

BENZYLAMINE DERIVATIVES AS INHIBITORS OF PLASMA KALLIKREIN

This invention relates to benzylamine derivatives and to pharmaceutical compositions containing and the uses of, such derivatives.

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Background to the Invention

The benzylamine derivatives of the present invention are inhibitors of plasma kallikrein and have a number of therapeutic applications, particularly in the treatment of retinal vascular permeability associated with diabetic retinopathy and diabetic macular oedema.

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Plasma kallikrein is a trypsin-like serine protease that can liberate kinins from kininogens (see K. D. Bhoola *et al.*, "Kallikrein-Kinin Cascade", *Encyclopedia of Respiratory Medicine*, p483-493; J. W. Bryant *et al.*, "Human plasma kallikrein-kinin system: physiological and biochemical parameters" *Cardiovascular and haematological agents in medicinal chemistry*, 7, p234-250, 2009; K. D. Bhoola *et al.*, *Pharmacological Rev.*, 1992, **44**, 1; and D. J. Campbell, "Towards understanding the kallikrein-kinin system: insights from the measurement of kinin peptides", *Brazilian Journal of Medical and Biological Research* 2000, **33**, 665-677). It is an essential member of the intrinsic blood coagulation cascade although its role in this cascade does not involve the release of bradykinin or enzymatic cleavage. Plasma prekallikrein is encoded by a single gene and synthesized in the liver. It is secreted by hepatocytes as an inactive plasma prekallikrein that circulates in plasma as a heterodimer complex bound to high molecular weight kininogen which is activated to give the active plasma kallikrein. Kinins are potent mediators of inflammation that act through G protein-coupled receptors and antagonists of kinins (such as bradykinin antagonists) have previously been investigated as potential therapeutic agents for the treatment of a number of disorders (F. Marceau and D. Regoli, *Nature Rev., Drug Discovery*, 2004, **3**, 845-852).

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Plasma kallikrein is thought to play a role in a number of inflammatory disorders. The major inhibitor of plasma kallikrein is the serpin C1 esterase inhibitor. Patients who present with a genetic deficiency in C1 esterase inhibitor suffer from hereditary angioedema (HAE) which results in intermittent swelling of face, hands, throat, gastro-intestinal tract and genitals. Blisters formed during acute episodes contain high levels of plasma kallikrein which cleaves high

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molecular weight kininogen liberating bradykinin leading to increased vascular permeability. Treatment with a large protein plasma kallikrein inhibitor has been shown to effectively treat HAE by preventing the release of bradykinin which causes increased vascular permeability (A. Lehmann "Ecallantide (DX-88), a plasma kallikrein inhibitor for the treatment of hereditary angioedema and the prevention of blood loss in on-pump cardiothoracic surgery" *Expert Opin. Biol. Ther.* 8, p1187-99).

The plasma kallikrein-kinin system is abnormally abundant in patients with advanced diabetic macular oedema. It has been recently published that plasma kallikrein contributes to retinal vascular dysfunctions in diabetic rats (A. Clermont *et al.* "Plasma kallikrein mediates retinal vascular dysfunction and induces retinal thickening in diabetic rats" *Diabetes*, 2011, 60, p1590-98). Furthermore, administration of the plasma kallikrein inhibitor ASP-440 ameliorated both retinal vascular permeability and retinal blood flow abnormalities in diabetic rats. Therefore a plasma kallikrein inhibitor should have utility as a treatment to reduce retinal vascular permeability associated with diabetic retinopathy and diabetic macular oedema.

Synthetic and small molecule plasma kallikrein inhibitors have been described previously, for example by Garrett *et al.* ("Peptide aldehyde...." *J. Peptide Res.* 52, p62-71 (1998)), T. Griesbacher *et al.* ("Involvement of tissue kallikrein but not plasma kallikrein in the development of symptoms mediated by endogenous kinins in acute pancreatitis in rats" *British Journal of Pharmacology* 137, p692-700 (2002)), Evans ("Selective dipeptide inhibitors of kallikrein" WO03/076458), Szelke *et al.* ("Kininogenase inhibitors" WO92/04371), D. M. Evans *et al.* (*Immunopharmacology*, 32, p115-116 (1996)), Szelke *et al.* ("Kininogen inhibitors" WO95/07921), Antonsson *et al.* ("New peptides derivatives" WO94/29335), J. Stürzbecher *et al.* (*Brazilian J. Med. Biol. Res* 27, p1929-34 (1994)), Kettner *et al.* (US 5,187,157), N. Teno *et al.* (*Chem. Pharm. Bull.* 41, p1079-1090 (1993)), W. B. Young *et al.* ("Small molecule inhibitors of plasma kallikrein" *Bioorg. Med. Chem. Letts.* 16, p2034-2036 (2006)), Okada *et al.* ("Development of potent and selective plasmin and plasma kallikrein inhibitors and studies on the structure-activity relationship" *Chem. Pharm. Bull.* 48, p1964-72 (2000)), Steinmetzer *et al.* ("Trypsin-like serine protease inhibitors and their preparation and use" WO08/049595), Zhang *et al.* ("Discovery of highly potent small molecule kallikrein inhibitors" *Medicinal Chemistry* 2, p545-553 (2006)), Sinha *et al.* ("Inhibitors of plasma kallikrein" WO08/016883), and Brandl *et al.* ("N-((6-amino-pyridin-3-yl)methyl)-heteroaryl-carboxamides as inhibitors of plasma

kallikrein” WO2012/017020). Also, Steinmetzer *et al.* (“Serine protease inhibitors” WO2012/004678) describes cyclized peptide analogs which are inhibitors of human plasmin and plasma kallikrein.

- 5 To date, no small molecule synthetic plasma kallikrein inhibitor has been approved for medical use. The molecules described in the known art suffer from limitations such as poor selectivity over related enzymes such as KLK1, thrombin and other serine proteases, and poor oral availability. The large protein plasma kallikrein inhibitors present risks of anaphylactic reactions, as has been reported for Ecallantide. Thus there remains a need for compounds that selectively
- 10 inhibit plasma kallikrein, that do not induce anaphylaxis and that are orally available. Furthermore, the molecules in the known art feature a highly polar and ionisable guanidine or amidine functionality. It is well known that such functionalities may be limiting to gut permeability and therefore to oral availability.
- 15 Other complications of diabetes such as cerebral haemorrhage, neuropathy, cardiomyopathy and neuropathy, all of which have associations with plasma kallikrein may also be considered as targets for a plasma kallikrein inhibitor.

Summary of the Invention

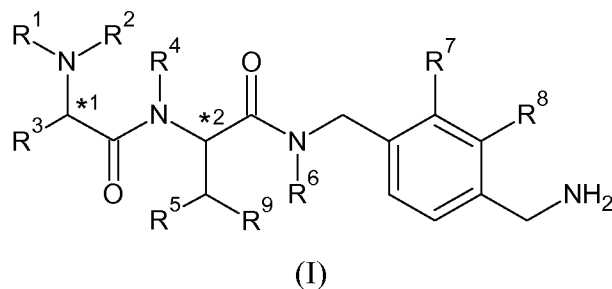
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The present invention relates to a series of benzylamines that are inhibitors of plasma kallikrein. These compounds demonstrate good selectivity for plasma kallikrein and are potentially useful in the treatment of impaired visual acuity, diabetic retinopathy, macular oedema, hereditary angioedema, diabetes, pancreatitis, cerebral haemorrhage, neuropathy, cardiomyopathy,

25 neuropathy, inflammatory bowel disease, arthritis, inflammation, septic shock, hypotension, cancer, adult respiratory distress syndrome, disseminated intravascular coagulation, cardiopulmonary bypass surgery and bleeding from post operative surgery. The invention further relates to pharmaceutical compositions of the inhibitors, to the use of the compositions as therapeutic agents, and to methods of treatment using these compositions.

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In an aspect, the present invention provides compounds of formula I



wherein:

R^1 is selected from H, alkyl, -COalkyl, -COaryl, -COheteroaryl, -CO₂alkyl, - $(CH_2)_aOH$, - $(CH_2)_bCOOR^{10}$, - $(CH_2)_cCONH_2$, -SO₂alkyl, -SO₂aryl, -SO₂ $(CH_2)_hR^{13}$, -CO $(CH_2)_iR^{14}$, -COcycloalkyl, -COCH=CHR¹⁵, -CO $(CH_2)_jNHCO(CH_2)_kR^{16}$ and -CONR¹⁷R¹⁸;

R^2 is selected from H and alkyl;

R^3 is selected from H, alkyl, - $(CH_2)_daryl$, - $(CH_2)_eheteroaryl$, - $(CH_2)_fcycloalkyl$, - $(CH_2)_gheterocycloalkyl$, -CH(cycloalkyl)₂, -CH(heterocycloalkyl)₂ and - $(CH_2)_laryl-O-(CH_2)_m-aryl$;

R^4 and R^6 are independently selected from H and alkyl;

R^5 is selected from H, alkyl, alkoxy and OH;

or R^4 and R^5 , together with the atoms to which they are attached, may join to form a 5- or 6-membered azacycloalkyl structure;

R^7 and R^8 are independently selected from H, alkyl, alkoxy, CN, halo and CF₃;

R^9 is aryl or heteroaryl;

R^{10} is H or alkyl;

a, b, c, d, e, f, g, h, i, j, l and m are independently 1, 2 or 3;

k is 0, 1, 2 or 3;

*1 and *2 denote chiral centres;

alkyl is a linear saturated hydrocarbon having up to 10 carbon atoms (C₁-C₁₀) or a branched saturated hydrocarbon of between 3 and 10 carbon atoms (C₃-C₁₀); alkyl may optionally be substituted with 1 or 2 substituents independently selected from (C₃-C₁₀)cycloalkyl, (C₁-C₆)alkoxy, OH, CN, CF₃, COOR¹¹, fluoro and NR¹¹R¹²;

cycloalkyl is a mono- or bi-cyclic saturated hydrocarbon of between 3 and 10 carbon atoms; cycloalkyl may optionally be fused to an aryl group; or cycloalkyl is adamantyl;

heterocycloalkyl is a C-linked or N-linked 3 to 10 membered saturated, mono- or bi-cyclic ring, wherein said heterocycloalkyl ring contains, where possible, 1, 2 or 3 heteroatoms independently selected from N, NR¹¹ and O;

alkoxy is a linear O-linked hydrocarbon of between 1 and 6 carbon atoms (C_1 - C_6) or a branched O-linked hydrocarbon of between 3 and 6 carbon atoms (C_3 - C_6); alkoxy may optionally be substituted with 1 or 2 substituents independently selected from (C_3 - C_{10})cycloalkyl, OH, CN, CF_3 , $COOR^{11}$, fluoro and $NR^{11}R^{12}$;

5 aryl is phenyl, biphenyl or naphthyl; aryl may be optionally substituted with up to 5 substituents independently selected from alkyl, alkoxy, OH, halo, CN, $COOR^{11}$, CF_3 and $NR^{11}R^{12}$;

heteroaryl is a 5, 6, 9 or 10 membered mono- or bi-cyclic aromatic ring, containing, where possible, 1, 2 or 3 ring members independently selected from N, NR^{11} , S and O;

10 heteroaryl may be optionally substituted with 1, 2 or 3 substituents independently selected from alkyl, alkoxy, OH, halo, CN, $COOR^{11}$, CF_3 , $NR^{11}R^{12}$ and NHR^{19} ;

R^{11} and R^{12} are independently selected from H and alkyl;

R^{13} is aryl or heteroaryl;

R^{14} is aryl, heteroaryl, cycloalkyl or heterocycloalkyl;

15 R^{15} is H, alkyl, aryl, heteroaryl, cycloalkyl or heterocycloalkyl;

R^{16} is H, aryl or heteroaryl;

R^{17} is H, alkyl, aryl, heteroaryl or heterocycloalkyl;

R^{18} is $-(CH_2)_mR^{21}$, where m is 0, 1, 2 or 3 and R^{21} is H, aryl or heteroaryl;

R^{19} -COalkyl, -COaryl or -COheteroaryl;

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and tautomers, isomers, stereoisomers (including enantiomers, diastereoisomers and racemic and scalemic mixtures thereof), pharmaceutically acceptable salts and solvates thereof.

In another aspect the present invention provides a prodrug of a compound of formula (I) as
25 herein defined, or a pharmaceutically acceptable salt thereof.

In yet another aspect the present invention provides an N-oxide of a compound of formula (I) as herein defined, or a prodrug or pharmaceutically acceptable salt thereof.

30 It will be understood that certain compounds of the present invention may exist in solvated, for example hydrated, as well as unsolvated forms. It is to be understood that the present invention encompasses all such solvated forms.

In an aspect, the invention comprises a subset of the compounds of formula (I) wherein:

R^1 is selected from H, alkyl, -COalkyl, -COaryl, -COheteroaryl, -CO₂alkyl, -(CH₂)_aOH, -(CH₂)_bCOOR¹⁰, -(CH₂)_cCONH₂, -SO₂alkyl and -SO₂aryl;

R^2 is selected from H and alkyl;

5 R^3 is selected from H, alkyl, -(CH₂)_daryl, -(CH₂)_eheteroaryl, -(CH₂)_fcycloalkyl, -(CH₂)_gheterocycloalkyl, -CH(cycloalkyl)₂ and -CH(heterocycloalkyl)₂;

R^4 and R^6 are independently selected from H and alkyl;

R^5 is selected from H, alkyl, alkoxy and OH;

10 or R^4 and R^5 , together with the atoms to which they are attached, may join to form a 5- or 6-membered azacycloalkyl structure;

R^7 and R^8 are independently selected from H, alkyl, alkoxy, CN and halo;

R^9 is aryl or heteroaryl;

R^{10} is H or alkyl;

a, b, c, d, e, f and g are independently 1, 2 or 3;

15 *1 and *2 denote chiral centres;

alkyl is a linear saturated hydrocarbon having up to 10 carbon atoms (C₁-C₁₀) or a branched saturated hydrocarbon of between 3 and 10 carbon atoms (C₃-C₁₀); alkyl may optionally be substituted with 1 or 2 substituents independently selected from (C₃-C₁₀)cycloalkyl, (C₁-C₆)alkoxy, OH, CN, CF₃, COOR¹¹, fluoro and NR¹¹R¹²;

20 cycloalkyl is a mono- or bi-cyclic saturated hydrocarbon of between 3 and 10 carbon atoms; cycloalkyl may optionally be fused to an aryl group;

heterocycloalkyl is a C-linked or N-linked 3 to 10 membered saturated, mono- or bi-cyclic ring, wherein said heterocycloalkyl ring contains, where possible, 1, 2 or 3 heteroatoms independently selected from N, NR¹¹ and O;

25 alkoxy is a linear O-linked hydrocarbon of between 1 and 6 carbon atoms (C₁-C₆) or a branched O-linked hydrocarbon of between 3 and 6 carbon atoms (C₃-C₆); alkoxy may optionally be substituted with 1 or 2 substituents independently selected from (C₃-C₁₀)cycloalkyl, OH, CN, CF₃, COOR¹¹, fluoro and NR¹¹R¹²;

30 aryl is phenyl, biphenyl or naphthyl; aryl may be optionally substituted with up to 5 substituents independently selected from alkyl, alkoxy, OH, halo, CN, COOR¹¹, CF₃ and NR¹¹R¹²;

heteroaryl is a 5, 6, 9 or 10 membered mono- or bi-cyclic aromatic ring, containing, where possible, 1, 2 or 3 ring members independently selected from N, NR¹¹, S and O;

heteroaryl may be optionally substituted with 1, 2 or 3 substituents independently selected from alkyl, alkoxy, OH, halo, CN, COOR¹¹, CF₃ and NR¹¹R¹²;

R¹¹ and R¹² are independently selected from H and alkyl;

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and tautomers, isomers, stereoisomers (including enantiomers, diastereoisomers and racemic and scalemic mixtures thereof), pharmaceutically acceptable salts and solvates thereof.

In another aspect, the invention comprises a subset of the compounds of formula (I) wherein:

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R¹ is selected from H, alkyl, -COalkyl, -COaryl, -CO₂alkyl, -CH₂CH₂OH, -CH₂COOR¹⁰, -CH₂CONH₂, -SO₂alkyl and -SO₂aryl;

R² is selected from H and alkyl;

R³ is selected from alkyl, -CH₂aryl, -CH₂cycloalkyl and -CH(cycloalkyl)₂;

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R⁴ and R⁶ are independently selected from H and alkyl;

R⁵ is selected from H, alkyl, and OH;

or R⁴ and R⁵, together with the atoms to which they are attached, may join to form a 5- or 6-membered azacycloalkyl structure;

R⁷ and R⁸ are independently selected from H, F, and Cl;

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R⁹ is aryl;

R¹⁰ is H or alkyl;

*1 and *2 denote chiral centres;

alkyl is a linear saturated hydrocarbon having up to 6 carbon atoms (C₁-C₆) or a branched saturated hydrocarbon of between 3 and 6 carbon atoms (C₃-C₆); alkyl may optionally be substituted with 1 or 2 substituents independently selected from (C₃-C₁₀)cycloalkyl, (C₁-C₆)alkoxy, OH, CN, CF₃, COOR¹¹, fluoro and NR¹¹R¹²;

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cycloalkyl is a mono- or bi-cyclic saturated hydrocarbon of between 3 and 10 carbon atoms;

alkoxy is a linear O-linked hydrocarbon of between 1 and 6 carbon atoms (C₁-C₆) or a branched O-linked hydrocarbon of between 3 and 6 carbon atoms (C₃-C₆); alkoxy may optionally be substituted with 1 or 2 substituents independently selected from (C₃-C₁₀)cycloalkyl, OH, CN, CF₃, COOR¹¹, fluoro and NR¹¹R¹²;

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aryl is phenyl, biphenyl or naphthyl; aryl may be optionally substituted with up to 5 substituents independently selected from alkyl, alkoxy, OH, halo, CN, COOR¹¹, CF₃ and NR¹¹R¹²;

R¹¹ and R¹² are independently selected from H and alkyl;

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and tautomers, isomers, stereoisomers (including enantiomers, diastereoisomers and racemic and scalemic mixtures thereof), pharmaceutically acceptable salts and solvates thereof.

The present invention also comprises the following aspects and combinations thereof:

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In an aspect of the invention, R¹ is selected from H, alkyl, -COalkyl, -COaryl, -(CH₂)_aOH, -(CH₂)_bCOOR¹⁰, -(CH₂)_cCONH₂, -SO₂alkyl and -SO₂aryl.

In an aspect of the invention, R¹ is selected from H, alkyl, -COalkyl, -COaryl, -(CH₂)_aOH, -CH₂COOR¹⁰, -CH₂CONH₂, -SO₂alkyl and -SO₂aryl; wherein a is 1 or 2.

15 In an aspect of the invention, R¹ is selected from H, -COaryl, -COalkyl, -CH₂COOH, -SO₂Ph and -SO₂CH₃.

In an aspect of the invention, R¹ is selected from H, -COethyl, methyl, methylsulfonyl, -COPhenyl, phenylsulfone, -CH₂COOH, -COⁱpropyl, propyl, -CH₂COOCH₃, -CH₂CONH₂, -CH₂CH₂OH and -CONaphthyl.

20 In an aspect of the invention, R¹ is selected from -COalkyl and -COPhenyl.

In an aspect of the invention, R¹ is selected from H, -COaryl, COheteroaryl, -COalkyl, -CH₂COOH, -SO₂Ph and -SO₂CH₃.

In an aspect of the invention, R¹ is selected from -COalkyl, COheteroaryl and -COaryl.

25 In an aspect of the invention, R² is selected from H and methyl.

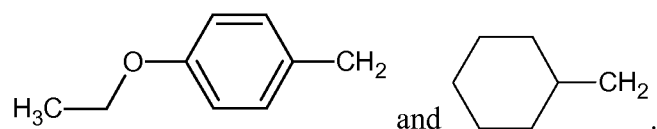
In an aspect of the invention, R² is H.

In an aspect of the invention, R³ is selected from alkyl, -(CH₂)_daryl, -(CH₂)_fcycloalkyl, and -CH(cycloalkyl)₂; wherein d and f are, independently, 1 or 2.

30 In an aspect of the invention, R³ is selected from alkyl, -CH₂aryl, -CH₂cycloalkyl, and -CH(cycloalkyl)₂.

In an aspect of the invention, R³ is selected from -CH₂aryl, -CH₂cycloalkyl, and -CH(cycloalkyl)₂.

In an aspect of the invention, R^3 is selected from:



In an aspect of the invention, R^4 is selected from H and methyl.

5 In an aspect of the invention, R^4 is H.

In an aspect of the invention, R^5 is selected from H, alkyl and OH.

In an aspect of the invention, R^5 is selected from H and OH.

In an aspect of the invention, R^5 is H.

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In an aspect of the invention, R^4 and R^5 , together with the atoms to which they are attached, join to form a pyrrolidine moiety.

In an aspect of the invention, R^4 and R^5 , together with the atoms to which they are attached, join to form a piperidine moiety.

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In an aspect of the invention, R^6 is selected from H and methyl.

In an aspect of the invention, R^6 is H.

In an aspect of the invention, R^7 is selected from H, methyl and halo.

20 In an aspect of the invention, R^7 is selected from H, F and Cl.

In an aspect of the invention, R^7 is H.

In an aspect of the invention, R^8 is selected from H, methyl and halo.

In an aspect of the invention, R^8 is selected from H, F and Cl.

25 In an aspect of the invention, R^8 is selected from H and F.

In an aspect of the invention, R^8 is H.

In an aspect of the invention, R^9 is aryl.

30 In an aspect of the invention, R^9 is selected from phenyl and naphthyl, wherein phenyl may be optionally substituted with up to 3 substituents independently selected from alkyl, alkoxy, OH, halo, CN, COOR¹¹, CF₃ and NR¹¹R¹².

In an aspect of the invention, R⁹ is phenyl, wherein phenyl may be optionally substituted with up to 2 substituents independently selected from alkyl, halo and CF₃.

In an aspect of the invention, R⁹ is selected from phenyl, 1-naphthalene, 2,4-dichlorophenyl, 3,4-dichlorophenyl, 3,4-difluorophenyl, 4-chlorophenyl, 4-trifluoromethylphenyl and 4-ethoxyphenyl.

In an aspect of the invention, R⁹ is selected from phenyl, heteroaryl and naphthyl, wherein phenyl may be optionally substituted with up to 3 substituents independently selected from alkyl, alkoxy, OH, halo, CN, COOR¹¹, CF₃ and NR¹¹R¹².

In an aspect of the invention, R⁹ is selected from phenyl, 1-naphthalene, 3,4-dichlorophenyl, 3,4-difluorophenyl, 4-chlorophenyl, 4-fluorophenyl, 3-fluorophenyl, 4-trifluoromethylphenyl, pyrid-3-yl, pyrid-2-yl, pyrid-4-yl, benzothiophen-3-yl, thiophen-2-yl, thiophen-3-yl, indol-3-yl, and thiazol-4yl.

In an aspect of the invention, R¹⁰ is H or methyl.

In an aspect of the invention, the stereochemical configuration about chiral centre *1 is *R*.

In an aspect of the invention, the stereochemical configuration about chiral centre *2 is *S*.

In an aspect of the invention, a is 2 and b, c, d, e, f and g are 1.

In an aspect of the invention, a is 2 and b, c, d, e, f, g, h, j, l and m are 1.

In an aspect of the invention, k is 0 or 1.

In an aspect, the invention comprises a compound selected from:

(S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

{(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino}-acetic acid;

(S)-N-(4-Aminomethyl-3-fluoro-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide;

(S)-N-(4-Aminomethyl-2-chloro-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide;

(S)-N-(4-Aminomethyl-benzyl)-3-(3,4-dichloro-phenyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-propionamide;

(S)-N-(4-Aminomethyl-3-chloro-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide;

(S)-N-(4-Aminomethyl-benzyl)-2- {[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionyl]-methyl-amino} -3-phenyl-propionamide;

((R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethyl)-methyl-amino)-acetic acid;

(S)-N-(4-Aminomethyl-3-fluoro-benzyl)-2- {[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionyl]-methyl-amino} -3-phenyl-propionamide;

N-[(R)-1- {[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-methyl-carbamoyl} -2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1- {[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-methyl-carbamoyl} -2-(4-ethoxy-phenyl)-ethyl]-isobutyramide;

Naphthalene-1-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-chloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2,4-dichloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,4-difluoro-benzamide;

(R)-2-Amino-N-[(1S,2S)-1-(4-aminomethyl-benzylcarbamoyl)-2-hydroxy-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-nicotinamide;

(2S,3S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-hydroxy-3-phenyl-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;

Thiophene-3-carboxylic acid-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Cyclohexanecarboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Isoxazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Benzo[b]thiophene-2-carboxylic acid[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-2-(4-chloro-benzenesulfonylamino)-3-(4-ethoxy-phenyl)-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3-chloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-chloro-benzamide

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3-trifluoromethyl-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,4-dichloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide;

(S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-(2-phenylacetyl-amino-acetyl-amino)-propionyl-amino]-3-phenyl-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-fluoro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-6-methyl-nicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-methyl-nicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2,6-dichloro-nicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-5,6-dichloro-nicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2,3,6-trifluoro-isonicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,3,3-trifluoro-propionamide;

2,4-Dimethyl-thiazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

2-Methyl-thiazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

4-Methyl-thiazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Furan-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Methyl-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-methoxy-isonicotinamide;

3-Methyl-1H-pyrrole-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-propoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(4-chloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(4-fluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(4-methoxy-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3-fluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiophen-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-3-chloro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiophen-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-3-chloro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-difluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;

Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-chloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethyl]-3-(4-ethoxy-phenyl)-2-propionylamino-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethyl]-3-(4-ethoxy-phenyl)-2-propionylamino-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,3,3-trifluoro-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-chloro-benzamide;

- Isoxazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-difluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- 3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(1H-indol-3-yl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- 5 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;
- 3-Acetylamino-thiophene-2-carboxylic acid-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(2-fluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- 10 3-Methyl-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-3-methyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- 15 3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- 3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide;
- 20 3-Methyl-1H-pyrrole-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- 3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- 25 3-Acetylamino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3-methyl-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-methyl-benzamide;

- 5 3,5-Dimethyl-1H-pyrrole-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-3-methyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

- 10 3-Acetylamino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Acetylamino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

- 15 3-Chloro-thiophene-2-carboxylic acid [(R)-1-{[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-methyl-carbamoyl}-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(1S,2R)-1-(4-Aminomethyl-benzylcarbamoyl)-2-hydroxy-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

- 20 3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(1S,2R)-1-(4-aminomethyl-benzylcarbamoyl)-2-hydroxy-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-{(R,S)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-ethyl}-benzamide;

and pharmaceutically acceptable salts and solvates thereof.

- 25 In an aspect, the invention comprises a compound selected from:

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

Naphthalene-1-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-chloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2,4-dichloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,4-difluoro-benzamide;

and pharmaceutically acceptable salts and solvates thereof.

In an aspect, the invention comprises a compound selected from:

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

Naphthalene-1-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

5 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-chloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2,4-dichloro-benzamide;

10 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-nicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,4-difluoro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;

15 Thiophene-3-carboxylic acid-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

20 Cyclohexanecarboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Isoxazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

25 (R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-2-(4-chloro-benzenesulfonylamino)-3-(4-ethoxy-phenyl)-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide;

- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-phenyl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-3,4-dichloro-benzamide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-phenyl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-3-chloro-benzamide;
- 5 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-phenyl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-phenyl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-4-fluoro-benzamide;
- 3-Methyl-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyle)-2-phenyl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- 10 3-Methyl-1H-pyrrole-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyle)-2-phenyl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- 3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyle)-2-phenyl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- 15 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-phenyl-ethylcarbamoyle]-2-(4-propoxy-phenyl)-ethyl]-benzamide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-pyridin-2-yl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-(3,4-dichloro-phenyl)-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- 20 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-(4-fluoro-phenyl)-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-pyridin-3-yl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- 25 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-thiophen-2-yl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-thiophen-3-yl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-benzo[b]thiophen-3-yl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- 30 N-[(R)-1-[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyle)-2-phenyl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-3-chloro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiophen-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

- 5 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-3-chloro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

- 10 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide;

Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide;

- 15 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-difluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

- 20 3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(1H-indol-3-yl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;

- 25 3-Acetyl-amino-thiophene-2-carboxylic acid-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Methyl-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

and pharmaceutically acceptable salts and solvates thereof.

30

Therapeutic Applications

As previously mentioned, the compounds of the present invention are potent and selective inhibitors of plasma kallikrein. They are therefore useful in the treatment of disease conditions for which over-activity of plasma kallikrein is a causative factor.

- 5 Accordingly, the present invention provides a compound of formula (I) for use in medicine.

The present invention also provides for the use of a compound of formula (I) in the manufacture of a medicament for the treatment or prevention of a disease or condition in which plasma kallikrein activity is implicated.

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The present invention also provides a compound of formula (I) for use in the treatment or prevention of a disease or condition in which plasma kallikrein activity is implicated.

- 15 The present invention also provides a method of treatment of a disease or condition in which plasma kallikrein activity is implicated comprising administration to a subject in need thereof a therapeutically effective amount of a compound of formula (I).

- 20 In one aspect, the disease or condition in which plasma kallikrein activity is implicated is selected from Diseases or conditions in which plasma kallikrein activity is implicated include impaired visual acuity, diabetic retinopathy, diabetic macular oedema, hereditary angioedema, diabetes, pancreatitis, cerebral haemorrhage, nephropathy, cardiomyopathy, neuropathy, inflammatory bowel disease, arthritis, inflammation, septic shock, hypotension, cancer, adult respiratory distress syndrome, disseminated intravascular coagulation, cardiopulmonary bypass surgery and bleeding from post operative surgery.

25

In another aspect, the disease or condition in which plasma kallikrein activity is implicated is retinal vascular permeability associated with diabetic retinopathy and diabetic macular oedema.

Combination Therapy

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The compounds of the present invention may be administered in combination with other therapeutic agents. Suitable combination therapies include a compound of formula (I) combined with one or more agents selected from agents that inhibit platelet-derived growth factor (PDGF),

endothelial growth factor (VEGF), integrin $\alpha 5 \beta 1$, steroids, other agents that inhibit plasma kallikrein and other inhibitors of inflammation. Specific examples of therapeutic agents that may be combined with the compounds of the present invention include those disclosed in EP2281885A and by S. Patel in *Retina*, 2009 Jun;29(6 Suppl):S45-8.

5

When combination therapy is employed, the compounds of the present invention and said combination agents may exist in the same or different pharmaceutical compositions, and may be administered separately, sequentially or simultaneously.

10 Definitions

The term "alkyl" includes saturated hydrocarbon residues including:

- linear groups up to 10 carbon atoms (C_1 - C_{10}), or of up to 6 carbon atoms (C_1 - C_6), or of up to 4 carbon atoms (C_1 - C_4). Examples of such alkyl groups include, but are not limited, to C_1 - methyl, C_2 - ethyl, C_3 - propyl and C_4 - n-butyl.
 - branched groups of between 3 and 10 carbon atoms (C_3 - C_{10}), or of up to 7 carbon atoms (C_3 - C_7), or of up to 4 carbon atoms (C_3 - C_4). Examples of such alkyl groups include, but are not limited to, C_3 - iso-propyl, C_4 - sec-butyl, C_4 - iso-butyl, C_4 - tert-butyl and C_5 - neo-pentyl.
- each optionally substituted as stated above.

20

The term "alkoxy" includes O-linked hydrocarbon residues including:

- linear groups of between 1 and 6 carbon atoms (C_1 - C_6), or of between 1 and 4 carbon atoms (C_1 - C_4). Examples of such alkoxy groups include, but are not limited to, C_1 - methoxy, C_2 - ethoxy, C_3 - n-propoxy and C_4 - n-butoxy.
 - branched groups of between 3 and 6 carbon atoms (C_3 - C_6) or of between 3 and 4 carbon atoms (C_3 - C_4). Examples of such alkoxy groups include, but are not limited to, C_3 - iso-propoxy, and C_4 - sec-butoxy and tert-butoxy.
- each optionally substituted as stated above.

30 Unless otherwise stated, halo is selected from Cl, F, Br and I.

Cycloalkyl is as defined above. Cycloalkyl groups may contain from 3 to 10 carbon atoms, or from 4 to 10 carbon atoms, or from 5 to 10 carbon atoms, or from 4 to 6 carbon atoms.

Examples of suitable monocyclic cycloalkyl groups include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl and cycloheptyl. Examples of suitable bicyclic cycloalkyl groups include decahydronaphthalene and octahydro-1*H*-indene. Examples of suitable cycloalkyl groups, when fused with aryl, include indanyl and 1,2,3,4-tetrahydronaphthyl.

5

Heterocycloalkyl is as defined above. Examples of suitable heterocycloalkyl groups include oxiranyl, aziridinyl, azetidiny, tetrahydrofuranyl, pyrrolidinyl, tetrahydropyranyl, piperidinyl, N-methylpiperidinyl, morpholinyl, N-methyl morpholinyl, piperazinyl, N-methylpiperazinyl, azepanyl, oxazepanyl and diazepanyl.

10

Aryl is as defined above. Typically, aryl will be optionally substituted with 1, 2 or 3 substituents. Optional substituents are selected from those stated above. Examples of suitable aryl groups include phenyl and naphthyl (each optionally substituted as stated above).

15 Heteroaryl is as defined above. Examples of suitable heteroaryl groups include thienyl, furanyl, pyrrolyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, triazolyl, oxadiazolyl, thiadiazolyl, tetrazolyl, pyridinyl, pyridazinyl, pyrimidinyl, pyrazinyl, indolyl, benzimidazolyl, benzotriazolyl, quinolinyl and isoquinolinyl (optionally substituted as stated above).

20 The term "C-linked", such as in "C-linked heterocycloalkyl", means that the heterocycloalkyl group is joined to the remainder of the molecule *via* a ring carbon atom.

The term "N-linked", such as in "N-linked heterocycloalkyl", means that the heterocycloalkyl group is joined to the remainder of the molecule *via* a ring nitrogen atom.

25

The term "O-linked", such as in "O-linked hydrocarbon residue", means that the hydrocarbon residue is joined to the remainder of the molecule *via* an oxygen atom.

In groups such as -COalkyl and -(CH₂)₆COOR¹⁰, "-" denotes the point of attachment of the
30 substituent group to the remainder of the molecule.

"Pharmaceutically acceptable salt" means a physiologically or toxicologically tolerable salt and includes, when appropriate, pharmaceutically acceptable base addition salts and

pharmaceutically acceptable acid addition salts. For example (i) where a compound of the invention contains one or more acidic groups, for example carboxy groups, pharmaceutically acceptable base addition salts that can be formed include sodium, potassium, calcium, magnesium and ammonium salts, or salts with organic amines, such as, diethylamine, *N*-methyl-
5 glucamine, diethanolamine or amino acids (e.g. lysine) and the like; (ii) where a compound of the invention contains a basic group, such as an amino group, pharmaceutically acceptable acid addition salts that can be formed include hydrochlorides, hydrobromides, sulfates, phosphates, acetates, citrates, lactates, tartrates, mesylates, succinates, oxalates, phosphates, esylates, tosylates, benzenesulfonates, naphthalenedisulphonates, maleates, adipates, fumarates,
10 hippurates, camphorates, xinafoates, *p*-acetamidobenzoates, dihydroxybenzoates, hydroxynaphthoates, succinates, ascorbates, oleates, bisulfates and the like.

Hemisalts of acids and bases can also be formed, for example, hemisulfate and hemicalcium salts.

15

For a review of suitable salts, see "Handbook of Pharmaceutical Salts: Properties, Selection and Use" by Stahl and Wermuth (Wiley-VCH, Weinheim, Germany, 2002).

"Prodrug" refers to a compound which is convertible *in vivo* by metabolic means (e.g. by
20 hydrolysis, reduction or oxidation) to a compound of the invention. Suitable groups for forming pro-drugs are described in 'The Practice of Medicinal Chemistry, 2nd Ed. pp561-585 (2003) and in F. J. Leinweber, *Drug Metab. Res.*, 1987, **18**, 379. .

The compounds of the invention can exist in both unsolvated and solvated forms. The term
25 'solvate' is used herein to describe a molecular complex comprising the compound of the invention and a stoichiometric amount of one or more pharmaceutically acceptable solvent molecules, for example, ethanol. The term 'hydrate' is employed when the solvent is water.

Where compounds of the invention exist in one or more geometrical, optical, enantiomeric,
30 diastereomeric and tautomeric forms, including but not limited to *cis*- and *trans*-forms, *E*- and *Z*-forms, *R*-, *S*- and *meso*-forms, keto-, and enol-forms. Unless otherwise stated a reference to a particular compound includes all such isomeric forms, including racemic and other mixtures thereof. Where appropriate such isomers can be separated from their mixtures by the application

or adaptation of known methods (e.g. chromatographic techniques and recrystallisation techniques). Where appropriate such isomers can be prepared by the application or adaptation of known methods (e.g. asymmetric synthesis).

- 5 In the context of the present invention, references herein to "treatment" include references to curative, palliative and prophylactic treatment.

General Methods

- 10 The compounds of formula (I) should be assessed for their biopharmaceutical properties, such as solubility and solution stability (across pH), permeability, *etc.*, in order to select the most appropriate dosage form and route of administration for treatment of the proposed indication. They may be administered alone or in combination with one or more other compounds of the invention or in combination with one or more other drugs (or as any combination thereof).
- 15 Generally, they will be administered as a formulation in association with one or more pharmaceutically acceptable excipients. The term 'excipient' is used herein to describe any ingredient other than the compound(s) of the invention which may impart either a functional (i.e., drug release rate controlling) and/or a non-functional (i.e., processing aid or diluent) characteristic to the formulations. The choice of excipient will to a large extent depend on
- 20 factors such as the particular mode of administration, the effect of the excipient on solubility and stability, and the nature of the dosage form.

- Compounds of the invention intended for pharmaceutical use may be administered as a solid or liquid, such as a tablet, capsule or solution. Pharmaceutical compositions suitable for the
- 25 delivery of compounds of the present invention and methods for their preparation will be readily apparent to those skilled in the art. Such compositions and methods for their preparation may be found, for example, in Remington's Pharmaceutical Sciences, 19th Edition (Mack Publishing Company, 1995).

- 30 Accordingly, the present invention provides a pharmaceutical composition comprising a compound of formula (I) and a pharmaceutically acceptable carrier, diluent or excipient.

For the treatment of conditions such as retinal vascular permeability associated with diabetic retinopathy and diabetic macular oedema, the compounds of the invention may be administered in a form suitable for injection into the ocular region of a patient, in particular, in a form suitable for intra-vitreous injection. It is envisaged that formulations suitable for such use will take the form of sterile solutions of a compound of the invention in a suitable aqueous vehicle. The compositions may be administered to the patient under the supervision of the attending physician.

The compounds of the invention may also be administered directly into the blood stream, into subcutaneous tissue, into muscle, or into an internal organ. Suitable means for parenteral administration include intravenous, intraarterial, intraperitoneal, intrathecal, intraventricular, intraurethral, intrasternal, intracranial, intramuscular, intrasynovial and subcutaneous. Suitable devices for parenteral administration include needle (including microneedle) injectors, needle-free injectors and infusion techniques.

Parenteral formulations are typically aqueous or oily solutions. Where the solution is aqueous, excipients such as sugars (including but not restricted to glucose, mannitol, sorbitol, etc.), salts, carbohydrates and buffering agents (preferably to a pH of from 3 to 9), but, for some applications, they may be more suitably formulated as a sterile non-aqueous solution or as a dried form to be used in conjunction with a suitable vehicle such as sterile, pyrogen-free water.

Parenteral formulations may include implants derived from degradable polymers such as polyesters (i.e., polylactic acid, polylactide, polylactide-co-glycolide, polycaprolactone, polyhydroxybutyrate), polyorthoesters and polyanhydrides. These formulations may be administered via surgical incision into the subcutaneous tissue, muscular tissue or directly into specific organs.

The preparation of parenteral formulations under sterile conditions, for example, by lyophilisation, may readily be accomplished using standard pharmaceutical techniques well known to those skilled in the art.

The solubility of compounds of formula (I) used in the preparation of parenteral solutions may be increased by the use of appropriate formulation techniques, such as the incorporation of co-

solvents and/or solubility-enhancing agents such as surfactants, micelle structures and cyclodextrins.

In one embodiment, the compounds of the invention may be administered orally. Oral administration may involve swallowing, so that the compound enters the gastrointestinal tract, and/or buccal, lingual, or sublingual administration by which the compound enters the blood stream directly from the mouth.

Formulations suitable for oral administration include solid plugs, solid microparticulates, semi-solid and liquid (including multiple phases or dispersed systems) such as tablets; soft or hard capsules containing multi- or nano-particulates, liquids, emulsions or powders; lozenges (including liquid-filled); chews; gels; fast dispersing dosage forms; films; ovules; sprays; and buccal/mucoadhesive patches.

Formulations suitable for oral administration may also be designed to deliver the compounds of the invention in an immediate release manner or in a rate-sustaining manner, wherein the release profile can be delayed, pulsed, controlled, sustained, or delayed and sustained or modified in such a manner which optimises the therapeutic efficacy of the said compounds. Means to deliver compounds in a rate-sustaining manner are known in the art and include slow release polymers that can be formulated with the said compounds to control their release.

Examples of rate-sustaining polymers include degradable and non-degradable polymers that can be used to release the said compounds by diffusion or a combination of diffusion and polymer erosion. Examples of rate-sustaining polymers include hydroxypropyl methylcellulose, hydroxypropyl cellulose, methyl cellulose, ethyl cellulose, sodium carboxymethyl cellulose, polyvinyl alcohol, polyvinyl pyrrolidone, xanthum gum, polymethacrylates, polyethylene oxide and polyethylene glycol.

Liquid (including multiple phases and dispersed systems) formulations include emulsions, solutions, syrups and elixirs. Such formulations may be presented as fillers in soft or hard capsules (made, for example, from gelatin or hydroxypropylmethylcellulose) and typically comprise a carrier, for example, water, ethanol, polyethylene glycol, propylene glycol, methylcellulose, or a suitable oil, and one or more emulsifying agents and/or suspending agents.

Liquid formulations may also be prepared by the reconstitution of a solid, for example, from a sachet.

The compounds of the invention may also be used in fast-dissolving, fast-disintegrating dosage forms such as those described in Liang and Chen, Expert Opinion in Therapeutic Patents, 2001, 11 (6), 981-986.

The formulation of tablets is discussed in Pharmaceutical Dosage Forms: Tablets, Vol. 1, by H. Lieberman and L. Lachman (Marcel Dekker, New York, 1980).

10

For administration to human patients, the total daily dose of the compounds of the invention is typically in the range 0.01 mg and 1000 mg, or between 0.1 mg and 250 mg, or between 1 mg and 50 mg depending, of course, on the mode of administration. For example, if administered by intra-vitreal injection it is envisaged that the compound of the invention will be dosed infrequently, e.g. once a month. In this circumstance, a dose of between 0.5 mg and 20 mg, such as between 1 mg and 10 mg, is envisaged. If dosed more frequently, e.g. once daily, a much lower dose of between 0.005 mg and 0.02 mg is envisaged.

20 The total dose may be administered in single or divided doses and may, at the physician's discretion, fall outside of the typical range given herein. These dosages are based on an average human subject having a weight of about 60kg to 70kg. The physician will readily be able to determine doses for subjects whose weight falls outside this range, such as infants and the elderly.

25

Synthetic Methods

The compounds of the present invention can be prepared according to the procedures of the following schemes and examples, using appropriate materials, and are further exemplified by the specific examples provided herein below. Moreover, by utilising the procedures described herein, one of ordinary skill in the art can readily prepare additional compounds that fall within the scope of the present invention claimed herein. The compounds illustrated in the examples

are not, however, to be construed as forming the only genus that is considered as the invention. The examples further illustrate details for the preparation of the compounds of the present invention. Those skilled in the art will readily understand that known variations of the conditions and processes of the following preparative procedures can be used to prepare these
5 compounds.

The compounds of the invention may be isolated in the form of their pharmaceutically acceptable salts, such as those described previously herein above.

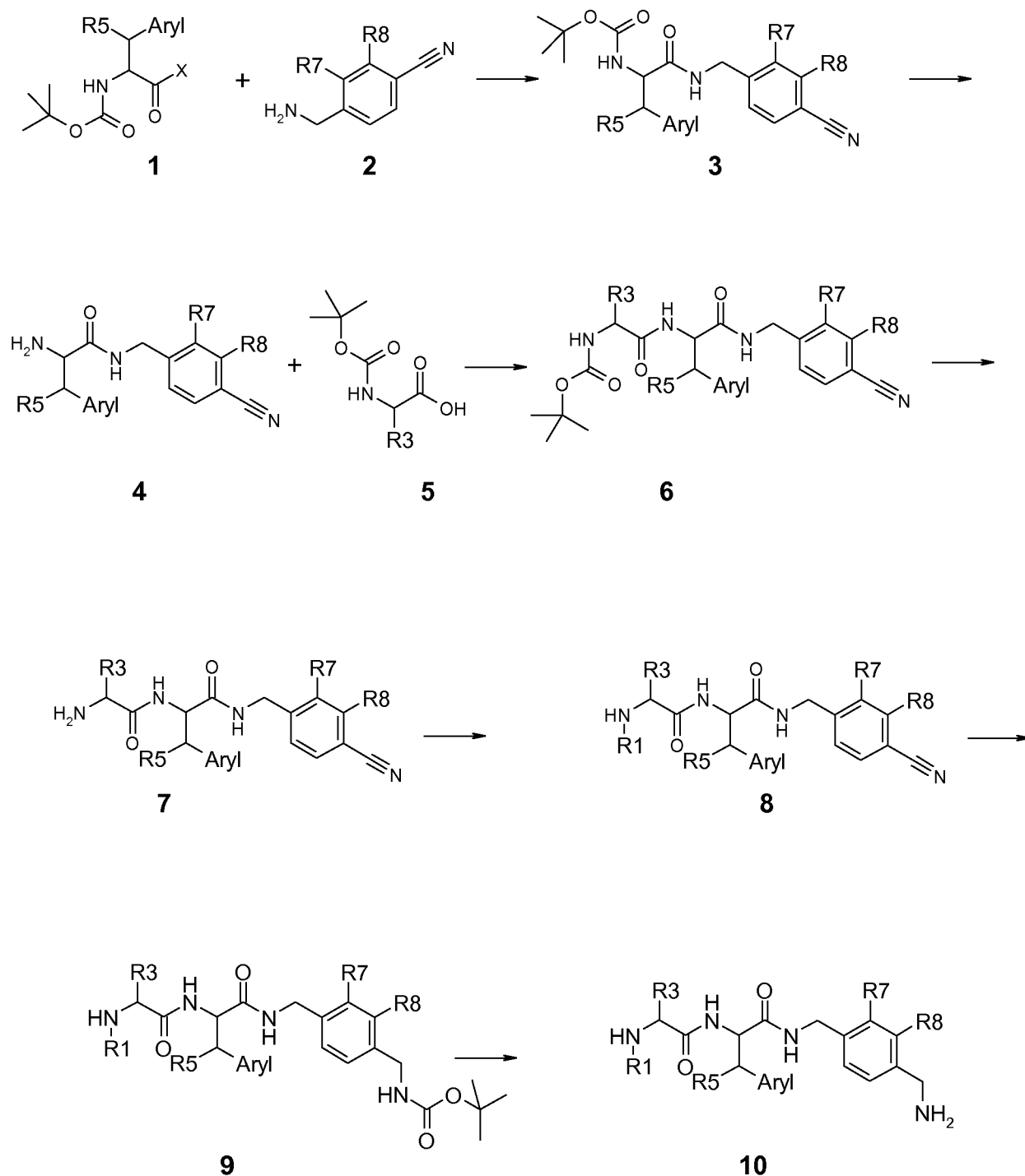
10 It may be necessary to protect reactive functional groups (e.g. hydroxy, amino, thio or carboxy) in intermediates used in the preparation of compounds of the invention to avoid their unwanted participation in a reaction leading to the formation of the compounds. Conventional protecting groups, for example those described by T. W. Greene and P. G. M. Wuts in "Protective groups in organic chemistry" John Wiley and Sons, 4th Edition, 2006, may be used. For example, a
15 common amino protecting group suitable for use herein is tert-butoxy carbonyl (Boc), which is readily removed by treatment with an acid such as trifluoroacetic acid or hydrogen chloride in an organic solvent such as dichloromethane. Alternatively the amino protecting group may be a benzyloxycarbonyl (Z) group which can be removed by hydrogenation with a palladium catalyst under a hydrogen atmosphere or 9-fluorenylmethyloxycarbonyl (Fmoc) group which can be
20 removed by solutions of secondary organic amines such as diethylamine or piperidine in an organic solvents. Carboxyl groups are typically protected as esters such as methyl, ethyl, benzyl or tert-butyl which can all be removed by hydrolysis in the presence of bases such as lithium or sodium hydroxide. Benzyl protecting groups can also be removed by hydrogenation with a palladium catalyst under a hydrogen atmosphere whilst tert-butyl groups can also be removed by
25 trifluoroacetic acid. Alternatively a trichloroethyl ester protecting group is removed with zinc in acetic acid. A common hydroxy protecting group suitable for use herein is a methyl ether, deprotection conditions comprise refluxing in 48% aqueous HBr for 1-24 hours, or by stirring with borane tribromide in dichloromethane for 1-24 hours. Alternatively where a hydroxy group is protected as a benzyl ether, deprotection conditions comprise hydrogenation with a palladium
30 catalyst under a hydrogen atmosphere.

The compounds according to general formula **I** can be prepared using conventional synthetic methods for example, but not limited to, the route outlined in Scheme 1. In a typical first step the

amine (**2**) is coupled using standard peptide coupling conditions to an activated alpha amino acid (**1**) suitably amino-protected with a standard protecting group such as tert-butyloxycarbonyl (Boc), benzyloxycarbonyl (Z) or 9-fluorenylmethyloxycarbonyl (Fmoc). The activating group (X) may be N-hydroxysuccinimide. The use of such groups is well known in the art. Where R⁵ or R⁹ (shown as 'Aryl' in Scheme 1) has a reactive functional group such as an amine or a carboxylic acid, this group will also be protected. Other standard peptide coupling methods include the reaction of acids with amines in the presence of hydroxybenzotriazole and carbodiimide such as water soluble carbodiimide, or 2-(1H-benzotriazole-1-yl)-1,1,3,3-tetramethylaminium hexafluorophosphate or benzotriazole-1-yl-oxy-tris-pyrrolidino-phosphonium hexafluorophosphate or bromo-trispyrrolidino-phosphonium hexafluorophosphate in the presence of organic bases such as triethylamine, diisopropylethylamine or N-methylmorpholine. In a typical second step the protecting group is removed using standard methods as previously described.

15 The route exemplified in Scheme 1 then proceeds in the third step via a further standard peptide coupling and in the fourth step via removal of the Boc protecting group using standard conditions as previously described. The amine revealed in **7** may, in a fifth step, then typically be alkylated or acylated with the group R₁. Acylation may be carried out by treatment with an acylating agent such as an acyl chloride, for example acetyl chloride or benzoyl chloride, in the presence of a base, typically a tertiary amine base such as triethylamine or diisopropylethylamine. Alkylation may typically be carried by treatment with an alkyl halide or by reductive alkylation. Typically, in a reductive alkylation procedure the amine is allowed to react with an aldehyde or ketone in the presence of a suitable reducing agent such as sodium cyanoborohydride or sodium acetoxyborohydride in a suitable solvent such as methanol, at room temperature. The resulting nitrile compound **8** may then be reduced by hydrogenation. Conversion from **8** to **10** may be achieved in a single step either by direct reduction of the nitrile by hydrogenation in a suitable solvent such as methanol in the presence of a suitable catalyst such as palladium on charcoal in the presence of an acid such as hydrochloric acid or reduction with a suitable borohydride in the presence of a suitable transition metal such as cobalt or nickel chloride in a suitable solvent such as methanol at room temperature. Alternatively, the *tert*-butoxycarbonyl (Boc) protected amine **9** may be isolated (using, for example, the method as described in S. Caddick et al., *Tetrahedron Lett.*, 2000, **41**, 3513) and subsequently deprotected by standard means described previously to give the amine **10**.

Scheme 1



5 Brief description of drawings

Figure 1 shows the inhibitory effect of Example 3 and CH-3457 (positive control; plasma kallikrein inhibitor) upon CA-I stimulated RVP in Sprague Dawley rats.

Figure 2 shows the ocular tissue concentrations of Example 3 following IVT administration of 4.2 µg/mL (210ng/eye).

Examples

5

The invention is illustrated by the following non-limiting examples in which the following abbreviations and definitions are used:

Cha	3-Cyclohexylalanine
DMF	N,N-Dimethylformamide
DMSO	Dimethyl sulfoxide
Et	Ethyl
EtOAc	Ethyl Acetate
hrs	Hours
HOBt	Hydroxybenzotriazole
LCMS	Liquid chromatography mass spectrometry
Me	Methyl
MeCN	Acetonitrile
MeOH	Methanol
min	Minutes
MS	Mass spectrum
m/z	Mass charge ratio (of parent, MH^+ , ion unless otherwise stated)
NMR	Nuclear magnetic resonance spectrum – NMR spectra were recorded at a frequency of 400MHz unless otherwise indicated
Pet. Ether	Petroleum ether fraction boiling at 60-80°C
Ph	Phenyl
Phe	Phenylalanine
n-Pr	n-Propyl
THF	Tetrahydrofuran
TFA	Trifluoroacetic acid

10 All reactions were carried out under an atmosphere of nitrogen unless specified otherwise.

¹H NMR spectra were recorded on a Bruker Avance III (400MHz) spectrometer with reference to deuterium solvent and at room temperature.

Molecular ions were obtained using LCMS which was carried out using a Chromolith Speedrod RP-18e column, 50 x 4.6 mm, with a linear gradient 10% to 90% 0.1% HCO₂H/MeCN into 0.1% HCO₂H/H₂O over 11 min, flow rate 1.5 mL/min. Data was collected using a Thermofinnigan Surveyor MSQ mass spectrometer with electrospray ionisation in conjunction with a Thermofinnigan Surveyor LC system.

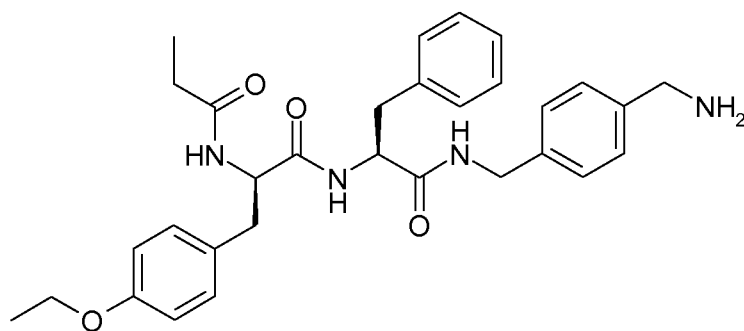
Chemical names were generated using the Autonom software provided as part of the ISIS draw package from MDL Information Systems.

Where products were purified by flash chromatography, 'silica' refers to silica gel for chromatography, 0.035 to 0.070 mm (220 to 440 mesh) (e.g. Merck silica gel 60), and an applied pressure of nitrogen up to 10 p.s.i accelerated column elution. Reverse phase preparative HPLC purifications were carried out using a Waters 2525 binary gradient pumping system at flow rates of typically 20 mL/min using a Waters 2996 photodiode array detector.

All solvents and commercial reagents were used as received.

EXAMPLE 1

(S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide



A. (S)-2-[(R)-2-tert-Butoxycarbonylamino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionic acid methyl ester

H-Phe-OMe.HCl (2.3g, 10.7mmol) was dissolved in CH₂Cl₂ (100 mL) and DMF (10 mL). This solution was cooled to 0°C. (R)-2-Butoxyloxycarbonylamino-3-(4-ethoxy-phenyl)-propionic acid (3.0g, 9.7mmol) was added followed by HOBt (1.57g, 11.6mmol) and triethylamine (2.9g, 29.0mmol). Water soluble carbodiimide (2.04g, 10.6mmol) was then added. After 18 hrs at 0°C to room temperature reaction mixture was diluted with chloroform (100 mL) and washed with NaHCO₃ (1x30 mL), water (1x30 mL), brine (1x30 mL), dried (Na₂SO₄) and evaporated in vacuo giving a yellow oil. The residue was purified by flash chromatography (silica), eluent 20% Pet. Ether (60-80°C), 80% EtOAc, fractions combined and evaporated in vacuo to give a colourless oil identified as (S)-2-[(R)-2-tert-butoxycarbonylamino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionic acid methyl ester (4.25g, 9.03mmol, 93%).
[M+H]⁺ = 471.27.

B. (S)-2-[(R)-2-tert-Butoxycarbonylamino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionic acid

(S)-2-[(R)-2-tert-Butoxycarbonylamino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionic acid methyl ester (2.5g, 5.3mmol) was dissolved in THF (100 mL). Lithium hydroxide monohydrate (668mg, 15.9mmol) in water (10 mL) was added. The reaction mixture was stirred at room temperature for 18 hrs after which time the reaction mixture was diluted with EtOAc (150 mL). This solution was washed with 0.3M KHSO₄ (1x50 mL), water (1x30 mL), brine (1x30 mL), dried (Na₂SO₄) and evaporated in vacuo to give a white solid identified as (S)-2-[(R)-2-tert-butoxycarbonylamino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionic acid (2.095g, 4.58mmol, 86%).
[M+H]⁺ = 457.25.

C. [(R)-1-[(S)-1-(4-Cyano-benzylcarbamoyle)-2-phenyl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-carbamic acid tert-butyl ester

4-(Aminomethyl)benzonitrile hydrochloride (303mg, 1.80mmol) was dissolved in CH₂Cl₂ (50 mL) and DMF (5 mL). This solution was cooled to 0°C. (S)-2-[(R)-2-tert-butoxycarbonylamino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionic acid (745mg, 1.63mmol) was added followed by HOBt (265mg, 1.96mmol) and triethylamine (495mg, 4.9mmol). Water soluble carbodiimide (344mg, 1.8mmol) was then added. After 18 hrs at 0°C to room temperature

reaction mixture was diluted with chloroform (100 mL) and washed with NaHCO₃ (1x30 mL), water (1x30 mL), brine (1x30 mL), dried (Na₂SO₄) and evaporated in vacuo giving a yellow oil. The residue was purified by flash chromatography (silica), eluent 20%Pet. Ether (60-80°C), 80% EtOAc, fractions combined and evaporated in vacuo to give a colourless oil identified as [(R)-1-
 5 [(S)-1-(4-cyano-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-
 carbamic acid tert-butyl ester (493mg, 0.86mmol, 53%).

$$[M+H]^+ = 571.29$$

**D. (R)-2-Amino-N-[(S)-1-(4-cyano-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-
 10 propionamide Hydrochloride**

[(R)-1-[(S)-1-(4-Cyano-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-
 ethyl]-carbamic acid tert-butyl ester (225mg, 0.39mmol) was treated with 4M HCl/dioxan (50
 mL). After one hour at room temperature the solvent was removed to give a white solid
 identified as (R)-2-amino-N-[(S)-1-(4-cyano-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-
 15 phenyl)-propionamide hydrochloride (200mg, 0.39mmol, 100%).

$$[M+H]^+ = 471.26$$

**E. (S)-N-(4-Cyano-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-
 3-phenyl-propionamide**

(R)-2-Amino-N-[(S)-1-(4-cyano-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-
 propionamide hydrochloride (200mg, 0.37mol) was dissolved in dichloromethane (50 mL), this
 solution was cooled to 0°C. Triethylamine (111mg, 1.1mmol) was added followed by propionyl
 chloride (39mg, 0.40mmol). After 18 hrs at 0°C to room temperature the reaction mixture was
 diluted with CHCl₃ (50 mL), this solution was washed with sat. NaHCO₃ (1x20 mL), water
 25 (1x20 mL), brine (1x20 mL), dried (Na₂SO₄) and evaporated in vacuo. The residue was purified
 by flash chromatography (silica), eluent 2% MeOH, 98% CHCl₃, fractions combined and
 evaporated in vacuo to give a colourless oil identified as (S)-N-(4-cyano-benzyl)-2-[(R)-3-(4-
 ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide (189mg, 0.36mmol,
 98%).

$$[M+H]^+ = 527.27$$

**F. [4-((S)-2-[(R)-3-(4-Ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-
 propionylamino}-methyl)-benzyl]-carbamic acid tert-butyl ester**

(S)-N-(4-Cyano-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide (100mg, 0.19mmol) was dissolved in methanol (50 mL). This solution was cooled to 0°C. Nickel (II) chloride hexahydrate (4.5mg, 0.0192mmol) and di-tertbutyl dicarbonate (83mg, 0.38mmol) were added followed by sodium borohydride (50mg, 1.33mmol) 5 portionwise. The reaction mixture was stirred at 0°C to room temp for 18hrs. The methanol was removed by evaporation. The residue was dissolved in CHCl₃ (70 mL), washed with sat NaHCO₃ (1x30 mL), water (1x30 mL), brine (1x30 mL), dried (Na₂SO₄) and evaporated in vacuo to give a yellow oil. Purified by flash chromatography, eluant 1% MeOH, 99% CHCl₃ to give a colourless oil identified as [4-({(S)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionylamino}-methyl)-benzyl]-carbamic acid tert-butyl ester 10 (89mg, 0.14mmol, 74%).

$$[M+H]^+ = 631.39$$

G. (S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide Trifluoroacetate 15

[4-({(S)-2-[(R)-3-(4-Ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionylamino}-methyl)-benzyl]-carbamic acid tert-butyl ester (89mg, 0.13mmol) was dissolved in trifluoroacetic acid (20 mL). This solution was stirred at room temperature for one hour after which time the solvent was removed in vacuo to give a yellow oil. The residue was 20 purified by Prep HPLC (Sunfire prep C18 OBD column. 19x250mm, 10μ). 10 to 90% 0.1% TFA/MeCN into 0.1%TFA/H₂O over 35 min at 20 mL/min. Fractions combined and freeze dried to give a white solid identified as (S)-N-(4-aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide trifluoroacetate (38mg, 0.056mmol, 42%)

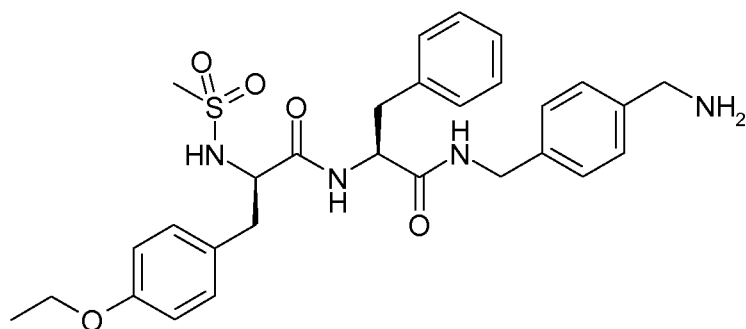
25 $[M+H]^+ = 531.31$

¹H NMR: (CD₃OD) 1.02 (3H, t, J=7.7Hz), 1.42 (3H, t, J=7.0Hz), 2.13-2.21 (2H, m), 2.71-2.77 (1H, m), 2.81-2.92 (2H, m), 3.12-3.16 (1H, m), 4.05 (2H, q, J=6.9Hz), 4.13 (2H, s), 4.37-4.50 (3H, m), 4.57-4.69 (1H, m), 6.82 (2H, d, J=8.6Hz), 7.05 (2H, d, J=8.6Hz), 7.17-7.19 (2H, m), 7.24-7.31 (5H, m), 7.41 (2H, d, J= 8.1Hz).

30

EXAMPLE 2

(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-2-methanesulfonylamino-propionamide



A. (R)-N-[(S)-1-(4-Cyano-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-2-methanesulfonylamino-propionamide

(R)-2-Amino-N-[(S)-1-(4-cyano-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-propionamide hydrochloride (150mg, 0.30mmol) was dissolved in CH_2Cl_2 (20 mL). This solution was cooled to 0°C . Methanesulfonyl chloride (37mg, 0.33mmol) was added followed triethylamine (90mg, 0.89mmol). After 18 hrs at 0°C to room temperature reaction mixture was diluted with chloroform (50 mL) and washed with NaHCO_3 (1x20 mL), water (1x20 mL), brine (1x20 mL), dried (Na_2SO_4) and evaporated in vacuo giving a yellow oil. The residue was purified by flash chromatography (silica), eluant 2%MeOH, 98% CHCl_3 , fractions combined and evaporated in vacuo to give a white solid identified as (R)-N-[(S)-1-(4-cyano-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-2-methanesulfonylamino-propionamide (110mg, 0.20mmol, 68%).

$[\text{M}+\text{H}]^+ = 549.11$

B. [4-((S)-2-[(R)-3-(4-Ethoxy-phenyl)-2-methanesulfonylamino-propionylamino]-3-phenyl-propionylamino)-methyl]-benzyl]-carbamic acid tert-butyl ester

(R)-N-[(S)-1-(4-Cyano-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-2-methanesulfonylamino-propionamide (110mg, 0.20mmol) was dissolved in methanol (50 mL). This solution was cooled to 0°C . Nickel (II) chloride hexahydrate (4.8mg, 0.02mmol) and di-tertbutyl dicarbonate (88mg, 0.4mmol) were added followed by sodium borohydride (53mg, 1.4mmol) portionwise. The reaction mixture was stirred at 0°C to room temp for 18hrs. The MeOH was removed by evaporation. The residue was dissolved in CHCl_3 (70 mL), washed with sat NaHCO_3 (1x30 mL), water (1x30 mL), brine (1x30 mL), dried (Na_2SO_4) and evaporated in vacuo to give a yellow oil. Purified by flash chromatography, eluant 2% MeOH, 98% CHCl_3 to give white solid identified as [4-((S)-2-[(R)-3-(4-ethoxy-phenyl)-2-methanesulfonylamino-

propionylamino]-3-phenyl-propionylamino}-methyl)-benzyl]-carbamic acid tert-butyl ester (86mg, 0.13mmol, 66%).

$[M+H]^+ = 653.23, 675.19 (M+Na)$.

5 **C. (R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-2-methanesulfonylamino-propionamide Trifluoroacetate**

[4-({(S)-2-[(R)-3-(4-Ethoxy-phenyl)-2-methanesulfonylamino-propionylamino]-3-phenyl-propionylamino}-methyl)-benzyl]-carbamic acid tert-butyl ester (86mg, 0.13mmol) was treated with trifluoroacetic acid (20 mL). After one hour at room temp the solvent was evaporated in vacuo. The residue was purified by Prep HPLC (Sunfire prep C18 OBD column. 19x250mm, 10 μ). 10 to 90% 0.1% TFA/MeCN into 0.1% TFA/H₂O over 35 min at 20 mL/min. Fractions combined and freeze dried to give a white solid identified as (R)-N-[(S)-1-(4-aminomethyl-benzylcarbamoyle)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-2-methanesulfonylamino-propionamide trifluoroacetate (28mg, 0.042mmol, 32%)

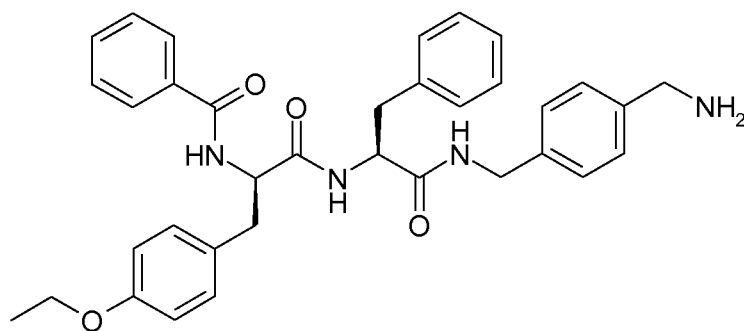
15 $[M+H]^+ = 553.08$

¹H NMR: (CD₃OD) 1.41 (3H, t, J=7.0Hz), 2.60 (3H, s), 2.69-2.75 (1H, m), 2.81-2.91 (2H, m), 3.09 (1H, dd, J=13.7, 6.5Hz), 4.04 (2H, q, J=7.0Hz), 4.13 (3H, m), 4.39 (2H, s), 4.62 (1H, dd, J=8.1, 6.6Hz), 6.87 (2H, d, J=8.6Hz), 7.13 (2H, d, J=8.6Hz), 7.23 (2H, t, J=6.6Hz), 7.25-7.32 (5H, m), 7.41 (2H, d, J= 8.1Hz).

20

EXAMPLE 3

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyle)-2-phenyl-ethylcarbamoyle]-2-(4-ethoxy-phenyl)-ethyl]-benzamide



25

A. {(S)-1-[4-(tert-Butoxycarbonylamino-methyl)-benzylcarbamoyle]-2-phenyl-ethyl}-carbamic acid benzyl ester

(S)-2-Benzyloxycarbonylamino-3-phenyl-propionic acid 2,5-dioxo-pyrrolidin-1-yl ester (4.25g, 10.72mmol) was dissolved in CH₂Cl₂ (100 mL). This solution was cooled to 0°C. 1-(N-Boc-Aminomethyl)-4-(aminomethyl)benzene (2.79g, 11.79mmol) was added followed by triethylamine (3.25g, 32.16mmol). After 18 hrs at 0°C to room temperature reaction mixture was diluted with chloroform (100 mL) and washed with NaHCO₃ (1x30 mL), water (1x30 mL), brine (1x30 mL), dried (Na₂SO₄) evaporated in vacuo giving a yellow oil. The residue was triturated with Pet. Ether (60-80°C) and EtOAc to give a white solid identified as {(S)-1-[4-(tert-butoxycarbonylamino-methyl)-benzylcarbamoyl]-2-phenyl-ethyl}-carbamic acid benzyl ester (3.88g, 7.49mmol, 70%).

10 $[M+H]^+ = 518.28, 540.32 (M+Na).$

B. {4-[(S)-2-Amino-3-phenyl-propionylamino)-methyl]-benzyl}-carbamic acid tert-butyl ester

{(S)-1-[4-(tert-Butoxycarbonylamino-methyl)-benzylcarbamoyl]-2-phenyl-ethyl}-carbamic acid benzyl ester (3.66g, 7.08mmol) was dissolved in methanol (200 mL). This solution was hydrogenated over 10% Pd/C (500mg) at atmospheric pressure and room temperature for one hour after which time the catalyst was filtered off through celite and the residue washed with methanol (30 mL), the combined filtrates were evaporated in vacuo to give a white solid identified as {4-[(S)-2-amino-3-phenyl-propionylamino)-methyl]-benzyl}-carbamic acid tert-

20 butyl ester (2.627g, 6.85mmol, 97%).

$[M+H]^+ = 384.37$

C. (R)-2-Amino-3-(4-ethoxy-phenyl)-propionic acid

(R)-2-Butoxycarbonylamino-3-(4-ethoxy-phenyl)-propionic acid (4.0g, 12.93mmol) was dissolved in 4M HCl in dioxan (150 mL). After one hour at room temperature the solvent was removed in vacuo to give a white solid identified as (R)-2-amino-3-(4-ethoxy-phenyl)-propionic acid hydrochloride (3.18g, 12.9mmol, 100%).

$[M+H]^+ = 210.18$

30 **D. (R)-2-Benzyloxycarbonylamino-3-(4-ethoxy-phenyl)-propionic acid**

(R)-2-Amino-3-(4-ethoxy-phenyl)-propionic acid hydrochloride (3.17g, 12.9mmol) was dissolved in a solution of sodium hydroxide (1.14g, 28.38mmol) in water (100 mL). Benzyl chloroformate (2.64g, 15.48mmol) in dioxan (100 mL) was added. The reaction mixture was

stirred at room temperature for 18 hrs after which time the dioxan was removed in vacuo. The aqueous residue was washed with diethyl ether (1x 100 mL), acidified to pH 2 with 1M HCl and extracted with chloroform (2x200 mL). The combined extracts were washed with water (1x50 mL), brine (1x50 mL), dried (Na₂SO₄) and evaporated in vacuo to give a white solid identified as
5 (R)-2-benzyloxycarbonylamino-3-(4-ethoxy-phenyl)-propionic acid (4.0g, 11.65mmol, 90%).
[M+H]⁺ = 344.20.

E. [(R)-1-[(S)-1-[4-(tert-Butoxycarbonylamino-methyl)-benzylcarbamoyl]-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-carbamic acid benzyl ester

10 {4-[(S)-2-Amino-3-phenyl-propionylamino]-methyl]-benzyl}-carbamic acid tert-butyl ester (2.63g, 6.86mmol) was dissolved in CH₂Cl₂ (100 mL) and DMF (5 mL). This solution was cooled to 0 °C. (R)-2-Benzyloxycarbonylamino-3-(4-ethoxy-phenyl)-propionic acid (2.59g, 7.54mmol) was added followed by HOBt (1.11g, 8.23mmol) and triethylamine (2.08g, 20.57mmol). Water soluble carbodiimide (1.45g, 7.54mmol) was then added. After 18 hrs at 0 °C
15 to room temperature reaction mixture was diluted with chloroform (200 mL) and washed with NaHCO₃ (1x50 mL), water (1x50 mL), brine (1x50 mL), dried (Na₂SO₄) and evaporated in vacuo giving a yellow oil. The residue was triturated with ethyl acetate and Pet. Ether (60-80°C) to give a white solid identified as [(R)-1-[(S)-1-[4-(tert-butoxycarbonylamino-methyl)-benzylcarbamoyl]-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-carbamic acid benzyl
20 ester (3.55g, 5.01mmol, 73%).
[M+H]⁺ = 709.34.

F. [4-[(S)-2-[(R)-2-Amino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionylamino]-methyl]-benzyl]-carbamic acid tert-butyl ester

25 [(R)-1-[(S)-1-[4-(tert-Butoxycarbonylamino-methyl)-benzylcarbamoyl]-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-carbamic acid benzyl ester (3.55g, 5.00mmol) was dissolved in methanol (200 mL). This solution was hydrogenated over 10% Pd/C (500mg) at atmospheric pressure and room temperature for one hour after which time the catalyst was filtered off through Celite and the residue washed with methanol (30 mL), the combined filtrates
30 were evaporated in vacuo to give a white solid identified as [4-[(S)-2-[(R)-2-amino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionylamino]-methyl]-benzyl]-carbamic acid tert-butyl ester (2.8g, 4.87mmol, 97%).
[M+H]⁺ = 575.37.

G. [4-((S)-2-[(R)-2-Benzoylamino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionylamino}-methyl)-benzyl]-carbamic acid tert-butyl ester

[4-((S)-2-[(R)-2-Amino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionylamino}-methyl)-benzyl]-carbamic acid tert-butyl ester (3.45g, 5.99mmol) was dissolved in dichloromethane (150 mL). Benzoyl chloride (1.01g, 7.19mmol) was added followed by triethylamine (1.82g, 17.98mmol). The reaction mixture was stirred at room temperature for 5 hrs and diluted with CHCl_3 (150 mL), this solution was washed with 0.3M KHSO_4 (1x50 mL), sat. NaHCO_3 (1x50 mL), water (1x50 mL), brine (1x50 mL), dried (Na_2SO_4) and evaporated in vacuo. The residue was triturated with Pet Ether (60-80°C) and EtOAc to give a white solid identified as [4-((S)-2-[(R)-2-benzoylamino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionylamino}-methyl)-benzyl]-carbamic acid tert-butyl ester (3.06g, 4.51mmol, 75%).
[M+H]⁺ = 679.34.

H. N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide hydrochloride

[4-((S)-2-[(R)-2-Benzoylamino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionylamino}-methyl)-benzyl]-carbamic acid tert-butyl ester (2.86g, 4.21mmol) was dissolved in 4M HCl in dioxan (150 mL). After one hour at room temperature the solvent was removed in vacuo. The residue was precipitated from ethanol to give a white solid identified as N-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide hydrochloride (2.1g, 3.41mmol, 81%).

[M+H]⁺ = 579.34

¹H NMR: (CD_3OD), 1.40 (3H, t, J= 6.9 Hz), 2.91-2.99 (3H, m), 3.14-3.19 (1H, m), 4.02 (2H, q, J= 6.9 Hz), 4.08 (2H, s), 4.41 (1H, d, J= 15.5 Hz), 4.51 (1H, d, J= 15.5 Hz), 4.66-4.69 (2H, m), 6.82 (2H, d, J= 8.4 Hz), 7.10 (2H, d, J= 8.2 Hz), 7.18-7.20 (2H, m), 7.25-7.38 (7H, m), 7.44-7.59 (3H, m), 7.72 (2H, d, J= 7.8 Hz).

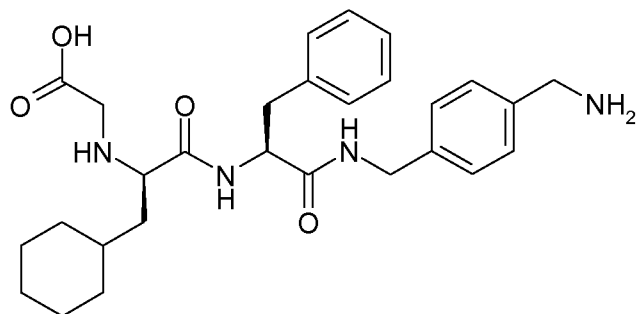
EXAMPLE 3b

30 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide hydrochloride

[4-({(S)-2-[(R)-2-Benzoylamino-3-(4-ethoxy-phenyl)-propionylamino]-3-phenyl-propionylamino}-methyl)-benzyl]-carbamic acid tert-butyl ester (10.0g, 14.7mmol) was stirred in hydrogen chloride / ethyl acetate (3.7M, 250 mL) at room temperature. After two hours the mixture was filtered, washed with ethyl acetate (2 x 50 mL) and dried to afford a solid (7.9g). A portion of the solid (0.106g) was suspended in a mixture of acetonitrile (2.1 mL) and water (0.32 mL), stirred, and heated to 77°C. Additional aliquots of water (0.05 mL) were added successively to the mixture until dissolution was observed. The stirred mixture was then cooled to room temperature overnight. The resulting solid was isolated by filtration and dried in vacuo at 40°C to afford N-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide hydrochloride (0.067g, 3.41mmol, 81%). ¹H NMR (CD₃OD) was identical to that of Example 3, Step H.

EXAMPLE 4

{(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino}-acetic acid



A. [(S)-1-(4-Cyano-benzylcarbamoyl)-2-phenyl-ethyl]-carbamic acid tert-butyl ester

4-Aminomethylbenzonitrile hydrochloride (1.53g, 9.1mmol) was dissolved in CH₂Cl₂ (100 mL). This solution was cooled to 0°C. (S)-2-tert-Butoxycarbonylamino-3-phenylpropionic acid 2,5-dioxo-pyrrolidin-1-yl ester (3.00g, 8.3mmol) was added followed triethylamine (2.51g, 25mmol). After 18 hrs at 0°C to room temperature reaction mixture was diluted with chloroform (100 mL) and washed with NaHCO₃ (1x30 mL), water (1x30 mL), brine (1x30 mL), dried (Na₂SO₄) and evaporated in vacuo giving a yellow oil. The residue was crystallised from EtOAc/Pet. Ether (60-80°C) to give a white solid identified as [(S)-1-(4-cyano-benzylcarbamoyl)-2-phenyl-ethyl]-carbamic acid tert-butyl ester (2.71g, 7.1mmol, 86%).

[M+H]⁺ = 380.13

B. (S)-2-Amino-N-(4-cyano-benzyl)-3-phenyl-propionamide Hydrochloride

[(S)-1-(4-Cyano-benzylcarbamoyl)-2-phenyl-ethyl]-carbamic acid tert-butyl ester

(2.71g, 7.1mmol) was treated with 4M HCl/dioxan (150 mL). After one hour at room temperature the solvent was removed to give a white solid identified as (S)-2-amino-N-(4-cyano-benzyl)-3-phenyl-propionamide hydrochloride (2.24g, 7.1mmol, 99%).

$$[M+H]^+ = 280.14$$

C. {(R)-1-[(S)-1-(4-Cyano-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethyl}-carbamic acid tert-butyl ester

(S)-2-Amino-N-(4-cyano-benzyl)-3-phenyl-propionamide hydrochloride (500mg, 1.58mmol) was dissolved in CH₂Cl₂ (30 mL) and DMF (3 mL). This solution was cooled to 0 °C. Boc-DCha-OH (473mg, 1.74mmol) was added followed by HOBt (257mg, 1.74mmol) and triethylamine (481mg, 4.75mmol). Water soluble carbodiimide (339mg, 1.74mmol) was then added. After 18 hrs at 0 °C to room temperature reaction mixture was diluted with chloroform (100 mL) and washed with NaHCO₃ (1x30 mL), water (1x30 mL), brine (1x30 mL), dried (Na₂SO₄) and evaporated in vacuo giving a yellow oil. The residue was purified by flash chromatography (silica), eluent 60% cyclohexane, 40% EtOAc, fractions combined and evaporated in vacuo to give a foamy white solid identified as {(R)-1-[(S)-1-(4-cyano-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethyl}-carbamic acid tert-butyl ester (799mg, 1.50mmol, 95%).

$$[M+H]^+ = 533.18$$

D. (R)-2-Amino-N-[(S)-1-(4-cyano-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-propionamide Hydrochloride

{(R)-1-[(S)-1-(4-Cyano-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethyl}-carbamic acid tert-butyl ester (799mg, 1.5mmol) was treated with 4M HCl/dioxan (50 mL). After one hour at room temperature the solvent was removed to give a white solid identified as (R)-2-amino-N-[(S)-1-(4-cyano-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-propionamide hydrochloride (703mg, 1.5mmol, 100%).

$$[M+H]^+ = 433.06$$

E. {(R)-1-[(S)-1-(4-Cyano-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino}-acetic acid tert-butyl ester

(R)-2-Amino-N-[(S)-1-(4-cyano-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-propionamide hydrochloride (290mg, 0.62mmol) was dissolved in acetonitrile (10 mL). tert-Butylbromoacetate (144mg, 0.74mmol) was added followed diisopropylethylamine (160mg, 1.24mmol). The reaction mixture was stirred at 60°C for 2 days after which time it was diluted with chloroform (100 mL), washed with water (1x30 mL), brine (1x30 mL), dried (Na₂SO₄) and evaporated in vacuo giving a yellow oil. The residue was purified by flash chromatography (silica), eluent 25%Pet. Ether (60-80°C), 75% EtOAc, fractions combined and evaporated in vacuo to give a colourless oil identified as {(R)-1-[(S)-1-(4-cyano-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino}-acetic acid tert-butyl ester (240mg, 0.44mmol, 71%).

10 [M+H]⁺ = 547.30.

F. ((R)-1-[(S)-1-[4-(tert-Butoxycarbonylamino-methyl)-benzylcarbamoyl]-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino)-acetic acid tert-butyl ester

{(R)-1-[(S)-1-(4-Cyano-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino}-acetic acid tert-butyl ester (240mg, 0.44mmol) was dissolved in methanol (25 mL). This solution was cooled to 0°C. Nickel (II) chloride hexahydrate (10.4mg, 0.44mmol) and di-tertbutyl dicarbonate (192mg, 0.88mmol) were added followed by sodium borohydride (116mg, 3.1mmol) portionwise. The reaction mixture was stirred at 0°C to room temp for 3 days. The MeOH was removed by evaporation. The residue was dissolved in CHCl₃ (70 mL), washed with sat NaHCO₃ (1x30 mL), water (1x30 mL), brine (1x30 mL), dried (Na₂SO₄) and evaporated in vacuo to give a yellow oil. Purified by flash chromatography, eluant 40% Pet. Ether (60-80°C), 60% EtOAc to give white solid identified as ((R)-1-[(S)-1-[4-(tert-butoxycarbonylamino-methyl)-benzylcarbamoyl]-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino)-acetic acid tert-butyl ester (65mg, 0.10mmol, 23%).

25 [M+H]⁺ = 651.44.

G. {(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino}-acetic acid Ditrifluoroacetate

((R)-1-[(S)-1-[4-(tert-Butoxycarbonylamino-methyl)-benzylcarbamoyl]-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino)-acetic acid tert-butyl ester (65mg, 0.1mmol) was treated with trifluoroacetic acid (4 mL) and CH₂Cl₂ (2 mL). After one hour at room temp the solvent was evaporated in vacuo. The residue was purified by Prep HPLC (Sunfire prep C18 OBD column. 19x250mm, 10μ). 10 to 90% 0.1% TFA/MeCN into 0.1% TFA/H₂O over 35 min

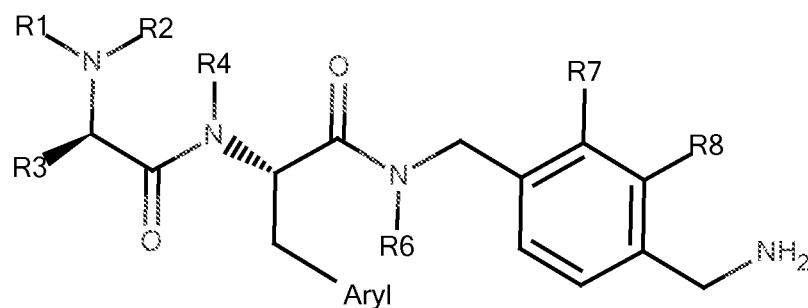
at 20 mL/min. Fractions combined and freeze dried to give a white solid identified as {(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino}-acetic acid ditrifluoroacetate (46mg, 0.064mmol, 64%).

$$[M+H]^+ = 495.28$$

- 5 ^1H NMR: (CD_3OD) 0.78-0.98 (2H, m), 1.10-1.25 (4H, m), 1.53-1.70 (7H, m), 2.97 (1H, dd, $J=14.0, 10.5\text{Hz}$), 3.25 (1H, dd, $J=14.1, 5.2\text{Hz}$), 3.74 (2H, s), 4.01 (1H, dd, $J=8.1, 6.1\text{Hz}$), 4.15 (2H, s), 4.47 (2H, s), 4.76 (1H, dd, $J=10.5, 5.2\text{Hz}$), 7.28-7.38 (7H, m), 7.45 (2H, d, $J= 8.2\text{Hz}$), 8.83 (1H, t, $J= 5.9\text{Hz}$).

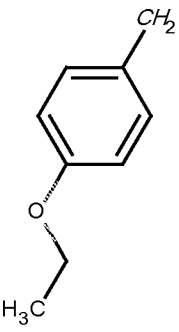
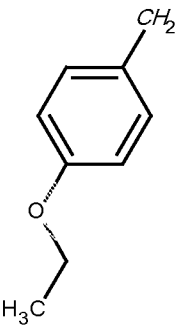
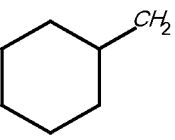
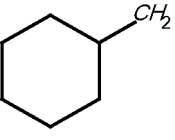
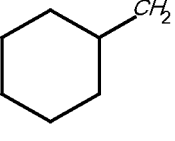
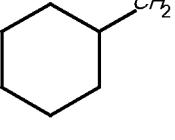
- 10 Compounds in Tables 1 to 5 were synthesised as described for Examples 1 to 4 (above) and 199 to 201 (below).

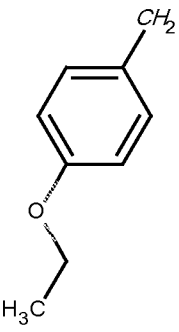
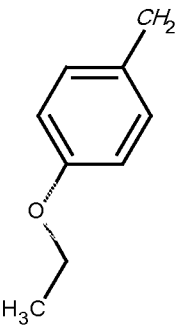
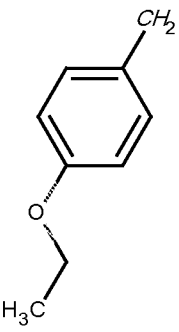
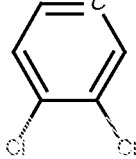
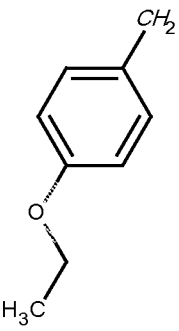
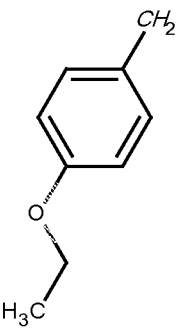
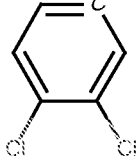
Table 1

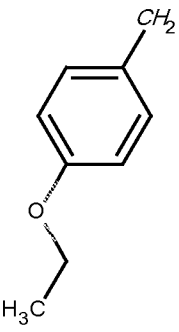
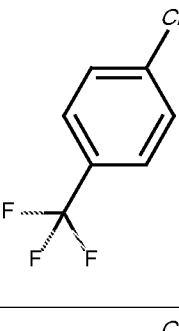
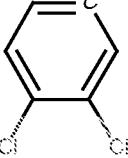
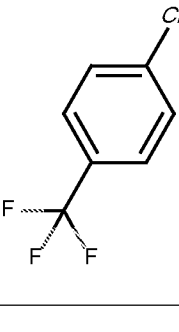
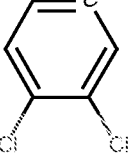
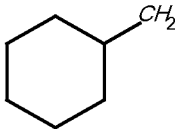
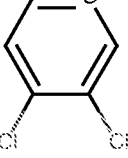
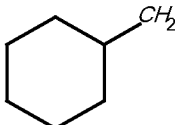
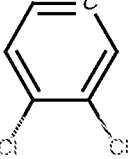


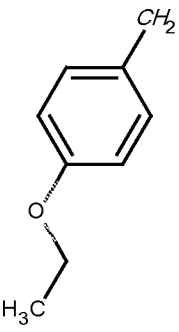
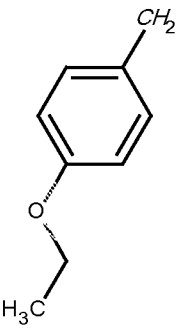
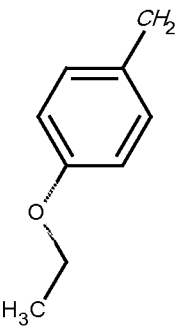
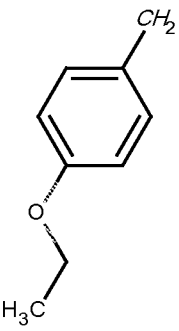
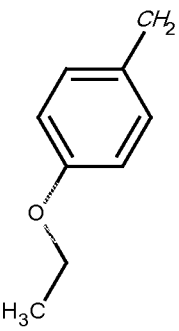
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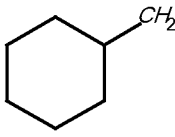
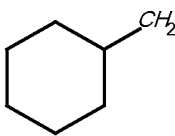
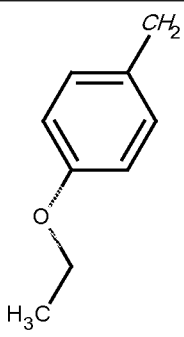
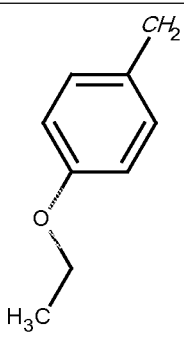
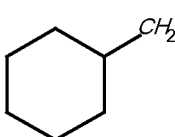
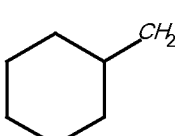
Ex. No.	R1	R2	R3	R4	Aryl	R6	R7	R8	m/z
5	H	H		H	phenyl	H	H	H	475.3

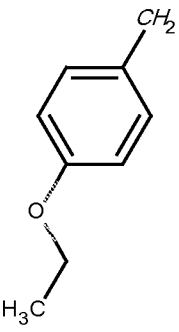
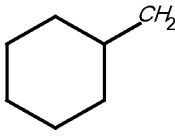
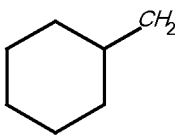
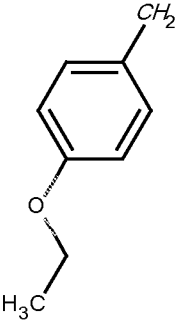
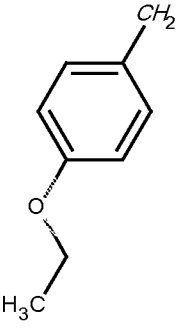
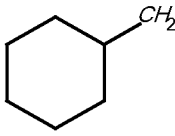
6	H	H		H	phenyl	H	H	F	493.3
7	CH ₃ CH ₂ CO	H		H	phenyl	H	H	F	549.3
8	H	H		H	phenyl	H	H	H	315.2
9	CH ₃	CH ₃		H	phenyl	H	H	H	465.3
10	CH ₃ CH ₂ CO	H		H	phenyl	H	H	H	493.2
11	CH ₃ SO ₂	H		H	phenyl	H	H	H	515.2

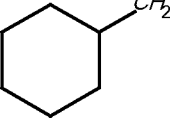
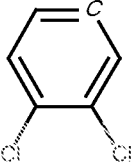
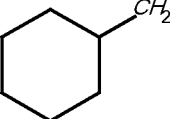
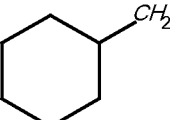
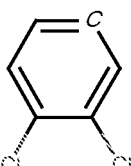
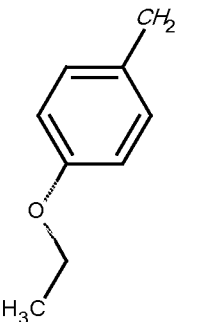
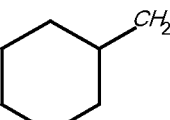
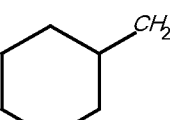
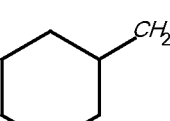
12	$\text{CH}_3\text{CH}_2\text{CO}$	H		H	phenyl	H	Cl	H	565.1
13	CH_3	CH_3		H	phenyl	H	H	H	503.2
14	H	H		H		H	H	H	543.2 and 545.2
15	PhSO_2	H		H	phenyl	H	H	H	615.2
16	$\text{CH}_3\text{CH}_2\text{CO}$	H		H		H	H	H	599.1 and 601.2

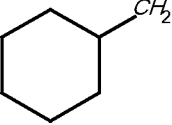
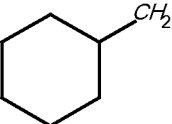
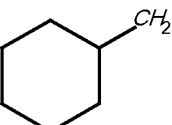
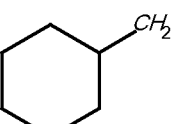
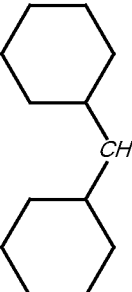
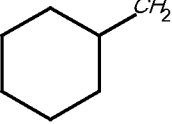
17	HOOCCH ₂	H		H	phenyl	H	H	H	533.2
18	H	H		H		H	H	H	567.1 and 569.1
19	CH ₃ CH ₂ CO	H		H		H	H	H	623.2 and 625.2
20	H	H		H		H	H	H	505.2 and 507.2
21	CH ₃ CH ₂ CO	H		H		H	H	H	561.2 and 563.2
22	PhCO	H	CH ₃	H	phenyl	H	H	H	459.2

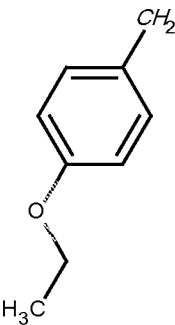
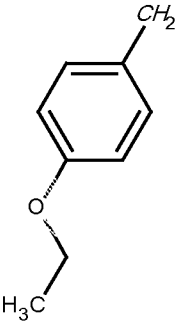
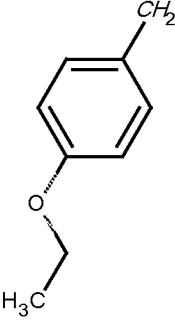
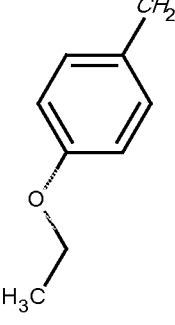
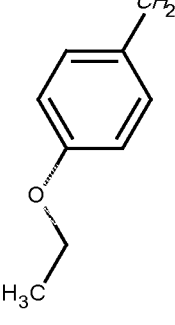
23	CH ₃ CH ₂ CO	H		H	phenyl	H	H	Cl	565.1
24	CH ₃ CH ₂ CO	H		H	phenyl	CH ₃	H	H	545.3
25	H	H		CH 3	phenyl	H	H	H	489.2
26	CH ₃ CH ₂ CO	H		CH 3	phenyl	H	H	H	545.2
27	CH ₃	H		H	phenyl	H	H	H	489.3

28	HOOCCH ₂	H	(CH ₃) ₂ CHCH ₂	H	phenyl	H	H	H	455.3
29	HOOCCH ₂	CH ₃		H	phenyl	H	H	H	509.3
30	HOOCCH ₂	H		CH 3	phenyl	H	H	H	509.3
31	CH ₃ CH ₂ CO	CH ₃		H	phenyl	H	H	H	545.3
32	CH ₃ CH ₂ CO	H		CH 3	phenyl	H	H	F	563.3
33	CH ₃	H		H	phenyl	H	H	H	451.4
34	HOOCCH ₂	H		CH 3	phenyl	H	H	F	527.3

35	n-Pr	H		CH 3	phenyl	H	H	H	531.4
36	HOOCCH ₂	H		H	phenyl	CH ₃	H	H	509.3
37	n-Pr	H		H	phenyl	H	H	H	479.4
38	PhCO	H		CH 3	phenyl	H	H	H	593.3
39	CH ₃ CO	H		CH 3	phenyl	H	H	H	531.3
40	CH ₃	H		H	phenyl	H	H	F	469.3

41	CH ₃	H		H		H	H	F	537.2 and 539.2
42	HOOCCH ₂	CH ₃		H	phenyl	H	H	F	527.3
43	HOOCCH ₂	CH ₃		H		H	H	F	595.3 and 597.3
44	(CH ₃) ₂ CHCO	H		CH 3	phenyl	H	H	H	559.3
45	n-Pr	CH ₃		H	phenyl	H	H	H	493.4
46	CH ₃ OCOCH ₂	H		H	phenyl	H	H	H	509.3
47	CH ₃ OCOCH ₂	CH ₃		H	phenyl	H	H	H	523.3

48	NH ₂ COCH ₂	CH ₃		H	phenyl	H	H	H	508.3
49	HOCH ₂ CH ₂	CH ₃		H	phenyl	H	H	H	495.4
50	NH ₂ COCH ₂	H		H	phenyl	H	H	H	494.3
51	HOCH ₂ CH ₂	H		H	phenyl	H	H	H	481.3
52	H	H		H	phenyl	H	H	H	519.3
53	n-Pr	n-Pr		H	phenyl	H	H	H	521.3

54	1-NaphCO	H		H	phenyl	H	H	H	629.3
55	4-Cl-PhCO	H		H	phenyl	H	H	H	613.3
56	4-CF ₃ -PhCO	H		H	phenyl	H	H	H	647.3
57	4-Ph-PhCO	H		H	phenyl	H	H	H	655.3
58	2,4-diCl-PhCO	H		H	phenyl	H	H	H	647.3

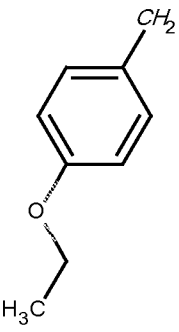
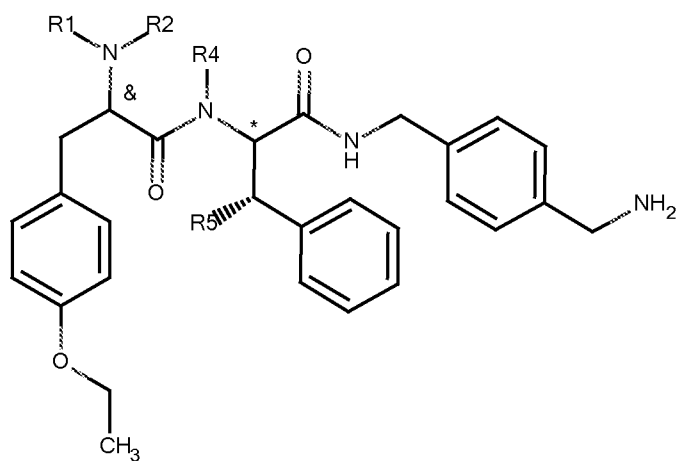
59	3,4-diF- PhCO	H		H	phenyl	H	H	H	615.3
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Table 2



5

Example No.	*	&	R1	R2	R4	R5	m/z
60	separated but not confirmed	R	H	H	H	OH	491.2
61		R	H	H	H	OH	491.2
62	separated but not confirmed	R	CH ₃ CH ₂ CO	H	H	OH	547.2
63		R	CH ₃ CH ₂ CO	H	H	OH	547.3
64	separated but not confirmed	R	CH ₃ CH ₂ CO	H	CH ₃ CH ₂	H	559.3
65		R	CH ₃ CH ₂ CO	H	CH ₃ CH ₂	H	559.3
66	S	S	CH ₃ CH ₂ CO	CH ₃	H	H	545.3

Table 3

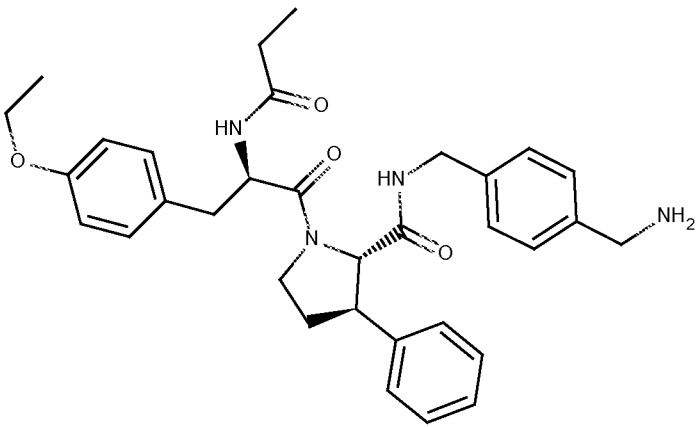
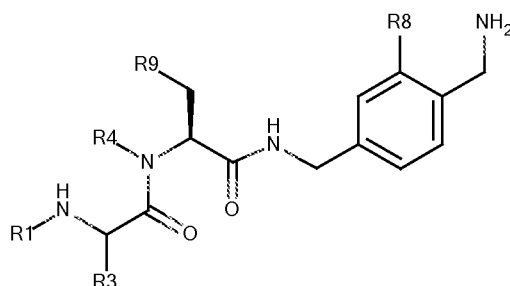
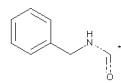
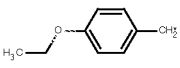
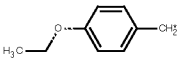
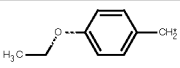
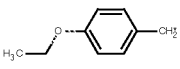
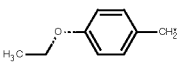
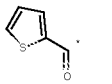
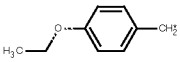
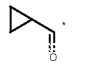
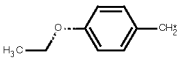
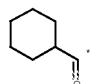
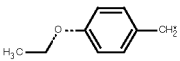
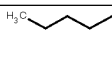
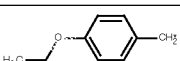
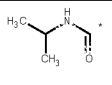
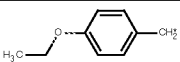
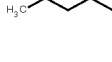
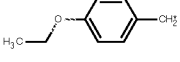
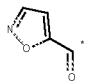
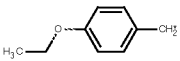
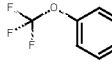
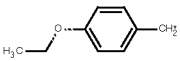
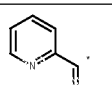
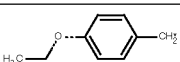
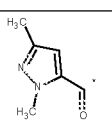
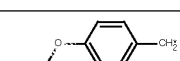
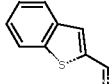
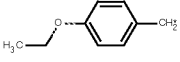
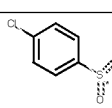
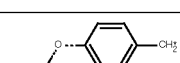
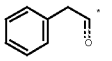
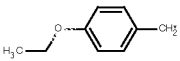
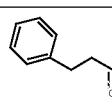
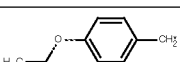
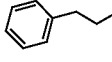
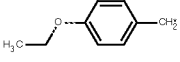
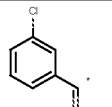
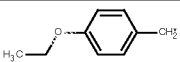
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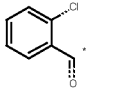
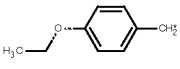
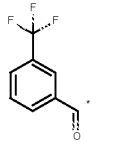
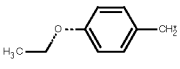
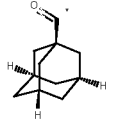
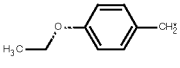
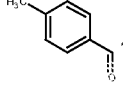
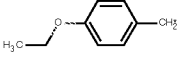
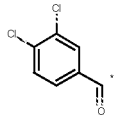
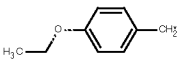
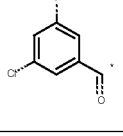
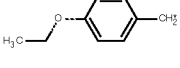
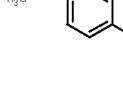
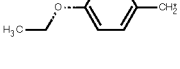
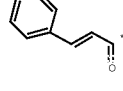
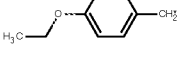
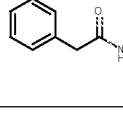
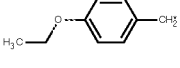
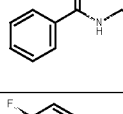
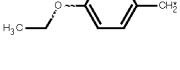
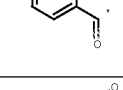
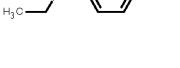
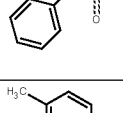
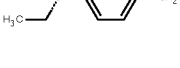
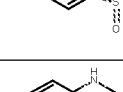
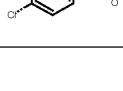
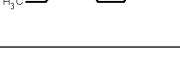
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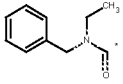
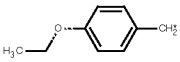
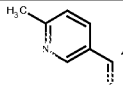
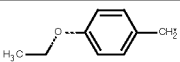
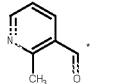
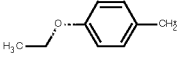
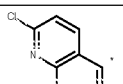
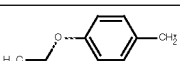
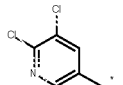
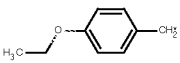
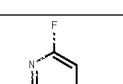
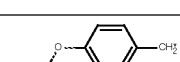
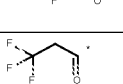
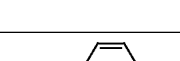
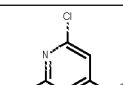
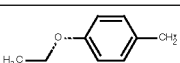
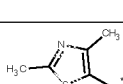
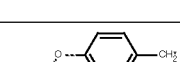
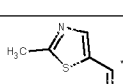
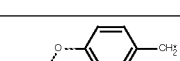
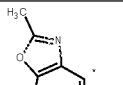
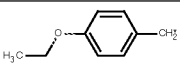
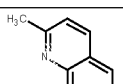
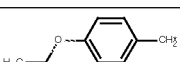
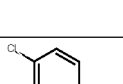
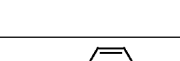
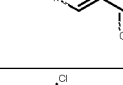


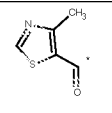
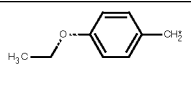
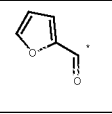
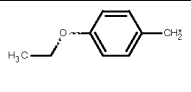
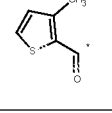
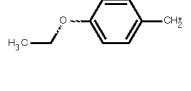
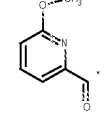
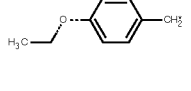
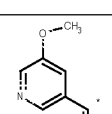
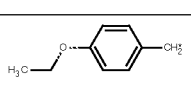
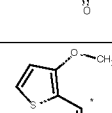
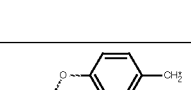
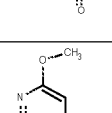
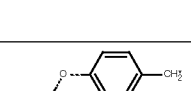
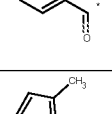
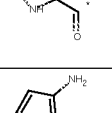
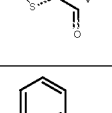
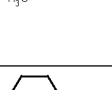
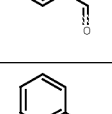
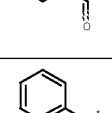
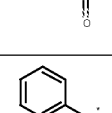
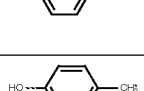
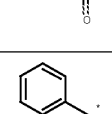
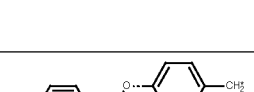
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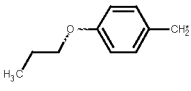
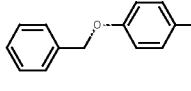
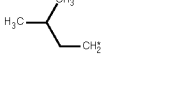
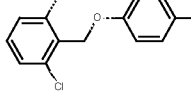
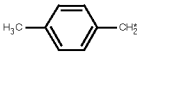
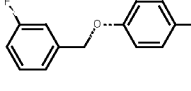
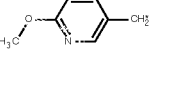
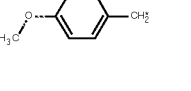
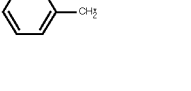
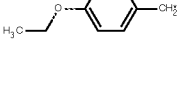
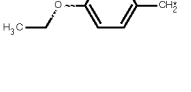
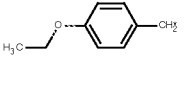
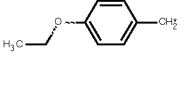
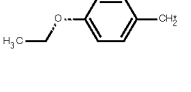
Example No	R1	R3	R4	R8	R9	m/z
68			H	H		607.99
69			H	H		594.04
70			H	H		580.19
71			H	H		580.18
72			H	H		585.15

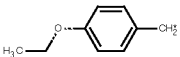
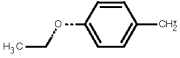
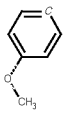
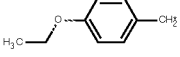
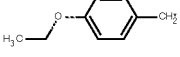
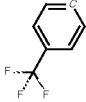
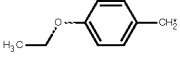
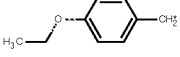
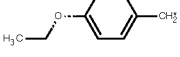
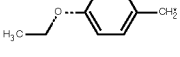
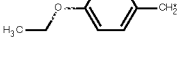
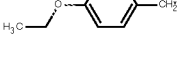
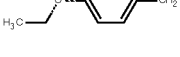
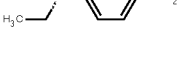
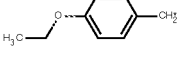
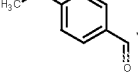
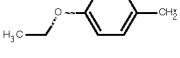
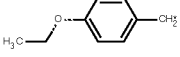
73			H	H		585.15
74			H	H		542.95
75			H	H		585
76			H	H		572.99
77			H	H		560.28
78			H	H		602.26
79			H	H		570.13
80			H	H		663.12
81			H	H		580.18
82			H	H		597.2
83			H	H		635.14
84			H	H		649.13
85			H	H		594.10
86			H	H		607.18
87			H	H		621.2
88			H	H		613.1

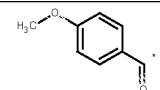
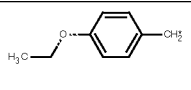
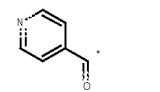
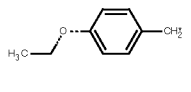
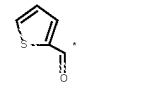
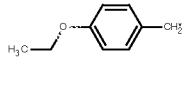
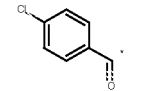
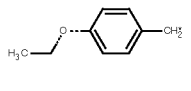
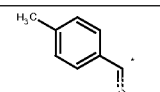
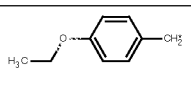
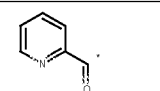
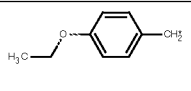
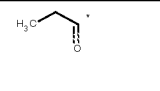
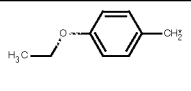
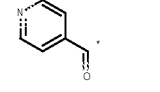
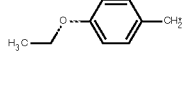
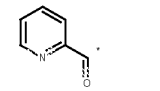
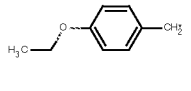
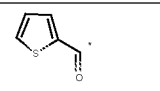
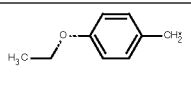
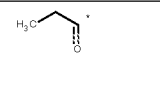
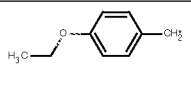
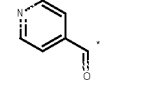
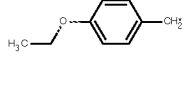
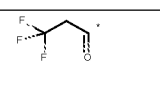
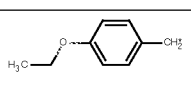
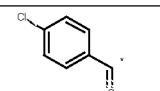
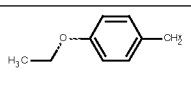
89			H	H		613.1
90			H	H		647.1
91			H	H		637.21
92			H	H		594.12
93			H	H		647.05
94			H	H		647.05
95			H	H		610.11
96			H	H		605.17
97			H	H		650.16
98			H	H		636.16
99			H	H		597.14
100			H	H		629.15
101			H	H		686.14
102			H	H		628.1

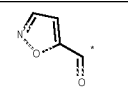
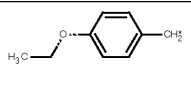
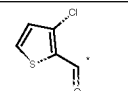
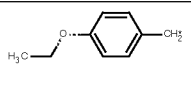
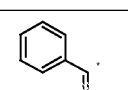
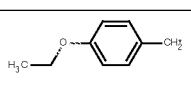
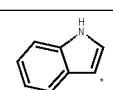
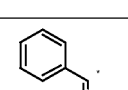
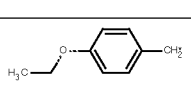
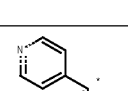
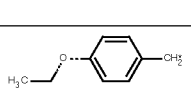
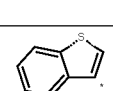
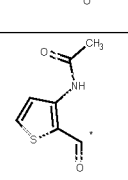
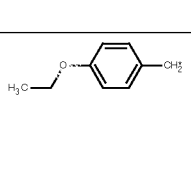
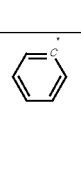
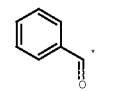
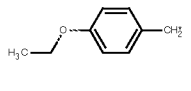
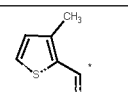
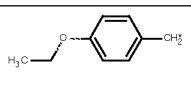
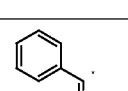
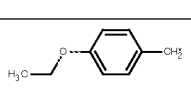
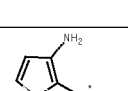
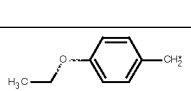
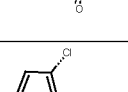
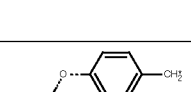
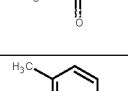
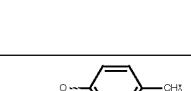
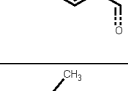
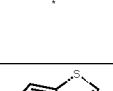
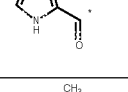
103			H	H		636.1
104			H	H		594.04
105			H	H		594.06
106			H	H		648.09
107			H	H		648.07
108			H	H		633.87
109			H	H		585.12
110			H	H		649.92 (M+Na)
111			H	H		613.95
112			H	H		599.92
113			H	H		597.75
114			H	H		644.13
115			H	H		664.06
116			H	H		618.97

117			H	H		600.06
118			H	H		569.04
119			H	H		599.04
120			H	H		610.01
121			H	H		610.06
122			H	H		615.05
123			H	H		610.07
124			H	H		582.06
125			H	H		600.57
126			H	H		541.21
127			H	H		569.11
128			H	H		675.1
129			H	H		551.15
130			H	H		659.12

131			H	H		594.05
132			H	H		642.12
133			H	H		501.21
134			H	H		709.12
135			H	H		548.94
136			H	H		659.12
137			H	H		566.09
138			H	H		565.1
139			H	H		535.11
140			H	H		580.06
141			H	H		647.04 and 649.06
142			H	H		612.94
143			H	H		596.93
144			H	H		646.96

145			H	H		580.13
146			H	H		608.97
147			H	H		579.95
148			H	H		646.97
149			H	H		618.93 (M+Na)
150			H	H		606.88 (M+Na)
151			H	H		584.98
152			H	H		586.03
153			H	H		635.05
154			H	F		596.93
155			H	Cl		612.92
156			H	CF ₃		646.9
157			H	H		586.26
158			H	H		610.02
159			H	Cl		614.29

160			H	H		610.35
161			H	H		616.03
162			H	H		585.98
163			H	H		614.05
164			H	H		616.05
165			H	H		648.04 and 650.04
166			H	H		532.04
167			H	F		598.03
168			H	F		598.04
169			H	H		653.00 and 654.99
170			H	H		532.04
171			H	H		648.02 and 650.04
172			H	H		653.02
173			H	H		614.30

174			H	H		571.02
175			H	H		620.26
176			H	H		641.37 (M+Na)
177			H	H		569.48
178			H	H		636.36
179			H	H		642.38
180			H	H		597.35
181			H	H		600.32
182			H	Me		593.29
183			H	H		607.26
184			H	H		625.22
185			H	H		600.32
186			H	H		638.19
187			H	H		588.09

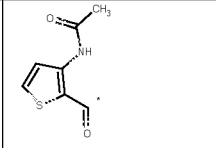
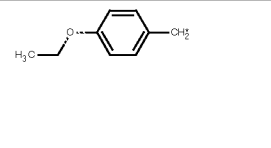
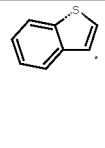
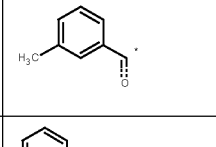
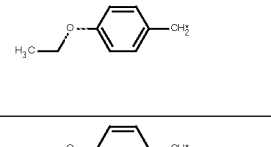
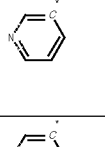
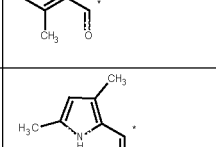
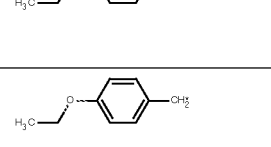
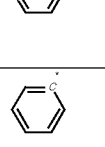
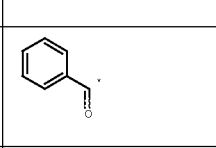
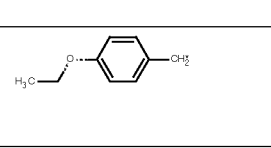
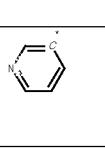
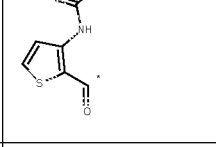
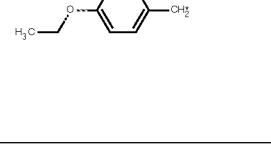
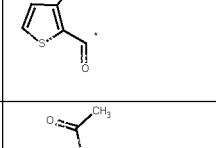
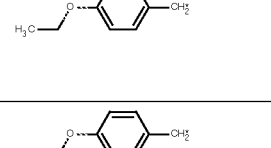
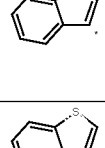
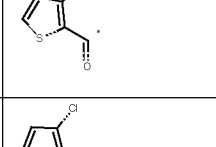
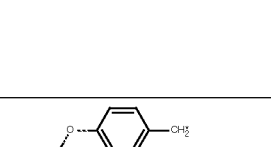
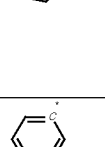
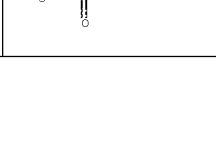
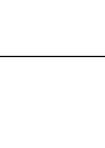
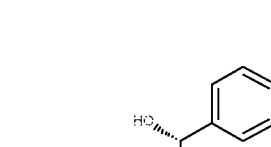
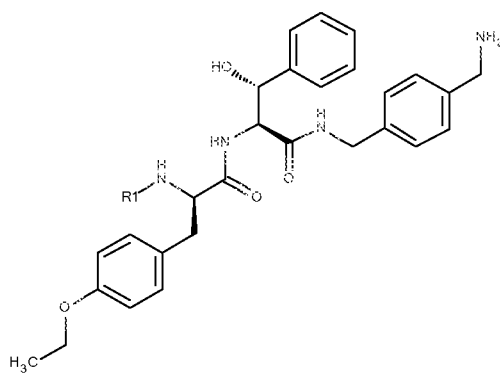
188			H	H		698.15
189			H	H		594.02
190			H	H		594.07
191			H	H		596.18
192			H	Me		594.05
193			H	H		648.16
194			H	H		656.47
195			H	H		605.52
196			Me	H		633.18

Table 5



Example No.	R1	m/z
197		594.48
198		635.06

Table 6

Example No	Name
5	(R)-2-Amino-N-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-propionamide
6	(R)-2-Amino-N-[(S)-1-(4-aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-propionamide
7	(S)-N-(4-Aminomethyl-3-fluoro-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide
8	(R)-2-Amino-N-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-propionamide
9	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-2-dimethylamino-propionamide
10	(S)-N-(4-Aminomethyl-benzyl)-2-((R)-3-cyclohexyl-2-propionylamino-propionylamino)-3-phenyl-propionamide
11	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-2-methanesulfonylamino-propionamide
12	(S)-N-(4-Aminomethyl-2-chloro-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide
13	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-2-dimethylamino-3-(4-ethoxy-phenyl)-propionamide
14	(R)-2-Amino-N-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethyl]-3-(4-ethoxy-phenyl)-propionamide

15	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-2-benzenesulfonylamino-3-(4-ethoxy-phenyl)-propionamide
16	(S)-N-(4-Aminomethyl-benzyl)-3-(3,4-dichloro-phenyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-propionamide
17	[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethylamino]-acetic acid
18	(R)-2-Amino-N-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethyl]-3-(4-trifluoromethyl-phenyl)-propionamide
19	((R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethyl]-2-propionylamino-3-(4-trifluoromethyl-phenyl)-propionamide
20	(R)-2-Amino-N-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethyl]-3-cyclohexyl-propionamide
21	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethyl]-3-cyclohexyl-2-propionylamino-propionamide
22	N- {(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-ethyl} -benzamide
23	(S)-N-(4-Aminomethyl-3-chloro-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide
24	(S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-N-methyl-3-phenyl-propionamide
25	(R)-2-Amino-N-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-N-methyl-propionamide
26	(S)-N-(4-Aminomethyl-benzyl)-2- {[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionyl]-methyl-amino} -3-phenyl-propionamide
27	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-2-methylamino-propionamide
28	{(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-3-methyl-butylamino}-acetic acid
29	((R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethyl)-methyl-amino)-acetic acid
30	((R)-1- {[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-methyl-carbamoyl}-2-cyclohexyl-ethylamino)-acetic acid

31	(S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-(methyl-propionyl-amino)-propionylamino]-3-phenyl-propionamide
32	(S)-N-(4-Aminomethyl-3-fluoro-benzyl)-2- {[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionyl]-methyl-amino }-3-phenyl-propionamide
33	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-2-methylamino-propionamide
34	((R)-1- {[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethyl]-methyl-carbamoyl }-2-cyclohexyl-ethylamino)-acetic acid
35	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-N-methyl-2-propylamino-propionamide
36	((R)-1- {[(S)-1-[(4-Aminomethyl-benzyl)-methyl-carbamoyl]-2-phenyl-ethylcarbamoyl }-2-cyclohexyl-ethylamino)-acetic acid
37	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-2-propylamino-propionamide
38	N-[(R)-1- {[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-methyl-carbamoyl }-2-(4-ethoxy-phenyl)-ethyl]-benzamide
39	(R)-2-Acetylamino-N-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-N-methyl-propionamide
40	(R)-N-[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-2-methylamino-propionamide
41	(R)-N-[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethyl]-3-cyclohexyl-2-methylamino-propionamide
42	({ (R)-1-[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethyl }-methyl-amino)-acetic acid
43	({ (R)-1-[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-cyclohexyl-ethyl }-methyl-amino)-acetic acid
44	N-[(R)-1- {[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-methyl-carbamoyl }-2-(4-ethoxy-phenyl)-ethyl]-isobutyramide
45	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-2-(methyl-propyl-amino)-propionamide
46	{ (R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino }-acetic acid methyl ester

47	{(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino}-acetic acid methyl ester
48	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-2-(carbamoylmethyl-methyl-amino)-3-cyclohexyl-propionamide
49	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-2-[(2-hydroxy-ethyl)-methyl-amino]-propionamide
50	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-2-(carbamoylmethyl-amino)-3-cyclohexyl-propionamide
51	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-2-(2-hydroxy-ethylamino)-propionamide
52	(R)-2-Amino-N-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3,3-dicyclohexyl-propionamide
53	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-cyclohexyl-2-dipropylamino-propionamide
54	Naphthalene-1-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
55	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-chloro-benzamide
56	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-trifluoromethyl-benzamide
57	Biphenyl-4-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
58	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2,4-dichloro-benzamide
59	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,4-difluoro-benzamide
60	(R)-2-Amino-N-[(1R,2S)-1-(4-aminomethyl-benzylcarbamoyl)-2-hydroxy-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-propionamide
61	(R)-2-Amino-N-[(1S,2S)-1-(4-aminomethyl-benzylcarbamoyl)-2-hydroxy-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-propionamide
62	(2R,3S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-hydroxy-3-phenyl-propionamide

63	(2S,3S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-hydroxy-3-phenyl-propionamide
64	(R)-N-(4-Aminomethyl-benzyl)-2- {[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionyl]-ethyl-amino} -3-phenyl-propionamide
65	(S)-N-(4-Aminomethyl-benzyl)-2- {[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionyl]-ethyl-amino} -3-phenyl-propionamide
66	(S)-N-(4-Aminomethyl-benzyl)-2-[(S)-3-(4-ethoxy-phenyl)-2-(methyl-propionyl-amino)-propionylamino]-3-phenyl-propionamide
67	(2S,3R)-1-[(R)-3-(4-Ethoxy-phenyl)-2-propionylamino-propionyl]-3-phenyl-pyrrolidine-2-carboxylic acid 4-aminomethyl-benzylamide
68	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-2-(3-benzyl-ureido) -3-(4-ethoxy-phenyl)-propionamide
69	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-2-(3-phenyl-ureido)-propionamide
70	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-nicotinamide
71	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide
72	Thiophene-3-carboxylic acid-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
73	Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
74	Cyclopropanecarboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
75	Cyclohexanecarboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
76	Hexanoic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
77	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-2-(3-isopropyl-ureido)-propionamide
78	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-2-(3-hexyl-ureido)-propionamide

79	Isoxazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
80	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-trifluoromethoxy-benzamide
81	Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
82	2,5-Dimethyl-2H-pyrazole-3-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
83	Benzo[b]thiophene-2-carboxylic acid[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
84	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-2-(4-chloro-benzenesulfonylamino)-3-(4-ethoxy-phenyl)-propionamide
85	(S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-phenylacetyl-amino-propionyl-amino]-3-phenyl-propionamide
86	(S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-(3-phenyl-propionyl-amino)-propionyl-amino]-3-phenyl-propionamide
87	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-phenyl-butyramide
88	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3-chloro-benzamide
89	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-chloro-benzamide
90	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3-trifluoromethyl-benzamide
91	Adamantane-1-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
92	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide
93	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,4-dichloro-benzamide
94	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,5-dichloro-benzamide

95	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide
96	(E)-N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3-phenyl-acrylamide
97	(S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-(2-phenylacetyl-amino-acetyl-amino)-propionyl-amino]-3-phenyl-propionamide
98	N- {[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethylcarbamoyl]-methyl} -benzamide
99	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-fluoro-benzamide
100	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-2-phenylmethanesulfonylamino-propionamide
101	(S)-N-(4-Aminomethyl-benzyl)-2- { (R)-3-(4-ethoxy-phenyl)-2-[2-(toluene-4-sulfonylamino)-acetyl-amino]-propionyl-amino } -3-phenyl-propionamide
102	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-2-[3-(4-chloro-phenyl)-ureido]-3-(4-ethoxy-phenyl)-propionamide
103	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-2-(3-benzyl-3-ethyl-ureido)-3-(4-ethoxy-phenyl)-propionamide
104	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-6-methyl-nicotinamide
105	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-methyl-nicotinamide
106	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2,6-dichloro-nicotinamide
107	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-5,6-dichloro-nicotinamide
108	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2,3,6-trifluoro-isonicotinamide
109	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,3,3-trifluoro-propionamide
110	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-chloro-6-methyl-isonicotinamide

111	2,4-Dimethyl-thiazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
112	2-Methyl-thiazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
113	2,5-Dimethyl-oxazole-4-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
114	2-Methyl-quinoline-6-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
115	7-Chloro-quinoline-3-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
116	3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
117	4-Methyl-thiazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
118	Furan-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
119	3-Methyl-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
120	6-Methoxy-pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
121	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-5-methoxy-nicotinamide
122	3-Methoxy-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
123	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-methoxy-isonicotinamide
124	3-Methyl-1H-pyrrole-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
125	3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
126	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethyl]-benzamide

127	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-chloro-phenyl)-ethyl]-benzamide
128	N-{(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-[4-(4-chloro-benzyloxy)-phenyl]-ethyl}-benzamide
129	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-hydroxy-phenyl)-ethyl]-benzamide
130	N-{(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-[4-(4-fluoro-benzyloxy)-phenyl]-ethyl}-benzamide
131	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-propoxy-phenyl)-ethyl]-benzamide
132	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-benzyloxy-phenyl)-ethyl]-benzamide
133	N-{(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-3-methyl-butyl}-benzamide
134	N-{(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-[4-(2,6-dichloro-benzyloxy)-phenyl]-ethyl}-benzamide
135	N-{(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-p-tolyl-ethyl}-benzamide
136	N-{(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-[4-(3-fluoro-benzyloxy)-phenyl]-ethyl}-benzamide
137	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(6-methoxy-pyridin-3-yl)-ethyl]-benzamide
138	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-methoxy-phenyl)-ethyl]-benzamide
139	N-{(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-phenyl-ethyl}-benzamide
140	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
141	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
142	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(4-chloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide

143	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(4-fluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
144	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3-trifluoromethyl-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
145	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
146	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(4-methoxy-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
147	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
148	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(4-trifluoromethyl-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
149	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3-fluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
150	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiophen-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
151	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
152	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
153	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
154	N-[(R)-1-[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
155	N-[(R)-1-[(S)-1-(4-Aminomethyl-3-chloro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
156	N-[(R)-1-[(S)-1-(4-Aminomethyl-3-trifluoromethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
157	Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiophen-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
158	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide

159	Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-3-chloro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
160	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide
161	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-difluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide
162	Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
163	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-chloro-benzamide
164	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide
165	Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
166	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethyl]-3-(4-ethoxy-phenyl)-2-propionylamino-propionamide
167	N-[(R)-1-[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide
168	Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
169	Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
170	(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethyl]-3-(4-ethoxy-phenyl)-2-propionylamino-propionamide
171	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide
172	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,3,3-trifluoro-propionamide
173	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-chloro-benzamide
174	Isoxazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide

175	3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
176	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(1H-indol-3-yl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
177	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(1H-imidazol-4-yl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
178	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide
179	3-Acetylamino-thiophene-2-carboxylic acid-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
180	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(2-fluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
181	3-Methyl-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
182	N-[(R)-1-[(S)-1-(4-Aminomethyl-3-methyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
183	3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
184	3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
185	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide
186	3-Methyl-1H-pyrrole-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
187	3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide

188	3-Acetylamino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
189	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3-methyl-benzamide
190	N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-methyl-benzamide
191	3,5-Dimethyl-1H-pyrrole-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
192	N-[(R)-1-[(S)-1-(4-Aminomethyl-3-methyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
193	3-Acetylamino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
194	3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
195	3-Acetylamino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide
196	3-Chloro-thiophene-2-carboxylic acid [(R)-1-{[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-methyl-carbamoyl}-2-(4-ethoxy-phenyl)-ethyl]-amide
197	N-[(R)-1-[(1S,2R)-1-(4-Aminomethyl-benzylcarbamoyl)-2-hydroxy-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide
198	3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(1S,2R)-1-(4-aminomethyl-benzylcarbamoyl)-2-hydroxy-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide

Table 7

NMR data of examples

Example No	Solvent	NMR (ppm)
5	CD ₃ OD	δ 1.42 (3H, t, J=7.0Hz), 2.80-2.91 (2H, m), 2.94-2.99 (1H, m), 3.08-3.13 (1H, m), 3.96-4.10 (3H, m), 4.14 (2H, s), 4.31-4.36 (1H, m), 4.47-4.52 (1H, m), 4.66-4.70 (1H, m), 6.88 (2H, d, J=8.6Hz), 7.02 (2H, d, J=8.6Hz), 7.25-7.40 (7H, m), 7.42 (2H, d, J= 8.1Hz), 8.73-8.76(1H, m)
6	CD ₃ OD	δ 1.42 (3H, t, J=7.0Hz), 2.80-2.89 (2H, m), 2.95-3.00 (1H, m), 3.08-3.13 (1H, m), 4.03-4.13 (3H, m), 4.21 (2H, s), 4.32-4.38 (1H, m), 4.46-4.51 (1H, m), 4.65-4.69 (1H, m), 6.88 (2H, d, J=8.6Hz), 7.03 (2H, d, J= 8.6Hz), 7.05-7.09 (2H, m), 7.26-7.37 (5H, m), 7.41-7.47 (1H, m), 8.78-8.80 (1H, m)
7	CD ₃ OD	δ 1.02 (3H, t, J=7.7Hz), 1.42 (3H, t, J=7.0Hz), 2.13-2.21 (2H, m), 2.71-2.77 (1H, m), 2.81-2.92 (3H, m), 3.12-3.16 (1H, m), 4.05 (2H, q, J=6.9Hz), 4.13 (2H, s), 4.37-4.50 (3H, m), 4.57-4.69 (1H, m), 6.82 (2H, d, J=8.6Hz), 7.05 (2H, d, J= 8.6Hz), 7.17-7.27 (4H, m), 7.30-7.35 (3H, m), 7.46-7.49 (1H, m)
8	CD ₃ OD	δ 0.82-0.94 (2H, m), 1.19-1.28 (4H, m), 1.42-1.48 (2H, m), 1.64-1.71 (5H, m), 2.92-2.98 (1H, m), 3.23-3.28 (1H, m), 3.88 (1H, q, J=7.6Hz), 4.15 (2H, s), 4.39 (1H, d, J=15.3Hz), 4.52 (1H, d, J=15.3Hz), 4.76-4.79 (1H, m), 7.33-7.37 (7H, m), 7.44 (2H, d, J= 8.1Hz)
9	CD ₃ OD	δ 0.84-1.03 (4H, m), 1.13-1.23 (2H, m), 1.52-1.77 (7H, m), 2.93 (6H, s), 2.96-3.02 (1H, m), 3.20-3.30 (1H, m), 3.81 (1H, q, J=3.9Hz), 4.15 (2H, s), 4.47 (2H, q, J=15.4Hz), 4.76 (1H, dd, J=5.3, 10.5Hz), 7.29-7.40 (7H, m), 7.45 (2H, d, J= 8.2Hz)
10	CD ₃ OD	δ 0.84-0.92 (2H, m), 1.03 (3H, t, J=7.7Hz), 1.11-1.20 (4H, m), 1.35-1.39 (2H, m), 1.64-1.67 (5H, m), 2.15-2.26 (2H, m), 2.87-2.93 (1H, m), 3.39-3.42 (1H, m), 4.14 (2H, s), 4.21-4.26 (1H, m), 4.41-4.42 (1H, m), 4.44-4.57 (1H, m), 4.69-4.75 (1H, m), 7.27-7.44 (9H, m), 8.08(1H, d, J=5.8Hz), 8.49 (1H, d, J= 8.3Hz), 8.63-8.70 (1H, m)
11	CD ₃ OD	δ 0.85-0.95 (2H, m), 1.21-1.40 (6H, m), 1.70-1.79 (5H, m), 2.85 (3H, s), 2.95 (1H, dd, J=10.0, 14.0Hz), 3.29 (1H, dd, J=5.5, 14.0Hz), 3.98 (1H, dd, J= 5.6, 8.8Hz), 4.14 (2H, s), 4.43 (2H, s), 4.71 (1H, dd, J=5.4, 10.0Hz), 7.27-7.35 (7H, m), 7.42 (2H, d, J= 8.1Hz)

12	CD ₃ OD	δ 1.04 (3H, t, J=7.5Hz), 1.40 (3H, t, J=7.0Hz), 2.15-2.23 (2H, m), 2.74-2.90 (3H, m), 2.97-3.21(2H, m), 3.98-4.13 (2H, m), 4.14 (2H, s), 4.35-4.58 (3H, m), 4.61-4.68 (1H, m), 6.82 (2H, d, J=8.6Hz), 7.04-7.08 (1H, m), 7.12-7.42 (9H, m), 7.55 (1H, d, J= 1.6Hz)
13	CD ₃ OD	δ 1.42 (3H, t, J=7.0Hz), 2.61-2.66 (1H, m), 2.83-2.88 (1H, m), 3.00 (6H, s), 3.02-3.08 (1H, m), 3.29 (1H, dd, J= 12.9, 4.9Hz), 3.98-4.06 (3H, m), 4.14 (2H, s), 4.19-4.21 (1H, m), 4.33-4.45 (2H, m), 6.87 (2H, d, J=8.6Hz), 6.89-7.05 (2H, m), 7.12 (2H, d, J= 8.6Hz), 7.22-7.27 (5H, m), 7.40 (2H, d, J= 8.2Hz), 8.61-8.64 (1H, m)
14	CD ₃ OD	δ 1.42 (3H, t, J=7.0Hz), 2.82-2.92 (2H, m), 3.03-3.09 (2H, m), 4.04-4.13 (3H, m), 4.15 (2H, s), 4.30-4.35 (1H, m), 4.48-4.53 (1H, m), 4.65-4.69 (1H, m), 6.87-6.91 (2H, m), 7.02-7.06 (2H, m), 7.15 (1H, dd, J=8.3, 2.0Hz), 7.29 (2H, d, J= 8.1Hz), 7.43-7.46 (4H, m), 8.74 (1H, t, J= 5.8Hz)
15	CD ₃ OD	δ 1.42 (3H, t, J=7.0Hz), 2.54-2.60 (1H, m), 2.67-2.72 (1H, m), 2.78-2.83 (1H, m), 3.06-3.11 (1H, m), 3.91-3.95 (1H, m), 3.98-4.04 (2H, m), 4.14 (2H, s), 4.36-4.49 (2H, m), 4.54-4.58 (1H, m), 6.69-6.72 (2H, m), 6.91 (2H, d, J= 8.6Hz), 7.19-7.21 (2H, m), 7.24-7.32 (5H, m), 7.41-7.45 (4H, m), 7.55-7.59 (1H, m), 7.64-7.66 (2H, m).
16	CD ₃ OD	δ 1.03 (3H, t, J=7.6Hz), 1.42 (3H, t, J=7.0Hz), 2.14-2.24 (2H, m), 2.75-2.94 (3H, m), 3.07-3.12 (1H, m), 4.04 (2H, q, J=7.0Hz), 4.14 (2H, s), 4.36-4.51 (3H, m), 4.61-4.65 (1H, m), 6.81-6.84 (2H, m), 7.03-7.09 (3H, m), 7.33 (2H, d, J= 8.1Hz), 7.38-7.43 (4H, m)
17	CD ₃ OD	δ 1.17 (3H, t, J=7.0Hz), 2.49-2.54 (1H, m), 2.74-2.83 (3H, m), 3.32 (2H, s), 3.81 (2H, q, J=7.0Hz), 3.88 (2H, s), 3.90-3.94 (1H, m), 4.08-4.21 (2H, m), 4.30-4.34 (1H, m), 6.60-6.65 (2H, m), 6.77-6.82 (2H, m), 6.94-6.96 (2H, m), 7.01-7.11 (5H, m), 7.17 (2H, d, J= 8.2Hz), 8.43 (1H, t, J= 6.0Hz)
18	CD ₃ OD	δ 2.82-2.88 (1H, m), 3.02-3.09 (2H, m), 3.20-3.26 (1H, m), 4.15 (2H, s), 4.21-4.34 (2H, m), 4.49-4.54 (1H, m), 4.68-4.72 (1H, m), 7.19 (1H, dd, J=8.2, 2.0Hz), 7.29 (2H, d, J=8.1Hz), 7.34 (2H, d, J=8.0Hz), 7.43-7.48 (4H, m), 7.68 (2H, d, J=8.2Hz), 8.76 (1H, t, J=5.8Hz)

19	CD ₃ OD	δ 1.02 (3H, t, J=7.6Hz), 2.19 (2H, q, J=7.6Hz), 2.86-2.94 (2H, m), 3.06-3.16 (2H, m), 4.14 (2H, s), 4.37-4.50 (2H, m), 4.61-4.70 (2H, m), 7.13 (1H, dd, J= 8.3, 2.0Hz), 7.32-7.37 (4H, m), 7.42-7.45 (4H, m), 7.60 (2H, d, J=8.2Hz)
20	CD ₃ OD	δ 0.82-0.99 (2H, m), 1.15-1.30 (4H, m), 1.44-1.54 (2H, m), 1.62-1.73 (5H, m), 2.91-2.97 (1H, m), 3.22-3.27 (1H, m), 3.90 (1H, t, J=7.1Hz), 4.16 (2H, s), 4.38-4.55 (2H, m), 4.77-4.81 (1H, m), 7.26 (1H, dd, J=8.2, 2.0Hz), 7.37 (2H, d, J=8.1Hz), 7.43-7.52 (4H, m)
21	CD ₃ OD	δ 0.85-0.94 (2H, m), 1.02-1.22 (7H, m), 1.34-1.42 (2H, m), 1.62-1.69 (5H, m), 2.16-2.27 (2H, m), 2.85-2.92 (1H, m), 3.39-3.44 (1H, m), 4.14 (2H, s), 4.18-4.24 (1H, m), 4.40-4.45 (1H, m), 4.53-4.58 (1H, m), 4.73-4.79 (1H, m), 7.24 (1H, dd, J=8.2, 2.0Hz), 7.38-7.51 (6H, m), 8.11 (1H, d, J=5.6Hz), 8.59 (1H, d, J=8.5Hz), 8.69 (1H, t, J=6.0Hz)
22	CD ₃ OD	δ 1.30 (3H, d, J=7.1Hz), 2.92-2.98 (1H, m), 3.40 (1H, d, J=4.9Hz), 4.09 (2H, s), 4.41-4.48 (2H, m), 4.54-4.60 (1H, m), 4.68-4.74 (1H, m), 7.24-7.34 (5H, m), 7.38 (4H, s), 7.43-7.49 (2H, m), 7.56-7.60 (1H, m), 7.80-7.82 (2H, m), 8.51 (1H, d, J=8.2Hz), 8.56 (1H, d, J=5.3Hz), 8.66 (1H, t, J=6.0Hz)
23	CD ₃ OD	δ 1.03 (3H, t, J=7.6Hz), 1.42 (3H, t, J=7.0Hz), 2.16-2.22 (2H, m), 2.73-2.93 (3H, m), 3.11-3.16 (1H, m), 4.05 (2H, q, J=6.9Hz),), 4.28 (2H, s), 4.37-4.48 (3H, m), 4.60-4.64 (1H, m), 6.82 (2H, d, J=8.6Hz), 7.05 (2H, d, J= 8.7Hz), 7.17 (2H, t, J=6.6Hz), 7.29-7.33 (4H, m), 7.46 (2H, dd, J=9.6,1.4Hz)
24	CD ₃ OD	δ 1.08 (3H, t, J=7.7Hz), 1.38-1.42 (3H, m), 2.18-2.25 (2H, m), 2.71-3.07 (7H, m), 4.00-4.07 (2H, m), 4.14 (2H, s), 4.54-4.58 (2H, m), 4.62-4.66 (1H, m), 5.09-5.15 (1H, m), 6.83-6.88 (2H, m), 7.08-7.21 (4H, m), 7.26-7.38 (4H, m), 7.42 (2H, d, J= 8.1Hz), 8.29 (1H, d, J= 7.8Hz), 8.35 (1H, d, J= 8.3Hz)
25	CD ₃ OD	δ 1.41 (3H, t, J= 7.0Hz), 2.56-2.68 (2H, m), 2.94-2.98 (1H, m), 3.01 (3H, s), 3.39-3.41 (1H, m), 4.05 (2H, q, J= 7.0Hz), 4.14 (2H, s), 4.33-4.38 (1H, m), 4.49-4.55 (2H, m), 5.52-5.56 (1H, m), 6.86-6.89 (2H, m), 7.02 (2H, d, J= 8.6Hz), 7.28-7.39 (7H, m), 7.43 (2H, d, J= 8.1Hz), 8.53 (1H, t, J= 5.8Hz)

26	CD ₃ OD	δ 1.05 (3H, t, J= 7.6Hz), 1.39 (3H, t, J= 7.0Hz), 2.15-2.23 (2H, m), 2.51-2.63 (2H, m), 2.83-2.90 (1H, m), 2.96 (3H, s), 3.39-3.41 (1H, m), 4.01 (2H, q, J= 7.0Hz), 4.13 (2H, s), 4.38-4.50 (2H, m), 4.83-4.87 (1H, m), 5.47-5.51 (1H, m), 6.78-6.82 (2H, m), 7.01 (2H, d, J= 8.6Hz), 7.22-7.40 (7H, m), 7.43 (2H, d, J= 8.1Hz), 8.19 (1H, d, J= 6.8Hz), 8.34 (1H, t, J= 6.0Hz)
27	CD ₃ OD	δ 1.42 (3H, t, J= 7.0Hz), 2.66 (3H, s), 2.78-2.90 (1H, m), 3.01-3.15 (3H, m), 4.02-4.10 (3H, m), 4.14 (2H, s), 4.34-4.47 (2H, m), 4.59-4.70 (1H, m), 6.83-6.88 (2H, m), 6.97-7.02 (2H, m), 7.20-7.22 (2H, m), 7.26-7.34 (5H, m), 7.42 (2H, d, J= 8.1Hz), 8.69 (1H, t, J= 5.8Hz)
28	CD ₃ OD	δ 0.81-0.83 (6H, m), 1.10-1.18 (1H, m), 1.48-1.55 (1H, m), 1.58-1.65 (1H, m), 2.92-2.98 (1H, m), 3.25-3.30 (1H, m), 3.75 (2H, s), 3.92-3.96 (1H, m), 4.15 (2H, s), 4.48-4.53 (2H, m), 4.75-4.79 (1H, m), 7.27-7.40 (7H, m), 7.46 (2H, d, J= 8.2Hz), 8.83 (1H, t, J= 5.9Hz)
29	CD ₃ OD	δ 0.70-1.03 (6H, m), 1.37-1.69 (7H, m), 2.81-2.87 (4H, m), 3.06-3.11 (1H, m), 3.72-3.96 (3H, m), 3.99 (2H, s), 4.26-4.37 (2H, m), 4.58-4.62 (1H, m), 7.13-7.24 (7H, m), 7.30 (2H, d, J= 8.1Hz), 8.72 (1H, t, J= 5.9Hz)
30	CD ₃ OD	δ 0.75-0.93 (3H, m), 1.20-1.31 (5H, m), 1.54-1.81 (5H, m), 3.02 (3H, s), 3.08-3.17 (1H, m), 3.43-3.48 (1H, m), 3.81 (2H, s), 4.15 (2H, s), 4.45-4.53 (3H, m), 5.65-5.71 (1H, m), 7.27-7.39 (5H, m), 7.41 (2H, d, J= 8.1Hz), 7.48 (2H, d, J= 8.2Hz), 8.67 (1H, t, J= 5.9Hz)
31	CD ₃ OD	δ 0.67-0.71 and 0.77-0.80 (3H, m), 1.19-1.24 (3H, m), 1.99-2.11 (2H, m), 2.65-2.83 (5H, m), 2.91-3.09 (2H, m), 3.81-3.88 (2H, m), 3.95 (2H, s), 4.17-4.32 (2H, m), 4.43-4.54 (1H, m), 4.77-4.82 (1H, m), 6.62-6.69 (2H, m), 6.90-6.94 (2H, m), 6.98-7.20 (7H, m), 7.21-7.26 (2H, m), 7.69 (1H, d, J= 7.9Hz), 8.26-8.29 and 8.40-8.43 (1H, m)
32	CD ₃ OD	δ 1.02-1.10 (3H, m), 1.40 (3H, t, J= 7.0Hz), 2.11-2.24 (2H, m), 2.51-2.63 (2H, m), 2.82-2.90 (1H, m), 2.95 (3H, s), 3.36-3.41 (1H, m), 3.98-4.03 (2H, m), 4.20 (2H, s), 4.38-4.51 (2H, m), 4.85 (1H, t, J= 7.3Hz), 5.46-5.50 (1H, m), 6.77-6.85 (2H, m), 7.00-7.08 (2H, m), 7.11-7.18 (2H, m), 7.21-7.36 (5H, m), 7.42-7.48 (1H, m), 8.43 (1H, t, J= 6.1Hz)

33	CD ₃ OD	δ 0.63-0.80 (2H, m), 0.87-1.08 (4H, m), 1.32-1.54 (7H, m), 2.50 (3H, s), 2.79-2.85 (1H, m), 3.08-3.13 (1H, m), 3.66-3.70 (1H, m), 3.99 (2H, s), 4.27-4.36 (2H, m), 4.59-4.65 (1H, m), 7.13-7.24 (7H, m), 7.30 (2H, d, J= 8.2Hz), 8.71 (1H, t, J= 5.9Hz), 8.80 (1H, d, J= 7.6Hz)
34	CD ₃ OD	δ 0.75-0.91 (3H, m), 1.20-1.33 (5H, m), 1.55 (1H, d, J= 12.3Hz), 1.62-1.71 (3H, m), 1.80 (1H, d, J= 12.5Hz), 3.02 (3H, s), 3.08-3.18 (1H, m), 3.45-3.50 (1H, m), 3.83-3.87 (2H, m), 4.23 (2H, s), 4.46-4.52 (3H, m), 5.66-5.72 (1H, m), 7.21-7.41 (7H, m), 7.52 (1H, t, J= 7.8Hz), 8.77 (1H, t, J= 6.0Hz)
35	CD ₃ OD	δ 0.83 (3H, t, J= 7.4Hz), 1.24-1.30 (3H, m), 1.51-1.62 (2H, m), 2.64 (3H, s), 2.65-2.89 (5H, m), 3.09-3.14 (1H, m), 3.85-3.96 (2H, m), 3.98 (2H, s), 4.16-4.22 (1H, m), 4.28-4.37 (1H, m), 4.47-4.50 (1H, m), 5.29 (1H, t, J= 8.1Hz), 6.67-6.75 (2H, m), 6.87-6.90 (2H, m), 7.09 (2H, d, J= 8.2Hz), 7.15-7.29 (7H, m), 8.42 (1H, t, J= 6.0Hz)
36	CD ₃ OD	δ 1.15-1.30 (6H, m), 1.52-1.76 (7H, m), 2.89-3.02 (4H, m), 3.10-3.19 (1H, m), 3.73-3.80 (2H, m), 3.99-4.05 (1H, m), 4.12-4.14 (2H, m), 4.48-4.59 (1H, m), 4.71-4.80 (1H, m), 5.08-5.16 (1H, m), 7.19-7.49 (9H, m)
37	CD ₃ OD	δ 0.62-1.08 (9H, m), 1.34-1.66 (9H, m), 2.71-2.86 (3H, m), 3.07-3.11 (1H, m), 3.69-3.73 (1H, m), 4.00 (2H, s), 4.27-4.37 (2H, m), 4.59-4.63 (1H, m), 7.13-7.27 (7H, m), 7.30 (2H, d, J= 8.1Hz), 8.68 (1H, t, J= 5.9Hz)
38	CD ₃ OD	δ 1.39 (3H, t, J= 7.0Hz), 2.66-2.71 (1H, m), 2.79-2.94 (2H, m), 3.01 (3H, s), 3.38-3.43 (1H, m), 3.97-4.03 (2H, m), 4.07-4.14 (2H, m), 4.40-4.50 (2H, m), 5.03-5.09 (1H, m), 5.52-5.56 (1H, m), 6.78-6.86 (2H, m), 7.05-7.18 (2H, m), 7.20-7.71 (12H, m), 7.73-7.81 (2H, m), 8.37 (1H, t, J= 6.0Hz), 8.58 (1H, d, J= 6.7Hz)
39	CD ₃ OD	δ 1.39 (3H, t, J= 7.0Hz), 1.93 (3H, s), 2.50-2.63 (2H, m), 2.81-2.90 (1H, m), 2.95 (3H, s), 3.34-3.41 (1H, m), 3.95-4.05 (2H, m), 4.13 (2H, s), 4.33-4.51 (2H, m), 4.82-4.89 (1H, m), 5.46-5.50 (1H, m), 6.78-6.85 (2H, m), 7.00-7.05 (2H, m), 7.09-7.36 (7H, m), 7.39-7.43 (2H, m), 8.29 (1H, t, J= 6.0Hz)

40	CD ₃ OD	δ 0.61-0.81 (2H, m), 0.88-1.05 (4H, m), 1.34-1.54 (7H, m), 2.51 (3H, s), 2.80-2.86 (1H, m), 3.08-3.13 (1H, m), 3.67-3.70 (1H, m), 4.07 (2H, s), 4.27-4.37 (2H, m), 4.59-4.63 (1H, m), 6.99-7.04 (2H, m), 7.13-7.24 (5H, m), 7.34 (1H, t, J= 7.9Hz), 8.73 (1H, t, J= 6.0Hz)
41	CD ₃ OD	δ 0.73-1.25 (6H, m), 1.50-1.69 (7H, m), 2.67 (3H, s), 2.94-3.00 (1H, m), 3.24-3.29 (1H, m), 3.81-3.85 (1H, m), 4.23 (2H, s), 4.44-4.54 (2H, m), 4.76-4.80 (1H, m), 7.18-7.23 (2H, m), 7.31 (1H, dd, J= 8.3, 2.0 Hz), 7.45-7.56 (3H, m), 8.91 (1H, t, J= 5.9Hz), 8.97 (1H, d, J= 7.7Hz)
42	CD ₃ OD	δ 0.83-1.19 (6H, m), 1.53-1.85 (7H, m), 2.94-3.08 (4H, m), 3.21-3.26 (1H, m), 3.87-4.08 (3H, m), 4.22 (2H, s), 4.38-4.55 (2H, m), 4.72-4.76 (1H, m), 7.18-7.22 (2H, m), 7.28-7.39 (5H, m), 7.47-7.51 (1H, m), 8.93 (1H, t, J= 6.0Hz), 9.05 (1H, d, J= 7.2Hz)
43	CD ₃ OD	δ 0.78-0.97 (4H, m), 1.03-1.20 (2H, m), 1.47-1.50 (1H, m), 1.54-1.63 (5H, m), 1.75-1.82 (1H, m), 2.92-2.95 (1H, m), 2.98 (3H, s), 3.19-3.24 (1H, m), 3.98-4.13 (3H, m), 4.19 (2H, s), 4.38-4.52 (2H, m), 4.70-4.74 (1H, m), 7.17-7.20 (2H, m), 7.27 (1H, dd, J= 8.3, 2.0 Hz), 7.37-7.52 (3H, m), 8.92 (1H, t, J= 5.9Hz), 9.08 (1H, d, J= 7.3Hz)
44	CD ₃ OD	δ 1.01-1.05 (6H, m), 1.40 (3H, t, J=7.0Hz), 2.36-2.62 (3H, m), 2.85-2.91 (1H, m), 2.96 (3H, s), 3.37-3.42 (1H, m), 3.98-4.04 (2H, m), 4.13 (2H, s), 4.39-4.50 (2H, m), 4.82-4.86 (1H, m), 5.48-5.52 (1H, m), 6.77-6.85 (2H, m), 6.99-7.04 (2H, m), 7.22-7.36 (7H, m), 7.39-7.44 (2H, m), 8.37 (1H, t, J= 6.0Hz)
45	CD ₃ OD	δ 0.71-1.03 (9H, m), 1.36-1.63 (9H, m), 2.75 (3H, s), 2.81-3.11 (4H, m), 3.71-3.74 (1H, m), 4.00 (2H, s), 4.31-4.33 (2H, m), 4.59-4.63 (1H, m), 7.14-7.24 (7H, m), 7.30 (2H, d, J= 8.2Hz), 8.71 (1H, t, J= 5.8Hz)
46	CD ₃ OD	δ 0.78-0.97 (2H, m), 1.09-1.25 (4H, m), 1.54-1.69 (7H, m), 2.93-2.99 (1H, m), 3.24-3.29 (1H, m), 3.87 (3H, s), 3.94 (2H, d, J= 2.5Hz), 3.99-4.04 (1H, m), 4.15 (2H, s), 4.43-4.53 (2H, m), 4.73-4.77 (1H, m), 7.25-7.38 (7H, m), 7.46 (2H, d, J= 8.1Hz), 8.83 (1H, t, J= 5.9Hz)
47	CD ₃ OD	δ 0.72-1.14 (6H, m), 1.40-1.73 (7H, m), 2.85-2.91 (4H, m), 3.11-3.16 (1H, m), 3.77 (3H, s), 3.93-3.96 (1H, m), 4.04 (2H, s), 4.07-4.18 (2H, m), 4.36 (2H, s), 4.63-4.67 (1H, m), 7.17-7.27 (7H, m), 7.35 (2H, d, J= 8.1Hz), 8.73 (1H, t, J= 5.9Hz), 8.92 (1H, d, J= 7.3Hz)

48	CD ₃ OD	δ 0.87-1.17 (6H, m), 1.54-1.79 (7H, m), 2.89 (3H, s), 2.95-3.02 (1H, m), 3.21-3.27 (1H, m), 3.79-3.99 (3H, m), 4.15 (2H, s), 4.47 (2H, s), 4.74-4.78 (1H, m), 7.29-7.39 (7H, m), 7.46 (2H, d, J= 8.1Hz), 8.84 (1H, t, J= 5.8Hz)
49	CD ₃ OD	δ 0.75-1.08 (6H, m), 1.41-1.76 (7H, m), 2.85-2.91 (4H, m), 3.11-3.36 (3H, m), 3.78-3.90 (3H, m), 4.04 (2H, s), 4.31-4.41 (2H, m), 4.63-4.67 (1H, m), 7.18-7.29 (7H, m), 7.35 (2H, d, J= 8.2Hz), 8.76(1H, t, J= 5.9Hz)
52	CD ₃ OD	δ , 0.74-0.88 (4H 0.78-2.10 (23H, m), 2.92-3.08 (1H, m), 3.12-3.30 (1H, m), 4.07 (1H, d, 4.2Hz), 4.15 (2H, s), 4.42 (1H, dd, J=15.3, 5.5Hz), 4.52 (1H, dd, J=15.3, 6.0Hz), 4.66 (1H, dd, J=10.2, 5.2Hz), 7.20-7.48 (9H, m), 8.87 (1H, t, J=6.0Hz)
53	CD ₃ OD	δ 0.83-1.19 (12H, m), 1.50 (1H, d, J=12.2 Hz), 1.60-1.79 (10H, m), 2.98 (1H, dd, J=10.6, 14.2 Hz), 3.14-3.28 (5H, m), 3.91 (1H, dd, J=3.4, 11.7 Hz), 4.15 (2H, s), 4.47-4.49 (2H, m), 4.77 (1H, dd, J=5.2, 10.5 Hz), H, s7.36-7.41(6H (7H, m), 7.45 (2H, d, J=8.1 Hz), 8.89-8.92 (1H, m)
54	CD ₃ OD	δ 1.44 (3H, t, J= 7.0 Hz), 2.81-2.87 (1H, m), 2.95-3.02 (2H, m), 3.19-3.24 (1H, m), 4.03-4.09 (4H, m), 4.46 (2H, d, J = 6.0 Hz), 4.84-4.89 (1H, m), 6.88 (2H, d, J = 8.5 Hz), 7.04 (2H, d, J = 8.5 Hz), 7.17-7.51 (13H, m), 7.85 (1H, d, J= 8.5 Hz), 7.93 (1H, d, J= 8.5 Hz), 7.90 and 8.00 (total 1H, each dd, J= each 7.2 Hz), 8.43 (1H, d, J= 8.1 Hz), 8.63 (1H, t, J= 5.9 Hz), 8.75 (1H, t, J= 7.0 Hz)
55	CD ₃ OD	δ 1.25 (3H, t, J= 7.0 Hz), 2.77-2.83 (H3H, m), 2.96-3.02 (1H, m), 3.85-3.97 (4H, m), 4.22-4.26 (1H, m), 4.27-4.39 (1H, m), 4.48-4.54 (2H, m), 6.66-6.68 (2H, m), 6.94 (2H, d, J = 8.6 Hz), 7.01-7.03 (2H, m), 7.11-7.29 (9H, m), 7.31 (1H, d, J= 1.8 Hz), 7.53 (1H, d, J= 8.6 Hz), 8.22 (1H, d, J= 8.0 Hz), 8.36-8.44 (2H, m)
56	CD ₃ OD	δ 1.25 (3H, t, J= 7.0 Hz), 2.75-2.83 (4H, m), 2.87-2.99 (1H, m), 3.85-3.89 (2H, m), 3.97 (2H, s), 4.22-4.39 (2H, m), 4.53-4.57 (H1H, m), 6.67 (2H, d, J = 8.5 Hz), 6.95 (2H, d, J = 8.5 Hz), 7.02-7.04 (2H, m), 7.11-7.22 (7H, m), 7.60 (2H, d, J= 8.3 Hz), 7.71 (2H, d, J= 8.2 Hz), 8.25 (1H, d, J= 8.0 Hz), 8.42 (1H, t, J= 5.8 Hz)

57	CD ₃ OD	δ 1.41 (3H, t, J= 7.0 Hz), 2.91-3.04 (4H, m), 3.15-3.20 (1H, m), 4.01-4.07 (4H, m), 4.40 (1H, d, J= 15.4 Hz), 4.53 (1H, d, J= 15.4 Hz), 4.67-4.71 (2H, m), 6.84 (2H, d, J = 8.7 Hz), 7.11 (2H, d, J = 8.6 Hz), 7.19-7.24 (2H, m), 7.27-7.37 (9H, m), 7.44-7.54 (2H, m), 7.70-7.73 (4H, m), 7.80-7.82 (1H, m)
58	CD ₃ OD	δ 1.42 (3H, t, J= 7.0 Hz), 2.80-2.98 (4H, m), 3.02-3.06 (1H, m), 4.05 (2H, d, J = 7.0 Hz), 4.12 (2H, s), 4.39-4.46 (2H, m), 4.68-4.76 (2H, m), 6.85 (2H, d, J = 8.6 Hz), 7.12 (2H, d, J = 8.6 Hz), 7.20-7.22 (2H, m), 7.26-7.32 (6H, m), 7.39-7.41 (3H, m), 7.54 (1H, d, J = 1.9 Hz), 8.54 (1H, t, J = 5.9 Hz)
59	CD ₃ OD	δ 1.25 (3H, t, J= 7.0 Hz), 2.77-2.81 (4H, m), 2.97-3.02 (1H, m), 3.87 (2H, q, J = 7.0 Hz), 3.93 (2H, s), 4.26-4.30 (1H, m), 4.35-4.40 (1H, m), 4.48-4.55 (2H, m), 6.68 (2H, d, J = 8.6 Hz), 6.94 (2H, d, J = 8.6 Hz), 7.01-7.03 (2H, m), 7.09-7.22 (8H, m), 7.42-7.45 (2H, m), 8.22 (1H, d, J= 8.1 Hz), 8.41 (1H, t, J= 5.8 Hz)
60	CD ₃ OD	δ 1.23 (3H, t, J= 7.0Hz), 2.81-2.87 (1H, m), 3.01-3.06 (1H, m), 3.85-3.91 (2H, m), 3.94 (2H, s), 3.96-4.04 (2H, m), 4.18-4.24 (1H, m), 4.53 (1H, d, J= 7.0Hz), 4.84 (1H, d, J= 7.1Hz), 6.70-6.74 (2H, m), 6.88 (2H, d, J= 8.0Hz), 7.01-7.04 (2H, m), 7.13-7.23 (5H, m), 7.27-7.31 (2H, m)
61	CD ₃ OD	δ 1.43 (3H, t, J= 7.0Hz), 2.68-2.74 (1H, m), 2.98-3.03 (1H, m), 4.06 (2H, q, J= 7.0Hz), 4.14 (2H, s), 4.21-4.24 (1H, m), 4.42-4.50 (2H, m), 4.75 (1H, d, J= 4.4Hz), 5.28 (1H, d, J= 4.4Hz), 6.85-6.89 (2H, m), 6.97-7.00 (2H, m), 7.30-7.35 (4H, m), 7.38-7.43 (3H, m), 7.5 (2H, d, J= 7.3Hz)
62	CD ₃ OD	δ 0.93 (3H, t, J= 7.7Hz), 1.25 (3H, t, J= 7.0Hz), 2.01-2.11 (2H, m), 2.60-2.66 (1H, m), 2.79-2.84 (1H, m), 3.86 (2H, q, J= 7.0Hz), 3.99 (2H, s), 4.22-4.32 (2H, m), 4.41-4.43 (1H, m), 4.46 (1H, d, J= 4.0Hz), 5.13 (1H, d, J= 4.0Hz), 6.64-6.67 (2H, m), 6.91-6.95 (2H, m), 7.14-7.33 (9H, m)

63	CD ₃ OD	δ 1.00 (3H, t, J= 7.6Hz), 1.43 (3H, t, J= 7.0Hz), 2.13-2.19 (2H, ,m), 2.61-2.66 (1H, m), 2.78-2.83 (1H, m), 4.04 (2H, q, J= 7.0Hz), 4.14 (2H, s), 4.45-4.65 (4H, m), 5.37 (1H, d, J= 3.0Hz), 6.77-6.81 (2H, m), 7.01 (2H, d, J= 8.6Hz), 7.28-7.45 (9H, ,m), 8.00 (1H, d, J= 6.9Hz), 8.23 (1H, d, J= 8.4Hz), 8.51 (1H, t, J= 6.0Hz)
64	CD ₃ OD	δ 0.98 (3H, t, J= 7.2Hz), 1.06-1.13 (3H, m), 1.40 (3H, t, J= 7.0Hz), 2.19-2.24 (2H, m), 2.71-2.82 (2H, ,m), 2.98-3.05 (1H, m), 3.14-3.22 (1H, m), 3.29-3.35 (1H, ,m), 3.51-3.60 (1H, m), 3.98-4.04 (2H, m), 4.13 (2H, s), 4.35-4.48 (2H, m), 4.78-4.89 (1H, m), 5.12-5.21 (1H, m), 6.79-6.83 (2H, m), 7.07-7.16 (2H, m), 7.26-7.38 (7H, m), 7.41 (2H, d, J= 8.1Hz), 8.21 (1H, t, J= 6.1Hz)
65	CD ₃ OD	δ 0.96-1.10 (6H, m), 1.33-1.44 (3H, m), 2.10-2.27 and 2.41-2.46 (total 3H, m), 2.74-2.97 (2H, m), 3.12-3.23 (1H, m), 3.28-3.34 (1H, m), 3.41-3.50 (1H, m), 3.92-4.08 (2H, m), 4.12 (2H, s), 4.34-4.50 (2H, m), 4.60-4.64 and 4.78-4.82 (1H, m), 4.94-4.98 and 5.08-5.12(total 1H, m), 6.80-6.89 (2H, m), 7.09-7.42 (11H, m), 8.11 (1H, t, J=6.0 Hz), 8.20 (1H, d, J=6.5 Hz), 8.45 (1H, d, J=7.2 Hz), 8.77 (1H, t, J=5.8 Hz)
66	CD ₃ OD	δ 0.82-0.86 and 0.96-1.00 (total 3H, m), 1.38-1.43 (3H, m), 2.18-2.30 (2H, m), 2.63 (3H, s), 2.84-3.02 (2H, m), 3.11-3.28 (2H, m), 3.99-4.06 (2H, m), 4.15 (2H, s), 4.39-4.51 (2H, m), 4.64-4.79 (1H, m), 5.22-5.26 (1H, m), 6.81-6.87 (2H, m), 7.09-7.13 (2H, m), 7.22-7.37 (7H, m), 7.42-7.46 (2H, m), 7.76 (1H, d, J= 7.9Hz), 8.45-8.47 and 8.59-8.62 (total 1H, m)
67	CD ₃ OD	δ 1.00 (3H, t, J=7.7Hz), 1.44 (3H, t, J=7.0Hz), 1.64-1.72 (1H, m), 2.08-2.31 (3H, m), 2.98-3.09 (2H, m), 3.30-3.36 (1H, m), 3.39-3.44 (1H, m), 3.91-3.97 (1H, m), 4.03-4.07 (2H, m), 4.14 (2H, s), 4.10-4.16 (1H, m), 4.31-4.34 (1H, m), 4.37 (1H, d, J=4.9Hz), 4.51-4.57 (1H, m), 4.82-4.86 (1H, m), H, s6.91-6.97 (4H, m), 7.25-7.36 (7H, m), 7.42 (2H, d, J=8.2Hz), 8.35-8.50 (1H, m)

68	d ⁶ - DMSO	δ: 1.29 (3H, t, J = 7.0Hz), 2.43 (1H, dd, J = 13.7, 8.0Hz), 2.63 (1H, dd, J = 13.8, 4.7Hz), 2.77 (1H, dd, J = 13.6, 10.4Hz), 3.06 (1H, dd, J = 13.7, 4.4Hz), 3.91-3.96 (4H, m), 4.07 (1H, dd, J = 15.4, 5.8Hz), 4.15 (1H, dd, J = 15.4, 6.0Hz), 4.23-4.26 (2H, m), 4.32-4.37 (1H, m), 4.44-4.50 (1H, m), 6.09 (1H, d, J = 7.7Hz), 6.58 (1H, t, J = 6.0Hz), 6.67 (2H, d, J = 8.7Hz), 6.76 (2H, d, J = 8.6Hz), 7.08-7.38 (11H, m), 7.45 (2H, d, J = 7.5Hz), 8.28 (3H, br s), 8.48 (1H, d, J = 8.5Hz), 8.62 (1H, t, J = 6.0Hz).
69	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.0Hz), 2.47-2.50 (1H, m), 2.67 (1H, dd, J = 13.9, 4.6Hz), 2.79 (1H, dd, J = 13.6, 10.5Hz), 3.07 (1H, dd, J = 13.7, 4.3Hz), 3.90-3.97 (4H, m), 4.23-4.33 (2H, m), 4.43-4.53 (2H, m), 6.24 (1H, d, J = 7.7Hz), 6.66 (2H, d, J = 8.7Hz), 6.72 (2H, d, J = 8.7Hz), 6.86 (1H, t, J = 7.4Hz), 7.13-7.32 (11H, m), 7.36 (2H, d, J = 8.1Hz), 8.23 (2H, br s), 8.58-8.63 (2H, m), 8.77 (1H, s).
70	d ⁶ - DMSO	δ: 1.26 (3H, t, J = 7.0Hz), 2.66 (1H, dd, J = 13.5, 10.6Hz), 2.73 (1H, dd, J = 13.8, 4.0Hz), 2.82 (1H, dd, J = 13.5, 10.2Hz), 3.05 (1H, dd, J = 13.6, 4.9Hz), 3.89-4.01 (4H, m), 4.27 (1H, dd, J = 15.4, 5.8Hz), 4.34 (1H, dd, J = 15.4, 6.1Hz), 4.55-4.68 (2H, m), 6.75 (2H, d, J = 8.6Hz), 7.07-7.32 (8H, m), 7.39 (2H, d, J = 8.1Hz), 7.73 (1H, dd, J = 7.9, 5.2Hz), 8.13-8.59 (4H, m), 8.63 (2H, m), 8.81 (1H, dd, J = 5.1, 1.3Hz), 8.99-9.01 (2H, m).
71	d ⁶ - DMSO	δ: 1.26 (3H, t, J = 6.9Hz), 2.64 (1H, dd, J = 13.5, 10.8Hz), 2.72 (1H, dd, J = 13.6, 3.8Hz), 2.82 (1H, dd, J = 13.5, 10.2Hz), 3.05 (1H, dd, J = 13.6, 4.7Hz), 3.80-4.01 (4H, m), 4.27 (1H, dd, J = 15.4, 5.8Hz), 4.34 (1H, dd, J = 15.4, 6.0Hz), 4.58-4.68 (2H, m), 6.75 (2H, d, J = 8.6Hz), 7.14 (2H, d, J = 8.7Hz), 7.17-7.27 (6H, m), 7.38 (2H, d, J = 8.1Hz), 7.85 (2H, d, J = 5.9Hz), 8.14-8.50 (3H, m), 8.64 (1H, d, J = 6.2Hz), 8.67 (1H, d, J = 8.6Hz), 8.80 (2H, d, J = 6.0Hz), 9.03 (1H, d, J = 8.3Hz).

72	d ⁶ - DMSO	δ: 1.26 (3H, t, J = 7.0Hz), 2.62-2.64 (2H, m), 2.81 (1H, dd, J = 13.6, 10.0Hz), 3.03 (1H, dd, J = 13.4, 4.5Hz), 3.92 (2H, q, J = 7.0Hz), 3.96 (2H, s), 4.25-4.35 (2H, m), 4.54-4.61 (2H, m), 6.74 (2H, d, J = 8.6Hz), 7.13 (2H, d, J = 8.7Hz), 7.16-7.25 (7H, m), 7.35 (2H, d, J = 8.1Hz), 7.43 (1H, dd, J = 5.0, 1.3Hz), 7.53 (1H, dd, J = 5.0, 3.0Hz), 8.11 (1H, dd, J = 2.8, 1.0), 8.20 (2H, br s), 8.33 (1H, d, J = 8.2Hz), 8.54 (1H, d, J = 8.7Hz), 8.60 (1H, t, J = 6.0Hz).
73	d ⁶ - DMSO	δ: 1.26 (3H, t, J = 7.0Hz), 2.48-2.68 (2H, m), 2.81 (1H, dd, J = 13.6, 10.1Hz), 3.03 (1H, dd, J = 13.6, 4.7Hz), 3.89-3.96 (4H, m), 4.24-4.34 (2H, m), 4.53-4.62 (2H, m), 6.74 (2H, d, J = 8.6Hz), 7.10-7.27 (9H, m), 7.37 (2H, d, J = 8.1Hz), 7.71 (1H, dd, J = 4.9, 0.9Hz), 7.82 (1H, dd, J = 3.6, 2.8Hz), 8.20 (3H, br s), 8.56 (2H, dd, J = 10.5, 9.1Hz), 8.62 (1H, t, J = 6.0Hz).
74	d ⁶ - DMSO	δ: 0.48-0.57(4H,m), 1.29(3H,t,J= 7.0Hz), 1.56-1.62(1H,m), 2.43-2.49(1H,m), 2.54-2.59(1H,m), 2.73-2.79(1H,m), 3.00-3.05(1H,m), 3.92-3.97(4H,m), 4.20-4.33(2H,m), 4.38-4.43(1H,m), 4.47-4.53(1H,m), 6.73(2H,d,J= 8.6Hz), 6.98(2H,d,J= 8.6Hz), 7.16-7.26(7H,m), 7.38(2H,d,J= 8.1Hz), 8.23(1H,d,J= 7.7Hz), 8.28(2H,s), 8.48(1H,d,J= 8.6Hz), 8.54(1H,t,J=6.0Hz).
75	d ⁶ - DMSO	δ: 0.96-1.15(5H,m), 1.20(3H,t,J= 7.0Hz), 1.36-1.55(5H,m), 1.96-2.01(1H,m), 2.34-2.41(1H,m), 2.51-2.56(1H,m), 2.67-2.73(1H,m), 2.92-2.97(1H,m), 3.86(2H,q,J= 7.0Hz), 3.90(2H,s), 4.15-4.24(2H,m), 4.27-4.32(1H,m), 4.41-4.47(1H,m), 6.64(2H,d,J= 8.6Hz), 6.88(2H,d,J= 8.6Hz), 7.08-7.18(7H,m), 7.30(2H,d,J= 8.1Hz), 7.66(1H,d,J= 8.0Hz), 8.17(2H,s), 8.29(1H,d,J= 8.5Hz), 8.51(1H,t,J=6.0Hz).
76	d ⁶ - DMSO	δ: 0.78(3H,t,J= 7.3Hz), 0.99-1.05(2H,m), 1.12-1.21(2H,m), 1.26-1.36(5H,m), 1.91-2.03(2H,m), 2.38-2.44(1H,m), 2.56-2.61(1H,m), 2.74-2.80(1H,m), 2.99-3.04(1H,m), 3.93(2H,q,J= 7.0Hz), 3.97(2H,s), 4.23-4.33(2H,m), 4.39-4.44(1H,m), 4.50-4.56(1H,m), 6.72(2H,d,J= 8.6Hz), 6.98(2H,d,J= 8.6Hz), 7.15-7.26(7H,m), 7.38(2H,d,J= 8.1Hz), 7.88(1H,d,J= 8.0Hz), 8.22(2H,s), 8.42(1H,d,J= 8.6Hz), 8.58(1H,t,J=6.0Hz).

77	d ⁶ - DMSO	δ: 0.93 (3H, d, J = 4.2Hz), 0.95 (3H, d, J = 4.1Hz), 1.28 (3H, t, J = 7.0Hz), 2.41 (1H, dd, J = 13.7, 8.0Hz), 2.58 (1H, dd, J = 13.8, 4.7Hz), 2.77 (1H, dd, J = 13.6, 10.5Hz), 3.06 (1H, dd, J = 13.7, 4.2Hz), 3.50-3.58 (1H, m), 3.70-4.01 (4H, m), 4.25-4.30 (3H, m), 4.41-4.47 (1H, m), 5.82 (1H, d, J = 7.6Hz), 5.99 (1H, d, J = 7.6Hz), 6.67 (2H, d, J = 8.6Hz), 6.76 (2H, d, J = 8.6Hz), 7.16-7.26 (7H, m), 7.39 (2H, d, J = 8.1Hz), 8.37 (2H, br s), 8.46 (1H, d, J = 8.5Hz), 8.65 (1H, t, J = 6.0Hz).
78	d ⁶ - DMSO	δ: 0.83 (3H, t, J = 7.0Hz), 1.16-1.30 (11H, m), 2.41 (1H, dd, J = 13.8, 8.0Hz), 2.59 (1H, dd, J = 13.6, 4.6Hz), 2.77 (1H, dd, J = 13.7, 10.5Hz), 2.83-2.90 (2H, m), 3.05 (1H, dd, J = 13.7, 4.2Hz), 3.93 (2H, q, J = 7.0Hz), 3.95 (2H, m), 4.26-4.32 (3H, m), 4.42-4.46 (1H, m), 5.91 (1H, d, J = 7.6Hz), 6.09 (1H, t, J = 5.6Hz), 6.67 (2H, d, J = 8.7Hz), 6.76 (2H, d, J = 8.6Hz), 7.16-7.26 (7H, m), 7.39 (2H, d, J = 8.1Hz), 8.35 (2H, br s), 8.45 (1H, d, J = 8.5Hz), 8.65 (1H, t, J = 6.0Hz).
79	d ⁶ - DMSO	δ: 1.34(3H,t,J= 7.0Hz), 2.65-2.79(2H,m), 2.85-2.91(1H,m), 3.09-3.13(1H,m), 3.99(2H,q,J= 7.0Hz), 4.04(2H,s), 4.32-4.42(2H,m), 4.65-4.73(2H,m), 6.80(2H,d,J= 8.6Hz), 7.15(2H,d,J= 1.9Hz), 7.16(1H,s), 7.22-7.34(7H,m), 7.44(2H,d,J= 8.1Hz), 8.29(2H,s), 8.68-8.72(2H,m), 8.77(1H,d,J= 1.8Hz), 8.96(1H,d,J= 8.6Hz).
80	d ⁶ - DMSO	δ: 1.26(3H,t,J= 7.0Hz), 2.61-2.72(2H,m), 2.79-2.84(1H,m), 3.02-3.07(1H,m), 3.92(2H,q,J= 7.0Hz), 3.96(2H,s), 4.25-4.37(2H,m), 4.56-4.63(2H,m), 6.75(2H,d,J= 8.6Hz), 7.14(2H,d,J= 8.6Hz), 7.16-7.25(7H,m), 7.37(2H,d,J= 8.1Hz), 7.42(2H,d,J= 8.2Hz), 7.85-7.88(2H,m), 8.21(2H,s), 8.56-8.64(3H,m).
81	d ⁶ - DMSO	δ: 1.25 (3H, t, J = 7.0Hz), 2.68 (1H, m), 2.78-2.85 (2H, m), 3.06 (1H, dd, J = 13.6, 4.4Hz), 3.86-4.01 (4H, m), 4.28 (2H, d, J = 6.0Hz), 4.50-4.56 (1H, m), 4.74-4.79 (1H, m), 6.59 (2H, d, J = 8.7Hz), 6.65 (2H, d, J = 8.8Hz), 7.17-7.38 (8H, m), 7.59-7.61 (1H, m), 7.96-7.99 (2H, m), 8.33 (3H, br s), 8.48 (1H, d, J = 8.2Hz), 8.61 (1H, dt, J = 4.7, 1.2Hz), 8.66 (1H, t, J = 6.0Hz), 8.71 (1H, d, J = 8.5Hz).

82	d ⁶ - DMSO	δ: 1.27 (3H, t, J = 6.9Hz), 2.13 (3H, s), 2.57-2.69 (2H, m), 2.82 (1H, dd, J = 13.6, 10.0Hz), 3.05 (1H, dd, J = 13.6, 4.7Hz), 3.78 (3H, s), 3.91 (2H, q, J = 7.0Hz), 3.92-3.99 (2H, m), 4.25-4.36 (2H, m), 4.50-4.62 (2H, m), 6.63 (1H, s), 6.74 (2H, d, J = 8.6Hz), 7.11 (2H, d, J = 8.7Hz), 7.17-7.25 (7H, m), 7.37 (2H, d, J = 8.2Hz), 8.25 (2H, br s), 8.35 (1H, d, J = 8.4Hz), 8.55 (1H, d, J = 8.6Hz), 8.60 (1H, t, J = 6.0Hz).
83	d ⁶ - DMSO	δ: 1.25 (3H, t, J = 7.0Hz), 2.61-2.70 (2H, m), 2.82 (1H, dd, J = 13.5, 10.1Hz), 3.05 (1H, dd, J = 13.6, 4.7Hz), 3.91 (2H, q, J = 7.0Hz), 3.94 (2H, s), 4.25-4.35 (2H, m), 4.58-4.64 (2H, m), 6.75 (2H, d, J = 8.6Hz), 7.14-7.25 (9H, m), 7.38 (2H, d, J = 8.2Hz), 7.42-7.45 (2H, m), 7.93-7.99 (2H, m), 8.18 (1H, s), 8.27 (2H, br s), 8.63-8.67 (2H, m), 8.84 (1H, d, J = 8.4Hz).
84	d ⁶ - DMSO	δ: 1.31(3H,t,J= 7.0Hz), 2.21-2.27(1H,m), 2.35-2.39(1H,m), 2.67-2.72(1H,m), 2.90-2.95(1H,m), 3.94(2H,q,J= 7.0Hz), 3.99(3H,s), 4.22-4.27(1H,m), 4.38-4.44(1H,m), 4.49-4.55(1H,m), 6.63(2H,d,J= 8.6Hz), 6.93(2H,d,J= 8.6Hz), 7.13-7.25(7H,m), 7.31-7.34(2H,m), 7.38-7.42(4H,m), 8.10(1H,s), 8.20(2H,s), 8.48(1H,d,J= 8.8Hz), 8.56(1H,t,J = 6.0Hz).
85	d ⁶ - DMSO	δ: 1.30 (3H, t, J= 6.96Hz), 2.38-2.42 (1H, m), 2.58-2.62 (1H, m), 2.73-2.79 (1H,m), 2.99-3.03 (1H,m), 3.91-3.95 (4H, m), 4.24-4.25 (2H, m), 4.41-4.46 (1H,m), 4.51-4.57 (1H,m), 6.67 (2H, d, J= 8.69Hz), 6.89 (2H, d, J= 8.64Hz), 7.07 (2H, d, J= 1.8Hz), 7.17-7.25 (10H,m), 7.37 (2H, t, J= 8.08Hz), 8.14 (1H, t, J= 8.12Hz), 8.26 (3H,s, br), 8.49 (1H, d, J = 8.57Hz), 8.56-8.59 (1H, m).
86	d ⁶ - DMSO	δ: 1.28 (3H, t, J= 6.92Hz), 2.26-2.35 (2H, m), 2.39-2.45 (1H, m), 2.49 (2H, s), 2.56-2.65 (2H,m), 2.74-2.80 (1H, m), 3.00-3.05 (1H, m), 3.93-3.96 (3H, m), 4.25-4.32 (2H, m), 4.41-4.46 (1H,m), 4.49-4.55 (1H, m), 6.70 (2H, d, J= 8.48Hz), 6.92 (2H, d, J= 8.48Hz), 7.08 (2H, d, J= 7.16Hz), 7.18-7.24 (10H, m), 7.38 (2H, t, J= 8.0Hz), 8.00 (1H, d, J= 8.00Hz), 8.28 (3H,s, br), 8.46 (1H, d, J = 8.48Hz), 8.57 (1H, t, J= 5.92Hz).

87	d ⁶ - DMSO	δ: 1.28 (3H, t, J= 6.92Hz), 1.51-1.58 (2H, m), 1.73-1.86 (2H, m), 2.39-2.45 (2H, m), 2.49 (2H, s), 2.56-2.65 (2H, m), 2.95-3.03 (1H, m), 3.86-3.97 (4H, m), 4.29-4.34 (3H, m), 4.47-4.53 (1H, m), 6.66 (2H, d, J= 8.21Hz), 6.96 (2H, d, J= 8.17Hz), 7.17-7.34 (10H, m), 7.36-7.40 (2H, m), 8.02-8.11 (1H, m), 8.26 (3H,s, br), 8.34-8.56 (2H, m).
88	d ⁶ - DMSO	δ: 1.33 (3H,t,J= 6.93Hz), 2.71-2.75 (2H, m), 2.86-2.92 (1H, m), 3.09-3.14 (1H,m), 3.98-4.03 (4H, m), 4.32-4.43 (2H, m), 4.63-4.69 (2H,m), 6.82 (2H, d, J= 8.52Hz), 7.21 (2H, d, J= 8.68Hz), 7.27-7.31 (7H,m), 7.44 (2H, d, J = 8.07Hz), 7.51-7.54 (1H,m), 7.63-7.65 (1H,m), 7.77 (1H, d, J= 7.84Hz), 7.86 (1H, d, J= 1.68Hz), 8.35 (3H,s, br), 8.63-8.74 (3H, m).
89	d ⁶ - DMSO	δ: 1.29 (3H, t, J= 6.88Hz), 2.52-2.58 (1H, m), 2.71-2.79 (1H, m), 2.81-2.84 (1H, m), 3.02-3.07 (1H,m), 3.95-3.98 (4H, m), 4.24-4.36 (2H, m), 4.58-4.63 (2H,m), 6.76 (2H, d, J= 8.56Hz), 7.08 (2H, d, J= 8.56Hz), 7.14-7.25 (9H, m), 7.33-7.43 (4H,m), 8.285 (3H,s, br), 8.47 (2H, t, J= 7.84Hz), 8.62 (1H, t, J= 5.764Hz).
90	d ⁶ - DMSO	δ: 1.26 (3H,t,J= 6.89Hz), 2.61-2.74 (2H, m), 2.79-2.85 (1H, m), 3.02-3.07 (1H, m), 3.91-3.96 (4H, m), 4.25-4.36 (2H, m), 4.58-4.67 (2H,m), 6.74 (2H, d, J= 8.56Hz), 7.15-7.24 (9H, m), 7.37 (2H, d, J= 8.12Hz), 7.68 (1H, t, J= 7.80Hz), 7.87 (1H, d, J= 7.89Hz), 8.03 (1H, d, J= 7.89Hz), 8.09 (1H, s) 8.28 (3H,s, br), 8.48-8.65 (2H, m), 8.82 (1H, d, J= 8.28Hz).
91	d ⁶ - DMSO	δ: 1.27 (3H, t, J= 6.88Hz), 1.55-1.61 (11H, m), 1.89 (2H, s), 2.49 (2H, s), 2.62-2.72 (2H, m), 2.79-2.84 (1H, m), 3.01-3.05 (1H,m), 3.91-3.96 (4H, m), 4.23-4.39 (3H, m), 4.49-4.54 (1H,m), 6.69 (2H, d, J= 8.36Hz), 6.88 (2H, d, J= 8.37Hz), 7.19-7.25 (8H, m), 7.39 (2H, d, J = 7.84Hz), 8.32-8.38 (3H,m), 8.65-8.68 (1H, m).
92	d ⁶ - DMSO	δ: 1.26 (3H, t, J= 6.93Hz), 2.32 (3H, s), 2.67 (2H,d, J = 7.20Hz), 2.78-2.84 (1H,m), 3.02-3.07 (1H,m), 3.91-3.97 (4H, m), 4.27-4.36 (2H, m), 4.57-4.60 (2H,m), 6.73 (2H, d, J= 8.57Hz), 7.12 (2H, d, J= 8.57Hz), 7.21-7.24 (9H,m), 7.36 (2H, t, J= 8.08Hz), 7.65 (2H, t, J= 8.17Hz), 8.25 (3H,s, br), 8.37 (1H, d, J = 8.08Hz), 8.54 (1H, d, J= 8.60Hz), 8.59-8.62 (1H, m).

93	d ⁶ - DMSO	δ : 1.33 (3H,t,J= 6.97Hz), 2.67-2.79 (2H,m), 2.86-2.91 (1H,m), 3.093-3.14 (1H, m), 3.98-4.04 (4H, m), 4.32-4.44 (2H, m), 4.63-4.70 (2H, m), 6.81 (2H, d, J= 8.57Hz), 7.21 (2H, d, J= 8.60Hz), 7.27-7.32 (8H,m), 7.44 (2H, t, J= 8.08Hz), 7.79 (2H, s), 8.06 (1H, s), 8.38 (3H,s, br), 8.63-8.70 (1H, m), 8.83 (1H, d, J= 8.28Hz).
94	d ⁶ - DMSO	δ : 1.26 (3H, t, J= 6.93Hz), 2.61-2.71 (2H, m), 2.71-2.84 (1H, m), 3.02-3.06 (1H, m), 3.92-3.96 (4H, m), 4.25-4.35 (2H, m), 4.57-4.64 (2H, m), 6.75 (2H, d, J= 8.64Hz), 7.14 (2H, d, J= 8.56Hz), 7.21-7.25 (8H,m), 7.36 (2H, d, J= 8.08Hz), 7.76-7.79 (3H,m), 8.24 (3H,s, br), 8.59 (1H, t, J= 6.32Hz), 8.78 (1H, d, J= 8.24Hz).
95	d ⁶ - DMSO	δ : 1.26 (3H, t, J= 6.97Hz), 2.65-2.69 (2H, m), 2.79-2.84 (1H, m), 3.03-3.07 (1H, m), 3.797 (3H, s), 3.91-3.95 (4H, m), 4.25-4.34 (2H, m), 4.53-4.59 (2H,m), 6.73 (2H, d, J= 8.57Hz), 6.95 (2H, d, J= 8.88Hz), 7.13 (2H, d, J= 8.64Hz), 7.18-7.24 (7H,m), 7.37 (2H, d, J= 8.12H), 7.74 (2H, t, J= 8.85Hz), 8.29 (3H,s, br), 8.34 (1H, d, J = 8.60Hz), 8.54 (1H, d, J= 8.60Hz), 8.61 (1H, t, J= 5.96Hz).
96	d ⁶ - DMSO	δ : 1.27 (3H, t, J= 6.92Hz), 2.61-2.66 (2H, m), 2.76-2.82 (1H, m), 3.01-3.06 (1H, m), 3.90-3.95 (4H, m), 4.28-4.31 (2H, m), 4.52-4.62 (2H, m), 6.67-6.73 (3H, m), 6.97 (2H, d, J= 8.53Hz), 7.18-7.27 (10H, m), 7.35-7.41 (3H, m), 7.50 (2H, d, J= 6.73Hz), 8.21 (4H, d, J= 8.00Hz), 8.59 (2H, t, J= 7.24Hz).
97	d ⁶ - DMSO	δ : 1.28 (3H, t, J= 6.88Hz), 2.40-2.46 (1H, m), 2.59-2.64 (1H, m), 2.73-2.79 (1H,m), 2.98-3.03 (1H,m), 3.43 (2H, s), 3.51-3.57 (1H,m), 3.67-3.73 (1H, m), 3.92-3.97 (4H, m), 4.24-4.33 (2H, m), 4.43-4.54 (2H,m), 6.68 (2H, d, J= 8.56Hz), 6.88 (2H, d, J= 8.57Hz), 7.20-7.26 (12H, m), 7.37 (2H, d, J= 8.08Hz), 7.96 (1H, d, J= 8.21Hz), 8.20-8.27 (4H, m), 8.47 (1H, d, J= 8.50Hz), 8.57 (1H, t, J= 5.97Hz).

98	d ⁶ - DMSO	δ: 1.27 (3H, t, J= 6.93Hz), 2.47-2.49 (1H, m), 2.62-2.66 (1H, m), 2.76-2.82 (1H, m), 3.01-3.06 (1H, m), 3.72-3.77 (1H, m), 3.85-3.91 (3H, m), 3.94-3.98 (2H, m), 4.26-4.30 (2H, m), 4.44-4.54 (2H,m), 6.61 (2H, d, J= 8.64Hz), 6.83 (2H, d, J= 8.60Hz), 7.18 (2H, d, J= 8.05Hz), 7.26-7.27 (5H,m), 7.36 (2H, d, J= 8.12Hz), 7.46 (2H, d, J= 7.68Hz), 7.51-7.55 (1H,m), 7.84 (2H, t, J= 7.126Hz), 7.92 (1H, d, J= 8.08Hz), 8.30 (3H,s, br), 8.50 (1H, d, J = 8.53Hz), 8.57 (1H, t, J= 5.93Hz), 8.73 (1H, t, J = 5.84).
99	d ⁶ - DMSO	δ: 1.26 (3H, t, J= 6.93Hz), 2.65-2.67 (2H, m), 2.78-2.84 (1H, m), 3.03-3.07 (1H, m), 3.91-3.96 (4H, m), 4.25-4.36 (2H, m), 4.55-4.61 (2H, m), 6.74 (2H, d, J= 8.60Hz), 7.14 (2H, d, J= 8.64Hz), 7.21-7.28 (9H,m), 7.36 (2H, d, J = 8.12Hz), 7.80-7.83 (2H, m), 8.23 (3H,s, br), 8.53-8.583 (3H, m).
100	d ⁶ - DMSO	δ: 1.36(3H,t,J= 7.0Hz), 2.50-2.56(1H,m), 2.59-2.63(1H,m), 2.82-2.88(1H,m), 3.02-3.07(1H,m), 3.79-3.85(1H,m), 3.97(2H,s), 4.03(2H,q,J= 7.0Hz), 4.16-4.21(2H,m), 4.32(2H,d,J= 5.8Hz), 4.63-4.69(1H,m), 6.87(2H,d,J= 8.6Hz), 7.15(2H,d,J= 8.6Hz), 7.20-7.35(14H,m), 7.47-7.57(1H,m), 7.73-7.81(2H,m), 8.61-8.66(2H,m).
101	d ⁶ - DMSO	δ: 1.28 (3H, t, J= 6.92Hz), 2.36 (3H, s), 2.38-2.42 (1H, m), 2.56-2.60 (1H, m), 2.73-2.78 (1H, m), 2.98-3.03 (1H, m), 3.23-3.28 (1H, m), 3.92-3.98 (4H, m), 4.27-4.32 (2H, m), 4.41-4.46 (1H, m), 4.49-4.55 (1H, m), 6.66 (2H, d, J= 8.61Hz), 6.79 (2H, d, J= 8.60Hz), 7.20 (2H, d, J= 6.21Hz), 7.26-7.29 (6H,m), 7.32-7.40 (4H,m), 7.62 (2H, d, J= 8.28Hz), 7.80-7.88 (2H, m), 8.29 (3H,s, br), 8.52 (1H, d, J = 8.52Hz), 8.59-8.62 (1H, m).
102	d ⁶ - DMSO	δ: 1.27(3H,t,J= 7.0Hz), 2.42-2.47(1H,m), 2.64-2.69(1H,m), 2.75-2.81(1H,m), 3.04-3.08(1H,m), 3.92(2H,q,J= 7.0Hz), 3.96(2H,s), 4.28(2H,d,J= 5.8Hz), 4.43-4.53(2H,m), 6.23(1H,d,J= 7.7Hz), 6.65(4H,s), 7.17-7.23(5H,m), 7.26-7.36(8H,m), 8.19(2H,s), 8.63-8.65(2H,m), 8.91(1H,s).

103	d ⁶ - DMSO	δ: 1.26 (3H, t, J= 6.93Hz), 1.31-1.43 (7H, m), 2.68 (2H, d, J= 7.47Hz), 2.79-2.84 (1H, m), 3.02-3.07 (1H, m), 3.90-3.95 (2H, m), 4.06 (2H, d, J= 6.40Hz), 4.25-4.32 (2H, m), 4.56-4.63 (2H,m), 6.75 (2H, d, J= 8.52Hz), 7.13-7.22 (12H, m), 7.32-7.35 (1H, m), 7.40-7.43 (2H, m), 7.59-7.63 (1H, m), 7.74 (2H, d, J = 7.90Hz), 8.45-8.52 (3H, m).
104	d ⁶ - DMSO	δ:1.26 (3H,t,J= 6.93Hz), 2.65-2.75 (2H, m + 3H, s), 2.82-2.86 (1H, m), 3.04-3.08 (1H,m), 3.91-3.96 (4H, m), 4.24-4.36 (2H, m), 4.57-4.69 (2H,m), 6.73 (2H, d, J= 8.52Hz), 7.14 (2H, d, J= 8.68Hz), 7.17-7.24 (6H,m), 7.39 (2H, d, J = 8.07Hz), 7.79 (1H, d, J= 7.84Hz), 8.44 (3H,s, br), 8.49-8.52 (1H, m), 8.65-8.69 (2H, m), 9.00 (1H, d, J= 1.68Hz), 9.14 (1H, d, J= 1.68Hz).
105	d ⁶ - DMSO	δ: 1.29 (3H,t,J= 6.93Hz), 2.42 (3H, s), 2.75-2.86 (3H, m), 3.05-3.09 (1H,m), 3.95-3.97 (4H, m), 4.25-4.37 (2H, m), 4.61-4.74 (2H,m), 6.78 (2H, d, J= 8.52Hz), 7.09 (2H, d, J= 8.68Hz), 7.21-7.27 (6H,m), 7.40 (2H, d, J = 8.07Hz), 7.76-7.80 (1H, m), 8.07-8.13 (1H, m), 8.42 (3H,s, br), 8.67-8.73 (3H, m), 8.94 (1H, d, J= 1.68Hz).
106	d ⁶ - DMSO	δ: 1.13 (3H,t,J= 6.93Hz), 2.70-2.84 (3H, m), 3.01-3.06 (1H,m), 3.95-3.98 (4H, m), 4.24-4.36 (2H, m), 4.59-4.69 (2H,m), 6.76 (2H, d, J= 8.52Hz), 7.04 (2H, d, J= 8.68Hz), 7.16-7.26 (6H,m), 7.38 (2H, d, J = 8.07Hz), 7.61 (2H, s), 8.33 (3H,s, br), 8.58-8.60 (1H, m), 8.64-8.67 (1H, m), 8.78 (1H, d, J= 1.68Hz).
107	d ⁶ - DMSO	δ: 1.24 (3H,t,J= 6.93Hz), 2.56-2.62 (1H, m), 2.69-2.73 (1H, m), 2.78-2.84 (1H, m), 3.02-3.07 (1H,m), 3.89-3.95 (4H, m), 4.24-4.37 (2H, m), 4.58-4.67 (2H,m), 6.75 (2H, d, J= 8.52Hz), 7.13 (2H, d, J= 8.68Hz), 7.16-7.25 (8H, m), 7.36 (2H, d, J = 8.07Hz), 8.26 (3H,s, br), 8.59-8.64 (1H, m), 8.65 (1H, d, J= 1.68Hz), 8.94 (1H, d, J= 1.68Hz).
108	d ⁶ - DMSO	δ: 1.28 (3H,t,J= 6.93Hz), 2.69-2.82 (2H, m), 3.02-3.09 (2H,m), 3.92-3.98 (4H, m), 4.26-4.33 (2H, m), 4.59-4.62 (2H,m), 6.74 (2H, d, J= 8.52Hz), 6.98 (2H, d, J= 8.68Hz), 7.23-7.29 (7H, m), 7.37-7.40 (2H, m), 8.05 (1H,s, br), 8.23 (3H,s, br), 8.65-8.69 (1H, m), 8.85 (1H, d, J= 1.68Hz).

109	d ⁶ - DMSO	δ: 1.28 (3H,t,J= 6.93Hz), 2.36-2.40 (1H, m), 2.59-2.63 (1H, m), 2.73-2.79 (1H, m), 3.00-3.03 (1H, m), 3.19-3.27 (1H, m), 3.91-3.97 (4H, m), 4.28-4.36 (2H, m), 4.52-4.54 (2H,m), 6.68 (2H, d, J= 8.52Hz), 6.84 (2H, d, J= 8.68Hz), 7.19-7.24 (7H,m), 7.38 (2H, d, J = 8.07Hz), 8.35 (3H,s, br), 8.59-8.63 (2H, m).
110	d ⁶ - DMSO	δ: 1.26 (3H, t, J= 6.93Hz), 2.47 (3H, s), 2.49-2.50 (1H, m), 2.59-2.63 (1H, m), 2.69-2.74 (1H, m), 3.02-3.07 (1H,m), 3.91-3.95 (4H, m), 4.28-4.37 (2H, m), 4.57-4.66 (2H,m), 4.79-4.80 (2H,m), 6.74 (2H, d, J= 8.52Hz), 7.14 (2H, d, J= 8.68Hz), 7.16-7.23 (6H, m), 7.38 (2H, d, J = 8.07Hz), 7.55 (2H, d, J= 7.84Hz), 8.40 (2H,s, br), 8.63-8.67 (1H, m), 8.91 (1H, d, J= 1.68Hz).
111	d ⁶ - DMSO	δ: 1.27 (3H, t, J= 6.93Hz), 2.30 (3H, s), 2.59 (3H, s), 2.61-2.72 (2H, m), 2.79-2.84 (1H, m), 3.04-3.09 (1H,m), 3.91-3.96 (4H, m), 4.29-4.35 (2H, m), 4.50-4.60 (2H, m), 6.73 (2H, d, J= 8.52Hz), 7.03 (2H, d, J= 8.68Hz), 7.15-7.27 (7H, m), 7.39 (2H, d, J = 8.07Hz), 8.07 (1H, d, J= 7.84Hz), 8.42 (2H,s, br), 8.62-8.66 (2H, m).
112	d ⁶ - DMSO	δ: 1.26 (3H, t, J= 6.93Hz), 2.66 (3H, s), 2.77-2.84 (3H, m), 3.03-3.08 (1H, m), 3.87-3.97 (4H, m), 4.28 (2H, d), 4.49-4.55 (1H, m), 4.70-4.75 (1H, m), 6.61 (2H, d, J= 8.52Hz), 6.67 (2H, d, J= 8.68Hz), 7.19-7.30 (7H, m), 7.38 (2H, d, J = 8.07Hz), 7.89 (1H, d, J= 7.84Hz), 8.06 (1H, s), 8.41 (2H,s, br), 8.65-8.70 (2H, m).
113	d ⁶ - DMSO	δ: 1.27 (3H, t, J= 6.93Hz), 2.36 (3H, s), 2.45 (3H, s), 2.63-2.68 (1H, m), 2.75-2.84 (2H, m), 3.04-3.08 (1H,m), 3.88-3.96 (4H, m), 4.29 (2H, d), 4.48-4.54 (1H, m), 4.64-4.69 (1H, m), 6.62 (2H, d, J= 8.52Hz), 6.65 (2H, d, J= 8.68Hz), 7.17-7.24 (6H, m), 7.39 (2H, d, J = 8.07Hz), 7.54 (1H, d, J= 7.84Hz), 8.40 (3H,s, br), 8.64-8.66 (2H, m).
114	d ⁶ - DMSO	δ: 1.26 (3H, t, J= 7.07Hz), 2.09 (3H, s), 2.67-2.77 (2H, m), 2.78—2.88 (1H, m), 3.05-3.09 (1H, m), 3.92-3.97 (4H, m), 4.31-4.40 (2H, m), 4.59-4.63 (1H, m), 4.64-4.74 (1H, m), 6.77 (2H, d, J= 8.34Hz), 7.17-7.29 (10H, m), 7.39 (2H, d, J = 8.36Hz), 7.87 (1H, d, J= 8.55Hz), 8.26-8.36 (4H, m), 8.62 (1H, s), 8.68-8.71 (2H, m), 8.88-8.95 (1H, m).

115	d ⁶ - DMSO	δ: 1.26 (3H, t, J= 6.36Hz), 2.66-2.87 (3H, m), 3.05-3.10 (1H,m), 3.92-3.97 (4H, m), 4.31-4.38 (2H, m), 4.38-4.40 (2H, m), 4.59-4.63 (1H,m), 4.69-4.75 (1H,m), 6.77 (2H, d, J= 7.95Hz), 7.14 (2H, d, J= 7.95Hz), 7.16-7.23 (6H, m), 7.38 (2H, d, J = 7.82Hz), 7.55 (2H, d, J= 7.95Hz), 7.74 (1H, d, J= 11.13Hz), 8.40 (3H,s, br), 8.63-8.67 (1H, m), 8.91 (1H, d, J= 2.69Hz), 9.05 (1H, J = 8.68Hz), 9.20 (1H, d, J = 2.2Hz).
116	d ⁶ - DMSO	δ: 1.28(3H, t, J= 6.57Hz), 2.61-2.67 (1H, m), 2.79-2.85 (2H, m), 3.08-3.12 (1H,m), 3.92-3.98 (4H, m), 4.31 (2H, d, J = 6.57Hz), 4.53-4.58 (1H,m), 4.67-4.71 (1H, m), 6.66 (2H, d, J= 8.09Hz), 6.74 (2H, d, J= 8.23Hz), 7.14 (1H, d, J= 5.47Hz), 7.22-7.25 (2H, m), 7.30-7.35 (4H, m), 7.39 (2H, d, J = 8.09Hz), 7.82 (1H, d, J= 7.66Hz), 7.86 (1H, d, J= 5.47Hz), 8.35 (3H,s, br), 8.69(1H, t, J = 6.57Hz), 8.76 (1H, d, J= 8.21Hz).
117	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.0 Hz), 2.38 (3H, s), 2.62 (1H, d, J = 10.3 Hz), 2.70 (1H, dd, J = 14.3 Hz, 4.2 Hz), 2.82 (1H, dd, J = 13.5 Hz, 10.1 Hz), 3.07 (1H, dd, J = 13.6 Hz, 4.5 Hz), 3.95 (2H, q, J = 7.0 Hz), 3.98 (2H, d, J = 4.8 Hz), 4.26 - 4.37 (2H, m), 4.52 - 4.62 (2H, m), 6.75 (2H, d, J = 8.7 Hz), 7.07 (2H, d, J = 8.6 Hz), 7.18 - 7.12 (1H, m), 7.23 - 7.28 (6H, m), 7.38 (2H, d, J = 8.1 Hz), 8.21 (1H, d, J = 8.3 Hz), 8.26 (2H, br.s), 8.62 (2H, d, J = 7.6 Hz), 9.01 (1H, s)
118	d ⁶ - DMSO	δ: 1.27 (3H, t, J = 7.0 Hz), 2.64 - 2.67 (2H, m), 2.81(1H, dd, J = 13.5 Hz, 10.1 Hz), 3.04 (1H, dd, J = 13.5, 4.6 Hz), 3.93 (2H, q, J = 7.0 Hz), 3.97 (2H, d, J = 5.4 Hz), 4.30 (2H, dd, J = 5.5 Hz, 3.5 Hz), 4.54 - 4.63 (2H, m), 6.60 (1H, dd, J = 3.5 Hz, 1.9 Hz), 6.7 (2H, d, J = 8.6 Hz), 7.01 (2H, d, J = 8.6 Hz), 7.11 (1H, d, J = 3.5 Hz), 7.16 - 7.28 (7H, m), 7.38 (2H, d, J = 8.1 Hz), 7.81 (1H, d, J = 1.1 Hz), 8.18 (1H, d, J = 8.4 Hz), 8.28 (2H, br.s), 8.60 (1H, d, J = 8.6 Hz), 8.64 (1H, t, J = 6.0 Hz)

119	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 6.9 Hz), 2.22 (3H, s), 2.65 (1H, dd, J = 13.6 Hz, 9.3 Hz), 2.72 (1H, dd, J = 13.8 Hz, 4.2 Hz), 2.82 (1H, dd, J = 13.5 Hz, 10.2 Hz), 3.08 (1H, dd, J = 13.5 Hz, 4.5 Hz), 3.94 (2H, q, J = 7.0 Hz), 3.98 (2H, br.s), 4.31 (2H, d, J = 5.8 Hz), 4.52 - 4.59 (2H, m), 6.73 (2H, d, J = 8.6 Hz), 6.91(1H, d, J = 5.0 Hz), 6.98 (2H, d, J = 8.6 Hz), 7.18 - 7.21 (1H, m), 7.23 - 7.30 (6H, m), 7.38 (2H, d, J = 7.9 Hz), 7.55 (1H, d, J = 5.0 Hz), 7.72 (1H, d, J = 7.8 Hz), 8.23 (2H, br.s), 8.62 (2H, d, J = 9.6 Hz)
120	d ⁶ - DMSO	δ: 1.27(3H,t,J= 7.0Hz), 2.70-2.75(1H,m), 2.80-2.86(2H,m), 3.07-3.11(1H,m), 3.88(3H,s), 3.92(2H,q,J= 6.9Hz), 3.97(2H,q,J= 5.7Hz), 4.31(2H,d,J= 6.1Hz), 4.52-4.58(1H,m), 4.67-4.72(1H,m), 6.65(2H,d,J= 8.6Hz), 6.76(2H,d,J= 8.4Hz), 7.02-7.04(1H,m), 7.21-7.34(7H,m), 7.37(2H,d,J= 8.1Hz), 7.53-7.55(1H,m), 7.85-7.88(1H,m), 8.23(2H,s), 8.27(1H,d,J= 8.1Hz), 8.67(1H,t,J= 5.9Hz), 8.71(1H,d,J= 8.4Hz).
121	d ⁶ - DMSO	δ: 1.28(3H,t,J= 6.9Hz), 2.66-2.75(2H,m), 2.80-2.86(1H,m), 3.03-3.08(1H,m), 3.87(3H,s), 3.91-4.00(4H,m), 4.30-4.38(2H,m), 4.59-4.68(2H,m), 6.77(2H,d,J= 8.6Hz), 7.16(1H,s), 7.18-7.21(2H,m), 7.23-7.27(6H,m), 7.38(2H,d,J= 8.2Hz), 7.70(1H,t,J= 4.2Hz), 8.24(2H,s), 8.44(1H,d,J= 2.9Hz), 8.51(1H,d,J= 1.6Hz),, 8.62-8.64(2H,m), 8.79(1H,d,J= 8.4Hz).
122	d ⁶ - DMSO	δ: 1.29(3H,t,J= 7.0Hz), 2.56-2.62(1H,m), 2.73-2.84(2H,m), 3.07-3.11(1H,m), 3.89(3H,s), 3.94(2H,q,J= 7.0Hz), 3.98(2H,s), 4.30(2H,d,J= 6.0Hz), 4.49-4.55(1H,m), 4.65-4.70(1H,m), 6.67(4H,s), 7.11(1H,d,J= 5.4Hz), 7.20-7.25(3H,m), 7.29-7.35(4H,m), 7.37(2H,d,J= 8.1Hz), 7.47(1H,d,J= 7.4Hz), 7.72(1H,d,J= 5.5Hz), 8.18(2H,s), 8.63(1H,t,J= 6.0Hz), 8.67(1H,d,J= 8.5Hz).
123	d ⁶ - DMSO	δ: 1.28(3H,t,J= 7.0Hz), 2.58-2.70(2H,m), 2.79-2.85(1H,m), 3.03-3.08(1H,m), 3.87(3H,s), 3.93(2H,q,J= 7.0Hz), 3.99(2H,q,J= 5.5Hz), 4.26-4.38(2H,m), 4.59-4.65(2H,m), 6.76(2H,d,J= 8.6Hz), 7.08(1H,s), 7.15(2H,d,J= 8.3Hz), 7.18-7.27(8H,m), 7.36(2H,d,J= 8.0Hz), 8.19(2H,s), 8.25(1H,d,J= 5.2Hz), 8.59-8.62(2H,m), 8.72(1H,d,J= 8.1Hz).

124	d ⁶ - DMSO	δ: 1.28(3H, t, J = 6.84Hz), 2.13 (3H, s), 2.56-2.59 (1H, m), 2.69-2.74 (1H, m), 2.78-2.84 (1H, m), 3.05-3.10 (1H,m), 3.91-4.00 (4H, m), 4.30 (2H, d, J = 5.82Hz), 4.52-4.64 (2H,m), 5.91 (1H, s) 6.71 (2H, d, J = 8.66Hz), 6.77 (1H, t, J = 2.36Hz), 6.92 (2H, d, J = 7.78Hz), 7.18-7.28 (8H, m), 7.34 (2H, d, J = 8.49Hz), 8.09 (3H,s, br), 8.60-8.64(2H, m).
125	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.0 Hz), 2.67 - 2.69 (2H, m), 2.82 (1H, dd, J = 13.5 Hz, 10.0 Hz), 3.05 (1H, dd, J = 13.6 Hz, 6.4 Hz), 3.93 (2H, q, J = 7.0 Hz), 3.97 (2H, d, J = 5.6 Hz), 4.31 (2H, d, J = 5.8 Hz), 4.47 - 4.57 (2H, m), 6.57 (1H, d, J = 5.4 Hz), 6.71 (2H, d, J = 8.6 Hz), 6.97 (2H, d, J = 8.6 Hz), 7.18 - 7.27 (10H m), 7.38 (2H, d, J = 6.0 Hz), 7.40 (1H, d, J = 3.1 Hz), 8.28 (2H, br.s), 8.51 (1H, d, J = 8.5 Hz), 8.61 (1H, t, J = 6.0 Hz)
126	d ⁶ - DMSO	δ: 0.80-0.85(2H,m), 1.02-1.12(4H,m), 1.22-1.32(1H,m), 1.41-1.48(1H,m), 1.53-1.62(5H,m), 2.76-2.82(1H,m), 3.09-3.13(1H,m), 3.95(2H,s), 4.28-4.35(2H,m), 4.37-4.43(1H,m), 4.46-4.52(1H,m), 7.15-7.27(7H,m), 7.34-7.37(2H,m), 7.39-7.46(2H,m), 7.48-7.54(1H,m), 7.80-7.83(2H,m), 8.21(2H,s), 8.51(2H,d,J= 7.8Hz), 8.59(1H,t,J= 6.1Hz).
127	d ⁶ - DMSO	δ: 2.62-2.69 (2H, m), 2.71-2.77 (1H, m), 2.96-3.01(1H, m), 3.87 (2H, d, J = 4.56Hz), 4.22-4.25 (2H, m), 4.49-4.61 (2H,m), 7.10-7.22 (10H, m), 7.30-7.36 (4H, m), 7.40-7.48 (1H, m), 7.67 (2H, d, J = 6.61Hz), 8.30 (3H,s, br), 8.47 (1H, d, J = 8.82Hz), 8.56-8.62 (2H, m).
128	d ⁶ - DMSO	δ: 2.70 (2H, d, J = 7.21Hz), 2.78-2.85 (1H, m), 3.03-3.07 (1H, m), 3.95 (2H, d, J = 8.78Hz), 4.27-4.36 (2H, m), 4.55-4.63 (2H,m), 6.84 (2H, d, J = 8.52Hz), 7.05-7.29 (12H, m), 7.37-7.44 (6H, m), 7.74 (2H, d, J = 7.52Hz), 8.34 (3H,s, br), 8.50 (1H, d, J = 7.57Hz), 8.58 (1H, d, J = 8.77Hz), 8.63-8.66 (2H, m).
129	d ⁶ - DMSO	δ: 2.61-2.63 (2H, m), 2.70-2.77 (1H, m), 2.95-3.00 (1H, m), 3.88 (2H, d, J = 6.26Hz), 4.18-4.29 (2H, m), 4.47-4.54 (2H,m), 4.91 (1H, s), 6.76 (2H, d, J = 8.18Hz), 7.07-7.18 (9H, m), 7.26-7.36 (4H, m), 7.66-7.68 (2H, m), 8.24 (3H,s, br), 8.42 (1H, d, J = 8.17Hz), 8.50 (1H, d, J = 8.02Hz), 8.53-8.58 (1H, m).

130	d ⁶ - DMSO	δ: 2.65 (2H, d, J = 5.97Hz), 2.79-2.85 (1H, m), 3.03-3.06 (1H, m), 3.95 (2H, s), 4.31 (2H, s), 4.55 (2H, s, br), 6.58 (2H, d, J= 7.55Hz), 7.01 (2H, d, J= 7.12Hz), 7.18-7.24 (10H, m), 7.37-7.50 (7H, m), 7.74 (2H, d, J= 7.10Hz), 8.32 (3H,s, br), 8.46 (1H, d, J= 9.60Hz), 8.53 (1H, d, J= 9.60Hz), 8.61 (1H, s).
131	d ⁶ - DMSO	δ: 1.43 (3H, t, J= 6.97Hz), 2.88 (2H, d, J= 6.67Hz), 2.95-3.06 (2H, m), 3.14-3.23 (2H, m), 4.06-4.12 (2H, m), 4.39-4.49 (2H, m), 4.75-4.78 (2H, m), 6.91 (2H, d, J= 8.95Hz), 7.31 (2H, m), 7.37-7.46 (9H, m), 7.54-7.67 (5H, m), 7.89-7.98 (2H, m), 8.46 (2H, s, br), 8.65-8.81 (3H, m).
132	d ⁶ - DMSO	δ: 2.68 (2H, d, J = 7.36Hz), 2.79-2.98 (1H, m), 3.03-3.06 (1H, m), 3.96 (2H, s), 4.21-4.32 (2H, m), 4.59-4.61 (2H,m), 5.00 (2H, s), 6.84 (2H, d, J= 8.15Hz), 7.14-7.24 (12H, m), 7.35-7.40 (5H, m), 7.67-7.76 (2H, m), 7.82-7.88 (2H, m), 8.24 (1H,s, br), 8.47-8.64 (4H, m).
133	d ⁶ - DMSO	δ: 0.81(3H,d,J= 6.3Hz), 0.87(3H,d,J= 6.3Hz), 1.27-1.39(2H,m), 1.45-1.52(1H,m), 2.81-2.87(1H,m), 3.20-3.24(1H,m), 4.02(2H,s), 4.35-4.44(3H,m), 4.52-4.58(1H,m), 7.22-7.35(7H,m), 7.43(2H,d,J= 8.1Hz), 7.48-7.52(2H,m), 7.58-7.64(1H,m), 7.87-7.89(2H,m), 8.30(2H,s), 8.59-8.69(3H,m).
134	d ⁶ - DMSO	δ: 2.70 (2H, d, J = 7.36Hz), 2.78-2.84 (1H, m), 3.03-3.06 (1H, m), 3.97 (2H, s), 4.11-4.34 (2H, m), 4.59-4.63 (2H,m), 5.14 (2H, s), 6.90 (2H, d, J= 8.21Hz), 7.11-7.30 (11H, m), 7.36-7.54 (5H, m), 7.67-7.75 (2H, m), 7.80-7.88 (2H, m), 8.47-8.58 (4H, m).
135	d ⁶ - DMSO	δ: 2.20 (3H, s), 2.71 (2H, d, J = 7.81Hz), 2.79-2.85 (1H, m), 3.03-3.07(1H,m), 3.93-3.96 (2H, m), 4.25-4.36 (2H, m), 4.55-4.65 (2H,m), 7.00 (2H, d, J= 9.01Hz), 7.12 (2H, d, J= 7.78Hz), 7.18 (2H, d, J= 7.79Hz), 7.22-7.24 (4H, m), 7.38 (2H, d, J= 6.49Hz), 7.42 (2H, d, J= 7.79Hz), 7.48-7.50 (1H, m), 7.74-7.76 (2H,m), 8.33 (3H,s, br), 8.51 (1H, d, J= 7.79Hz), 8.58 (1H, d, J= 7.79Hz), 8.62-8.65 (1H, m).

136	d ⁶ - DMSO	δ: 2.70 (2H, d, J = 7.28Hz), 2.74-2.85 (1H, m), 3.04-3.08 (1H, m), 3.95 (2H, s), 4.27-4.36 (2H, m), 4.58-4.62 (2H,m), 5.03 (2H, s), 6.85 (2H, d, J= 8.21Hz), 7.15-7.24 (11H, m), 7.37-7.52 (5H, m), 7.56-7.61 (1H, m), 7.75 (2H, d, J = 8.22Hz), 8.34 (3H, s, br), 8.50(1H, d, J= 7.02Hz), 8.58(1H, d, J= 9.38Hz), 8.62-8.65 (1H, m).
137	d ⁶ - DMSO	δ: 2.66-2.69 (2H, m), 2.81 (1H, dd, J = 13.5, 10.0Hz), 3.05 (1H, dd, J = 13.6, 4.7Hz), 3.69 (2H, s), 3.77 (3H, s), 4.23-4.33 (2H, m), 4.58-4.67 (2H, m), 6.68 (1H, d, J = 8.5Hz), 7.07-7.25 (11H, m), 7.43 (2H, t, J = 7.3Hz), 7.49-7.55 (2H, m), 7.75 (2H, d, J = 7.2Hz), 8.00 (1H, d, J = 1.8Hz),8.49-8.57 (3H, m).
138	d ⁶ - DMSO	δ: 2.69-2.71 (1H, m), 2.79-2.85 (1H, m), 3.03-3.07(1H,m), 3.67 (3H, s), 3.96 (2H, s), 4.26-4.32 (2H, m), 4.55-4.63 (2H,m), 6.76 (2H, d, J= 8.52Hz), 7.14-7.25 (9H, m), 7.36-7.44 (4H, m), 7.48-7.52 (1H,m), 7.74-7.76 (2H,m), 8.29 (3H,s, br), 8.56 (1H, d, J= 1.68Hz), 8.613-8.64 (2H, m).
139	d ⁶ - DMSO	δ: 2.82-2.85 (3H, m), 3.03-3.08 (1H,m), 3.93-3.98 (2H, m), 4.26-4.31 (2H, m), 4.56-4.69 (2H,m), 7.11-7.26 (12H, m), 7.37-7.43 (4H, m), 7.48-7.51 (1H,m), 7.73-7.75 (2H,m), 8.32 (3H,s, br), 8.53-8.65 (2H, m).
140	d ⁶ - DMSO	δ: 1.09 (3H, t, J = 7.0Hz), 2.50-2.66 (2H, m), 3.05 (1H, dd, J = 13.4, 10.4Hz), 3.26-3.32 (1H, m), 3.45-3.54 (2H, m), 3.74 (2H, q, J = 7.0Hz), 4.09-4.15 (2H, m), 4.33-4.38 (1H, m), 4.65-4.71 (1H, m), 6.57 (2H, d, J = 8.6Hz), 7.00 (2H, d, J = 8.5Hz), 7.09 (2H, d, J = 8.0Hz), 7.16-7.35 (5H, m), 7.53-7.58 (4H, m), 7.95-8.20 (3H, br m), 8.48-8.56 (3H, m), 8.64 (1H, d, J = 8.5Hz).
141	d ⁶ - DMSO	δ: 1.26(3H,t,J= 7.0Hz), 2.65-2.84(3H,m), 3.03-3.08(1H,m), 3.92(2H,q,J= 7.0Hz), 3.96(2H,s), 4.27-4.35(2H,m), 4.57-4.63(2H,m), 6.75(2H,d,J= 8.7Hz), 7.16(2H,d,J= 8.6Hz), 7.23(1H,dd,J= 8.3,1.7Hz), 7.27(2H,d,J= 8.1Hz), 7.37-7.51(6H,m), 7.55(1H,d,J= 1.8Hz), 7.73-7.75(2H,m), 8.24(2H,s), 8.50(1H,d,J= 8.1Hz), 8.67-8.71(2H,m).

142	d ⁶ - DMSO	δ: 1.26(3H,t,J= 6.9Hz), 2.67-2.83(3H,m), 3.01-3.05(1H,m), 3.89-3.95(4H,m), 4.27-4.35(2H,m), 4.53-4.63(2H,m), 6.75(2H,d,J= 8.5Hz), 7.14(2H,d,J= 8.5Hz), 7.20-7.26(5H,m), 7.37-7.44(4H,m), 7.50(1H,t,J= 7.3Hz), 7.66-7.72(1H,m), 7.75(2H,d,J= 7.4Hz), 8.31(2H,s), 8.53(1H,d,J= 8.1Hz), 8.62(1H,d,J= 8.7Hz), 8.69(1H,t,J= 5.9Hz).
143	d ⁶ - DMSO	δ: 1.26 (3H, t, J= 6.93Hz), 2.70 (2H, d, J= 7.52Hz), 2.89-2.95 (1H, m), 3.12-3.17 (1H, m), 3.89-3.94 (2H, m), 3.95 (2H, s), 4.31 (2H, d, , J= 5.38Hz), 4.56-4.66 (2H,m), 6.74 (2H, d, J= 8.52Hz), 7.12 (2H, d, J= 6.93Hz), 7.26 (2H, d, J= 9.72Hz), 7.38-7.44 (5H, m), 7.50-7.54 (3H, m), 7.75 (2H, d, J= 8.31Hz), 8.34 (3H, s, br), 8.51 (1H, d, J= 8.32Hz), 8.62 (1H, d, J= 8.32Hz), 8.68-8.71 (1H, m).
144	d ⁶ - DMSO	δ: 1.26 (3H, t, J= 6.93Hz), 2.65 (2H, d, J= 6.98Hz), 2.89-2.95 (1H, m), 3.16-3.21 (1H, m), 3.91-3.96 (2H, m), 4.32 (2H, J= 5.82Hz), 4.57-4.66 (2H, m), 6.73 (2H, d, J= 8.54Hz), 7.13 (2H, d, J= 9.08Hz), 7.27 (2H, d, J= 8.01Hz), 7.39-7.44 (8H, m), 7.50-7.57 (1H, m), 7.68-7.71 (1H, m), 7.73-7.75 (2H, m), 8.44-8.50 (4H, m), 8.72-8.76 (2H, m).
145	d ⁶ - DMSO	δ: 1.26 (3H, t, J = 7.0Hz), 2.64 (2H, d, J = 7.2Hz), 2.82 (1H, dd, J = 13.5, 10.3Hz), 3.07 (1H, dd, J = 13.6, 4.2Hz), 3.81 (2H, s), 3.92 (2H, q, J = 7.0Hz), 4.26-4.35 (2H, m), 4.55-4.65 (2H, m), 5.70-6.60 (2H, br s), 6.75 (2H, d, J = 8.6Hz), 7.14-7.34 (7H, m), 7.39-7.51 (3H, m), 7.62 (1H, d, J = 7.7Hz), 7.73 (2H, d, J = 7.2Hz), 8.39 (1H, dd, J = 4.8, 1.4Hz), 8.46 (1H, d, J = 1.6Hz), 8.52 (1H, d, J = 8.0Hz), 8.62 (1H, t, J = 6.0Hz), 8.68 (1H, d, J = 8.8Hz).
146	d ⁶ - DMSO	δ: 1.26(3H,t,J= 7.0Hz), 2.69-2.77(3H,m), 2.94-2.99(1H,m), 3.65(3H,s), 3.92(2H,q,J= 7.0Hz), 3.96(2H,s), 4.25-4.35(2H,m), 4.49-4.54(1H,m), 4.57-4.63(1H,m), 6.75(4H,dd,J= 8.6,1.7Hz), 7.13-7.18(4H,m), 7.25(2H,d,J= 8.1Hz), 7.36(2H,d,J= 8.1Hz), 7.40-7.44(2H,m),7.48-7.52(1H,m), 7.72-7.75(2H,m), 8.19(2H,s), 8.48-8.53(2H,m), 8.62(1H,t,J= 6.0Hz).

147	d ⁶ - DMSO	δ: 1.26 (3H, t, J = 7.0 Hz), 2.69 (2H, d, J = 7.2 Hz), 2.83 (1H, dd J = 13.6, 10.0 Hz), 3.06 (1H, dd, J = 13.7, 4.5 Hz), 3.87 (2H,s), 3.92 (2H, q, J = 7.0 Hz), 4.26-4.31(2H, m), 4.55-4.60 (1H, m), 4.62-4.67 (1H, m), 6.51-6.79 (1H, m), 6.75 (2H, d, J = 8.6Hz), 7.13-7.19 (3H, m), 7.20-7.22 (4H, m), 7.31 (1H, d, J = 8.1Hz), 7.40-7.44 (3H, m), 7.45-7.52 (1H, m), 7.71-7.75 (2H, m), 8.36-8.39 (2H, m), 8.50-8.61 (3H, m).
148	d ⁶ - DMSO	δ: 1.26 (3H, t, J= 6.93Hz), 2.70 (2H, d, J= 7.09Hz), 2.89-2.95 (1H, m), 3.12-3.17 (1H, m), 3.89-3.94 (2H, m), 3.95 (2H, s), 4.31-4.32 (2H, m), 4.56-4.66 (2H,m), 6.74 (2H, d, J= 7.19Hz), 7.12 (2H, d, J= 8.40Hz), 7.26 (2H, d, J= 8.40Hz), 7.38-7.44 (5H, m), 7.50-7.54 (3H, m), 7.74-7.76 (2H, m), 8.35 (3H, s, br), 8.51 (1H, d, J= 7.2Hz), 8.62 (1H, d, J= 8.40Hz), 8.68-8.71 (1H, m).
149	d ⁶ - DMSO	δ: 1.26 (3H, t, J= 6.93Hz), 2.67-2.69 (2H, m), 2.81-2.87 (1H, m), 3.06-3.10 (1H,m), 3.59 (2H, s, br), 3.89-3.98 (4H, m), 4.22-4.33 (2H, m), 4.57-4.62 (2H,m), 6.74 (2H, d, J= 8.52Hz), 7.13-7.16 (7H, m), 7.22-7.27 (2H,m), 7.35-7.41 (3H,m), 7.43-7.51 (1H, m), 7.72-7.81 (2H,m), 8.29-8.32 (1H,m), 8.48-8.53 (1H, m), 8.54-8.64 (1H, m).
150	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.0 Hz), 2.80 - 2.87 (2H, m), 3.07 (1H, dd, J = 14.7, 9.4 Hz), 3.27 (2H, dd, J = 14.8, 4.3 Hz), 3.32 (2H, br.s), 3.69 (1H, s), 3.94 (2H, q, J = 6.9 Hz), 4.27 - 4.30 (2H, m), 4.53 - 4.58 (1H, m), 4.60 - 4.66 (1H, m), 6.78 (2H, d, J = 8.6 Hz), 6.86 - 6.90 (2H, m), 7.12 - 7.24 (6H, m), 7.31 (1H, d, J = 5.0 Hz), 7.42 (2H, t, J = 7.4 Hz), 7.50 (1H, q, J = 7.3 Hz), 7.73 (2H, d, J = 7.1 Hz), 8.51 (1H, t, J = 5.5 Hz), 8.57 (2H, m)
151	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 6.9 Hz), 2.77 (2H, br.s), 2.89 (1H, dd, J = 14.1, 9.8 Hz), 3.07 (1H, dd, J = 14.1, 4.4 Hz), 3.33 (3H, br.s), 3.65 (1H, br.s), 3.94 (2H, q, J = 6.9 Hz), 4.28 (2H, t, J = 5.8 Hz), 4.52 - 4.58 (1H, m), 4.60 - 4.65 (1H, m), 6.78 (2H, d, J = 8.5 Hz), 6.98 (1H, d, J = 4.6 Hz), 7.15 (3H, br.s), 7.18 - 7.23 (4H, m), 7.38 - 7.41 (1H, m), 7.44 (2H, d, J = 7.6 Hz), 7.51 (1H, t, J = 7.2 Hz), 7.75 (2H, d, J = 7.2 Hz), 8.50 (1H, br.s), 8.57 (2H, d, J = 6.2 Hz)

152	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.0 Hz), 2.67 (1H, t, J = 1.6 Hz), 2.77 (1H, d, J = 7.3 Hz), 3.06 (1H, dd, J = 14.5 Hz, 9.7 Hz), 3.23 (1H, dd, J = 14.6 Hz, 4.7 Hz), 3.94 (2H, q, J = 7.0 Hz), 3.96 - 3.99 (2H, m), 4.31 (2H, t, J = 5.7 Hz), 4.58 (1H, q, J = 7.5 Hz), 4.70 (1H, td, J = 9.1 Hz, 4.6 Hz), 6.77 (2H, d, J = 8.6 Hz), 7.16 (2H, d, J = 8.6 Hz), 7.23 (2H, d, J = 8.1 Hz), 7.26 (1H, d, J = 1.5 Hz), 7.33 (2H, d, J = 8.2 Hz), 7.43 (2H, t, J = 7.5 Hz), 7.52 (1H, t, J = 7.4 Hz), 7.42 (2H, d, J = 7.1 Hz), 8.05 (2H, br.s), 8.55 (2H, d, J = 7.6 Hz), 8.59 (1H, d, J = 8.6 Hz), 8.95 (1H, d, J = 1.8 Hz)
153	d ⁶ - DMSO	δ: 1.27 (3H, t, J = 7.0 Hz), 2.66 - 2.72 (3H, m), 3.10 (1H, dd, J = 14.4 Hz, 9.8 Hz), 3.91 (2H, q, J = 7.0 Hz), 3.98 (2H, d, J = 5.2 Hz), 4.33 (2H, qd, J = 14.5 Hz, 6.0 Hz), 4.61 (1H, td, J = 8.7 Hz, 4.9 Hz), 4.72 (1H, td, J = 9.1 Hz, 4.7 Hz), 6.71 (2H, d, J = 8.6 Hz), 7.12 (2H, d, J = 8.6 Hz), 7.26 (2H, d, J = 8.2 Hz), 7.35 (3H, d, J = 7.7 Hz), 7.38 - 7.40 (2H, m), 7.42 (2H, d, J = 7.8 Hz), 7.49 (1H, d, J = 7.4 Hz), 7.73 (2H, d, J = 7.2 Hz), 7.97 (2H, d, J = 8.0 Hz), 8.06 (2H, br.s), 8.45 (1H, d, J = 7.8 Hz), 8.66 - 8.72 (2H, m)
154	d ⁶ - DMSO	δ: 1.26(3H,t,J= 7.0Hz), 2.70(2H,d,J= 7.3Hz), 2.79-2.85(1H,m), 3.03-3.08(1H,m), 3.94(2H,q,J= 7.0Hz), 4.00(2H,s), 4.27-4.39(2H,m), 4.54-4.62(2H,m), 6.75(2H,d,J= 8.6Hz), 7.07-7.25(9H,m), 7.41-7.52(4H,m), 7.70-7.74(2H,m), 8.26(2H,s), 8.49(1H,d,J= 8.1Hz), 8.58(1H,d,J= 8.6Hz), 8.66(1H,t,J= 6.0Hz).
155	d ⁶ - DMSO	δ: 1.26(3H,t,J= 6.9Hz), 2.72(2H,d,J= 7.2Hz), 2.80-2.86(1H,m), 3.03-3.08(1H,m), 3.92(2H,q,J= 6.9Hz), 4.06(2H,s), 4.26-4.37(2H,m), 4.53-4.62(2H,m), 6.74(2H,d,J= 8.5Hz), 7.13(2H,d,J= 8.5Hz), 7.15-7.23(6H,m), 7.38-7.43(3H,m), 7.48-7.53(2H,m), 7.75(2H,d,J= 7.3Hz), 8.49(2H,s), 8.52(1H,d,J= 8.2Hz), 8.59(1H,d,J= 8.5Hz), 8.71(1H,t,J= 5.9Hz).
156	d ⁶ - DMSO	δ: 1.26(3H,t,J= 7.0Hz), 2.70(2H,d,J= 7.2Hz), 2.79-2.85(1H,m), 3.03-3.07(1H,m), 3.92(2H,q,J= 7.0Hz), 4.11(2H,s), 4.33-4.46(2H,m), 4.54-4.62(2H,m), 6.74(2H,d,J= 8.6Hz), 7.12(2H,d,J= 8.6Hz), 7.16-7.25(7H,m), 7.39-7.43(2H,m), 7.47-7.56(2H,m), 7.67-7.74(4H,m), 8.48(1H,d,J= 8.0Hz), 8.59(1H,d,J= 8.5Hz), 8.75(1H,t,J= 6.0Hz).

157	d ⁶ - DMSO	δ: 1.27(3H,t,J= 7.0Hz), 2.81-2.86(1H,m), 2.93-2.97(1H,m), 3.06-3.12(1H,m), 3.24-3.29(1H,m), 3.90-3.99(4H,m), 4.30(2H,d,J= 5.8Hz), 4.50-4.56(2H,m), 4.78-4.83(2H,m), 6.69(2H,d,J= 8.6Hz), 6.86(2H,d,J= 8.6Hz), 6.91-6.94(2H,m), 7.23(2H,d,J= 8.1Hz), 7.35-7.38(3H,m), 7.60-7.63(1H,m), 7.96-8.00(2H,m), 8.20(2H,s), 8.56(1H,d,J= 8.3Hz), 8.63(1H,d,J= 4.8Hz), 8.67(1H,d,J= 6.0Hz), 8.74(1H,d,J= 8.3Hz).
158	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 6.9 Hz), 2.74 (2H, d, J = 7.2 Hz), 3.02 (2H, dd, J = 13.9 Hz, 9.4 Hz), 3.18 (1H, d, J = 13.9 Hz, 4.6 Hz), 3.66 (1H, s), 3.80 (3H, s), 3.93 (2H, q, J = 6.9 Hz), 4.27 (2H, d, J = 6.0 Hz), 4.52 (1H, q, J = 7.4 Hz), 4.73 - 4.79 (1H, m), 6.74 (2H, d, J = 8.6 Hz), 6.96 (2H, d, J = 8.8 Hz), 7.09 (2H, d, J = 8.2 Hz), 7.13 (2H, d, J = 8.6 Hz), 7.17 - 7.21 (3H, m), 7.34 (1H, d, J = 7.8 Hz), 7.63 (1H, td, J = 7.6 Hz, 1.7 Hz), 7.7 (2H, d, J = 8.8 Hz), 8.36 (1H, d, J = 4.0 Hz), 8.41 (2H, d, J = 6.6 Hz), 8.60 (1H, d, J = 8.4 Hz)
159	d ⁶ - DMSO	δ: 1.27(3H,t,J= 7.0Hz), 2.68-2.74(1H,m), 2.82-2.88(2H,m), 3.05-3.10(1H,m), 3.90(2H,q,J= 7.0Hz), 4.06(2H,q,J= 5.7Hz), 4.31(2H,d,J= 5.9Hz), 4.50-4.55(1H,m), 4.76-4.81(1H,m), 6.59-6.65(4H,m), 7.19-7.23(2H,m), 7.28(2H,t,J= 7.4Hz), 7.33(2H,d,J= 7.2Hz), 7.37(1H,d,J= 1.0Hz), 7.50(1H,d,J= 8.0Hz), 7.59-7.63(1H,m), 7.98-8.02(2H,m), 8.38(2H,s), 8.51(1H,d,J= 8.2Hz), 8.61(1H,d,J= 4.7Hz), 8.71-8.74(2H,m).
160	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.66Hz), 2.70 (2H, d, J = 7.60Hz), 2.94-3.00 (1H, m), 3.21-3.24 (1H, m), 3.80 (3H, s), 3.92-4.00 (4H, m), 4.33 (2H, d, J = 6.69Hz), 4.52 (1H, q, J = 7.45Hz), 4.66-4.69 (2H, m), 6.75 (2H, d, J = 7.79Hz), 6.96 (2H, d, J = 8.68Hz), 7.17 (2H, d, J = 8.45Hz), 7.27 (2H, d, J = 7.82Hz), 7.38 (2H, d, J = 7.82Hz), 7.67 (1H, s, br), 7.74 (1H, d, J = 9.03Hz), 8.11 (1H, s, br), 8.22 (2H, s, br), 8.45 (1H, d, J = 8.34Hz), 8.68-8.69 (4H, m).
161	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 6.94Hz), 2.62-2.84 (3H, m), 2.96-3.07 (1H, m), 3.90-4.02 (4H, m), 4.32 (2H, d, J = 5.8Hz), 4.60-4.67 (2H, m), 6.77(2H, d, J = 8.88Hz), 7.16 – 7.22 (1H, m), 7.26 (2H, d, J = 8.32Hz), 7.29-7.35(5H, m), 7.36 (2H, d, J = 8.32Hz), 7.64 (1H, d, J = 6.53Hz), 8.08 (3H,s, br), 8.58-8.71(3H, m), 8.82 (1H, d, J = 7.71Hz).

162	d ⁶ - DMSO	δ: 1.27 (3H, t, J = 7.0 Hz), 2.61 (2H, dd, J = 14.0 Hz, 4.3 Hz), 2.86 (1H, dd, J = 13.7 Hz, 10.3 Hz), 3.10 (1H, dd, J = 13.7 Hz, 4.5 Hz), 3.93 (2H, q, J = 7.0 Hz), 3.98 - 4.00 (2H, m), 4.3 (2H, d, J = 5.8 Hz), 4.56 (1H, td, J = 9.0 Hz, 5.0 Hz), 4.66 (1H, td, J = 9.0 Hz, 4.8 Hz), 6.75 (2H, d, J = 8.6 Hz), 7.12 (1H, dd J = 4.9, 3.7 Hz), 7.16 (2H, d, J = 8.6 Hz), 7.26 (2H, d, J = 8.3 Hz), 7.36 (3H, d, J = 8.1 Hz), 7.72 (1H, dd, J = 5.0 Hz, 0.7 Hz), 7.76 (1H, br.s), 7.81 (1H, d, J = 3.2 Hz), 8.07 (2H, br.s), 8.47 - 8.55 (3H, m), 8.64 (2H, t, J = 9.0 Hz)
163	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.0 Hz), 2.67 - 2.75 (2H, m), 3.03 (2H, dd, J = 13.8 Hz, 9.5 Hz), 3.21 (2H, dd, J = 14.0 Hz, 4.7 Hz), 3.93 (2H, q, J = 7.0 Hz), 3.99 (2H, q, J = 5.8 Hz), 4.20 - 4.40 (3H, m), 4.53 - 4.58 (1H, m), 4.77 - 4.83 (1H, m), 6.75 (2H, d, J = 8.6 Hz), 7.14 (2H, d, J = 8.6 Hz), 7.23 (2H, d, J = 8.1 Hz), 7.34 (2H, d, J = 8.2 Hz), 7.52 (2H, d, J = 8.6 Hz), 7.76 (2H, d, J = 8.6 Hz), 8.07 (2H, br.s), 8.44 (1H, d, J = 4.5 Hz), 8.53 (1H, br.s), 8.66 (2H, d, J = 7.6 Hz)
164	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.0 Hz), 2.34 (3H, s), 2.74 (2H, d, J = 8.2 Hz), 3.05 (1H, dd, J = 13.9, 9.5 Hz), 3.23 (1H, dd, J = 14.0, 4.7 Hz), 3.93 (2H, q, J = 7.0 Hz), 3.99 (2H, q, J = 5.8 Hz), 4.26 - 4.36 (2H, m), 4.54 (1H, dd, J = 14.5, 7.7 Hz), 4.79 (1H, td, J = 8.9 Hz, 4.8 Hz), 6.75 (2H, d, J = 8.6 Hz), 7.13 (2H, d, J = 8.6 Hz), 7.24 (4H, d, J = 8.0 Hz), 7.30 (1H, br.s), 7.34 (3H, d, J = 8.0 Hz), 7.66 (2H, d, J = 8.1 Hz), 7.76 (1H, br.s), 8.09 (2H, br.s), 8.45 (2H, t, J = 6.0 Hz), 8.54 (1H, t, J = 6.0 Hz), 8.64 (1H, d, J = 8.3 Hz)
165	d ⁶ - DMSO	δ: 1.28(3H,t,J= 6.9Hz), 2.71-2.92(3H,m), 3.06-3.10(1H,m), 3.92(2H,q,J= 6.9Hz), 3.96-4.01(2H,m), 4.31(2H,d,J= 5.8Hz), 4.53-4.59(1H,m), 4.75-4.80(1H,m), 6.57-6.72(4H,m), 7.27(2H,d,J= 8.0Hz), 7.30-7.33(1H,m), 7.37(2H,d,J= 8.0Hz), 7.55(1H,d,J= 8.2Hz), 7.59-7.63(1H,m), 7.64(1H,s), 7.98-8.01(2H,m), 8.13(2H,s), 8.47(1H,d,J= 8.2Hz), 8.62(1H,d,J= 4.7Hz), 8.65-8.67(1H,m), 8.77(1H,d,J= 8.2Hz).

166	d ⁶ - DMSO	δ: 0.85 (3H, t, J = 7.6 Hz), 1.30 (3H, t, J = 7.0 Hz), 2.02 (2H, q, J = 7.6 Hz), 2.63 - 2.68 (1H, m), 2.99 (2H, dd, J = 14.0 Hz, 9.6 Hz), 3.20 (1H, dd, J = 13.9, 4.8 Hz), 3.95 (2H, q, J = 7.0 Hz), 4.00 (2H, q, J = 6.0 Hz), 4.28 (2H, t, J = 7.2 Hz), 4.32 - 4.35 (1H, m), 4.74 (1H, td, J = 8.9 Hz, 4.8 Hz), 6.73 (2H, d, J = 8.6 Hz), 7.00 (2H, d, J = 8.6 Hz), 7.23 (2H, d, J = 8.1 Hz), 7.29 (2H, d, J = 7.24 Hz), 7.36 (2H, d, J = 8.12 Hz), 7.74 (1H, br.s), 7.97 (1H, d, J = 5.2 Hz), 8.08 (2H, br.s), 8.51 (3H, d, J = 4.2 Hz)
167	d ⁶ - DMSO	δ: 1.27(3H,t,J= 6.9Hz), 2.65-2.89(3H,m), 3.05-3.10(1H,m), 3.93(2H,q,J= 6.9Hz), 4.00(2H,q,J= 5.4Hz), 4.27-4.40(2H,m), 4.58-4.70(2H,m), 6.76(2H,d,J= 8.5Hz), 7.08(1H,s), 7.11(1H,d,J= 2.8Hz), 7.16-7.29(7H,m), 7.51(1H,t,J= 8.0Hz), 7.95(2H,s), 8.47(2H,s), 8.72-8.78(2H,m), 8.87(2H,s), 9.19(1H,d,J= 7.7Hz).
168	d ⁶ - DMSO	δ: 1.27(3H,t,J= 7.0Hz), 2.69-2.74(1H,m), 2.82-2.93(2H,m), 3.05-3.10(1H,m), 3.90(2H,q,J= 7.0Hz), 4.01(2H,q,J= 5.7Hz), 4.32(2H,d,J= 6.0Hz), 4.51-4.56(1H,m), 4.76-4.79(1H,m), 6.60-6.67(3H,m), 7.06-7.09(2H,m), 7.18-7.22(2H,m), 7.28(2H,t,J= 7.4Hz), 7.33(2H,d,J= 3.8Hz), 7.46(1H,t,J= 7.8Hz), 7.59-7.63(1H,m), 7.96-7.99(2H,m), 8.31(2H,s), 8.50(1H,d,J= 8.2Hz), 8.60-8.62(1H,m), 8.71-8.74(2H,m).
169	d ⁶ - DMSO	δ: 1.26-1.30(3H,m), 2.60-2.84(3H,m), 3.03-3.07(1H,m), 3.93(2H,q,J= 7.0Hz), 4.00(2H,q,J= 5.4Hz), 4.32(2H,d,J= 5.8Hz), 4.56-4.66(2H,m), 6.76(2H,d,J= 8.6Hz), 7.11-7.13(1H,m), 7.16(2H,d,J= 8.5Hz), 7.23(1H,dd,J= 8.3,1.8Hz), 7.28(2H,d,J= 8.0Hz), 7.38(2H,d,J= 8.0Hz), 7.48(1H,d,J= 8.2Hz) 7.55(1H,d,J= 1.7Hz), 7.73(1H,dd,J= 5.0,0.7Hz), 7.81(1H,d,J= 3.2Hz), 8.17(2H,s), 8.53(1H,d,J= 8.3Hz), 8.63-8.66(2H,m).

170	d ⁶ - DMSO	δ: 0.84 (3H, t, J = 7.6 Hz), 1.30 (3H, t, J = 7.0 Hz), 2.00 (2H, q, J = 7.6 Hz), 2.43 - 2.47 (1H, m), 2.61 (1H, dd, J = 13.7 Hz, 4.8 Hz), 2.94 (1H, dd, J = 13.6 Hz, 10.2 Hz), 3.20 (1H, dd, J = 13.8 Hz, 4.4 Hz), 3.97 (2H, q, J = 7.0 Hz), 3.99 (2H, q, J = 5.5 Hz), 4.30 (2H, d, J = 5.6 Hz), 4.35 (1H, td, J = 8.4, 5.2 Hz), 4.62 (1H, td, J = 9.3, 4.6 Hz), 6.73 (2H, d, J = 8.6 Hz), 7.02 (2H, d, J = 8.6 Hz), 7.26 (2H, d, J = 8.0 Hz), 7.40 (2H, d, J = 8.1 Hz), 7.71 (1H, br.s), 8.00 (1H, d, J = 7.6 Hz), 8.12 (1H, d, J = 6.4 Hz), 8.27 (2H, br.s), 8.58 (1H, d, J = 8.4 Hz), 8.65 (1H, br.s), 8.66 (2H, br.s)
171	d ⁶ - DMSO	δ: 1.28(3H,t,J= 7.0Hz), 2.62-2.68(1H,m), 2.75-2.85(2H,m), 3.03-3.08(1H,m), 3.94(2H,q,J= 7.0Hz), 3.97-4.02(2H,m), 4.28-4.38(2H,m), 4.63-4.67(2H,m), 6.77(2H,d,J= 8.6Hz), 7.17(2H,d,J= 8.5Hz), 7.22-7.25(1H,m), 7.28(2H,d,J= 8.1Hz), 7.38(2H,d,J= 8.0Hz), 7.50(1H,d,J= 8.2Hz), 7.55(1H,d,J= 1.4Hz), 7.65(2H,d,J= 5.8Hz), 7.71-7.78(1H,m), 8.12(2H,s), 8.62-8.73(3H,m), 8.81(1H,d,J= 8.4Hz).
173	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 6.82Hz), 2.61-2.71 (2H, m), 2.85-2.91 (1H, m), 3.02-3.14 (1H, m), 3.93-4.01 (4H, m), 4.32-4.34 (2H, m), 4.56-4.69 (2H, m), 6.76 (2H, d, J = 8.67Hz), 7.16 (2H, d, J = 8.67Hz), 7.26 (2H, d, J = 8.67Hz), 7.36 (2H, d, J = 7.51Hz), 7.40-7.47 (1H, m), 7.51 (2H, d, J = 8.48Hz), 7.56 (1H, d, J = 7.51Hz), 7.76 (2H, d, J = 8.09Hz), 8.12 (2H, s, br), 8.48-8.58 (2H, m), 8.62-8.68(3H, m).
174	d ⁶ - DMSO	δ: 1.28(3H, t, J= 7.05Hz), 2.65-2.71 (1H, m), 2.79-2.82 (1H, m), 3.04-3.07 (1H, m), 3.28-3.31 (1H,m), 3.91-3.99 (4H, m), 4.31 (2H, s), 4.58-4.60 (1H,m), 4.74-4.75 (1H,m), 6.75 (2H, d, J= 8.13Hz), 7.14 (2H, d, J= 9.73Hz), 7.27 (2H, d, J= 8.41Hz), 7.41 (2H, d, J = 8.41Hz), 7.85-7.88 (1H, m),8.34-8.42 (4H, m), 8.72-8.83 (5H, m), 9.08 (1H, d, J= 8.41Hz).
175	d ⁶ - DMSO	δ: 1.29 (3H, t, J = 6.75 Hz), 2.61-2.64 (1H, m), 2.78-2.83 (1H, m), 2.87-2.93 (1H, m), 3.14-3.19 (1H, m), 3.92-4.00 (4H, m), 4.32 (2H, d, J = 5.78Hz), 4.60-4.65 (2H, m), 6.71 (2H, d, J = 8.18 Hz), 6.87 (2H, d J = 8.56 Hz), 7.14 (1H, d, J = 5.45 Hz), 7.26 (2H, d, J = 8.56 Hz), 7.37 (2H, d, J = 7.78 Hz), 7.49-7.52 (1H, m), 7.83-7.90 (2H, m), 8.10 (3H, s, br), 8.54 (1H, d, J = 4.69Hz), 8.60 - 8.66 (2H, m), 8.77 (1H, d, J = 8.63 Hz).

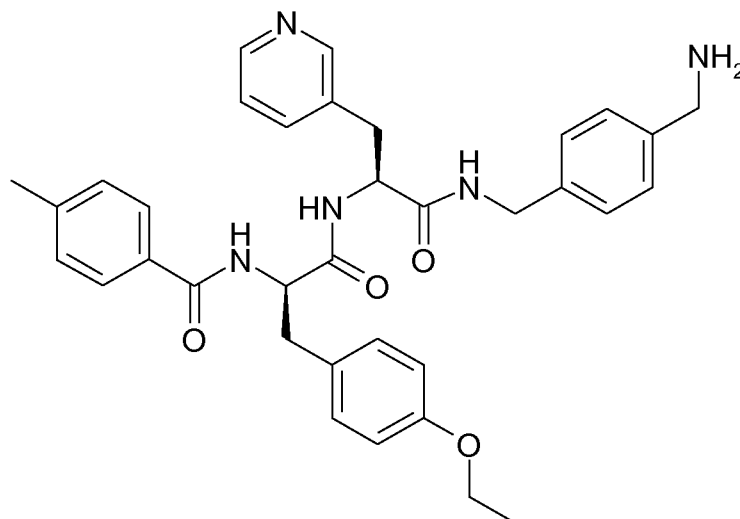
176	d ⁶ - DMSO	δ: 1.27 (3H, t, J = 7.0 Hz), 2.66 - 2.77 (2H, m), 2.98 (1H, dd, J = 14.5 Hz, 9.6 Hz), 3.19 (1H, dd, J = 14.5 Hz, 4.6 Hz), 3.91 (2H, q, J = 7.0 Hz), 3.98 (2H, q, J = 5.7 Hz), 4.32 (2H, d, J = 5.9 Hz), 4.62 (2H, qd, J = 9.3 Hz, 4.6 Hz), 6.69 (2H, d, J = 8.6 Hz), 6.99 (1H, t, J = 6.9 Hz), 7.08 (3H, d, J = 8.7 Hz), 7.16 (1H, d, J = 2.1 Hz), 7.26 (2H, d, J = 8.0 Hz), 7.33 (3H, dd, J = 8.0 Hz, 6.1 Hz), 7.42 (2H, t, J = 7.5 Hz), 7.51 (1H, t, J = 7.4 Hz), 7.66 (1H, d, J = 7.8 Hz), 7.73 (2H, d, J = 7.2 Hz), 8.06 (2H, br.s), 8.39 (1H, d, J = 7.8 Hz), 8.52 (1H, d, J = 6.5 Hz), 8.61 (1H, t, J = 4.9 Hz), 10.82 (1H, br.s)
177	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.0 Hz), 2.32 - 2.34 (1H, m), 2.66 (2H, t, J = 1.7 Hz), 2.85 (1H, d, J = 7.44 Hz), 2.91 - 3.03 (1H, m), 3.17 (1H, dd, J = 15.2 Hz, 5.0 Hz), 3.94 (2H, q, J = 7.0 Hz), 3.98 - 4.00 (1H, br.m), 4.20 (1H, d, J = 6.2 Hz), 4.55 (1H, q, J = 7.4 Hz), 4.62 - 4.68 (1H, m), 6.78 (2H, d, J = 8.5 Hz), 7.19 (2H, d, J = 8.5 Hz), 7.28 (2H, d, J = 8.4 Hz), 7.34 - 7.38 (2H, m), 7.44 (2H, q, J = 8.0 Hz), 7.51 - 7.56 (1H, m), 7.72 - 7.78 (2H, m), 8.08 (2H, br.s), 8.52 (1H, t, J = 6.6 Hz), 8.59 - 8.66 (2H, m), 8.97 (1H, br.s), 14.02 (1H, br.s), 14.18 (1H, br.s)
178	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.0 Hz), 2.66 (1H, d, J = 11.0 Hz), 2.73 (1H, dd, J = 13.5, 3.6 Hz), 3.10 (1H, dd, J = 14.4, 9.8 Hz), 3.29 (1H, d, J = 4.9 Hz), 3.66 (2H, s), 3.92 (2H, q, J = 7.0 Hz), 4.30 (2H, qd, J = 15.2, 5.7 Hz), 4.66 (1H, td, J = 9.1, 4.4 Hz), 4.75 (1H, td, J = 9.0, 5.0 Hz), 6.73 (2H, d, J = 8.6 Hz), 7.14 (5H, dd, J = 8.3, 5.1 Hz), 7.22 (2H, d, J = 8.1 Hz), 7.35 - 7.44 (4H, m), 7.61 (2H, dd, J = 4.5, 1.5 Hz), 7.97 (2H, t, J = 9.4 Hz), 8.62 (1H, br, s), 8.67 (2H, dd, J = 4.5, 1.5 Hz), 8.71 (1H, br,s), 8.80 (1H, br,s)
179	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.0 Hz), 2.02 (3H, s), 2.68 - 2.74 (2H, m), 2.83 (1H, dd, J = 13.6, 10.0 Hz), 3.05 (1H, dd, J = 13.5, 4.6 Hz), 3.93 (2H, q, J = 7.0 Hz), 3.98 (2H, d, J = 5.8 Hz), 4.32 (2H, qd, J = 15.3, 6.0 Hz), 4.55 - 4.64 (2H, m), 6.73 (2H, d, J = 8.5 Hz), 7.06 (2H, d, J = 8.5 Hz), 7.19 - 7.27 (7H, m), 7.38 (2H, d, J = 8.1 Hz), 7.69 (1H, d, J = 5.4 Hz), 7.85 (1H, d, J = 5.4 Hz), 8.11 (1H, d, J = 8.1 Hz), 8.22 (2H, br.s), 8.58 (1H, d, J = 8.5 Hz), 8.65 (1H, t, J = 5.7 Hz), 10.70 (1H, s)

180	d ⁶ - DMSO	δ: 1.27 (3H, t, J= 7.09Hz), 2.72-2.74 (2H, m), 2.84-2.90 (1H, m), 3.09-3.16 (1H,m), 3.59 (2H, s, br), 3.91-3.98 (4H, m), 4.29-4.33 (2H, m), 4.57-4.67 (2H,m), 6.76 (2H, d, J= 8.79Hz), 7.01-7.16 (7H, m), 7.22-7.27 (2H,m), 7.35-7.41 (3H,m), 7.43-7.51 (1H, m), 7.76 (2H, d, J = 7.19Hz), 8.37 (3H, s, br), 8.57 (1H, d, J = 7.97Hz), 8.64-8.69 (1H, m).
181	d ⁶ - DMSO	δ: 1.30 (3H, t, J = 7.21 Hz), 2.22 (3H, s), 2.56-2.78 (2H, m), 2.96-3.08 (1H, m), 3.23-3.28 (1H, m), 3.95-4.00 (4H, m), 4.32 (2H, d, J = 5.60Hz), 4.49-4.54 (1H, m), 4.66-4.72 (1H, m), 6.76 (2H, d, J = 8.32 Hz), 6.91 (1H, d J = 5.14 Hz), 7.05 (2H, d, J = 8.69Hz), 7.27 (2H, d, J = 7.90 Hz), 7.39 (2H, d, J = 8.30 Hz), 7.56 (1H, d, J = 5.14Hz), 7.72 (1H, s, br), 7.88 (1H, d, J = 7.61?Hz), 8.16 (1H, d, J = 7.61Hz), 8.27 (3H, s, br), 8.67 - 8.74 (3H, m).
182	d ⁶ - DMSO	δ: 1.28(3H,t,J= 7.0Hz), 2.29(3H,s), 2.69(2H,d,J= 7.2Hz), 2.80-2.86(1H,m), 3.04-3.09(1H,m), 3.91-3.97(4H,m), 4.24-4.34(2H,m), 4.56-4.63(2H,m), 6.76(2H,d,J= 8.7Hz), 7.08-7.31(10H,m), 7.41-7.45(2H,m), 7.50-7.53(1H,m), 7.74-7.76(2H,m), 8.15(2H,s), 8.48(1H,d,J= 8.0Hz), 8.57-8.61(2H,m)
183	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.0 Hz), 2.76 (2H, d, J = 5.8 Hz), 3.06 (2H, dd, J = 14.5, 9.5 Hz), 3.22 (1H, dd, J = 14.5, 4.5 Hz), 3.94 (2H, q, J = 7.0 Hz), 3.98 (1H, d, J = 5.7 Hz), 4.29 (2H, d, J = 6.0 Hz), 4.44 - 4.50 (1H, m), 4.65 (1H, td, J = 8.8, 4.9 Hz), 6.33 (2H, br.s), 6.55 (1H, d, J = 5.3 Hz), 6.74 (2H, d, J = 8.6 Hz), 7.04 (2H, d, J = 8.5 Hz), 7.22 (2H, d, J = 8.0 Hz), 7.27 (1H, d, J = 1.4 Hz), 7.29 (1H, s), 7.34 (2H, d, J = 8.0 Hz), 7.39 (1H, d, J = 5.4 Hz), 8.07 (2H, br.s), 8.52 (1H, s), 8.53 (1H, s), 8.97 (1H, d, J = 1.8 Hz)
184	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 7.0 Hz), 1.35 (1H, s), 1.89 (3H, s), 2.72 (1H, dd, J = 14.7, 7.9 Hz), 2.86 (1H, dd, J = 13.7, 4.7 Hz), 3.05 (1H, dd, J = 14.4, 9.8 Hz), 3.24 (1H, dd, J = 14.7, 4.6 Hz), 3.67 (2H, s), 3.94 (2H, q, J = 7.0 Hz), 4.25 (2H, d, J = 5.2 Hz), 4.65-4.70 (2H, m), 6.72 (2H, d, J = 8.6 Hz), 6.88 (2H, d, J = 8.7 Hz), 7.11 (2H, d, J = 7.9 Hz), 7.14 (1H, d, J = 5.2 Hz), 7.22 (2H, d, J = 7.8 Hz), 7.34 (1H, d, J = 1.7 Hz), 7.85 (1H, d, J = 5.2 Hz), 7.89 (2H, d, J = 7.5 Hz), 8.51 (1H, t, J = 6.0 Hz), 8.68 (1H, d, J = 8.6 Hz), 9.04 (1H, d, J = 1.9 Hz)

185	d ⁶ - DMSO	δ: 1.28(3H,t,J= 6.9Hz), 2.34(3H,s), 2.77(2H,d,J= 7.3Hz), 3.05(1H,dd,J=14.6,9.6Hz), 3.22(1H,dd,J=14.6,4.7Hz), 3.93(2H,q,J=7.0Hz), 3.97(2H,m), 4.3(2H,dd,J=5.0,4.9Hz), 4.56(1H,dd,J=14.8,7.6), 4.66-4.71(1H,m), 6.76(2H,d,J= 8.7Hz), 7.15(2H,d,J=8.6Hz), 7.22-7.26(4H,m), 7.34(2H,d,J=8.2Hz), 7.66(2H,d,J=8.2Hz), 8.08(3H,br s), 8.47(1H,d,J=7.8Hz), 8.54(1H,d,J=6.0), 8.58(1H,d,J= 8.3Hz), 8.95(1H,d,J=2.0Hz).
186	d ⁶ - DMSO	δ: 1.26 (3H, t, J = 7.0 Hz), 2.12 (3H, s), 2.26-2.67 (1H, m), 2.74 (1H, dd, J = 13.7, 4.7 Hz), 3.09 (1H, dd, J = 14.5, 10.1 Hz), 3.35 (1H, d, J = 5.6 Hz), 3.65 (2H, s), 3.89 (2H, q, J =7.0 Hz), 4.27 (2H, d, J = 6.0 Hz), 4.64 (2H, dd, J = 7.5, 4.7 Hz), 5.90 (1H, br.s), 6.59 (2H, d, J = 8.6 Hz), 6.76 (1H, t, J = 2.5 Hz), 6.86 (2H, d, J = 8.5 Hz), 7.13 (2H, d, J = 7.3 Hz), 7.20 (2H, d, J = 8.1 Hz), 7.24 (1H, br.s), 7.36-7.45 (3H, m). 7.97 (2H, d, J = 8.2 Hz), 8.64 (1H, br.s), 8.68 (1H, d, J = 7.2 Hz), 11.12 (1H, br.s)
187	d ⁶ - DMSO	δ: 1.29 (3H, t, J = 7.0 Hz), 2.14 (3H, s), 2.66 (1H, dd, J = 13.1, 9.0 Hz), 2.78 (1H, dd, J = 13.7, 4.5 Hz), 2.86 (1H, dd, J = 14.2, 10.0 Hz), 3.07 (1H, dd, J = 14.2, 4.4 Hz), 3.66 (2H, s), 3.95 (2H, q, J = 7.0 Hz), 4.26 (2H, d, J = 4.6 Hz), 4.51 (1H, td, J = 9.0, 4.8 Hz), 4.61-4.67 (1H, m), 5.91 (1H, s), 6.75 (2H, d, J = 8.6 Hz), 6.77 (1H, d, J = 2.7 Hz), 7.02 (3H, d, J = 8.5 Hz), 7.12 (2H, d, J = 7.6 Hz), 7.15 (2H, br.s), 7.21 (2H, d, J = 8.3 Hz), 7.29 (1H, d, J = 7.4 Hz), 7.42 (1H, dd, J = 4.8, 2.9 Hz), 8.32 (1H, s), 8.49 (1H, br.s), 8.57 (1H, d, J = 7.0 Hz), 11.11 (1H, s)
188	d ⁶ - DMSO	δ: 1.26 (3H, t, J = 7.0 Hz), 2.01 (3H, s), 2.08 (1H, s), 2.66-2.72 (1H, m), 2.76-2.80 (1H, m), 3.12 (1H, dd, J = 14.4, 9.6), 3.67 (1H, s), 3.90 (2H, q, J = 7.0 Hz), 4.10 (1H, dd, J = 17.3, 6.1 Hz), 4.28-4.31 (2H, m), 4.35 (1H, dd, J = 13.5, 6.8 Hz), 4.59 (1H, br.s), 4.71-4.76 (1H, m), 6.66 (2H, d, J = 8.6 Hz), 7.02 (2H, d, J = 8.5 Hz), 7.14 (3H, d, J = 7.6 Hz), 7.22 (1H, d, J = 8.2 Hz), 7.31-7.42 (4H, m), 7.67 (1H, d, J = 5.2 Hz), 7.84 (1H, d, J = 5.4 Hz), 7.97 (2H, dd, J = 12.2, 8.0 Hz), 8.05 (br.s, 1H), 8.60 (2H, br.d, J = 7.1 Hz), 10.66 (1H, br.s)

189	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 6.70Hz), 2.33 (3H, s), 2.69 (2H, d, J = 7.51Hz), 2.91-2.99 (1H, m), 3.13-3.18 (1H, m), 3.92-4.00 (4H, m), 4.33 (2H, d, J = 5.94Hz), 4.52-4.61 (1H, m), 4.63-4.69 (1H, m), 6.76 (2H, d, J = 8.67Hz), 7.15 (2H, d, J = 7.86Hz), 7.27 (2H, d, J = 8.096Hz), 7.31 (2H, d, J = 6.36Hz), 7.36 (2H, d, J = 7.51Hz), 7.47-7.53 (1H, m), 7.56 (1H, d, J = 7.51Hz), 7.87 (1H, d, J = 7.341Hz), 8.11 (3H, s, br), 8.45 (1H, d, J = 7.86Hz), 8.53 (2H, d, J = 4.72Hz), 8.57-8.65(2H, m).
190	d ⁶ - DMSO	δ: 1.31 (3H, t, J = 6.88Hz), 2.08 (3H, s), 2.70-2.71 (1H, m), 2.74-2.75 (1H, m), 2.88-2.94 (1H, m), 3.13-3.18 (1H, m), 3.90-4.01 (4H, m), 4.33 (2H, d, J = 6.21?Hz), 4.57-4.62 (1H, m), 4.66-4.71 (1H, m), 6.80 (2H, d, J = 8.06Hz), 7.09-7.11 (5H, m), 7.14-7.31 (2H, m), 7.36 (2H, d, J = 8.056Hz), 7.45-7.48 (1H, m), 7.84 (1H, d, J = 7.051Hz), 8.11 (3H, s, br), 8.34 (1H, d, J = 8.06Hz), 8.52-8.53(1H, m), 8.56 (1H, d, J = 5.021Hz), 8.63-8.65(2H, m).
191	d ⁶ - DMSO	δ: 1.29 (3H, t, J = 6.84Hz), 2.07 (3H, s), 2.14 (3H, s), 2.53-2.59 (1H, m), 2.69-2.74 (1H, m), 2.77-2.83 (1H, m), 3.05-3.09 (1H, m), 3.91-4.01 (4H, m), 4.30 (2H, d, J = 5.98Hz), 4.51-4.65 (2H, m), 5.63 (1H, d, J = 1.50Hz), 6.71 (2H, d, J = 8.88Hz), 6.90 (2H, t, J = 8.88Hz), 7.04 (1H, d, J = 8.17Hz), 7.18-7.28 (8H, m), 7.34 (2H, d, J = 7.35Hz), 8.10 (2H, s, br), 8.57-8.64(2H, m).
192	d ⁶ - DMSO	δ: 1.28 (3H, t, J = 6.82Hz), 2.30 (3H, s), 2.71-2.74 (2H, m), 2.80-2.86(1H, m), 3.04-3.09(1H, m), 3.91-3.95 (4H, m), 4.25-4.30 (2H, m), 4.54-4.59(1H, m), 4.69-4.74 (1H, m), 6.76 (2H, d, J = 7.887Hz), 7.08-7.31(10H, m), 7.41-7.45(2H, m), 7.50-7.53(1H, m), 7.74-7.76(2H, m), 8.31-8.39 (3H, m), 8.63(1H, d, J = 7.88Hz), 8.75-8.78(3H, m).
193	d ⁶ - DMSO	δ: 1.27 (3H, t, J = 7.0 Hz), 2.01 (3H, s), 2.32-2.34 (1H, m), 2.67 (1H, t, J = 1.7 Hz), 2.76-2.78 (2H, m), 2.88 (2H, dd, J = 14.1, 9.6 Hz), 3.04 (2H, dd, J = 14.2, 4.6 Hz), 3.67 (1H, s), 3.93 (2H, q, J = 7.0 Hz), 4.28 (2H, t, J = 6.4 Hz), 4.56-4.62 (2H, m), 6.75 (2H, d, J = 8.6 Hz), 6.99 (1H, d, J = 5.8 Hz), 7.12 (3H, t, J = 8.3Hz), 7.15 (1H, d, J = 3.5 Hz), 7.22 (2H, d, J = 8.0 Hz), 7.39 (1H, dd, J = 4.8, 3.0 Hz), 7.69 (1H, d, J = 5.4 Hz), 7.85 (1H, d, J = 5.4 Hz), 8.15 (1H, d, J = 8.3 Hz), 8.50 (2H, t, J = 8.0 Hz), 10.69 (1H, br.s)

194	d ⁶ - DMSO	(DMSO) δ : 1.27 (3H, t, J = 7.0 Hz), 7.00 (2H, d, J = 7.0 Hz), 3.10 (2H, dd, J = 14.5, 10.0 Hz), 3.90 (2H, q, J = 7.0 Hz), 4.00 (2H, q, J = 5.5 Hz), 4.33 (2H, d, J = 6.1 Hz), 4.52 (1H, q, J = 6.6 Hz), 4.67 (1H, td, J = 9.6 Hz, 4.1 Hz), 6.35 (1H, br.s), 6.56 (1H, d, J = 5.4 Hz), 6.61 (2H, d, J = 8.6 Hz), 6.91 (2H, d, J = 8.6 Hz), 7.14 (1H, d, J = 7.7 Hz), 7.26 (2H, d, J = 7.5 Hz), 7.36 (2H, d, J = 8.1 Hz), 7.38 (2H, d, J = 5.3 Hz), 7.46-7.41 (3H, m), 7.98 (2H, d, J = 8.8 Hz), 8.06 (2H, br.s), 8.60 (1H, d, J = 8.12 Hz), 8.70 (1H, br.s)
195	d ⁶ - DMSO	δ : 1.28 (3H, t, J = 7.0 Hz), 2.76 (2H, d, J = 5.8 Hz), 3.06 (2H, dd, J = 14.5, 9.5 Hz), 3.22 (1H, dd, J = 14.5, 4.5 Hz), 3.94 (2H, q, J = 7.0 Hz), 3.98 (1H, d, J = 5.7 Hz), 4.29 (2H, d, J = 6.0 Hz), 4.44 - 4.50 (1H, m), 4.65 (1H, td, J = 8.8, 4.9 Hz), 6.33 (2H, br.s), 6.55 (1H, d, J = 5.3 Hz), 6.74 (2H, d, J = 8.6 Hz), 7.04 (2H, d, J = 8.5 Hz), 7.22 (2H, d, J = 8.0 Hz), 7.27 (1H, d, J = 1.4 Hz), 7.29 (1H, s), 7.34 (2H, d, J = 8.0 Hz), 7.39 (1H, d, J = 5.4 Hz), 8.07 (2H, br.s), 8.52 (1H, s), 8.53 (1H, s), 8.97 (1H, d, J = 1.8 Hz)
196	d ⁶ - DMSO	δ : 1.29(3H, t, J= 6.91Hz), 2.42-2.46 (1H, m), 2.76-3.00 (3H, m), 3.05 (3H, s), 3.27-3.31 (1H,m), 3.93-4.00 (4H, m), 4.29 (2H, d, J = 6.28Hz), 4.89-4.99 (1H, m), 5.34-5.38 (1H, m), 6.67 (2H, d, J= 8.78Hz), 6.74 (2H, d, J= 8.21Hz), 6.98-7.01 (1H, m), 7.14 (1H, d, J= 5.67Hz), 7.19-7.38 (7H, m), 7.86 (1H, d, J= 5.67Hz), 7.88 (1H, d, J= 7.09Hz), 8.09 (3H,s, br), 8.41-8.48 (1H, m).
197	d ⁶ - DMSO	δ : 1.28 (3H, t, J= 7.15Hz), 2.72-2.74 (2H, m), 3.92-3.99 (4H, m), 4.36 (2H, d, J = 6.40Hz), 4.47-4.50 (1H, m), 4.67-4.73 (1H, m), 5.22 (1H, s, br), 5.83 (1H, s, br), 6.76 (2H, d, J= 8.51Hz), 7.17-7.30 (6H, m), 7.37-7.45 (5H, m), 7.52-7.53 (1H, m), 8.30 (4H,s, br), 8.38-8.44 (2H, m), 8.60 (1H, d, J = 7.64Hz).
198	d ⁶ - DMSO	δ : 1.28(3H, t, J= 6.70Hz), 2.61-2.64 (1H, m), 2.79-2.83 (1H, m), , 3.91-4.00 (4H, m), 4.34-4.36 (2H, m), 4.45-4.48 (1H, m), 4.82-4.85 (1H, m), 5.26-5.28 (1H, m), 5.85-5.86 (1H, m), 6.54 (1H,s, br), 6.65 (2H, d, J= 8.21Hz), 6.71 (2H, d, J= 8.213Hz), 7.13 (1H, d, J= 5.47Hz), 7.23-7.36 (4H, m), 7.49 (2H, d, J = 7.11Hz), 7.82-7.86 (3H, m), 8.07 (3H,s, br), 8.36(1H, t, J = 5.81Hz), 8.53 (1H, d, J= 8.92Hz).

EXAMPLE 199**N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide****A. [4-(tert-Butoxycarbonylamino-methyl)-benzyl]-carbamic acid benzyl ester**

tert-Butyl 4-(Aminomethyl)benzylcarbamate (7.5g, 31.74mmol) was dissolved in dichloromethane (250mL). This solution was cooled to 0°C and triethylamine (9.63g, 93.2mmol) was added followed by carbonic acid benzyl ester 2,5-dioxo-pyrrolidin-1-yl ester (9.5g, 38.09mmol). The reaction mixture was stirred at 0°C to room temperature for 18 hours and diluted with CHCl₃ (200mL) the filtrate was washed with 0.3M KHSO₄ (1x50mL), sat. NaHCO₃ (1x50mL), water (1x50mL), brine (1x50mL), dried (Na₂SO₄) and evaporated in vacuo to give a white solid. The solid were triturated with EtOAc/Pet Ether 60-80°C to give a white solid identified as [4-(tert-butoxycarbonylamino-methyl)-benzyl]-carbamic acid benzyl ester (11.3g, 30.5mmol, 96%).

$$[M+H]^+ = 392.98 (M+ Na)$$

20 B. (4-Aminomethyl-benzyl)-carbamic acid benzyl ester hydrochloride

[4-(tert-Butoxycarbonylamino-methyl)-benzyl]-carbamic acid benzyl ester (10.8g, 29.15mmol) was dissolved in 4M HCl in dioxan (400mL). After one hour at room temperature the solvent was removed in vacuo. The residue was slurried with acetone and the solid was filtered off to

give a white solid identified as (4-aminomethyl-benzyl)-carbamic acid benzyl ester hydrochloride (11.9g, 30.135mmol, 99%).

$$[M+H]^+ = 359.15$$

5 **C. {(S)-1-[4-(Benzyloxycarbonylamino-methyl)-benzylcarbamoyl]-2-pyridin-3-yl-ethyl}-carbamic acid tert-butylester**

(S)-2-tert-Butoxycarbonylamino-3-pyridin-3-yl-propionic acid (2.12g, 7.96mmol) was dissolved in CH₂Cl₂(100mL), HBTU (3.29g, 8.68mmol) and triethylamine (2.20g, 21.71mmol) were added. After 20 mins at room temperature the reaction mixture was cooled to 0°C and (4-aminomethyl-benzyl)-carbamic acid benzyl ester hydrochloride (1.96g, 7.24mmol) was added. After 2 hours at 0°C the reaction mixture was diluted with CHCl₃ (200mL), this solution was washed with 0.3M KHSO₄ (1x50mL), sat NaHCO₃ (1x50mL), water (1x50mL), brine (1x50mL), dried (Na₂SO₄) and evaporated in vacuo to give a white solid. The solid was triturated with EtOAc/Pet Ether 60-80°C to give a white solid identified as {(S)-1-[4-(benzyloxycarbonylamino-methyl)-benzylcarbamoyl]-2-pyridin-3-yl-ethyl}-carbamic acid tert-butylester (2.53g, 4.88mmol, 67%).

$$[M+H]^+ = 519.16$$

20 **D. {4-[(S)-2-Amino-3-pyridin-3-yl-propionylamino)-methyl]-benzyl}-carbamic acid benzyl ester Dihydrochloride**

{(S)-1-[4-(Benzyloxycarbonylamino-methyl)-benzylcarbamoyl]-2-pyridin-3-yl-ethyl}-carbamic acid tert-butylester (2.52g, 4.89mmol) was treated with 4M HCl/dioxan (50 mL). After one hour at room temperature the solvent was removed to give a white solid identified as {4-[(S)-2-amino-3-pyridin-3-yl-propionylamino)-methyl]-benzyl}-carbamic acid benzyl ester dihydrochloride (2.31g, 4.71mmol, 97%).

$$[M+H]^+ = 419.18$$

¹H NMR: (d⁶-DMSO) δ: 9.38 (1H, t, J = 5.7Hz), 8.87 (1H, s), 8.81 (1H, d, J = 5.4Hz), 8.42-8.49 (2H, br s), 8.41 (1H, d, J = 8.0Hz), 7.93 (1H, dd, J = 7.9, 5.8Hz), 7.87 (1H, t, J = 6.2Hz), 7.28-7.38 (4H, m), 7.16-7.25 (4H, m), 5.03 (2H, s), 4.22-4.43 (4H, m), 4.18 (2H, d, J = 6.1Hz), 3.39 (1H, dd, J = 14, 5.6Hz), 3.26 (1H, dd, J = 14.0, 8.2Hz).

E. [(R)-1-{(S)-1-[4-(Benzyloxycarbonylamino-methyl)-benzylcarbamoyl]-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl}-carbamic acid tert-butyl ester

(R)-2-Benzoyloxycarbonylamino-3-(4-ethoxy-phenyl)-propionic acid (870mg, 2.80mmol) was dissolved in CH₂Cl₂(100mL), HBTU (1.21g, 3.20mmol) and triethylamine (1.35g, 13.33mmol) were added. After 20 mins at room temperature the reaction mixture was cooled to 0°C and {4-(((S)-2-amino-3-pyridin-3-yl-propionylamino)-methyl)-benzyl}-carbamic acid benzyl ester
 5 dihydrochloride (1.31g, 2.67mmol) was added. After 2 hours at 0°C the reaction mixture was diluted with CHCl₃ (200mL), this solution was washed with 0.3M KHSO₄ (1x50mL), sat NaHCO₃ (1x50mL), water (1x50mL), brine (1x50mL), dried (Na₂SO₄) and evaporated in vacuo to give a white solid. The solid was triturated with EtOAc/Pet Ether 60-80°C to give a white solid identified as [(R)-1-((S)-1-[4-(benzyloxycarbonylamino-methyl)-benzylcarbamoyl]-2-
 10 pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-carbamic acid tert-butyl ester (2.40g, 1.70mmol, 90%).

$$[M+H]^+ = 710.18$$

F. [4-((S)-2-[(R)-2-Amino-3-(4-ethoxy-phenyl)-propionylamino]-3-pyridin-3-yl-propionylamino}-methyl)-benzyl]-carbamic acid benzyl ester Dihydrochloride
 15

[(R)-1-((S)-1-[4-(Benzyloxycarbonylamino-methyl)-benzylcarbamoyl]-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-carbamic acid tert-butyl ester (1.70, 2.42mmol) was treated with 4M HCl/dioxan (100 mL). After one hour at room temperature the solvent was removed to give a white solid identified as [4-((S)-2-[(R)-2-amino-3-(4-ethoxy-phenyl)-
 20 propionylamino]-3-pyridin-3-yl-propionylamino}-methyl)-benzyl]-carbamic acid benzyl ester dihydrochloride (1.50g, 2.32mmol, 97%).

$$[M+H]^+ = 609.99$$

¹H NMR: (d⁶-DMSO) δ: 9.29 (1H, d, J = 8.4Hz), 8.96 (1H, t, J = 5.8Hz), 8.83 (1H, s), 8.77 (1H, d, J = 5.4Hz), 8.39 (1H, d, J = 8.2Hz), 8.28-7.98 (2H, br s), 7.92 (1H, dd, J = 8.0, 5.7Hz), 7.86
 25 (1H, t, J = 6.2Hz), 7.28-7.38 (4H, m), 7.11-7.20 (4H, m), 6.95 (2H, d, J = 8.6Hz), 6.79 (2H, d, J = 8.6Hz), 5.02 (2H, s), 4.68-4.75 (1H, m), 4.23-4.25 (2H, m), 4.16 (2H, d, J = 6.1Hz), 3.83-4.13 (4H, m), 3.22 (1H, dd, J = 14.0, 4.4Hz), 3.03 (1H, dd, J = 13.7, 9.7Hz), 2.84 (1H, dd, J = 14.0, 5.9Hz), 2.63 (1H, dd, J = 13.8, 6.1Hz), 1.29 (3H, t, J = 7.0Hz).

G. [4-((S)-2-[(R)-3-(4-Ethoxy-phenyl)-2-(4-methyl-benzoylamino)-propionylamino]-3-pyridin-3-yl-propionylamino}-methyl)-benzyl]-carbamic acid benzyl ester
 30

[4-((S)-2-[(R)-2-Amino-3-(4-ethoxy-phenyl)-propionylamino]-3-pyridin-3-yl-propionylamino}-methyl)-benzyl]-carbamic acid benzyl ester dihydrochloride (150mg, 0.23mol)

was dissolved in dichloromethane (50 mL), this solution was cooled to 0°C. Triethylamine (70mg, 0.70mmol) was added followed by p-toluoyl chloride (39mg, 0.26mmol). After 18 hrs at 0°C to room temperature the reaction mixture was diluted with CHCl₃ (50 mL), this solution was washed with sat. NaHCO₃ (1x20 mL), water (1x20 mL), brine (1x20 mL), dried (Na₂SO₄) and evaporated in vacuo. The residue was purified by flash chromatography (silica), eluent 2% MeOH, 98% CHCl₃, fractions combined and evaporated in vacuo to give a colourless oil identified as [4-({(S)-2-[(R)-3-(4-ethoxy-phenyl)-2-(4-methyl-benzoylamino)-propionylamino]-3-pyridin-3-yl-propionylamino}-methyl)-benzyl]-carbamic acid benzyl ester (130mg, 0.18mmol, 77%).

10 $[M+H]^+ = 728.14$

H. N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoylethyl)-4-methyl-benzamide Dihydrochloride

[4-({(S)-2-[(R)-3-(4-ethoxy-phenyl)-2-(4-methyl-benzoylamino)-propionylamino]-3-pyridin-3-yl-propionylamino}-methyl)-benzyl]-carbamic acid benzyl ester (98mg, 0.13mmol) was dissolved in methanol (100mL), 1M hydrochloric acid (0.263mL, 0.263mmol) was added and the reaction mixture was hydrogenated over 10% Pd/C (50mg) at atmospheric pressure for 2 hours after which time the catalyst was filtered off and washed with methanol (100mL), the combined filtrates were evaporated in vacuo to give a white solid which was recrystallised from ethanol to give a white solid identified as N-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoylethyl)-2-pyridin-3-yl-ethylcarbamoylethyl]-4-methyl-benzamide dihydrochloride.

Yield = 340mg, 0.498mmol, 57%

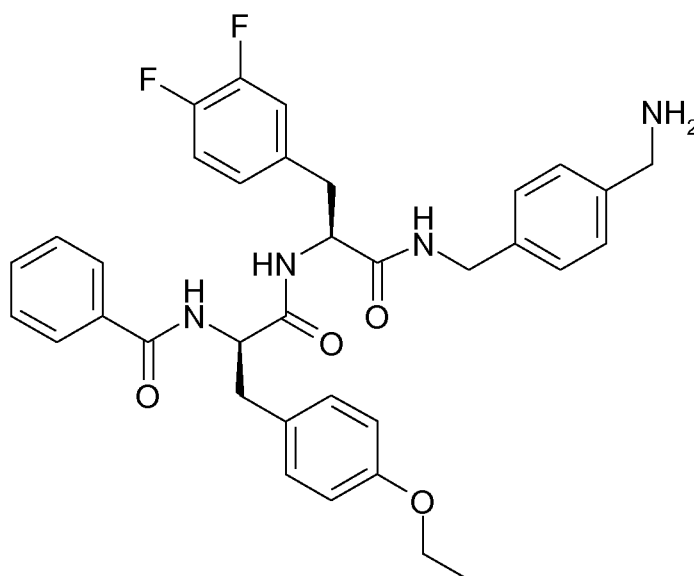
$[M+H]^+ = 593.99$

25 ¹H NMR: (d⁶-DMSO) δ: 1.28 (3H, t, J = 7.05Hz), 2.34 (3H, s), 2.72 (2H, d, J = 8.16Hz), 3.01-3.06 (1H, m), 3.25-3.28 (1H, m), 3.91-3.98 (4H, m), 4.32-4.38 (2H, m), 4.54-4.57 (1H, m), 4.70-4.73 (1H, m), 6.75 (2H, d, J = 6.83Hz), 7.18 (2H, d, J = 8.56Hz), 7.24 (2H, d, J = 7.56Hz), 7.25-7.27 (1H, m), 7.28 (2H, d, J = 6.78Hz), 7.39 (2H, d, J = 7.51Hz), 7.67 (1H, d, J = 7.51Hz), 7.76 (1H, s, br), 8.22 (1H, d, J = 7.56Hz), 8.33 (3H, s, br), 8.71-8.77(4H, m).

30

EXAMPLE 200

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoylethyl)-2-(3,4-difluoro-phenyl)-ethylcarbamoylethyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide



A. [4-(tert-Butoxycarbonylamino-methyl)-benzyl]-carbamic acid 9H-fluoren-9-ylmethyl ester

5 tert-Butyl 4-(Aminomethyl)benzylcarbamate (7.5g, 31.74mmol) was dissolved in dichloromethane (250mL). This solution was cooled to 0°C and triethylamine (9.63g, 93.2mmol) was added followed by carbonic acid 2,5-dioxo-pyrrolidin-1-yl ester 9H-fluoren-9-ylmethyl ester (12.85g, 38.09mmol). The reaction mixture was stirred at 0°C to room temperature for 3 hours and diluted with CHCl₃ (200mL) the filtrate was washed with 0.3M KHSO₄ (1x50mL), sat.

10 NaHCO₃ (1x50mL), water (1x50mL), brine (1x50mL), dried (Na₂SO₄) and evaporated in vacuo to give a white solid. The solid was triturated with EtOAc/Pet Ether 60-80°C to give a white solid identified as [4-(tert-butoxycarbonylamino-methyl)-benzyl]-carbamic acid 9H-fluoren-9-ylmethyl ester (13.96g, 30.44mmol, 96%).

[M+H]⁺ = 359.14 (M-Boc)

15

B. (4-Aminomethyl-benzyl)-carbamic acid 9H-fluoren-9-ylmethyl ester Hydrochloride

[4-(tert-Butoxycarbonylamino-methyl)-benzyl]-carbamic acid 9H-fluoren-9-ylmethyl ester (13.9g, 31.41mmol) was dissolved in 4M HCl in dioxan (400mL). After one hour at room temperature the solvent was removed in vacuo. The residue was triturated with acetone, the solid

20 was filtered off to give a white solid identified as (4-aminomethyl-benzyl)-carbamic acid 9H-fluoren-9-ylmethyl ester hydrochloride (11.9g, 30.135mmol, 99%)

[M+H]⁺ = 359.15

C. ((S)-2-(3,4-Difluoro-phenyl)-1-{4-[(9H-fluoren-9-ylmethoxycarbonylamino)-methyl]-benzylcarbamoyl}-ethyl)-carbamic acid tert-butyl ester

(S)-2-tert-Butoxycarbonylamino-3-(3,4-difluoro-phenyl)-propionic acid (2.1g, 6.96mmol) was dissolved in CH₂Cl₂(250mL) and DMF(25mL). This solution was cooled to 0°C. (4-Aminomethyl-benzyl)-carbamic acid 9H-fluoren-9-ylmethyl ester hydrochloride (2.5g, 6.33mmol) was added followed by HOBt (940mg, 6.96mmol) and triethylamine (1.92g, 18.99mmol). Water soluble carbodiimide (1.45g, 7.6mmol) was then added. After 18 hours at 0°C to room temperature reaction mixture was diluted with chloroform (400mL) washed with 0.3MKHSO₄ (1x30mL), NaHCO₃ (1x30mL), water (1x30mL), brine (1x30mL) and evaporated in vacuo giving a white solid. The residue was triturated with ethyl acetate/pet ether 60-80°C to give a white solid identified as ((S)-2-(3,4-difluoro-phenyl)-1-{4-[(9H-fluoren-9-ylmethoxycarbonylamino)-methyl]-benzylcarbamoyl}-ethyl)-carbamic acid tert-butyl ester (2.60g, 4.05mmol, 64%).

$$[M+H]^+ = 641.9, 664.07 (M+Na)$$

D. (4-{[(S)-2-Amino-3-(3,4-difluoro-phenyl)-propionylamino]-methyl}-benzyl)-carbamic acid 9H-fluoren-9-ylmethyl ester hydrochloride

((S)-2-(3,4-Difluoro-phenyl)-1-{4-[(9H-fluoren-9-ylmethoxycarbonylamino)-methyl]-benzylcarbamoyl}-ethyl)-carbamic acid tert-butyl ester (2.5g, 3.90mmol) was dissolved in 4M HCl in dioxan (150mL). After one hour at room temperature the solvent was removed in vacuo to give a white solid identified as (4-{[(S)-2-amino-3-(3,4-difluoro-phenyl)-propionylamino]-methyl}-benzyl)-carbamic acid 9H-fluoren-9-ylmethyl ester hydrochloride (2.25g, 3.89mmol, 100%).

$$[M+H]^+ = 542.12$$

E. [(R)-1-((S)-2-(3,4-Difluoro-phenyl)-1-{4-[(9H-fluoren-9-ylmethoxycarbonylamino)-methyl]-benzylcarbamoyl}-ethylcarbamoyl)-2-(4-ethoxy-phenyl)-ethyl]-carbamic acid tert-butyl ester

(R)-2-tert-Butoxycarbonylamino-3-(4-ethoxy-phenyl)-propionic acid (895mg, 2.90mmol) was dissolved in CH₂Cl₂(250mL) and DMF(25mL). This solution was cooled to 0°C. (4-{[(S)-2-Amino-3-(3,4-difluoro-phenyl)-propionylamino]-methyl}-benzyl)-carbamic acid 9H-fluoren-9-ylmethyl ester hydrochloride (1.5g, 2.63mmol) was added followed by HOBt (391mg, 2.90mmol) and triethylamine (800mg, 7.89mmol). Water soluble carbodiimide (605mg,

3.16mmol) was then added. After 18 hours at 0°C to room temperature reaction mixture was diluted with chloroform (200mL) washed with 0.3MKHSO₄ (1x30mL), NaHCO₃ (1x30mL), water (1x30mL), brine (1x30mL) and evaporated in vacuo giving a white solid. The residue was triturated with ethyl acetate/pet ether 60-80°C to give a white solid identified [(R)-1-((S)-2-(3,4-difluoro-phenyl)-1-{4-[(9H-fluoren-9-ylmethoxycarbonylamino)-methyl]-benzylcarbamoyl}-ethylcarbamoyl)-2-(4-ethoxy-phenyl)-ethyl]-carbamic acid tert-butyl ester (2.1g, 2.52mmol, 96%).

$[M+H]^+ = 733.15$ (M-Boc)

10 F. (4-{[(S)-2-[(R)-2-Amino-3-(4-ethoxy-phenyl)-propionylamino]-3-(3,4-difluoro-phenyl)-propionylamino]-methyl}-benzyl)-carbamic acid 9H-fluoren-9-ylmethyl ester Hydrochloride

[(R)-1-((S)-2-(3,4-Difluoro-phenyl)-1-{4-[(9H-fluoren-9-ylmethoxycarbonylamino)-methyl]-benzylcarbamoyl}-ethylcarbamoyl)-2-(4-ethoxy-phenyl)-ethyl]-carbamic acid tert-butyl ester (2.1g, 2.52mmol) was dissolved in 4M HCl in dioxan (150mL). After one hour at room temperature the solvent was removed in vacuo and the residue triturated with acetone to give a white solid identified as (4-{[(S)-2-[(R)-2-amino-3-(4-ethoxy-phenyl)-propionylamino]-3-(3,4-difluoro-phenyl)-propionylamino]-methyl}-benzyl)-carbamic acid 9H-fluoren-9-ylmethyl ester hydrochloride (1.9g, 2.47mmol, 98%).

20 $[M+H]^+ = 73.12$

G. (4-{[(S)-2-[(R)-2-Benzoylamino-3-(4-ethoxy-phenyl)-propionylamino]-3-(3,4-difluoro-phenyl)-propionylamino]-methyl}-benzyl)-carbamic acid 9H-fluoren-9-ylmethyl ester

(4-{[(S)-2-[(R)-2-Amino-3-(4-ethoxy-phenyl)-propionylamino]-3-(3,4-difluoro-phenyl)-propionylamino]-methyl}-benzyl)-carbamic acid 9H-fluoren-9-ylmethyl ester hydrochloride (410mg, 0.53mmol) was dissolved in dichloromethane (50mL). This solution was cooled to 0°C and triethylamine (162mg, 1.60mmol) was added followed by benzoyl chloride (82mg, 0.59mmol). The reaction mixture was stirred at 0°C to room temperature for 3 hours and diluted with CHCl₃ (100mL) the filtrate was washed with 0.3M KHSO₄ (1x30mL), sat. NaHCO₃ (1x30mL), water (1x30mL), brine (1x30mL), dried (Na₂SO₄) and evaporated in vacuo to give a white solid. The solid were triturated with hot ethanol, the cooled suspension was filtered to give a white solid identified as (4-{[(S)-2-[(R)-2-benzoylamino-3-(4-ethoxy-phenyl)-

propionylamino]-3-(3,4-difluoro-phenyl)-propionylamino]-methyl}-benzyl)-carbamic acid 9H-fluoren-9-ylmethyl ester (240mg, 0.34mmol, 99%).

$$[M+H]^+ = 697.18$$

5 **H. N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoylethyl)-2-(3,4-difluoro-phenyl)-ethyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide Hydrochloride**

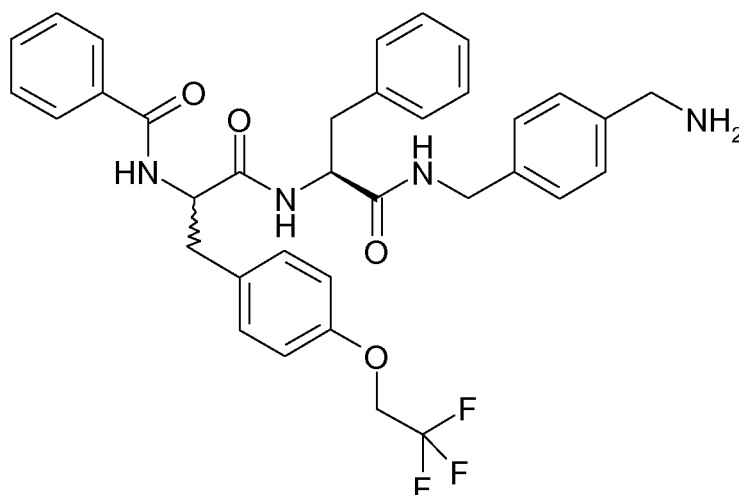
(4-{{[(S)-2-[(R)-2-Benzoylamino-3-(4-ethoxy-phenyl)-propionylamino]-3-(3,4-difluoro-phenyl)-propionylamino]-methyl}-benzyl)-carbamic acid 9H-fluoren-9-ylmethyl ester (180mg, 0.215mmol) was dissolved in diethylamine/THF (1:1, 100mL) the reaction mixture was stirred
10 at room temperature for 3 hours after which time the solvent was removed in vacuo and the residue was triturated with ethyl acetate/pet ether 60-80°C to give a white solid identified which was treated with 4M HCl in dioxan (20mL), the solvent was removed in vacuo and the residue recrystallised from EtOH to give a white solid identified as N-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoylethyl)-2-(3,4-difluoro-phenyl)-ethyl]-2-(4-ethoxy-phenyl)-ethyl]-
15 benzamide hydrochloride (62mg, 0.095mmol, 44%).

$$[M+H]^+ = 614.68$$

¹H NMR: (d⁶-DMSO) δ: 1.26(3H, t, J= 6.79Hz), 2.65-2.84(3H, m), 3.03-3.08(1H, m), 3.92(2H, q, J= 6.11Hz), 3.96(2H, s), 4.27-4.35(2H, m), 4.57-4.63(2H, m), 6.75(2H, d, J= 8.03Hz),
20 7.16(2H, d, J= 8.76Hz), 7.23-7.25(1H, m), 7.26-7.27(2H, m), 7.37-7.51(6H, m), 7.43(1H, d, J= 7.3Hz), 7.73-7.75(2H, m), 8.24 (2H, s), 8.50(1H, d, J= 7.40Hz), 8.67-8.71(2H, m).

EXAMPLE 201

25 **N-{(R,S)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoylethyl)-2-phenyl-ethyl]-2-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-ethyl}-benzamide**



A. (R,S)-2-Benzyloxycarbonylamino-3-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-propionic acid

(R,S)-2-Benzyloxycarbonylamino-3-(4-hydroxy-phenyl)-propionic acid (1.0g, 3.17mmol) was dissolved in THF (70mL), 2,2,2-trifluoroethyl trifluoromethanesulfonate (883mg, 3.81mmol) and cesium carbonate (3.1g, 9.51mmol) were added. The reaction mixture was stirred at 65°C for 18 hours after which time the solvent was removed in vacuo and the residue taken up in EtOAc (100mls), this solution was washed with 1M HCl (1x30mL), water (1x30mL), brine (1x30mL), dried (Na₂SO₄) and evaporated in vacuo. The residue was purified by flash chromatography (silica), eluent 1% AcOH, 5%MeOH, 94% CHCl₃, fractions combined and evaporated in vacuo to give a colourless oil identified as (R,S)-2-benzyloxycarbonylamino-3-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-propionic acid (380mg, 0.96mmol, 30%).

$[M+H]^+ = 395.11$

B. {(R,S)-1-[(S)-1-[4-(tert-Butoxycarbonylamino-methyl)-benzylcarbamoyl]-2-phenyl-ethylcarbamoyl]-2-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-ethyl}-carbamic acid benzyl ester

(R,S)-2-Benzyloxycarbonylamino-3-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-propionic acid (200mg, 0.50mmol) was dissolved in CH₂Cl₂(50mL) and DMF(2.5mL). This solution was cooled to 0°C. {4-[(S)-2-amino-3-phenyl-propionylamino)-methyl]-benzyl}-carbamic acid tert-butyl ester (231mg, 0.60mmol) was added followed by HOBt (75mg, 0.55mmol) and triethylamine (153mg, 1.51mmol). Water soluble carbodiimide (116mg, 0.60mmol) was then added. After 18 hours at 0°C to room temperature reaction mixture was diluted with chloroform (400mL) washed with 0.3M KHSO₄ (1x30mL), NaHCO₃ (1x30mL), water (1x30mL), brine (1x30mL), dried (Na₂SO₄) and evaporated in vacuo giving a yellow oil. The residue was purified by flash chromatography (silica), eluent 3%MeOH, 97% CHCl₃, fractions combined and evaporated in

vacuo to give a white solid identified as {(R,S)-1-[(S)-1-[4-(tert-butoxycarbonylamino-methyl)-benzylcarbamoyl]-2-phenyl-ethylcarbamoyl]-2-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-ethyl}-carbamic acid benzyl ester (350mg, 0.46mmol, 92%).

$[M+H]^+ = 663.43$ (M-Boc), 785.44 (M+Na)

5

C. 4-[(S)-2-[(R,S)-2-Amino-3-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-propionylamino]-3-phenyl-propionylamino)-methyl]-benzyl}-carbamic acid tert-butyl ester

{(R,S)-1-[(S)-1-[4-(tert-butoxycarbonylamino-methyl)-benzylcarbamoyl]-2-phenyl-ethylcarbamoyl]-2-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-ethyl}-carbamic acid benzyl ester (350mg, 0.46mmol) was dissolved in methanol (100mL). This solution was hydrogenated over 10% Pd/C (50mg) at atmospheric for 2 hours after which time the catalyst was filtered off and washed with methanol (100mls), the combined filtrates were evaporated in vacuo to give a white solid identified as 4-[(S)-2-[(R,S)-2-amino-3-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-propionylamino]-3-phenyl-propionylamino)-methyl]-benzyl}-carbamic acid tert-butyl ester

10 (270mg, 0.43mmol, 94%).

$[M+H]^+ = 629.40$

D. 4-[(S)-2-[(R,S)-2-Benzoylamino-3-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-propionylamino]-3-phenyl-propionylamino)-methyl]-benzyl}-carbamic acid tert-butyl ester

20 4-[(S)-2-[(R,S)-2-Amino-3-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-propionylamino]-3-phenyl-propionylamino)-methyl]-benzyl}-carbamic acid tert-butyl ester (250mg, 0.40mmol) was dissolved in dichloromethane (50mL). This solution was cooled to 0°C and triethylamine (121mg, 1.19mmol) was added followed by benzoyl chloride (61mg, 0.44mmol). The reaction mixture was stirred at 0°C to room temperature for 18 hours and diluted with CHCl₃ (100mls) 25 the filtrate was washed with 0.3M KHSO₄ (1x30mL), sat. NaHCO₃ (1x30mL), water (1x30mL), brine (1x30mL), dried (Na₂SO₄) and evaporated in vacuo to give a white solid. The solid was triturated with ethyl acetate/pet ether 60-80°C to give a white solid identified as 4-[(S)-2-[(R,S)-2-benzoylamino-3-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-propionylamino]-3-phenyl-propionylamino)-methyl]-benzyl}-carbamic acid tert-butyl ester (190mg, 0.26mmol, 65%).

30 $[M+H]^+ = 733.357, 755.49$ (M+Na)

E. N-[(R,S)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-ethyl]-benzamide Ditrifluoroacetate

{4-[(S)-2-[(R,S)-2-Benzoylamino-3-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-propionylamino]-3-phenyl-propionylamino)-methyl]-benzyl}-carbamic acid tert-butyl ester (190mg, 0.26mmol) was treated with 4M HCl/dioxan (50 mL). After one hour at room temperature the solvent was removed. The residue was purified by Prep HPLC (Sunfire prep C18 OBD column. 19x250mm, 10 μ). 10 to 90% 0.1% TFA/MeCN into 0.1%TFA/H₂O over 35 min at 20ml/min. Fractions were combined and freeze dried to give a white solid identified N-[(R,S)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-ethyl]-benzamide ditrifluoroacetate (56mg, 0.075mmol, 29%).

$[M+H]^+ = 632.51$

¹H NMR (d₆-DMSO) δ : 2.68 (1H, d, J = 7.44Hz), 2.82-3.08 (5H, m), 3.98 (2H, d, J= 5.92Hz), 4.31-4.34 (2H, m), 4.60-4.70 (4H, m), 6.90-6.94 (2H, m), 7.16-7.28 (6H, m), 7.34-7.37 (2H, m), 7.43-7.52 (3H, m), 7.74-7.79 (2H, m), 8.09 (3H, s, br), 8.47 (1H, d, J= 8.45Hz), 8.58-8.62 (2H, m).

15

Determination of the kinetic solubility in phosphate buffer

The solubility was determined turbidimetrically using standard published methods (Lipinski *et.al.* Advanced Drug Delivery Reviews 23 (1997) 3-25). A 10mM compound stock was prepared in DMSO, which was added to 25mM pH 7.0 sodium phosphate buffer to produce concentrations ranging from 12 to 235 μ M. After shaking for approximately 30 seconds the reduction in light transmission of these samples, at 650nm, was measured (Molecular Devices Spectromax UV/visible spectrophotometer). A second measurement was taken at approximately 30 seconds later. Absorbance of greater than 0.005 is taken to indicate that some precipitation of compound has occurred and therefore the compound is not soluble at that concentration.

25

Data acquired from these determinations are shown in Table 8 below:

Table 8

30

Example No	solubility in PO ₄ buffer (μM)	Example No	solubility in PO ₄ buffer (μM)	Example No	solubility in PO ₄ buffer (μM)
1	235	26	235	51	235
2	235	27	235	52	235
3	12	29	235	53	235
4	235	30	235	54	12
5	235	31	235	55	36
6	235	32	235	56	36
8	235	33	235	57	36
9	235	34	235	58	36
10	235	35	235	59	36
11	235	36	235	61	235
12	235	37	235	63	235
14	235	38	235	64	235
15	235	39	235	67	235
17	235	40	235		
18	235	41	235		
20	235	42	235		
21	12	43	235		
24	235	44	235		
25	235	45	235		

Determination of thermodynamic solubility in phosphate buffer

The thermodynamic solubility of compound was determined in Ammonium Phosphate Buffer (pH7.4, 290mOsm). Compounds were made up at a nominal concentration of 1 mg/mL, vortexed then placed on a shaker for 1h, 37°C at approximately 950 rpm. Following incubation samples were transferred to eppendorf tubes and centrifuged at 15,000 g (r.c.f.) for 10 minutes at 37°C. The concentration of compound in the supernatant was determined by LC-MS/MS analysis using a calibration line prepared from a DMSO stock.

Data acquired from these determinations are shown in Table 9 below:

Table 9

Example No	Solubility in PO ₄ buffer (µg/ml)
3	10
38	14.55
54	0.8
55	1.5
56	0.9
58	4.1
59	7.7
70	11.53
71	22.1
72	6.9
73	5.1
75	1.06
79	2.4
81	1.3
84	0.5
88	0.2
92	0.2
93	0.01
95	2.05
99	2.88
116	1.39
119	1.61
124	64.45
125	43.60
131	0.56
140	79.63
141	0.15
143	2.86

145	68.91
150	0.12
151	4.31
152	122.19
153	0.19
154	3.14
155	0.44
157	4.77
158	70.56
159	1.15
160	20.4
162	69.69
173	2.38
174	232.69
175	23.4
176	2.28
178	1.54
179	17.72
181	61.65
199	15.88
200	0.22

Biological Methods

The ability of the compounds of formula (I) to inhibit plasma kallikrein may be determined using
5 the following biological assay:

Determination of the IC₅₀ for plasma kallikrein

Plasma kallikrein inhibitory activity *in vitro* was determined using standard published methods
10 (see e.g. Johansen *et al.*, Int. J. Tiss. Reac. 1986, **8**, 185; Shori *et al.*, Biochem. Pharmacol., 1992, **43**, 1209; Stürzebecher *et al.*, Biol. Chem. Hoppe-Seyler, 1992, **373**, 1025). Human plasma kallikrein (Protogen) was incubated at 37°C with the fluorogenic substrate H-DPro-Phe-Arg-

AFC and various concentrations of the test compound. Residual enzyme activity (initial rate of reaction) was determined by measuring the change in optical absorbance at 410nm and the IC₅₀ value for the test compound was determined.

5 Data acquired from this assay are shown in Tables 10 and 11 below:

Table 10

Example No	IC50 (human PKal) μ M	Example No	IC50 (human PKal) μ M	Example No	IC50 (human PKal) μ M
1	0.089	24	0.45	47	2.1
2	0.82	25	0.34	48	4.7
3	0.022	26	0.081	49	2.2
4	0.23	27	1.3	50	4.3
5	0.46	28	8.9	51	1.0
6	1.0	29	0.30	52	0.86
7	0.074	30	0.58	53	0.65
8	0.74	31	4.3	54	0.083
9	2.0	32	0.20	55	0.031
10	7.2	33	1.1	56	0.50
11	1.7	34	1.4	57	10
12	0.29	35	2.0	58	0.17
13	10	36	8.2	59	0.078
14	0.40	37	0.91	60	10
15	0.47	38	0.047	61	0.22
16	0.053	39	0.31	62	10
17	5.8	40	1.8	63	0.052
18	8.8	41	1.3	64	3.4
19	10	42	0.53	65	10
20	0.73	43	1.3	66	10

Example No	IC50 (human PKal) μ M	Example No	IC50 (human PKal) μ M	Example No	IC50 (human PKal) μ M
21	5.1	44	0.18	67	1.6
22	10	45	0.79		
23	0.031	46	1.2		

Table 11

Example No	IC ₅₀ (human PKal)(μ M)
68	0.861
69	0.621
70	0.126
71	0.077
72	0.046
73	0.025
74	0.545
75	0.270
76	0.452
77	1.959
78	10.000
79	0.038
80	0.398
81	0.040
82	0.539
83	0.254
84	0.219
85	0.485
86	0.423
87	0.755

88	0.090
89	0.099
90	0.276
91	2.083
92	0.032
93	0.175
94	1.080
95	0.046
96	0.764
97	0.241
98	0.576
99	0.095
100	0.474
101	1.113
102	10.000
103	10.000
104	0.159
105	0.185
106	0.256
107	0.133
108	0.232
109	0.155
110	0.598
111	0.058
112	0.298
113	0.455
114	0.427
115	0.417
116	0.018
117	0.048
118	0.061
119	0.015

120	0.540
121	0.310
122	0.519
123	0.214
124	0.020
125	0.010
126	4.013
127	0.521
128	2.009
129	10.000
130	2.491
131	0.040
132	1.209
133	10.000
134	10.000
135	0.424
136	1.017
137	7.540
138	0.310
139	1.141
140	0.045
141	0.055
142	0.096
143	0.083
144	0.340
145	0.025
146	0.232
147	0.284
148	0.934
149	0.091
150	0.051
151	0.009

152	0.023
153	0.010
154	0.021
155	0.022
156	1.083
157	0.036
158	0.047
159	0.015
160	0.010
161	0.049
162	0.010
163	0.064
164	0.050
165	0.070
166	0.262
167	0.080
168	0.076
169	0.044
170	0.051
171	0.123
172	0.881
173	0.213
174	0.045
175	0.003
176	0.028
177	0.457
178	0.042
179	0.022
180	0.073
181	0.010
182	0.015
183	0.016

184	0.010
185	0.049
186	0.014
187	0.007
188	0.036
189	0.023
190	0.028
191	0.020
192	0.012
193	0.025
194	0.013
195	0.014
196	0.032
197	0.012
198	0.007
199	0.010
200	0.045
201	0.012

Selected compounds were further screened for inhibitory activity against the related enzyme KLK1. The ability of the compounds of formula (I) to inhibit KLK1 may be determined using the following biological assay:

5

Determination of the IC₅₀ for KLK1

KLK1 inhibitory activity *in vitro* was determined using standard published methods (see e.g. Johansen *et al.*, Int. J. Tiss. Reac. 1986, **8**, 185; Shori *et al.*, Biochem. Pharmacol., 1992, **43**, 1209; Stürzebecher *et al.*, Biol. Chem. Hoppe-Seyler, 1992, **373**, 1025). Human KLK1 (Calbiochem) was incubated at 37°C with the fluorogenic substrate H-DVal-Leu-Arg-AFC and various concentrations of the test compound. Residual enzyme activity (initial rate of reaction) was determined by measuring the change in optical absorbance at 410nm and the IC₅₀ value for the test compound was determined.

15

Data acquired from this assay are shown in Tables 12 and 13 below:

Table 12

Example No	IC ₅₀ (human KLK1) μ M	Example No	IC ₅₀ (human KLK1) μ M	Example No	IC ₅₀ (human KLK1) μ M
1	>1	24	>10	42	>10
3	>10	25	6.3	43	>10
4	>10	26	>10	44	>10
5	0.16	27	4.0	45	>10
6	>1	28	>10	46	>10
7	>1	29	>10	47	>10
8	>1	30	>10	48	>10
9	>1	31	9.6	49	>10
14	3.7	32	>10	50	5
15	2.2	33	>10	51	>10
16	4.9	34	7.7	52	9.5
17	>10	35	>10	53	>10
18	1.6	36	>10	66	8.4
19	>10	37	>10	67	>10
20	3.7	38	>10		
21	10	39	>10		
22	9.6	40	>10		
23	5.6	41	2.5		

5

Table 13

Example No	IC ₅₀ (human KLK1) (μ M)
68	>10

69	>10
70	7.5
71	>10
72	9.1
73	9.3
74	>10
75	>10
76	3.6
77	>10
78	>10
79	8.6
80	>10
81	>10
82	>10
83	>10
84	1.2
85	2.9
86	>10
87	3.4
88	>10
89	3.7
90	>10
91	>10
92	>10
93	>10
94	>10
95	>10
96	>10
97	4.7
98	5.1
99	>10
100	2.9

101	>10
102	9.0
103	>10
104	>10
105	>10
106	8.4
107	7.5
108	>10
109	>10
110	>10
111	>10
112	>10
113	>10
114	>10
115	9.0
116	6.3
117	6.1
118	>10
119	6.1
120	>10
121	>10
122	4.8
123	>10
124	8.4
125	7.0
126	>10
127	>10
128	>10
129	>10
130	>10
131	9.5
132	>10

133	>10
134	>10
135	>10
136	>10
137	>10
138	>10
139	>10
140	>10
141	>10
142	>10
143	>10
144	>10
145	>10
146	>10
147	>10
148	>10
149	>10
150	>10
151	>10
152	>10
153	>10
154	>10
155	>10
156	>10
157	>10
158	>10
159	>10
160	>10
161	6.9
162	>10
163	>10
164	>10

165	>10
166	>10
167	>10
168	>10
169	9.9
170	>10
171	9.4
172	>10
173	>10
174	>10
175	>10
176	>10
177	>10
178	>10
179	7.2
180	>10
181	>10
182	>10
183	>10
184	>10
185	>10
186	3.8
187	>10
188	>10
189	>10
190	>10
191	>10
192	>10
193	>10
194	7.1
195	>10
196	9.9

197	>10
198	9.1
199	>10
200	7.6
201	>10

Selected compounds were further screened for inhibitory activity against the related enzymes plasmin, thrombin, trypsin, Factor Xa and Factor XIIa. The ability of the compounds of formula (I) to these enzymes may be determined using the following biological assays:

5

Determination of enzyme selectivity

Human serine protease enzymes plasmin, thrombin, trypsin, Factor Xa and Factor XIIa were assayed for enzymatic activity using an appropriate fluorogenic substrate. Protease activity was measured by monitoring the accumulation of liberated fluorescence from the substrate over 5 minutes. The linear rate of fluorescence increase per minute was expressed as percentage (%) activity. The K_m for the cleavage of each substrate was determined by standard transformation of the Michaelis-Menten equation. The compound inhibitor assays were performed at substrate K_m concentration and activities were calculated as the concentration of inhibitor giving 50% inhibition (IC_{50}) of the uninhibited enzyme activity (100%).

15

Data acquired from these assays are shown in Table 14 below:

Table 14 (Selectivity data)

Example No	IC ₅₀ (μM)			
	Thrombin	Trypsin	Plasmin	Factor Xa
3	>40	10.8	3.5	>10

20

Table 15 (Selectivity data: Factor XIIa)

Example No	IC ₅₀ (Factor XIIa) (μM)
------------	-------------------------------------

3	>10
85	>10
91	>10
92	>10
93	>10
94	>10
95	>10
96	>10
157	>10
182	>10
183	>10
184	>10
185	>10
186	>10
187	>10
188	>10
189	>10
190	>10
191	>10
192	>10
193	>10
194	>10
195	>10
196	>10
197	>10
198	>10
199	>10
201	>10

Carbonic anhydrase I induced Retinal Vascular Permeability Model

The activity of Example 3 has been established using this *in vivo* model in the rat. Rats received an intravitreal injection (5 μ L) of phosphate buffered saline (PBS), CH-3457 (a plasma kallikrein inhibitor positive control) (10 μ M) or Example 3 (1 μ M) at time 0. After 30mins, a second intravitreal injection (5 μ L) of PBS or CA-I (200ng/eye) was given. After 15 minutes, 10% sodium fluorescein was infused and retinal vascular permeability (RVP) was measured by vitreous fluorophotometry 75 minutes following the initial IVT injections. Data for Example 3 are presented in Figure 1, in which the lower dashed line indicates basal RVP following PBS/PBS and upper dashed line indicates maximal stimulation. Intravitreal injection of 1 μ M Example 3 alone had no effect upon basal RVP compared to PBS alone (3.29 ± 0.21 vs. 3.64 ± 0.48). Intravitreal injection of Example 3 reduced RVP (stimulated by CA-I injection) by $53 \pm 21\%$.

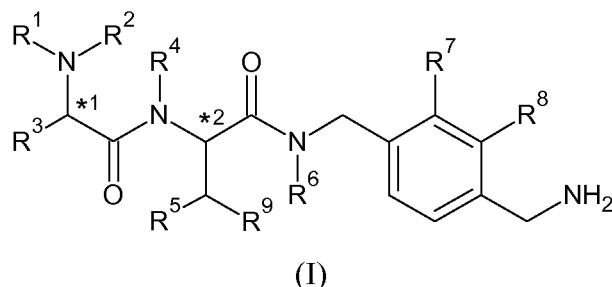
Pharmacokinetics

A pharmacokinetic study of Example 3 was performed to assess the ocular and systemic pharmacokinetics following a single IVT dose in pigmented (Dutch-belted) rabbits. Six rabbits per dose level were given a single, bilateral, IVT injection of 50 μ L of a 4.2 μ g/mL (210ng per eye) Example 3 formulated in phosphate buffered saline. One rabbit was euthanized at each time point (4, 8, 24, 48, 96 and 168 hours after IVT administration) and ocular tissue concentrations of Example 3 in the vitreous, retina/choroid and aqueous humour were measured. Serial blood samples were collected in surviving rabbits.

Ocular tissue concentration data are presented in Figure 2, in which the solid line for each ocular tissue concentration is the average of the left and right eye of each rabbit. The decline in ocular tissue concentrations of Example 3 was minimal over 7 days. Plasma concentrations of Example 3 following IVT administration were below 1 ng/mL at all time points.

Claims

1. A compound of formula I



5

wherein:

R^1 is selected from H, alkyl, -COalkyl, -COaryl, -COheteroaryl, -CO₂alkyl, -(CH₂)_aOH, -(CH₂)_bCOOR¹⁰, -(CH₂)_cCONH₂, -SO₂alkyl, -SO₂aryl, -SO₂(CH₂)_hR¹³, -CO(CH₂)_iR¹⁴, -COcycloalkyl, -COCH=CHR¹⁵, -CO(CH₂)_jNHCO(CH₂)_kR¹⁶ and -CONR¹⁷R¹⁸;

10 R^2 is selected from H and alkyl;

R^3 is selected from H, alkyl, -(CH₂)_daryl, -(CH₂)_eheteroaryl, -(CH₂)_fcycloalkyl, -(CH₂)_gheterocycloalkyl, -CH(cycloalkyl)₂, -CH(heterocycloalkyl)₂ and -(CH₂)_laryl-O-(CH₂)_m-aryl;

R^4 and R^6 are independently selected from H and alkyl;

15 R^5 is selected from H, alkyl, alkoxy and OH;

or R^4 and R^5 , together with the atoms to which they are attached, may join to form a 5- or 6-membered azacycloalkyl structure;

R^7 and R^8 are independently selected from H, alkyl, alkoxy, CN, halo and CF₃;

R^9 is aryl or heteroaryl;

20 R^{10} is H or alkyl;

a, b, c, d, e, f, g, h, i, j, l and m are independently 1, 2 or 3;

k is 0, 1, 2 or 3;

*1 and *2 denote chiral centres;

alkyl is a linear saturated hydrocarbon having up to 10 carbon atoms (C₁-C₁₀) or a branched

25 saturated hydrocarbon of between 3 and 10 carbon atoms (C₃-C₁₀); alkyl may optionally be substituted with 1 or 2 substituents independently selected from (C₃-C₁₀)cycloalkyl, (C₁-C₆)alkoxy, OH, CN, CF₃, COOR¹¹, fluoro and NR¹¹R¹²;

cycloalkyl is a mono- or bi-cyclic saturated hydrocarbon of between 3 and 10 carbon atoms;

cycloalkyl may optionally be fused to an aryl group; or cycloalkyl is adamantyl;

heterocycloalkyl is a C-linked or N-linked 3 to 10 membered saturated, mono- or bi-cyclic ring, wherein said heterocycloalkyl ring contains, where possible, 1, 2 or 3 heteroatoms independently selected from N, NR¹¹ and O;

alkoxy is a linear O-linked hydrocarbon of between 1 and 6 carbon atoms (C₁-C₆) or a branched O-linked hydrocarbon of between 3 and 6 carbon atoms (C₃-C₆); alkoxy may optionally be substituted with 1 or 2 substituents independently selected from (C₃-C₁₀)cycloalkyl, OH, CN, CF₃, COOR¹¹, fluoro and NR¹¹R¹²;

aryl is phenyl, biphenyl or naphthyl; aryl may be optionally substituted with up to 5 substituents independently selected from alkyl, alkoxy, OH, halo, CN, COOR¹¹, CF₃ and NR¹¹R¹²;

heteroaryl is a 5, 6, 9 or 10 membered mono- or bi-cyclic aromatic ring, containing, where possible, 1, 2 or 3 ring members independently selected from N, NR¹¹, S and O; heteroaryl may be optionally substituted with 1, 2 or 3 substituents independently selected from alkyl, alkoxy, OH, halo, CN, COOR¹¹, CF₃, NR¹¹R¹² and NHR¹⁹;

R¹¹ and R¹² are independently selected from H and alkyl;

R¹³ is aryl or heteroaryl;

R¹⁴ is aryl, heteroaryl, cycloalkyl or heterocycloalkyl;

R¹⁵ is H, alkyl, aryl, heteroaryl, cycloalkyl or heterocycloalkyl;

R¹⁶ is H, aryl or heteroaryl;

R¹⁷ is H, alkyl, aryl, heteroaryl or heterocycloalkyl;

R¹⁸ is -(CH₂)_mR²¹, where m is 0, 1, 2 or 3 and R²¹ is H, aryl or heteroaryl;

R¹⁹ -COalkyl, -COaryl or -COheteroaryl;

and tautomers, isomers, stereoisomers (including enantiomers, diastereoisomers and racemic and scalemic mixtures thereof), pharmaceutically acceptable salts and solvates thereof.

25

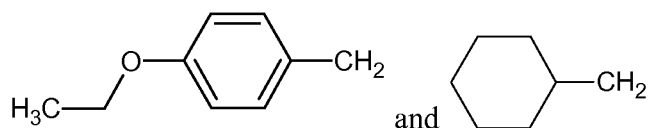
2. A compound as claimed in claim 1, wherein R⁹ is selected from phenyl and naphthyl, wherein phenyl may be optionally substituted with up to 3 substituents independently selected from alkyl, alkoxy, OH, halo, CN, COOR¹¹, CF₃ and NR¹¹R¹².

3. A compound as claimed in claim 1 or claim 2, wherein R⁹ is selected from phenyl, 1-naphthalene, 2,4-dichlorophenyl, 3,4-dichlorophenyl, 3,4-difluorophenyl, 4-chlorophenyl, 4-trifluoromethylphenyl and 4-ethoxyphenyl.

4. A compound as claimed in any one of claims 1 to 3, wherein R^1 is selected from H, -COaryl, -COalkyl, -CH₂COOH, -SO₂Ph and -SO₂CH₃.

5. A compound as claimed in any one of claims 1 to 4, wherein R^1 is selected from -COalkyl and -COaryl.

6. A compound as claimed in any one of claims 1 to 5, wherein R^3 is selected from:



7. A compound as claimed in any one of claims 1 to 6, wherein R^4 and R^6 are selected from H and CH₃.

8. A compound as claimed in any one of claims 1 to 7, wherein the stereochemical configuration about chiral centre *1 is *R*.

9. A compound as claimed in any one of claims 1 to 8, wherein the stereochemical configuration about chiral centre *2 is *S*.

10. A compound as claimed in any one of claims 1 to 9, wherein a is 2 and b, c, d, e, f, g, h, j, l and m are 1.

11. A compound as claimed in claim 1, selected from

(S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

{(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethylamino}-acetic acid;

(S)-N-(4-Aminomethyl-3-fluoro-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide;

(S)-N-(4-Aminomethyl-2-chloro-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide;

(S)-N-(4-Aminomethyl-benzyl)-3-(3,4-dichloro-phenyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-propionamide;

(S)-N-(4-Aminomethyl-3-chloro-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-phenyl-propionamide;

(S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionyl]-methyl-amino}-3-phenyl-propionamide;

((R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-cyclohexyl-ethyl)-methyl-amino)-acetic acid;

(S)-N-(4-Aminomethyl-3-fluoro-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionyl]-methyl-amino}-3-phenyl-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-methyl-carbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-methyl-carbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isobutyramide;

Naphthalene-1-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-chloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2,4-dichloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,4-difluoro-benzamide;

(R)-2-Amino-N-[(1S,2S)-1-(4-aminomethyl-benzylcarbamoyl)-2-hydroxy-2-phenyl-ethyl]-3-(4-ethoxy-phenyl)-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-nicotinamide;

(2S,3S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-propionylamino-propionylamino]-3-hydroxy-3-phenyl-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;

Thiophene-3-carboxylic acid-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Cyclohexanecarboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Isoxazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Benzo[b]thiophene-2-carboxylic acid[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-2-(4-chloro-benzenesulfonylamino)-3-(4-ethoxy-phenyl)-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3-chloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-chloro-benzamide

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3-trifluoromethyl-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,4-dichloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide;

(S)-N-(4-Aminomethyl-benzyl)-2-[(R)-3-(4-ethoxy-phenyl)-2-(2-phenylacetyl-amino-acetyl-amino)-propionyl-amino]-3-phenyl-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-fluoro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-6-methyl-nicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-methyl-nicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2,6-dichloro-nicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-5,6-dichloro-nicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2,3,6-trifluoro-isonicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,3,3-trifluoro-propionamide;

2,4-Dimethyl-thiazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

2-Methyl-thiazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

4-Methyl-thiazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Furan-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Methyl-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-methoxy-isonicotinamide;

3-Methyl-1H-pyrrole-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-propoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(4-chloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(4-fluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(4-methoxy-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3-fluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiophen-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-3-chloro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiophen-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-3-chloro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-difluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;

Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-chloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethyl]-3-(4-ethoxy-phenyl)-2-propionylamino-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethyl]-3-(4-ethoxy-phenyl)-2-propionylamino-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,3,3-trifluoro-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-chloro-benzamide;

- Isoxazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-difluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- 3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(1H-indol-3-yl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- 5 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;
- 3-Acetylamino-thiophene-2-carboxylic acid-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(2-fluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- 10 3-Methyl-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-3-methyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;
- 15 3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- 3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide;
- 20 3-Methyl-1H-pyrrole-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- 3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiazol-4-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;
- 25 3-Acetylamino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3-methyl-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2-methyl-benzamide;

- 5 3,5-Dimethyl-1H-pyrrole-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-3-methyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

- 10 3-Acetylamino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Acetylamino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

- 15 3-Chloro-thiophene-2-carboxylic acid [(R)-1-{[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-methyl-carbamoyl}-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(1S,2R)-1-(4-Aminomethyl-benzylcarbamoyl)-2-hydroxy-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

- 20 3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(1S,2R)-1-(4-aminomethyl-benzylcarbamoyl)-2-hydroxy-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-{(R,S)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-[4-(2,2,2-trifluoro-ethoxy)-phenyl]-ethyl}-benzamide;

and pharmaceutically acceptable salts and solvates thereof.

- 25 12. A compound as claimed in claim 1, selected from

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

Naphthalene-1-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-chloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-2,4-dichloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2- (4-ethoxy-phenyl)-ethyl]-nicotinamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,4-difluoro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2- (4-ethoxy-phenyl)-ethyl]-isonicotinamide;

Thiophene-3-carboxylic acid-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)- 2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Cyclohexanecarboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Isoxazole-5-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2- phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

(R)-N-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethyl]-2-(4-chloro-benzenesulfonylamino)-3-(4-ethoxy-phenyl)-propionamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3,4-dichloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-3-chloro-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-fluoro-benzamide;

3-Methyl-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Methyl-1H-pyrrole-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Amino-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-propoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-dichloro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(4-fluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiophen-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-3-fluoro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-3-chloro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-thiophen-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-2-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide;

Pyridine-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-3-chloro-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methoxy-benzamide;

Thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-4-methyl-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(3,4-difluoro-phenyl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

3-Chloro-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

5 N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-(1H-indol-3-yl)-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-benzamide;

N-[(R)-1-[(S)-1-(4-Aminomethyl-benzylcarbamoyl)-2-benzo[b]thiophen-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-isonicotinamide;

3-Acetyl-amino-thiophene-2-carboxylic acid-[(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-phenyl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

10 3-Methyl-thiophene-2-carboxylic acid [(R)-1-[(S)-1-(4-aminomethyl-benzylcarbamoyl)-2-pyridin-3-yl-ethylcarbamoyl]-2-(4-ethoxy-phenyl)-ethyl]-amide;

and pharmaceutically acceptable salts and solvates thereof.

15 13. A pharmaceutical composition comprising a compound as claimed in any one of claims 1 to 12 and a pharmaceutically acceptable carrier, diluent or excipient.

14. A compound as claimed in any one of claims 1 to 12 for use in medicine.

20 15. The use of a compound as claimed in any one of claims 1 to 12 in the manufacture of a medicament for the treatment or prevention of a disease or condition in which plasma kallikrein activity is implicated.

16. A method of treatment of a disease or condition in which plasma kallikrein activity is
25 implicated comprising administration to a subject in need thereof a therapeutically effective amount of a compound as claimed in any one of claims 1 to 12.

17. A compound as claimed in any one of claims 1 to 12 for use in a method of treatment of a disease or condition in which plasma kallikrein activity is implicated.

18. The use of claim 15, the method of claim 16 or a compound as claimed in claim 17,
5 wherein the disease or condition in which plasma kallikrein activity is implicated is selected from impaired visual acuity, diabetic retinopathy, diabetic macular oedema, hereditary angioedema, diabetes, pancreatitis, cerebral haemorrhage, nephropathy, cardiomyopathy, neuropathy, inflammatory bowel disease, arthritis, inflammation, septic shock, hypotension, cancer, adult respiratory distress syndrome, disseminated intravascular coagulation,
10 cardiopulmonary bypass surgery and bleeding from post operative surgery.

19. The use of claim 15, the method of claim 16 or a compound as claimed in claim 17, wherein the disease or condition in which plasma kallikrein activity is implicated is retinal vascular permeability associated with diabetic retinopathy and diabetic macular oedema.