C_{3-7} cycloalkyl, phenyl, thiophen-3-yl, phenylmethyl, or cyclopentylmethyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, C_{1-6} alkyl, OCH_3 , and/or CF_3 ; and

where m is 1 or 2; and

where R⁵ is phenyl, pyridin-2-yl, pyridin-3-yl, pyridin-4-yl, pyrimidin-5-yl, thiophen-3-yl, benzo[d]thiazol-4-yl, benzo[d]thiazol-2-yl, phenylcarbonyl, thiazol-2-yl, benzo[d]oxazol-2-yl, or benzo[d]thiazol-2-yl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C₁₋₆ alkyl, C₁₋₆ alkoxy, OCF₃, and/or CF₃,

with the proviso that R¹ is not benzyl or 3-phenylpropyl.

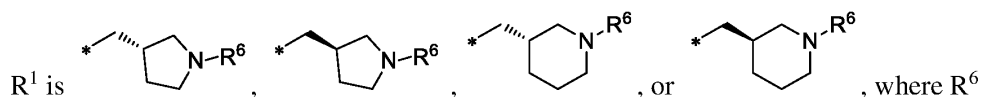
2. The compound of claim 1 wherein:

R¹ is (CH₂)_nR², wherein n is 1 or 2, and R² is cyclohexyl, cyclohexylmethyl, phenylethyl, 4-phenylcyclohexyl, 4-(trifluoromethoxy)cyclohexyl, spiro[2.5]octan-6-yl, spiro[3.5]nonan-7-yl, spiro[4.5]decan-8-yl, (5S,8s)-3,3-dimethyl-2-oxaspiro[4.5]decan-8-yl, 1,2,3,4-tetrahydronaphthalen-2-yl, 2,3-dihydro-1H-inden-2-yl, (adamantyl)methyl, (pyridine-2-yl)methyl, (benzo[d][1,3]dioxol-5-yl)methyl, (2,3-dihydrobenzo[b][1,4]dioxin-6-yl)methyl, ([1,1'-biphenyl]-4-yl)methyl, 1-(2,2,2-trifluoroethyl)piperidin-4-yl, 1-(pyridin-3-yl)piperidin-4-yl, 1-(pyridin-2-yl)piperidin-4-yl, 1-(benzo[d]thiazol-2-yl)piperidin-4-yl, 1-(cyclohexylcarbonyl)piperidin-4-yl, 1-(cyclohexylcarbamothioyl)piperidin-4-yl, 1-phenylpiperidin-4-yl, 1-cyclohexylazetidin-3-yl, thiophen-2-yl, 2-(thiophen-2-yl)methyl, or 2-(thiophen-3-yl)methyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C₁₋₆ alkyl, cyclopropyl, vinyl, difluoromethyl, 1,1-difluoroethyl-1-yl, 2-fluoropropan-2-yl, methoxymethyl, C₁₋₆ alkoxy, and/or CF₃; or

R¹ is benzyl, substituted from one up to the maximum number of substituents with one or more of F, Cl, C₁₋₆ alkyl, phenyl, pyrrolidin-1-yl, piperidin-1-yl, C₁₋₆ alkoxy, cyclopropylmethoxy, phenoxy, and/or CF₃; or

R¹ is phenylethyl, optionally substituted from one up to the maximum number of substituents with one or more of pyrrolidin-1-yl, piperidin-1-yl, 4-morpholino, cyclopropylmethoxy, (tetrahydrofuran-3-yl)oxy, (tetrahydro-2H-pyran-3-yl)oxy, (tetrahydro-2H-pyran-4-yl)oxy, phenoxy, (tetrahydrofuran-3-yl)methoxy, tetrahydro-2H-pyran-4-yl, 3,5-dimethylisoxazol-4-yl, 3,5-dimethyl-1H-pyrazol-4-yl, F, Cl, C₁₋₆ alkyl, cyclopropyl, propen-2-yl, OCH₃, and/or CF₃; or

R¹ is (1-formylpiperidin-4-yl)methyl, substituted on the formyl group with one of: C₁₋₆ alkyl, C₃₋₇ cycloalkyl, phenyl, thiophen-3-yl, phenylmethyl, or cyclopentylmethyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, C₁₋₆ alkyl, OCH₃, and/or CF₃; or



is selected from the group consisting of: phenyl, pyridin-2-yl, pyridin-3-yl, pyridin-4-yl, pyrimidin-5-yl, thiophen-3-yl, benzo[d]thiazol-4-yl, benzo[d]thiazol-2-yl, phenylcarbonyl, thiazol-2-yl, benzo[d]oxazol-2-yl, and benzo[d]thiazol-2-yl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C₁₋₆ alkyl, C₁₋₆ alkoxy, OCF₃, and/or CF₃,

with the proviso that R^1 is not benzyl or 3-phenylpropyl.

3. The compound of claim 1 wherein:

R^1 is cyclohexylmethyl, (4,4-dimethylcyclohexyl)methyl, (4,4-difluorocyclohexyl)methyl, (4,4-dichlorocyclohexyl)methyl, (4-ethylcyclohexyl)methyl, ((1s,4S)-4-vinylcyclohexyl)methyl, ((1s,4S)-4-isopropylcyclohexyl)methyl, ((1r,4R)-4-isopropylcyclohexyl)methyl, 4-(tert-butyl)cyclohexyl)methyl, ((1s,4S)-4-(tert-butyl)cyclohexyl)methyl, ((1r,4R)-4-(tert-butyl)cyclohexyl)methyl, ((1s,4S)-4-(trifluoromethyl)cyclohexyl)methyl, ((1r,4R)-4-(trifluoromethyl)cyclohexyl)methyl, ((1s,4S)-4-(2-fluoropropan-2-yl)cyclohexyl)methyl, ((1r,4R)-4-(2-fluoropropan-2-yl)cyclohexyl)methyl, ((1s,4S)-4-methoxycyclohexyl)methyl, ((1r,4R)-4-methoxycyclohexyl)methyl, (4-(methoxymethyl)cyclohexyl)methyl, ((1s,4S)-4-cyclopropylcyclohexyl)methyl, ((1r,4R)-4-cyclopropylcyclohexyl)methyl, (4-phenylcyclohexyl)methyl, (spiro[2.5]octan-6-yl)methyl, (spiro[3.5]nonan-7-yl)methyl, (spiro[4.5]decan-8-yl)methyl, 2-cyclohexylethyl, 2-(4,4-difluorocyclohexyl)ethyl, 2-((1s,4S)-4-(trifluoromethyl)cyclohexyl)ethyl, 2-((1r,4R)-4-(trifluoromethyl)cyclohexyl)ethyl, 2-(adamantan-1-yl)ethyl, 3-cyclohexylpropyl, phenethyl, 2-methylphenethyl, 2-methoxyphenethyl, 2-fluorophenethyl, 2-chlorophenethyl, 2,3-difluorophenethyl, 2,4-difluorophenethyl, 2,5-difluorophenethyl, 3,4-difluorophenethyl, 2-fluoro-4-methoxyphenethyl, 3-chloro-2-fluorophenethyl, 4-chloro-2-fluorophenethyl, 5-chloro-2-fluorophenethyl, 2,6-difluorophenethyl, 3-chloro-2,6-difluorophenethyl, 2,6-difluoro-4-(prop-1-en-2-yl)phenethyl, 2,6-difluoro-4-isopropylphenethyl, 2,6-difluoro-3-isopropylphenethyl, 4-cyclopropyl-2,6-difluorophenethyl, 2,6-difluoro-4-(trifluoromethyl)phenethyl, 2,6-difluoro-4-(pyrrolidin-1-yl)phenethyl, 2,6-difluoro-4-(piperidin-1-yl)phenethyl, 2,6-difluoro-4-morpholinophenethyl, 4-butoxy-2,6-difluorophenethyl, 4-(cyclopropylmethoxy)-2,6-difluorophenethyl, 4-((tetrahydrofuran-3-yl)oxy)phenethyl, 4-((tetrahydro-2H-pyran-3-

yl)oxy)phenethyl, 4-((tetrahydro-2H-pyran-4-yl)oxy)phenethyl, 4-phenoxyphenethyl, 4-((tetrahydrofuran-3-yl)methoxy)phenethyl, (R)-2-phenylpropyl, (S)-2-phenylpropyl, 2-([1,1'-biphenyl]-4-yl)ethyl, 2-(3,5-difluoro-[1,1'-biphenyl]-4-yl)ethyl, 2-(benzo[d][1,3]dioxol-5-yl)ethyl, 2-(6-fluorobenzo[d][1,3]dioxol-5-yl)ethyl, 2-(2,2-difluorobenzo[d][1,3]dioxol-5-yl)ethyl, 2-(2,3-dihydrobenzo[b][1,4]dioxin-6-yl)ethyl, 2-(thiophen-2-yl)ethyl, 2-(thiophen-3-yl)ethyl, 2-(pyridine-2-yl)ethyl, 3-(2-fluorophenyl)propyl, 3-(4-fluorophenyl)propyl, 3-(thiophen-2-yl)propyl, 3-(thiophen-3-yl)propyl, (1-phenylpiperidin-4-yl)methyl, (1-(2-fluorophenyl)piperidin-4-yl)methyl, (1-(3-fluorophenyl)piperidin-4-yl)methyl, (1-(4-fluorophenyl)piperidin-4-yl)methyl, (1-(4-(trifluoromethyl)phenyl)piperidin-4-yl)methyl, (4-methyl-1-phenylpiperidin-4-yl)methyl, (4-fluoro-1-phenylpiperidin-4-yl)methyl, 2-(1-phenylpiperidin-4-yl)ethyl, (1-(2,2,2-trifluoroethyl)piperidin-4-yl)methyl, (1-isobutyrylpiperidin-4-yl)methyl, (1-pivaloylpiperidin-4-yl)methyl, (1-butyrylpiperidin-4-yl)methyl, (1-(3-methylbutanoyl)piperidin-4-yl)methyl, (1-(3,3-dimethylbutanoyl)piperidin-4-yl)methyl, (1-(2-cyclopentylacetyl)piperidin-4-yl)methyl, (1-(cyclopropanecarbonyl)piperidin-4-yl)methyl, (1-(cyclobutanecarbonyl)piperidin-4-yl)methyl, (1-(cyclopentanecarbonyl)piperidin-4-yl)methyl, (1-(cyclohexanecarbonyl)piperidin-4-yl)methyl, (1-((1s,4s)-4-(tert-butyl)cyclohexanecarbonyl)piperidin-4-yl)methyl, (1-((1r,4r)-4-(tert-butyl)cyclohexanecarbonyl)piperidin-4-yl)methyl, (1-(4-methoxycyclohexanecarbonyl)piperidin-4-yl)methyl, (1-(4-(trifluoromethyl)cyclohexanecarbonyl)piperidin-4-yl)methyl, (1-benzoylpiperidin-4-yl)methyl, (1-(3-(trifluoromethyl)benzoyl)piperidin-4-yl)methyl, (1-(2-phenylacetyl)piperidin-4-yl)methyl, (1-(thiophene-3-carbonyl)piperidin-4-yl)methyl, ((5S,8s)-3,3-dimethyl-2-oxaspiro[4.5]decan-8-yl)methyl, (1,2,3,4-tetrahydronaphthalen-2-yl)methyl, (2,3-dihydro-1H-inden-2-yl)methyl, 2,6-difluoro-4-(tetrahydro-2H-pyran-4-yl)phenethyl, (1-(pyridin-3-yl)piperidin-4-yl)methyl, (1-(cyclohexylcarbamoyl)piperidin-4-yl)methyl, (1-(cyclohexylcarbamothioyl)piperidin-4-yl)methyl, (1-((1S,2R)-2-(trifluoromethyl)cyclohexyl)azetidin-3-yl)methyl, ((R)-1-phenylpyrrolidin-3-yl)methyl, ((R)-1-(o-tolyl)pyrrolidin-3-yl)methyl, ((R)-1-(2-(trifluoromethyl)phenyl)pyrrolidin-3-yl)methyl, ((S)-1-(2-(trifluoromethyl)phenyl)pyrrolidin-3-yl)methyl, (R)-1-(2-fluorophenyl)pyrrolidin-3-yl)methyl, (R)-1-(3-fluorophenyl)pyrrolidin-3-yl)methyl, ((R)-1-(2-(trifluoromethoxy)phenyl)pyrrolidin-3-yl)methyl, ((R)-1-(6-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl, ((R)-1-(3-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl, ((R)-1-(4-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl, ((R)-1-(pyridin-3-

yl)pyrrolidin-3-yl)methyl, ((R)-1-(4-methylpyridin-3-yl)pyrrolidin-3-yl)methyl, ((R)-1-(4-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl, ((R)-1-(5-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl, ((R)-1-(2-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl, ((R)-1-(4-(trifluoromethyl)pyrimidin-5-yl)pyrrolidin-3-yl)methyl, ((R)-1-(thiophen-3-yl)pyrrolidin-3-yl)methyl, ((R)-1-(benzo[d]thiazol-4-yl)pyrrolidin-3-yl)methyl, (S)-1-(4-(trifluoromethyl)benzoyl)pyrrolidin-3-yl)methyl, ((R)-1-(o-tolyl)piperidin-3-yl)methyl, ((R)-1-(2-fluorophenyl)piperidin-3-yl)methyl, ((R)-1-(3-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl, ((R)-1-(6-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl, ((R)-1-(4-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl, 3-fluorophenethyl, 4-fluorophenethyl, 3,4-dichlorophenethyl, 3-(trifluoromethyl)phenethyl, 4-(trifluoromethyl)phenethyl, ((R)-1-(benzo[d]thiazol-2-yl)pyrrolidin-3-yl)methyl, ((R)-1-(2-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl, 4-(3,5-dimethylisoxazol-4-yl)-2,6-difluorophenethyl, 4-(3,5-dimethyl-1H-pyrazol-4-yl)-2,6-difluorophenethyl, ((R)-1-(4-(trifluoromethyl)thiazol-2-yl)pyrrolidin-3-yl)methyl, ((R)-1-(benzo[d]oxazol-2-yl)pyrrolidin-3-yl)methyl, ((R)-1-(5-isopropylthiazol-2-yl)piperidin-3-yl)methyl, ((R)-1-(4-(trifluoromethyl)thiazol-2-yl)piperidin-3-yl)methyl, ((R)-1-(benzo[d]thiazol-2-yl)piperidin-3-yl)methyl, ((R)-1-(benzo[d]thiazol-4-yl)piperidin-3-yl)methyl, ((S)-1-(3-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl, ((S)-1-(4-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl, ((S)-1-(4-(trifluoromethyl)pyrimidin-5-yl)pyrrolidin-3-yl)methyl, ((S)-1-(benzo[d]thiazol-4-yl)pyrrolidin-3-yl)methyl, ((S)-1-(3-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl, ((S)-1-(4-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl, ((S)-1-(6-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl, ((S)-1-(benzo[d]thiazol-4-yl)piperidin-3-yl)methyl, ((R)-1-(5-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl, ((R)-1-(6-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl, ((R)-1-(3-(trifluoromethyl)pyridin-4-yl)pyrrolidin-3-yl)methyl, ((R)-1-(2-(trifluoromethyl)pyridin-4-yl)pyrrolidin-3-yl)methyl, ((S)-1-(2-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl, ((S)-1-(5-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl, ((S)-1-(4-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl, ((S)-1-(6-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl, ((S)-1-(5-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl, ((S)-1-(6-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl, ((S)-1-(3-(trifluoromethyl)pyridin-4-yl)pyrrolidin-3-yl)methyl, ((S)-1-(2-(trifluoromethyl)pyridin-4-yl)pyrrolidin-3-yl)methyl, ((R)-1-(5-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl, ((R)-1-(4-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl, ((R)-1-(5-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl, ((R)-1-(6-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl, ((R)-1-(3-(trifluoromethyl)pyridin-

4-yl)piperidin-3-yl)methyl, ((R)-1-(2-(trifluoromethyl)pyridin-4-yl)piperidin-3-yl)methyl, ((S)-1-(2-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl, ((S)-1-(5-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl, ((S)-1-(4-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl, ((S)-1-(5-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl, ((S)-1-(6-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl, ((S)-1-(3-(trifluoromethyl)pyridin-4-yl)piperidin-3-yl)methyl, ((S)-1-(2-(trifluoromethyl)pyridin-4-yl)piperidin-3-yl)methyl, 1-(3-chloro-2-fluorophenyl)propan-2-yl, 4-(3-chloro-2-fluorophenyl)butan-2-yl, ((1r,4R)-4-(difluoromethyl)cyclohexyl)methyl, ((1s,4S)-4-(difluoromethyl)cyclohexyl)methyl, ((1s,4S)-4-(1,1-difluoroethyl)cyclohexyl)methyl, ((1r,4R)-4-(1,1-difluoroethyl)cyclohexyl)methyl, ((1r,4R)-4-(trifluoromethoxy)cyclohexyl)methyl, (4,7-difluoro-2,3-dihydro-1H-inden-2-yl)methyl, (1-(cyclohexanecarbonyl)-4-fluoropiperidin-4-yl)methyl, (4-fluoro-1-(4-fluorophenyl)piperidin-4-yl)methyl, (4-fluoro-1-(2-(trifluoromethyl)phenyl)piperidin-4-yl)methyl, (4-fluoro-1-(3-(trifluoromethyl)pyridin-2-yl)piperidin-4-yl)methyl, (4-fluoro-1-(6-(trifluoromethyl)pyridin-2-yl)piperidin-4-yl)methyl, (1-(benzo[d]thiazol-2-yl)-4-fluoropiperidin-4-yl)methyl, 4-butoxybenzyl, 4-(pentyloxy)benzyl, 4-butoxy-2,6-difluorobenzyl, 4-(cyclopropylmethoxy)benzyl, 4-phenoxybenzyl, [1,1'-biphenyl]-4-ylmethyl, 4-(pyrrolidin-1-yl)benzyl, 4-(piperidin-1-yl)benzyl, or thiophen-2-ylmethyl.

4. The compound of claim 1 wherein the compound is:

(3S,4R,5R)-1-(((1r,4R)-4-(trifluoromethyl)cyclohexyl)methyl)piperidine-3,4,5-triol;
 (3S,4R,5R)-1-(((1s,4S)-4-(trifluoromethyl)cyclohexyl)methyl)piperidine-3,4,5-triol;
 (3S,4R,5R)-1-(((1s,4S)-4-(2-fluoropropan-2-yl)cyclohexyl)methyl)piperidine-3,4,5-triol;
 (3S,4r,5R)-1-(2-cyclohexylethyl)piperidine-3,4,5-triol;
 (3S,4r,5R)-1-(3-cyclohexylpropyl)piperidine-3,4,5-triol;
 (3S,4r,5R)-1-(3-chloro-2-fluorophenethyl)piperidine-3,4,5-triol;
 (3S,4r,5R)-1-(3-chloro-2,6-difluorophenethyl)piperidine-3,4,5-triol;
 (3S,4r,5R)-1-(3,4-dichlorophenethyl)piperidine-3,4,5-triol;
 (3S,4r,5R)-1-(4-(cyclopropylmethoxy)-2,6-difluorophenethyl)piperidine-3,4,5-triol;
 (3S,4r,5R)-1-(2,6-difluoro-4-(pyrrolidin-1-yl)phenethyl)piperidine-3,4,5-triol;
 (3S,4r,5R)-1-(2,6-difluoro-4-(piperidin-1-yl)phenethyl)piperidine-3,4,5-triol;
 (3S,4r,5R)-1-(2,6-difluoro-4-morpholinophenethyl)piperidine-3,4,5-triol;
 (3S,4r,5R)-1-(2,6-difluoro-4-(tetrahydro-2H-pyran-4-yl)phenethyl)piperidine-3,4,5-triol;
 (3S,4r,5R)-1-(4-(3,5-dimethylisoxazol-4-yl)-2,6-difluorophenethyl)piperidine-3,4,5-triol;
 (3S,4r,5R)-1-(4-(3,5-dimethyl-1H-pyrazol-4-yl)-2,6-difluorophenethyl)piperidine-3,4,5-triol;

(3S,4r,5R)-1-(1-(3-chloro-2-fluorophenyl)propan-2-yl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-(3-chloro-2-fluorophenyl)butan-2-yl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(o-tolyl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(3-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(4-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(4-(trifluoromethyl)thiazol-2-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(benzo[d]oxazol-2-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(benzo[d]thiazol-2-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(3-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(4-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(4-(trifluoromethyl)pyrimidin-5-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(4-(trifluoromethyl)thiazol-2-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(benzo[d]thiazol-4-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(3-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(6-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(2-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(5-isopropylthiazol-2-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(4-(trifluoromethyl)thiazol-2-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(benzo[d]thiazol-2-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(benzo[d]thiazol-4-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(3-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4S,5R)-1-(((S)-1-(6-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(4-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(4-(trifluoromethyl)thiazol-2-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(cyclohexylmethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((4,4-dimethylcyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((4,4-difluorocyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((4,4-dichlorocyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((4-ethylcyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1s,4S)-4-vinylcyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1s,4S)-4-isopropylcyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1r,4R)-4-isopropylcyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1s,4S)-4-(tert-butyl)cyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1r,4R)-4-(tert-butyl)cyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1r,4R)-4-(2-fluoropropan-2-yl)cyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1s,4S)-4-methoxycyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1r,4R)-4-methoxycyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((4-(methoxymethyl)cyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1s,4S)-4-cyclopropylcyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1r,4R)-4-cyclopropylcyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((4-phenylcyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(spiro[2.5]octan-6-ylmethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(spiro[3.5]nonan-7-ylmethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(spiro[4.5]decan-8-ylmethyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((5S,8s)-3,3-dimethyl-2-oxaspiro[4.5]decan-8-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((1,2,3,4-tetrahydronaphthalen-2-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((2,3-dihydro-1H-inden-2-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-(4,4-difluorocyclohexyl)ethyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(2-((1s,4S)-4-(trifluoromethyl)cyclohexyl)ethyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(2-((1r,4R)-4-(trifluoromethyl)cyclohexyl)ethyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(2-((3R,5R,7R)-adamantan-1-yl)ethyl)piperidine-3,4,5-triol;

(3S,4r,5R)-1-phenethylpiperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-methylphenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-methoxyphenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-fluorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-chlorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2,3-difluorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2,4-difluorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2,5-difluorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(3,4-difluorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-fluoro-4-methoxyphenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-chloro-2-fluorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(5-chloro-2-fluorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2,6-difluorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2,6-difluoro-4-(prop-1-en-2-yl)phenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2,6-difluoro-4-isopropylphenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2,6-difluoro-3-isopropylphenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-cyclopropyl-2,6-difluorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2,6-difluoro-4-(trifluoromethyl)phenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-butoxy-2,6-difluorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-((tetrahydrofuran-3-yl)oxy)phenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-((tetrahydro-2H-pyran-3-yl)oxy)phenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-((tetrahydro-2H-pyran-4-yl)oxy)phenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-phenoxyphenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-((tetrahydrofuran-3-yl)methoxy)phenethyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-((R)-2-phenylpropyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-((S)-2-phenylpropyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-([1,1'-biphenyl]-4-yl)ethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-(3,5-difluoro-[1,1'-biphenyl]-4-yl)ethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-(benzo[d][1,3]dioxol-5-yl)ethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-(6-fluorobenzo[d][1,3]dioxol-5-yl)ethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-(2,2-difluorobenzo[d][1,3]dioxol-5-yl)ethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-(2,3-dihydrobenzo[b][1,4]dioxin-6-yl)ethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-(thiophen-2-yl)ethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-(thiophen-3-yl)ethyl)piperidine-3,4,5-triol;

(3S,4r,5R)-1-(2-(pyridin-2-yl)ethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(3-(2-fluorophenyl)propyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(3-(4-fluorophenyl)propyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(3-(thiophen-2-yl)propyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(3-(thiophen-3-yl)propyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((1-phenylpiperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((1-(2-fluorophenyl)piperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((1-(3-fluorophenyl)piperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((1-(4-fluorophenyl)piperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((1-(4-(trifluoromethyl)phenyl)piperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((4-methyl-1-phenylpiperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((4-fluoro-1-phenylpiperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(2-(1-phenylpiperidin-4-yl)ethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((1-(pyridin-3-yl)piperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((1-(2,2,2-trifluoroethyl)piperidin-4-yl)methyl)piperidine-3,4,5-triol;
2-methyl-1-(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)propan-1-one;
2,2-dimethyl-1-(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)propan-1-one;
1-(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)butan-1-one;
3-methyl-1-(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)butan-1-one;
3,3-dimethyl-1-(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)butan-1-one;
2-cyclopentyl-1-(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)ethanone;
cyclopropyl(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)methanone;
cyclobutyl(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)methanone;
cyclopentyl(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)methanone;
cyclohexyl(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)methanone;
((1s,4S)-4-(tert-butyl)cyclohexyl)(4-(((3S,4R,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)methanone;
((1r,4R)-4-(tert-butyl)cyclohexyl)(4-(((3S,4R,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)methanone;

(4-methoxycyclohexyl)(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)methanone;

(4-(trifluoromethyl)cyclohexyl)(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)methanone;

phenyl(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)methanone;

(3-(trifluoromethyl)phenyl)(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)methanone;

2-phenyl-1-(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)ethanone;

thiophen-3-yl(4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)methanone;

N-cyclohexyl-4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidine-1-carboxamide;

N-cyclohexyl-4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidine-1-carbothioamide;

(3S,4S,5R)-1-((1-((1S,2R)-2-(trifluoromethyl)cyclohexyl)azetidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((R)-1-phenylpyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((R)-1-(2-(trifluoromethyl)phenyl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((R)-1-(2-fluorophenyl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((R)-1-(3-fluorophenyl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((R)-1-(2-(trifluoromethoxy)phenyl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((R)-1-(6-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((R)-1-(4-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((R)-1-(pyridin-3-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((R)-1-(4-methylpyridin-3-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((R)-1-(5-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((R)-1-(2-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((R)-1-(5-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(6-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(3-(trifluoromethyl)pyridin-4-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(2-(trifluoromethyl)pyridin-4-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(4-(trifluoromethyl)pyrimidin-5-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3R,4R,5S)-1-(((R)-1-(thiophen-3-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(benzo[d]thiazol-4-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(4-(trifluoromethyl)phenyl)((R)-3-(((3S,4R,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)pyrrolidin-1-yl)methanone;
(3S,4S,5R)-1-(((S)-1-(2-(trifluoromethyl)phenyl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(6-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(2-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(5-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(4-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(5-(trifluoromethyl)pyridin-2-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(6-(trifluoromethyl)pyridin-3-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(3-(trifluoromethyl)pyridin-4-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(2-(trifluoromethyl)pyridin-4-yl)pyrrolidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(o-tolyl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(2-fluorophenyl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;

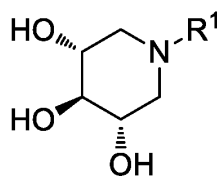
(3S,4R,5R)-1-(((R)-1-(4-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(5-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(4-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(5-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(6-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(3-(trifluoromethyl)pyridin-4-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((R)-1-(2-(trifluoromethyl)pyridin-4-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(2-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(5-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(4-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(5-(trifluoromethyl)pyridin-2-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(6-(trifluoromethyl)pyridin-3-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(3-(trifluoromethyl)pyridin-4-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4S,5R)-1-(((S)-1-(2-(trifluoromethyl)pyridin-4-yl)piperidin-3-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(3-fluorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-fluorophenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(3-(trifluoromethyl)phenethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-(trifluoromethyl)phenethyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1r,4R)-4-(difluoromethyl)cyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1s,4S)-4-(difluoromethyl)cyclohexyl)methyl)piperidine-3,4,5-triol;

(3S,4R,5R)-1-(((1s,4S)-4-(1,1-difluoroethyl)cyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1r,4R)-4-(1,1-difluoroethyl)cyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4R,5R)-1-(((1r,4R)-4-(trifluoromethoxy)cyclohexyl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((4,7-difluoro-2,3-dihydro-1H-inden-2-yl)methyl)piperidine-3,4,5-triol;
cyclohexyl(4-fluoro-4-(((3S,4r,5R)-3,4,5-trihydroxypiperidin-1-yl)methyl)piperidin-1-yl)methanone;
(3S,4r,5R)-1-((4-fluoro-1-(4-fluorophenyl)piperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((4-fluoro-1-(2-(trifluoromethyl)phenyl)piperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((4-fluoro-1-(3-(trifluoromethyl)pyridin-2-yl)piperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((4-fluoro-1-(6-(trifluoromethyl)pyridin-2-yl)piperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-((1-(benzo[d]thiazol-2-yl)-4-fluoropiperidin-4-yl)methyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-butoxybenzyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-(pentyloxy)benzyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-butoxy-2,6-difluorobenzyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-(cyclopropylmethoxy)benzyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-phenoxybenzyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-([1,1'-biphenyl]-4-ylmethyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-(pyrrolidin-1-yl)benzyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(4-(piperidin-1-yl)benzyl)piperidine-3,4,5-triol;
(3S,4r,5R)-1-(thiophen-2-ylmethyl)piperidine-3,4,5-triol;
or a pharmaceutically acceptable salt of any of the foregoing compounds.

5. The compound of claim 1 wherein the compound is a prodrug.
6. The compound of any one of claims 1 to 5 wherein the compound inhibits a non-lysosomal glucosylceramidase (GBA2).
7. The compound of any one of claims 1 to 6 wherein the compound specifically binds a GBA2.
8. The compound of any one of claims 1 to 7 wherein the compound decreases the enzyme activity levels of a GBA2.
9. The compound of any one of claims 6 to 8 wherein the GBA2 is a mammalian GBA2.

10. A pharmaceutical composition comprising the compound of any one of claims 1 to 9 or a pharmaceutically acceptable salt thereof in combination with a pharmaceutically acceptable carrier.

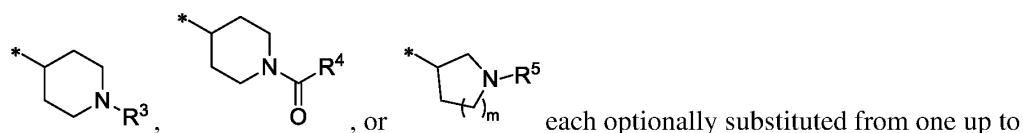
11. A method of inhibiting a GBA2 in a subject in need thereof, the method comprising administering to the subject an effective amount of a compound of Formula (I) or a pharmaceutically acceptable salt thereof:



(I)

wherein

R^1 is $(CH_2)_nR^2$, wherein n is 1, 2, or 3, and R^2 is cyclohexyl, phenyl, thiophen-2-yl, thiophen-3-yl, pyridine-2-yl, adamantyl, 2,3-dihydro-1H-inden-2-yl, 1,2,3,4-tetrahydronaphthalen-2-yl, spiro[2.5]octan-6-yl, spiro[3.5]nonan-7-yl, spiro[4.5]decan-8-yl, (5S,8s)-3,3-dimethyl-2-oxaspiro[4.5]decan-8-yl, (benzo[d][1,3]dioxol-5-yl)methyl, (2,3-dihydrobenzo[b][1,4]dioxin-6-yl)methyl, 1-cyclohexylazetidin-3-yl,



each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C_{1-6} alkyl, C_{2-6} alkenyl, cyclopropyl, difluoromethyl, 1,1-difluoroethyl-1-yl, 2-fluoropropan-2-yl, methoxymethyl, OCF_3 , CF_3 , phenyl, pyrrolidin-1-yl, piperidin-1-yl, 4-morpholino, C_{1-6} alkoxy, cyclopropylmethoxy, (tetrahydrofuran-3-yl)oxy, (tetrahydro-2H-pyran-3-yl)oxy, (tetrahydro-2H-pyran-4-yl)oxy, phenoxy, (tetrahydrofuran-3-yl)methoxy, tetrahydro-2H-pyran-4-yl, 3,5-dimethylisoxazol-4-yl, and/or 3,5-dimethyl-1H-pyrazol-4-yl;

where R^3 is C_{1-6} alkyl, phenyl, pyridine-2-yl, pyridin-3-yl, benzo[d]thiazol-2-yl, cyclohexylcarbamothioyl, or cyclohexylcarbamoyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C_{1-6} alkyl, C_{2-6} alkenyl, cyclopropyl, C_{1-6} alkoxy, OCF_3 , and/or CF_3 ; and

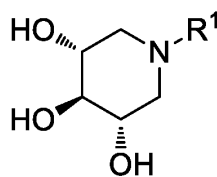
where R⁴ is C₁₋₆ alkyl, C₃₋₇ cycloalkyl, phenyl, thiophen-3-yl, phenylmethyl, or cyclopentylmethyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, C₁₋₆ alkyl, OCH₃, and/or CF₃; and

where m is 1 or 2; and

where R⁵ is phenyl, pyridin-2-yl, pyridin-3-yl, pyridin-4-yl, pyrimidin-5-yl, thiophen-3-yl, benzo[d]thiazol-4-yl, benzo[d]thiazol-2-yl, phenylcarbonyl, thiazol-2-yl, benzo[d]oxazol-2-yl, or benzo[d]thiazol-2-yl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C₁₋₆ alkyl, C₁₋₆ alkoxy, OCF₃, and/or CF₃,

with the proviso that R¹ is not benzyl or 3-phenylpropyl.

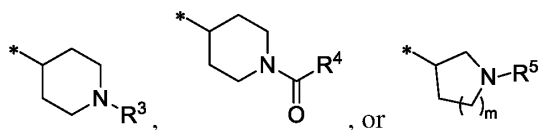
12. A method of reducing the GBA2 enzyme activity in a subject in need thereof, the method comprising administering to the subject an effective amount of a compound of Formula (I) or a pharmaceutically acceptable salt thereof:



(I)

wherein

R¹ is (CH₂)_nR², wherein n is 1, 2, or 3, and R² is cyclohexyl, phenyl, thiophen-2-yl, thiophen-3-yl, pyridine-2-yl, adamantyl, 2,3-dihydro-1H-inden-2-yl, 1,2,3,4-tetrahydronaphthalen-2-yl, spiro[2.5]octan-6-yl, spiro[3.5]nonan-7-yl, spiro[4.5]decan-8-yl, (5S,8s)-3,3-dimethyl-2-oxaspiro[4.5]decan-8-yl, (benzo[d][1,3]dioxol-5-yl)methyl, (2,3-dihydrobenzo[b][1,4]dioxin-6-yl)methyl, 1-cyclohexylazetidin-3-yl,



N^{R^3} , O^{R^3} , or $\text{C}^{\text{R}^3}_m$ each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C₁₋₆ alkyl, C₂₋₆ alkenyl, cyclopropyl, difluoromethyl, 1,1-difluoroethyl-1-yl, 2-fluoropropan-2-yl, methoxymethyl, OCF₃, CF₃, phenyl, pyrrolidin-1-yl, piperidin-1-yl, 4-morpholino, C₁₋₆ alkoxy, cyclopropylmethoxy, (tetrahydrofuran-3-yl)oxy, (tetrahydro-2H-pyran-3-yl)oxy, (tetrahydro-

2H-pyran-4-yl)oxy, phenoxy, (tetrahydrofuran-3-yl)methoxy, tetrahydro-2H-pyran-4-yl, 3,5-dimethylisoxazol-4-yl, and/or 3,5-dimethyl-1H-pyrazol-4-yl;

where R^3 is C_{1-6} alkyl, phenyl, pyridine-2-yl, pyridin-3-yl, benzo[d]thiazol-2-yl, cyclohexylcarbamothioyl, or cyclohexylcarbamoyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C_{1-6} alkyl, C_{2-6} alkenyl, cyclopropyl, C_{1-6} alkoxy, OCF_3 , and/or CF_3 ; and

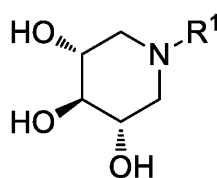
where R^4 is C_{1-6} alkyl, C_{3-7} cycloalkyl, phenyl, thiophen-3-yl, phenylmethyl, or cyclopentylmethyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, C_{1-6} alkyl, OCH_3 , and/or CF_3 ; and

where m is 1 or 2; and

where R^5 is phenyl, pyridin-2-yl, pyridin-3-yl, pyridin-4-yl, pyrimidin-5-yl, thiophen-3-yl, benzo[d]thiazol-4-yl, benzo[d]thiazol-2-yl, phenylcarbonyl, thiazol-2-yl, benzo[d]oxazol-2-yl, or benzo[d]thiazol-2-yl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C_{1-6} alkyl, C_{1-6} alkoxy, OCF_3 , and/or CF_3 ,

with the proviso that R^1 is not benzyl or 3-phenylpropyl.

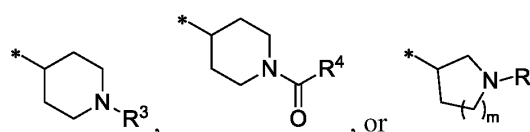
13. A method of treating a condition that is modulated by a GBA2, in a subject in need thereof, the method comprising administering to the subject an effective amount of a compound of Formula (I) or a pharmaceutically acceptable salt thereof:



(I)

wherein

R^1 is $(CH_2)_nR^2$, wherein n is 1, 2, or 3, and R^2 is cyclohexyl, phenyl, thiophen-2-yl, thiophen-3-yl, pyridine-2-yl, adamantyl, 2,3-dihydro-1H-inden-2-yl, 1,2,3,4-tetrahydronaphthalen-2-yl, spiro[2.5]octan-6-yl, spiro[3.5]nonan-7-yl, spiro[4.5]decan-8-yl, (5S,8s)-3,3-dimethyl-2-oxaspiro[4.5]decan-8-yl, (benzo[d][1,3]dioxol-5-yl)methyl, (2,3-dihydrobenzo[b][1,4]dioxin-6-yl)methyl, 1-cyclohexylazetidin-3-yl,


 each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C₁₋₆ alkyl, C₂₋₆ alkenyl, cyclopropyl, difluoromethyl, 1,1-difluoroethyl-1-yl, 2-fluoropropan-2-yl, methoxymethyl, OCF₃, CF₃, phenyl, pyrrolidin-1-yl, piperidin-1-yl, 4-morpholino, C₁₋₆ alkoxy, cyclopropylmethoxy, (tetrahydrofuran-3-yl)oxy, (tetrahydro-2H-pyran-3-yl)oxy, (tetrahydro-2H-pyran-4-yl)oxy, phenoxy, (tetrahydrofuran-3-yl)methoxy, tetrahydro-2H-pyran-4-yl, 3,5-dimethylisoxazol-4-yl, and/or 3,5-dimethyl-1H-pyrazol-4-yl;

where R³ is C₁₋₆ alkyl, phenyl, pyridine-2-yl, pyridin-3-yl, benzo[d]thiazol-2-yl, cyclohexylcarbamothioyl, or cyclohexylcarbamoyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C₁₋₆ alkyl, C₂₋₆ alkenyl, cyclopropyl, C₁₋₆ alkoxy, OCF₃, and/or CF₃; and

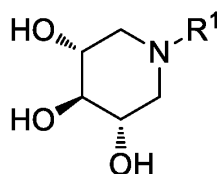
where R⁴ is C₁₋₆ alkyl, C₃₋₇ cycloalkyl, phenyl, thiophen-3-yl, phenylmethyl, or cyclopentylmethyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, C₁₋₆ alkyl, OCH₃, and/or CF₃; and

where m is 1 or 2; and

where R⁵ is phenyl, pyridin-2-yl, pyridin-3-yl, pyridin-4-yl, pyrimidin-5-yl, thiophen-3-yl, benzo[d]thiazol-4-yl, benzo[d]thiazol-2-yl, phenylcarbonyl, thiazol-2-yl, benzo[d]oxazol-2-yl, or benzo[d]thiazol-2-yl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C₁₋₆ alkyl, C₁₋₆ alkoxy, OCF₃, and/or CF₃,

with the proviso that R¹ is not benzyl or 3-phenylpropyl.

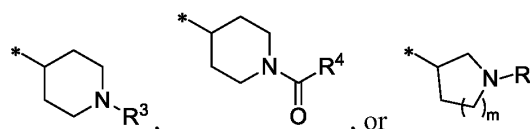
14. A method of treating a condition selected from a neurological disease, a lysosomal storage disease, and a liver disease, in a subject in need thereof, the method comprising administering to the subject an effective amount of a compound of Formula (I) or a pharmaceutically acceptable salt thereof:



(I)

wherein

R^1 is $(CH_2)_nR^2$, wherein n is 1, 2, or 3, and R^2 is cyclohexyl, phenyl, thiophen-2-yl, thiophen-3-yl, pyridine-2-yl, adamantyl, 2,3-dihydro-1H-inden-2-yl, 1,2,3,4-tetrahydronaphthalen-2-yl, spiro[2.5]octan-6-yl, spiro[3.5]nonan-7-yl, spiro[4.5]decan-8-yl, (5S,8s)-3,3-dimethyl-2-oxaspiro[4.5]decan-8-yl, (benzo[d][1,3]dioxol-5-yl)methyl, (2,3-dihydrobenzo[b][1,4]dioxin-6-yl)methyl, 1-cyclohexylazetidin-3-yl,

 each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C_{1-6} alkyl, C_{2-6} alkenyl, cyclopropyl, difluoromethyl, 1,1-difluoroethyl-1-yl, 2-fluoropropan-2-yl, methoxymethyl, OCF_3 , CF_3 , phenyl, pyrrolidin-1-yl, piperidin-1-yl, 4-morpholino, C_{1-6} alkoxy, cyclopropylmethoxy, (tetrahydrofuran-3-yl)oxy, (tetrahydro-2H-pyran-3-yl)oxy, (tetrahydro-2H-pyran-4-yl)oxy, phenoxy, (tetrahydrofuran-3-yl)methoxy, tetrahydro-2H-pyran-4-yl, 3,5-dimethylisoxazol-4-yl, and/or 3,5-dimethyl-1H-pyrazol-4-yl;

where R^3 is C_{1-6} alkyl, phenyl, pyridine-2-yl, pyridin-3-yl, benzo[d]thiazol-2-yl, cyclohexylcarbamothioyl, or cyclohexylcarbamoyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C_{1-6} alkyl, C_{2-6} alkenyl, cyclopropyl, C_{1-6} alkoxy, OCF_3 , and/or CF_3 ; and

where R^4 is C_{1-6} alkyl, C_{3-7} cycloalkyl, phenyl, thiophen-3-yl, phenylmethyl, or cyclopentylmethyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, C_{1-6} alkyl, OCH_3 , and/or CF_3 ; and

where m is 1 or 2; and

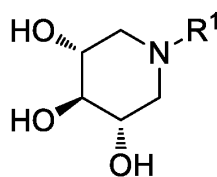
where R^5 is phenyl, pyridin-2-yl, pyridin-3-yl, pyridin-4-yl, pyrimidin-5-yl, thiophen-3-yl, benzo[d]thiazol-4-yl, benzo[d]thiazol-2-yl, phenylcarbonyl, thiazol-2-yl, benzo[d]oxazol-2-yl, or benzo[d]thiazol-2-yl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C_{1-6} alkyl, C_{1-6} alkoxy, OCF_3 , and/or CF_3 ,

with the proviso that R^1 is not benzyl or 3-phenylpropyl.

15. The method of claim 13 or 14 wherein the condition is Alzheimer's disease, Parkinson's disease, multiple sclerosis, Huntington's disease, amyotrophic lateral sclerosis (ALS), amyotrophic lateral sclerosis with cognitive impairment (ALSci), addiction, anxiety, argyrophilic grain dementia, ataxia-telangiectasia (A-T), attention deficit/hyperactivity disorder (ADHD), autism spectrum disorder (ASD), Becker muscular dystrophy (BMD), bipolar disorder (BD), Bluit disease, cerebellar ataxia, Charcot-Marie-Tooth disease (CMT), chronic fatigue syndrome, corticobasal degeneration (CBD), dementia pugilistica, dementia with Lewy bodies (DLB), Dejerine-Sottas disease, diffuse neurofibrillary tangles with calcification, Down's syndrome, Duchenne muscular dystrophy (DMD), epilepsy, essential tremor (ET), familial British dementia, familial Danish dementia, fibromyalgia, frontotemporal dementia with parkinsonism linked to chromosome 17 (FTDP-17), Friedreich's ataxia, Gerstmann-Straussler-Scheinker disease, glaucoma, Guadeloupean parkinsonism, Guillain-Barré syndrome, Hallevorden-Spatz disease (neurodegeneration with brain iron accumulation type 1), insomnia, Lambert-Eaton myasthenic syndrome (LEMS), major depressive disorder (MDD), migraine, mild cognitive impairment (MCI), multi-infarct dementia, multiple system atrophy (MSA), myasthenia gravis, myotonic dystrophy (including types DM1 and DM2), neuronal ceroid lipofuscinosis (including types 1, 2, 3, 4, 5, 6, 7, 8, 9, and 10), neuropathy (including peripheral neuropathy, autonomic neuropathy, neuritis, and diabetic neuropathy), oculopharyngeal muscular dystrophy, pain, pallido-ponto-nigral degeneration, parkinsonism-dementia complex of Guam, Pick's disease (PiD), post-encephalitic parkinsonism (PEP), primary lateral sclerosis (PLS), prion diseases (including Creutzfeldt-Jakob Disease (CJD), variant Creutzfeldt-Jakob Disease (vCJD), fatal familial insomnia, and kuru), progressive superecortical gliosis, progressive supranuclear palsy (PSP), Richardson's syndrome, schizophrenia, seizures, spinal cord injury, spinal muscular atrophy (SMA), spinocerebellar ataxia (including types 1, 2, 3, 4, 5, 6, 7, 8, 10, 11, 12, 13, 14, 16, 17, 18, 19, 20, 21, 22, 23, 25, 26, 27, 28, and 29), stroke, subacute sclerosing panencephalitis, tangle-only dementia, tardive dyskinesia, Tourette syndrome (TS), vascular dementia, Wilson's disease, Gaucher disease (including types I, II, and III), Niemann-Pick disease (including types A, B, and C), mucopolipidosis (including types I, II, III, IV, VI, and VII), cerebrotendineous xanthomatosis, Fabry disease, Farber disease, GM1 gangliosidosis, Krabbe disease, metachromatic leukodystrophy (MLD), multiple sulfatase deficiency, Pompe disease, Sandhoff disease, Tay-Sach's disease, non-alcoholic fatty liver disease (NAFLD), non-alcoholic steatohepatitis (NASH), Alagille syndrome, alcohol-related liver disease, alpha-1 antitrypsin deficiency, autoimmune hepatitis, autoimmune cholangitis, benign liver tumors,

biliary atresia, cirrhosis, Crigler-Najjar syndrome, drug-induced liver injury (DILI), galactosemia, Gilbert syndrome, hemochromatosis, hepatic encephalopathy, hepatocellular carcinoma (HCC), intrahepatic cholestasis of pregnancy (ICP), lysosomal acid lipase deficiency (LAL-D), liver cysts, liver cancer, newborn jaundice, primary biliary cholangitis (PBC), primary sclerosing cholangitis (PSC), Reye syndrome, type I glycogen storage disease, or viral hepatitis (including types A, B, C, D, and E).

16. The method of claim 13 or 14 wherein the condition is Parkinson's disease.
17. The method of claim 13 or 14 wherein the condition is Alzheimer's disease, Huntington's disease, amyotrophic lateral sclerosis (ALS), or multiple sclerosis.
18. The method of claim 13 or 14 wherein the condition is Niemann-Pick type C disease.
19. The method of claim 13 or 14 wherein the condition is Gaucher disease, mucopolipidosis type IV, neuronal ceroid lipofuscinosis, or Sandhoff disease.
20. The method of claim 13 or 14 wherein the condition is non-alcoholic steatohepatitis (NASH).
21. The method of any one of claims 11 to 20 wherein the compound is one or more of the compounds described in Table 1.
22. The method of any one of claims 11 to 21 wherein said administering decreases the level of GBA2 enzyme activity in the subject.
23. The method of any one of claims 11 to 22 wherein the subject is a human.
24. Use of an effective amount of a compound of Formula (I) or a pharmaceutically acceptable salt thereof:

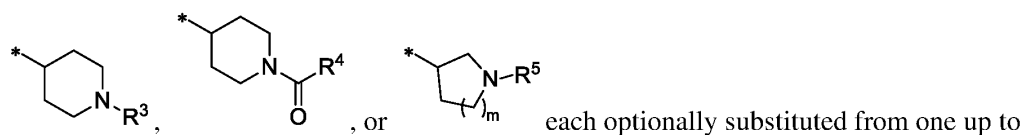


(I)

wherein

R^1 is $(CH_2)_nR^2$, wherein n is 1, 2, or 3, and R^2 is cyclohexyl, phenyl, thiophen-2-yl, thiophen-3-yl, pyridine-2-yl, adamantyl, 2,3-dihydro-1H-inden-2-yl, 1,2,3,4-tetrahydronaphthalen-2-yl, spiro[2.5]octan-6-yl, spiro[3.5]nonan-7-yl, spiro[4.5]decan-8-yl,

(5S,8s)-3,3-dimethyl-2-oxaspiro[4.5]decan-8-yl, (benzo[d][1,3]dioxol-5-yl)methyl, (2,3-dihydrobenzo[b][1,4]dioxin-6-yl)methyl, 1-cyclohexylazetidin-3-yl,



each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C₁₋₆ alkyl, C₂₋₆ alkenyl, cyclopropyl, difluoromethyl, 1,1-difluoroethyl-1-yl, 2-fluoropropan-2-yl, methoxymethyl, OCF₃, CF₃, phenyl, pyrrolidin-1-yl, piperidin-1-yl, 4-morpholino, C₁₋₆ alkoxy, cyclopropylmethoxy, (tetrahydrofuran-3-yl)oxy, (tetrahydro-2H-pyran-3-yl)oxy, (tetrahydro-2H-pyran-4-yl)oxy, phenoxy, (tetrahydrofuran-3-yl)methoxy, tetrahydro-2H-pyran-4-yl, 3,5-dimethylisoxazol-4-yl, and/or 3,5-dimethyl-1H-pyrazol-4-yl;

where R³ is C₁₋₆ alkyl, phenyl, pyridine-2-yl, pyridin-3-yl, benzo[d]thiazol-2-yl, cyclohexylcarbamothioyl, or cyclohexylcarbamoyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C₁₋₆ alkyl, C₂₋₆ alkenyl, cyclopropyl, C₁₋₆ alkoxy, OCF₃, and/or CF₃; and

where R⁴ is C₁₋₆ alkyl, C₃₋₇ cycloalkyl, phenyl, thiophen-3-yl, phenylmethyl, or cyclopentylmethyl, each optionally substituted from one up to the maximum number of substituents with one or more of F, C₁₋₆ alkyl, OCH₃, and/or CF₃; and

where m is 1 or 2; and

where R⁵ is phenyl, pyridin-2-yl, pyridin-3-yl, pyridin-4-yl, pyrimidin-5-yl, thiophen-3-yl, benzo[d]thiazol-4-yl, benzo[d]thiazol-2-yl, phenylcarbonyl, thiazol-2-yl, benzo[d]oxazol-2-yl, or benzo[d]thiazol-2-yl, each optionally substituted from one up to the maximum number of substituents with one or more of F, Cl, C₁₋₆ alkyl, C₁₋₆ alkoxy, OCF₃, and/or CF₃,

with the proviso that R¹ is not benzyl or 3-phenylpropyl, in the preparation of a medicament.

25. The use of claim 24 wherein said medicament is for inhibiting a GBA2, for decreasing the level of GBA2 enzyme activity, for treating a condition modulated by a GBA2, for treating a neurological disease, for treating a lysosomal storage disease, or for treating a liver disease.