

Acteli 81A/CR4

2,3,4,9-Tetrahydro-1*H*-carbazole derivatives as CRTH2 receptor antagonists

Field of the invention:

- 5 The present invention relates to 2,3,4,9-tetrahydro-1*H*-carbazole compounds of the Formula I and their use as potent “chemoattractant receptor-homologous molecule expressed on Th2 cells” (hereinafter called CRTH2) antagonists in the treatment of prostaglandin mediated diseases, to pharmaceutical compositions containing these compounds and to processes for their preparation. In particular, a compound of the Formula I may be used in pharmaceutical
- 10 compositions for the treatment of both chronic and acute allergic/immune disorders comprising allergic asthma, rhinitis, chronic obstructive pulmonary disease (COPD), dermatitis, inflammatory bowel disease, rheumatoid arthritis, allergic nephritis, conjunctivitis, atopic dermatitis, bronchial asthma, food allergy, systemic mast cell disorders, anaphylactic shock, urticaria, eczema, itching, inflammation, ischemia-
- 15 reperfusion injury, cerebrovascular disorders, pleuritis, ulcerative colitis, eosinophil-related diseases, such as Churg-Strauss syndrome and sinusitis, and basophil-related diseases, such as basophilic leukemia and basophilic leukocytosis, in humans and other mammals.

Background of the invention:

- 20 The complex response to allergen exposure results in a cascade of effects involving numerous mediators comprising different cell types like neutrophils, monocytes, eosinophiles, mast cells and T helper (Th) cells as well as cytokines and chemokines. For example, Th2 cells regulate allergic responses by producing Th2 cytokines, such as interleukin (IL)-4, IL-5 and IL-13. Among the events triggered by such mediators are Th2
- 25 cell differentiation, immunoglobulin (Ig)E synthesis, mast cell growth and differentiation, upregulation of CD23 expression, differentiation, recruitment, migration and activation of effector cells, such as eosinophils and basophils as well as the priming for their prolonged survival. Allergic or immune responses due to sustained release of above-mentioned mediators may culminate, if untreated, in severe inflammatory diseases with end-organ

damage, hyper-responsiveness, enhanced vascular permeability, edema, mucous hyper-secretion, airway hyperactivity, and bronchioconstriction.

Prostaglandin D2 (PGD2), histamine, cysteinyl leukotrienes (CysLTs) and thromboxane A2 (TxA2) are chemokines considered to act as proinflammatory key mediators in allergic responses. PGD2 is a major cyclooxygenase metabolite of arachidonic acid and is released in large amounts from activated mast cells during allergic attacks.

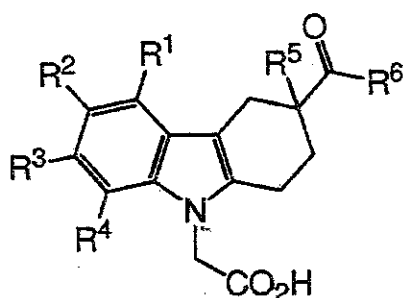
PGD2 is known to activate thromboxane A2 (TP) receptor, PGD2 (DP1) receptor and recently identified G-protein-coupled "chemoattractant receptor-homologous molecule expressed on Th2 cells" (CRTH2 or DP2) receptor. CRTH2 receptors are expressed on Th2 cells, eosinophiles and basophiles. PGD2 induces migration and activation of these cells via CRTH2 receptor activation.

A plethora of medications, such as anti-histamines, β 2-agonists, leukotriene modifiers, non-steroidal anti-inflammatory agents, cyclooxygenase-2 inhibitors, immunosuppressants, and monoclonal anti-IgE antibody have been used to treat symptoms of allergic conditions. Corticosteroids remain the most effective medication in the treatment of allergen-induced disorders, despite severe dose-limiting side effects that are due to non-specific inhibition of the transcription of several cytokines and chemokines. Such medications do not cure the disease and debilitating symptoms may relapse soon after treatment is stopped. Thus, there is still a need for new modalities to treat and/or to prevent allergic responses by suppressing specifically chemoattractant induced, e.g. PGD2 induced, tissue invasion of effector cells. Therefore, antagonists blocking CRTH2 receptor binding of PGD2 should be useful for the treatment of allergic conditions.

So far, few compounds having CRTH2 antagonistic activity have been reported in the patent literature. Bayer AG claims in GB Patent Specification No. 2388540 the use of Ramatroban ((3R)-3-(4-fluorobenzene-sulfonamido)-1,2,3,4-tetrahydrocarbazole-9-propionic acid) for the prophylaxis and treatment of allergic diseases, such as asthma, allergic rhinitis or allergic conjunctivitis. Oral bioavailability of Ramatroban and its ability to inhibit prostaglandin D2-induced eosinophil migration *in vitro* has been reported in *Journal of Pharmacology and Experimental Therapeutics*, 2003, 305(1), 347-352.

Description of the invention:

The disclosure relates to novel 2,3,4,9-tetrahydro-1*H*-carbazole compounds of the Formula I:



I

wherein

R^1 , R^2 , R^3 and R^4 independently represent hydrogen, alkyl (especially methyl or isopropyl), alkoxy (especially methoxy), halogen, nitro, cyano, trifluoromethyl, or formyl; preferably hydrogen, alkyl (especially methyl or isopropyl), alkoxy (especially methoxy), halogen, nitro, cyano, or trifluoromethyl; preferably R^1 represents hydrogen;

R^5 represents hydrogen, alkyl, or $-CF_3$; preferably hydrogen or alkyl, such as especially hydrogen, methyl, ethyl, or n-propyl;

R^6 represents alkoxy, aryl-alkoxy, or $-NR^7R^8$; especially phenyl-alkoxy or $-NR^7R^8$, most preferably $-NR^7R^8$;

R^7 and R^8 independently represent hydrogen, alkyl, cyano-alkyl, alkenyl, aryl, aryl-alkyl, phenylcarbonyl, cycloalkyl, pyridyl-alkyl (especially pyridyl-ethyl), thienyl-alkyl (especially thienyl-methyl), furanyl-alkyl, or imidazolyl-alkyl; or

R^7 and R^8 , together with the nitrogen atom to which they are attached, form a heterocyclic 5-, 6-, 7-, or 8-membered ring system with one to three heteroatoms which are selected from nitrogen, oxygen and sulfur (especially 1 or 2 nitrogen heteroatoms) and which ring system is optionally substituted with (i) one or two annellated benzene rings, which benzene rings are unsubstituted or substituted with one or two substituents independently selected from C1-C4 alkyl, C1-C4 alkoxy (especially methoxy), halogen, $-CF_3$, and $-OCF_3$; (ii) an unsubstituted phenyl ring; (iii) a mono- or di-substituted phenyl ring, wherein the substituents are independently selected from halogen, C1-C4 alkyl, C1-C4 alkoxy, $-CF_3$, and

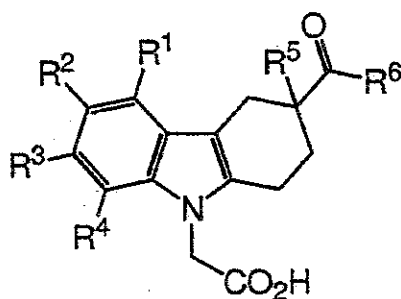
-OCF₃; or (iv) phenyl-alkyl wherein the alkyl moiety is substituted by phenyl, such as especially benzhydryl;

and optically pure enantiomers, mixtures of enantiomers such as racemates, optically pure diastereomers, mixtures of diastereomers, mixtures of enantiomers and diastereomers such as

5 diastereomeric racemates, meso forms, and geometric isomers;

prodrugs of such compounds in which a prodrug forming group is present, as well as solvates and morphological forms, and pharmaceutically acceptable salts thereof.

In a first aspect, there is provided a compound selected from the group consisting of 2,3,4,9-tetrahydro-1*H*-carbazoles of the Formula I:



I

wherein

R¹, R², R³ and R⁴ independently represent hydrogen, alkyl, alkoxy, halogen, nitro, cyano,

10 trifluoromethyl, or formyl;

R⁵ represents hydrogen, alkyl, or -CF₃;

R⁶ represents alkoxy, aryl-alkoxy, or -NR⁷R⁸;

R⁷ and R⁸ independently represent hydrogen, alkyl, cyano-alkyl, alkenyl, aryl, aryl-alkyl, phenylcarbonyl, cycloalkyl, pyridyl-alkyl, thienyl-alkyl, furanyl-alkyl, or imidazolyl-alkyl;

15 or

R⁷ and R⁸, together with the nitrogen atom to which they are attached, form a heterocyclic 5-, 6-, 7-, or 8-membered ring system with one to three heteroatoms which are selected from nitrogen, oxygen and sulfur and which ring system is optionally substituted with (i) one or two annellated benzene rings, which benzene rings are unsubstituted or substituted with one or two substituents independently selected from C1-C4 alkyl, C1-C4 alkoxy, halogen, -CF₃, and -OCF₃; (ii) an unsubstituted phenyl ring; (iii) a mono- or di-substituted phenyl ring, wherein the substituents are independently selected from halogen, C1-C4 alkyl, C1-C4 alkoxy, -CF₃, and -OCF₃; or (iv) phenyl-alkyl wherein the alkyl moiety is substituted by phenyl;

20

diastereomeric racemates, meso forms, and geometric isomers;
 prodrugs of such compounds in which a prodrug forming group is present, as well as
 solvates and morphological forms, and pharmaceutically acceptable salts thereof.

The compounds of the Formula I are CRTH2 receptor antagonists and may be used for the
 10 prevention and/or treatment of chronic and acute allergic immune disorders comprising
 allergic asthma, rhinitis, chronic obstructive pulmonary disease (COPD), dermatitis,
 inflammatory bowel disease, rheumatoid arthritis, allergic nephritis, conjunctivitis, atopic
 dermatitis, bronchial asthma, food allergy, systemic mast cell disorders, anaphylactic shock,
 urticaria, eczema, itching, inflammation, ischemia-reperfusion injury, cerebrovascular
 15 disorders, pleuritis, ulcerative colitis, eosinophil-related diseases comprising Churg-Strauss
 syndrome and sinusitis, and basophil-related diseases, comprising basophilic leukemia and
 basophilic leukocytosis, in humans and other mammals.

In one embodiment, the disclosure relates to a compound of Formula I, wherein
 R¹, R², R³ and R⁴ independently represent hydrogen, alkyl, alkoxy, halogen, nitro, cyano,
 20 trifluoromethyl, or formyl;
 R⁵ represents hydrogen, alkyl, or -CF₃;
 R⁶ represents alkoxy; aryl-alkoxy; mono-, di-, or tri-substituted aryl-alkoxy substituted
 independently in the aryl moiety with halogen, C1-C4 alkyl, C1-C4 alkoxy, -CF₃, -OCF₃; or
 -NR⁷R⁸;
 25 R⁷ and R⁸ independently represent hydrogen; alkyl; cyano-alkyl; alkenyl; aryl; mono-, di-,
 or tri- substituted aryl substituted independently in the aryl moiety with halogen, C1-C4
 alkyl, C1-C4 alkoxy, -CF₃, -OCF₃; aryl-alkyl; mono-, di-, or tri- substituted aryl-alkyl
 substituted independently in the aryl moiety with halogen, C1-C4 alkyl, C1-C4 alkoxy,
 -CF₃, -OCF₃; phenylcarbonyl; phenyl-C1-C4 alkyl optionally substituted with a
 30 methylenedioxy group at the phenyl ring; cycloalkyl; pyridyl-alkyl; thienyl-alkyl; furanyl-
 alkyl; or imidazolyl-alkyl; or
 R⁷ and R⁸, together with the nitrogen atom to which they are attached, form a heterocyclic
 5-, 6-, 7-, or 8-membered ring system with one to three heteroatoms which are selected from

nitrogen, oxygen and sulfur and which ring system is optionally substituted with (i) one or two annellated benzene rings, which benzene rings are unsubstituted or substituted with one or two substituents selected from C1-C4 alkyl, C1-C4 alkoxy, halogen, -CF₃, -OCF₃; (ii) an unsubstituted phenyl ring; or (iii) a mono- or di-substituted phenyl ring substituted with
5 halogen, C1-C4 alkyl, C1-C4 alkoxy, -CF₃, or -OCF₃.

In a preferred embodiment, R¹, R², R³ and R⁴ represent C₁-C₄ alkyl, C₁-C₄ alkoxy, halogen, nitro, cyano or trifluoromethyl. In a particularly preferred embodiment, R¹, R², R³ and R⁴ are selected from the group consisting of methyl, trifluoromethyl, methoxy, fluoro, chloro,
10 bromo or iodo.

In another preferred embodiment, R⁷ represents hydrogen, alkenyl, alkyl, aryl-alkyl, cycloalkyl, or heteroaryl-alkyl; and R⁸ represents aryl; aryl-alkyl; or heteroaryl-alkyl.

15 In a further preferred embodiment, R⁷ represents alkenyl, alkyl, aryl-C1-C4 alkyl, cycloalkyl, thienyl-C1-C4 alkyl, furanyl-C1-C4 alkyl, pyridyl-C1-C4 alkyl, or imidazolyl-C1-C4 alkyl; and R⁸ represents hydrogen, aryl, aryl-C1-C4 alkyl, furanyl-C1-C4 alkyl, pyridyl-C1-C4 alkyl, or thienyl-C1-C4 alkyl.

20 In a particularly preferred embodiment, R⁷ represents hydrogen, allyl, 2-cyano-ethyl, methyl, butyl, ethyl, isopropyl, benzyl, 1-phenyl-ethyl, 2-phenyl-ethyl, phenyl-propyl, cyclohexyl, or thiophen-3-ylmethyl; and R⁸ represents phenyl, 2-benzyl-phenyl, 2-methoxy-phenyl, 2-methyl-phenyl, 2-trifluoromethyl-phenyl, 3,4-dichloro-phenyl, 3-benzoyl-phenyl, 3-chloro-phenyl, 2-fluoro-phenyl, 3-fluoro-phenyl, 4-chloro-phenyl, 4-fluoro-phenyl, 4-
25 methoxy-biphenyl-3-yl, 4-trifluoromethoxy-phenyl, 5-chloro-2-methoxy-phenyl, naphthalen-1-yl, benzo[1,3]dioxol-5-ylmethyl, benzyl, diphenylmethyl, 1-phenyl-ethyl, 2-phenyl-ethyl, 2-pyridin-2-yl-ethyl, 4-phenyl-benzyl, 3,4-dichloro-benzyl, 2,4-dichloro-benzyl, difluoromethoxy-benzyl, 2-chloro-benzyl, 4-chloro-benzyl, 2-methylsulfanyl-benzyl, 2-fluoro-benzyl, 3-fluoro-benzyl, 4-fluoro-benzyl, 2-trifluoromethyl-benzyl, 3-
30 trifluoromethyl-benzyl, 2,4-difluoro-benzyl, 2,5-difluoro-benzyl, 2,6-difluoro-benzyl, 3,5-difluoro-benzyl, 4-chloro-2-fluoro-benzyl, (2-fluoro-phenyl)-ethyl, (3-fluoro-phenyl)-ethyl, (4-fluoro-phenyl)-ethyl, (4-chloro-phenyl)-ethyl, (2,6-dichloro-phenyl)-ethyl, naphthalene-1-ylmethyl, 1,2,3,4-tetrahydro-naphthalen-1-yl, indan-2-yl, or 2,2-diphenyl-ethyl.

- In a particular embodiment, R⁷ represents hydrogen, allyl, 2-cyano-ethyl, butyl, ethyl, isopropyl, 3-phenyl-propyl, benzyl, phenylethyl, cyclohexyl, or thiophen-3-ylmethyl; and R⁸ represents 2-benzyl-phenyl, 2-methoxy-phenyl, 2-methyl-phenyl, 2-trifluoromethyl-phenyl, 3,4-dichloro-phenyl, 3-benzoyl-phenyl, 3-chloro-phenyl, 3-fluoro-phenyl, 4-chloro-phenyl, 4-fluoro-phenyl, 4-methoxy-biphenyl-3-yl, 4-trifluoromethoxy-phenyl, 5-chloro-2-methoxy-phenyl, naphthalen-1-yl, (*R*)-1-phenyl-ethyl, (*S*)-1-phenyl-ethyl, benzo[1,3]dioxol-5-ylmethyl, benzyl, diphenylmethyl, phenylethyl, or 2-pyridin-2-yl-ethyl.
- 10 In a further embodiment, R⁷ represents allyl, 2-cyano-ethyl, butyl, ethyl, isopropyl, 3-phenyl-propyl, benzyl, phenylethyl, cyclohexyl, or thiophen-3-ylmethyl; and R⁸ represents 2-benzyl-phenyl, 2-methoxy-phenyl, 2-methyl-phenyl, 2-trifluoromethyl-phenyl, 3,4-dichloro-phenyl, 3-benzoyl-phenyl, 3-chloro-phenyl, 3-fluoro-phenyl, 4-chloro-phenyl, 4-fluoro-phenyl, 4-methoxy-biphenyl-3-yl, 4-trifluoromethoxy-phenyl, 5-chloro-2-methoxy-phenyl, naphthalen-1-yl, (*R*)-1-phenyl-ethyl, (*S*)-1-phenyl-ethyl, benzo[1,3]dioxol-5-ylmethyl, benzyl, diphenylmethyl, phenylethyl, or 2-pyridin-2-yl-ethyl; or
- 15 R⁸ represents hydrogen, aryl; aryl-C1-C4 alkyl, furanyl-C1-C4 alkyl, pyridyl-C1-C4 alkyl, or thienyl-C1-C4 alkyl.
- 20 In still another preferred embodiment, R⁷ and R⁸, together with the nitrogen atom to which they are attached, form a dihydro-dibenzo[*b,f*]azocine, dihydro-indole, dihydroisoquinoline, dihydroquinoline, or dibenzoazepine ring. Particularly preferred is 11,12-dihydro-6*H*-dibenzo[*b,f*]azocine-5-yl, 2,3-dihydro-indole-1-yl, 3,4-dihydro-1*H*-isoquinoline-2-yl, 3,4-dihydro-2*H*-quinoline-1-yl, 4-(4-fluoro-phenyl)-piperazine-1-yl, 6,11-dihydro-
- 25 dibenzo[*b,e*]azepine-5-yl, 6,7-dimethoxy-3,4-dihydro-1*H*-isoquinoline-2-yl, 7-trifluoromethyl-3,4-dihydro-2*H*-quinoline-1-yl, and dibenzo[*b,f*]azepine-5-yl. In a further particularly preferred embodiment R⁷ and R⁸, together with the nitrogen atom to which they are attached, form 11,12-dihydro-6*H*-dibenzo[*b,f*]azocine-5-yl, 2,3-dihydro-indole-1-yl, 3,4-dihydro-1*H*-isoquinoline-2-yl, 3,4-dihydro-2*H*-quinoline-1-yl, 4-(4-fluoro-phenyl)-
- 30 piperazine-1-yl, 6,11-dihydro-dibenzo[*b,e*]azepine-5-yl, 6,7-dimethoxy-3,4-dihydro-1*H*-isoquinoline-2-yl, 7-trifluoromethyl-3,4-dihydro-2*H*-quinoline-1-yl, dibenzo[*b,f*]azepine-5-yl, 1*H*,3*H*-benzo[*d,e*]isoquinoline-2-yl, 4-benzhydryl-piperazine-1-yl, or azocane-1-yl.

In a further preferred embodiment, R⁶ represents C₁-C₄ alkoxy or aryl-C₁-C₄ alkoxy, in particular (*R*)-1-phenyl-ethyloxy or benzyloxy.

The disclosure also relates to compounds of Formula I wherein the meanings of one
 5 or more of the substituents and symbols as defined for Formula I, or an embodiment of Formula I, are replaced by their preferred meanings as defined herein, such as those defined hereinabove.

In a very preferred embodiment, the disclosure relates to a compound of Formula I,
 10 wherein
 R¹, R², R³ and R⁴ independently represent hydrogen, alkyl, alkoxy, halogen, nitro, cyano, or trifluoromethyl; preferably R¹ represents hydrogen;
 R⁵ represents hydrogen or alkyl;
 R⁶ represents phenyl-alkoxy or -NR⁷R⁸;
 15 R⁷ and R⁸ independently represent hydrogen; alkyl; cyano-alkyl; alkenyl; phenyl optionally mono-substituted by halogen, alkyl, alkoxy, -CF₃, -OCF₃, phenyl-alkyl or phenyl-carbonyl; phenyl, di-substituted with a substituent independently selected from halogen, alkoxy and phenyl; phenyl-alkyl optionally substituted in the alkyl moiety by phenyl or optionally substituted in the phenyl ring by methylenedioxy; phenyl-alkyl which is di-substituted by
 20 halogen or mono-substituted by halogen, -CF₃, -OCHF₂, alkyl or alkylsulfanyl; naphthyl; naphthyl-alkyl; cycloalkyl, especially cyclopentyl or cyclohexyl, which is optionally substituted with an annellated benzene ring; pyridyl-alkyl; or thienyl-alkyl; or
 R⁷ and R⁸, together with the nitrogen atom to which they are attached, form a heterocyclic 5-, 6-, 7-, or 8-membered ring system with one or two heteroatoms selected from nitrogen
 25 and which ring system is optionally substituted with (i) one or two annellated benzene rings, which benzene rings are unsubstituted or substituted with one or two substituents independently selected from alkoxy and -CF₃; (ii) a mono-substituted phenyl ring substituted with halogen; or (iii) phenyl-alkyl, wherein the alkyl moiety is substituted with phenyl.

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Most preferred novel compounds of the disclosure include:

[3-methyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

{3-[(3-chloro-phenyl)-methyl-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic

acid;

[3-(2,3-dihydro-indole-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

[3-methyl-3-(phenyl-thiophen-3-ylmethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

5 [3-(benzyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

{3-[(4-fluoro-phenyl)-methyl-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

[3-(6,11-dihydro-dibenzo[*b,e*]azepine-5-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

10 [3-(benzyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

{3-[phenyl-(3-phenyl-propyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

[3-(2,3-dihydro-indole-1-carbonyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

{3-[(3,4-dichloro-phenyl)-methyl-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

15 [3-(allyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

{3-[benzyl-((*S*)-1-phenyl-ethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

and

{3-[methyl-(2-trifluoromethyl-phenyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid.

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Particularly preferred novel compounds of the disclosure include:

[3-(methyl-phenyl-carbamoyl)-6-trifluoromethyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

[3-methyl-3-(methyl-*o*-tolyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

25 [3-(2-benzyl-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

[3-(dibenzo[*b,f*]azepine-5-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

[3-(ethyl-naphthalen-1-yl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

[3-(benzhydryl-methyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

{3-methyl-3-[phenyl-(3-phenyl-propyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic

30 acid;

[3-(ethyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

[3-(phenyl-thiophen-3-ylmethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

[3-(benzyl-isopropyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

- {3-[4-(4-fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(benzyl-phenethyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(3,4-dihydro-2*H*-quinoline-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 5 [3-(cyclohexyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[(3-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-methyl-3-(phenethyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(2-methoxy-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(allyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 10 [3-(benzyl-phenethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(phenethyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(methyl-*o*-tolyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[benzyl-((*R*)-1-phenyl-ethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(methyl-phenyl-carbamoyl)-6-nitro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 15 [3-(7-trifluoromethyl-3,4-dihydro-2*H*-quinoline-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(3,4-dihydro-2*H*-quinoline-1-carbonyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[(4-fluoro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 20 [3-(4-methoxy-biphenyl-3-ylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [6-fluoro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [7-chloro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [6-chloro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[methyl-(2-pyridin-2-yl-ethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 25 {3-[(4-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- {(*R*)-3-[(4-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(isopropyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[(3,4-dichloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(benzhydryl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 30 [6-cyano-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[benzyl-((*R*)-1-phenyl-ethyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(2-benzyl-phenylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

- [6-isopropyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [6-methyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [3-(11,12-dihydro-6*H*-dibenzo[*b,f*]azocine-5-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-
 acetic acid;
- 5 [3-(methyl-phenethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [6-bromo-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [3-(3-benzoyl-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 (3-dibenzylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
 [3-(ethyl-naphthalen-1-yl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic
 10 acid;
 {3-[(3-fluoro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
 {3-[benzyl-(2-cyano-ethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
 (3-phenylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
 [3-(isopropyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 15 (3-diphenethylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
 [3-(benzhydryl-methyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [3-(5-chloro-2-methoxy-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [3-(butyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 {3-[ethyl-(4-trifluoromethoxy-phenyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic
 20 acid;
 [3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid; and
 [6-iodo-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid.

Preferred novel compounds of the disclosure include:

- 25 [6-methoxy-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 9-carboxymethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid (*R*)-1-phenyl-ethyl
 ester;
 {(*S*)-3-[(4-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
 [3-(6,11-dihydro-dibenzo[*b,e*]azepine-5-carbonyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-
 30 yl]-acetic acid;
 [8-chloro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 9-carboxymethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid benzyl ester;
 [3-(3,4-dihydro-1*H*-isoquinoline-2-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

- [3-(6,7-dimethoxy-3,4-dihydro-1*H*-isoquinoline-2-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 {3-[(benzo[1,3]dioxol-5-yl)methyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid; and
 5 [3-(cyclohexyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid.

Most preferred novel compounds of the disclosure further include:

- {3-[benzyl-((*R*)-1-phenyl-ethyl)-carbamoyl]-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
 10 [3-(benzhydryl-methyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 {3-[benzyl-((*S*)-1-phenyl-ethyl)-carbamoyl]-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
 [3-(benzyl-phenethyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [3-(2,3-dihydro-indole-1-carbonyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 15 [3-(benzyl-isopropyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [3-(azocane-1-carbonyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 (3-phenylcarbamoyl-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
 [3-(methyl-phenyl-carbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [6-fluoro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid; and
 20 {6-fluoro-3-[4-(4-fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid.

Particularly preferred novel compounds of the disclosure include:

- [3-(4-fluoro-phenylcarbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 25 {6-fluoro-3-[4-(4-fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
 [3-(benzhydryl-methyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 (3-{benzyl-[2-(2,6-dichloro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
 30 {3-[benzyl-((*S*)-1-phenyl-ethyl)-carbamoyl]-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
 [3-(3-fluoro-phenylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [3-(3,5-difluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

- [3-(3-fluoro-phenylcarbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(3-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(2,3-dihydro-indole-1-carbonyl)-3-ethyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-ethyl-3-(3-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
5 [3-ethyl-3-(4-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(2-chloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(benzyl-phenethyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-methyl-3-(2-methylsulfanyl-benzylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
10 (3-ethyl-3-phenylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
{3-[benzyl-((R)-1-phenyl-ethyl)-carbamoyl]-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
[3-ethyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(2-fluoro-phenylcarbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
15 [3-(2-fluoro-phenylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
{3-methyl-3-[(naphthalen-1-yl)methyl]-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
{3-[benzyl-(2-cyano-ethyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
20 [3-(benzyl-butyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(4-fluoro-phenylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(4-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(2,4-dichloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
{3-[benzyl-(4-fluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
25 [3-(benzyl-isopropyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(2-difluoromethoxy-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(2-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(benzyl-ethyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
30 (3-methyl-3-phenylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
[3-ethyl-3-(2-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(4-chloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid; and

(3-{benzyl-[2-(4-fluoro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid.

Preferred novel compounds of the disclosure include:

- 5 [3-methyl-3-(4-pentyl-benzylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 {3-[benzyl-(4-chloro-2-fluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
 [3-(azocane-1-carbonyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [3-(2-fluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 10 [3-(3,4-dichloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 (3-{benzyl-[2-(4-chloro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
 [3-(2,4-difluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [3-(benzhydryl-carbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 15 {8-chloro-3-[4-(4-fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
 [8-chloro-3-(2,3-dihydro-indole-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [3-(2,6-difluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 {3-[benzyl-(3-fluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 20 [3-(benzhydryl-methyl-carbamoyl)-8-chloro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 [3-(benzhydryl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 {3-[benzyl-(3-trifluoromethyl-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
 [3-(benzhydryl-carbamoyl)-3-ethyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 25 [3-(4-fluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 (3-{benzyl-[2-(3-fluoro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
 [3-(benzyl-isopropyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
 {3-[benzyl-(3,5-difluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 30 (3-{benzyl-[2-(2-fluoro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
 {3-[benzyl-(2-trifluoromethyl-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

{3-[benzyl-(4-fluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid; and

[3-(1*H*,3*H*-benzo[*de*]isoquinoline-2-carbonyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid.

5

Other preferred novel compounds of the disclosure include:

{3-[benzyl-(2,5-difluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

{3-[benzyl-(3-fluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

10 {3-methyl-3-[(*R*)-(1,2,3,4-tetrahydro-naphthalen-1-yl)carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

{3-[benzyl-(3,5-difluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

[3-(benzyl-methyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

15 (3-{benzyl-[2-(4-chloro-phenyl)-ethyl]-carbamoyl}-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;

(3-{benzyl-[2-(4-fluoro-phenyl)-ethyl]-carbamoyl}-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;

[8-chloro-3-(2,2-diphenyl-ethylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

20 {3-[benzyl-(3-trifluoromethyl-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

[3-(4-benzhydryl-piperazine-1-carbonyl)-8-chloro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

{3-[benzyl-(4-chloro-2-fluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

25

[3-(benzhydryl-carbamoyl)-8-chloro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

[3-(2,3-dihydro-indole-1-carbonyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

(3-{benzyl-[2-(3-fluoro-phenyl)-ethyl]-carbamoyl}-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;

30 (3-{benzyl-[2-(2,6-dichloro-phenyl)-ethyl]-carbamoyl}-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;

{3-[benzyl-(2-trifluoromethyl-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

{3-[benzyl-(2,5-difluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

[8-chloro-3-(indan-2-yl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

(3-{benzyl-[2-(2-fluoro-phenyl)-ethyl]-carbamoyl}-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;

[6-fluoro-3-((*R*)-1-phenyl-ethylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

[6-fluoro-3-((*R*)-1-phenyl-ethylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

[3-(benzyl-cyanomethyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

[3-(2,3-dichloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

10 and

[7-methyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid.

In a second aspect, there is provided a pharmaceutical composition comprising a compound according to any one of claims 1 to 12 and a pharmaceutically acceptable carrier.

In a third aspect, there is provided a compound according to any one of claims 1 to 12, or composition according to claim 13, for use as a medicament.

In a fourth aspect, there is provided use of a compound according to any one of claims 1 to 12 for the preparation of a pharmaceutical composition for the prevention and/or treatment of diseases selected from the group consisting of both chronic and acute allergic diseases or immune disorders comprising allergic asthma, rhinitis, chronic obstructive pulmonary disease, dermatitis, inflammatory bowel disease, rheumatoid arthritis, allergic nephritis, conjunctivitis, atopic dermatitis, bronchial asthma, food allergy, systemic mast cell disorders, anaphylactic shock, urticaria, eczema, itching, inflammation, ischemia-reperfusion injury, cerebrovascular disorders, pleuritis, ulcerative colitis, and eosinophil-related diseases, such as Churg-Strauss syndrome and sinusitis, and basophil-related diseases, such as basophilic leukemia and basophilic leukocytosis.

Unless explicitly stated otherwise, the general terms and names used hereinbefore and hereinafter preferably have within the context of this disclosure the following meanings:

15

The term "alkyl" as used herein, alone or in any combination, refers to a saturated aliphatic group including a straight or branched hydrocarbon chain containing 1-7, preferably 1-4 carbon atoms, i.e. C1-C4 alkyl. The alkyl group can optionally be substituted with one or more substituents, each independently selected from alkenyl, alkoxy, alkoxycarbonyl,

- 20 alkylcarbonyl, alkylcarbonyloxy, alkylendioxy, alkylsulfinyl, alkylsulfonyl, alkylthio, alkynyl, amino, aminocarbonyl, aryl, arylalkenyl, arylalkyloxy, aryloxy, aryloxycarbonyl, arylsulfinyl, arylsulfonyl, arylthio, carboxy, cyano, formyl, halogen, haloalkoxy, heterocyclyl, hydroxy, mercapto, and nitro, appended to any carbon atom of the alkyl moiety. If not explicitly indicated otherwise, the term "alkyl" preferably relates to an
- 25 unsubstituted alkyl group. Representative examples of alkyl include, but are not limited to, methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *tert*-butyl, *iso*-butyl (or 2-methylpropyl), cyclopropylmethyl, *n*-pentyl, *iso*-pentyl, *iso*-amyl, *n*-amyl, *n*-hexyl, *n*-heptyl, and *n*-octyl.

- The term "alkenyl" as used herein, alone or in any combination, refers to a straight or
- 30 branched hydrocarbon chain containing 2-7, preferably 2-4 carbon atoms with at least one carbon-carbon double bond ($R_aR_bC=CR_cR_d$). R_a - R_d refer to substituents, each individually and independently selected from hydrogen, alkyl, alkoxy, and alkoxyalkyl, preferably hydrogen and alkyl. Representative examples of alkenyl include, but are not limited to,

ethenyl, 2-propenyl, 2-methyl-2-propenyl, 3-butenyl, 4-pentenyl, and 5-hexenyl, especially 2-propenyl.

- The term "alkylendioxy" as used herein, alone or in any combination, refers to an
- 5 -O(CH₂)_nO- group, wherein n is preferably 1 or 2, and wherein the oxygen atoms are appended to two adjacent carbon atoms of the parent molecular moiety, preferably the two adjacent carbon atoms of a phenyl ring. Representative examples of alkylendioxy include, but are not limited to, methylenedioxy and ethylenedioxy.
- 10 The term "alkynyl" as used herein, alone or in any combination, refers to a straight or branched hydrocarbon chain containing 2-7 carbon atoms with at least one carbon-carbon triple bond (R_a-C≡C-R_b), R_a and R_b referring to substituents, each individually and independently selected from hydrogen, alkyl, alkenyl, alkoxy, and alkoxyalkyl, preferably hydrogen and alkyl. Representative examples of alkynyl include, but are not limited to,
- 15 acetylenyl, 1-propynyl, 2-propynyl, 1-butyne, 3-butyne, and 2-pentyne.

- The term "alkoxy" as used herein, alone or in any combination, refers to an alkyl group with 1 to 7, preferably 1 to 4, carbon atoms appended to the parent molecular moiety through an oxygen bridge. Representative examples of alkoxy include, but are not limited to, methoxy,
- 20 ethoxy, propoxy, 2-propoxy, butoxy, *tert*-butoxy, pentyloxy, and hexyloxy, especially methoxy.

- The term "alkoxyalkyl", as used herein, alone or in any combination, refers to an alkoxy group appended to the parent molecular moiety through an alkyl group. Representative
- 25 examples of alkoxyalkyl include, but are not limited to, *tert*-butoxymethyl, 2-ethoxyethyl, 2-methoxyethyl, and methoxymethyl.

- The term "alkoxycarbonyl", as used herein, alone or in any combination, refers to an alkoxy group appended to the parent molecular moiety through a carbonyl group. Representative
- 30 examples of alkoxycarbonyl include, but are not limited to, methoxycarbonyl, ethoxycarbonyl, and *tert*-butoxycarbonyl.

The term "alkoxycarbonylalkyl", as used herein, alone or in any combination, refers to an

alkoxycarbonyl group appended to the parent molecular moiety through an alkyl group. Representative examples of alkoxycarbonylalkyl include, but are not limited to, methoxycarbonylmethyl, methoxycarbonylpropyl, ethoxycarbonylbutyl, and 2-*tert*-butoxycarbonylethyl.

5

The term "alkylcarbonyl" or "acyl", as used herein, alone or in any combination, refers to an alkyl group appended to the parent molecular moiety through a carbonyl group. Representative examples of alkylcarbonyl include, but are not limited to, acetyl, 1-oxopropyl, 2,2-dimethyl-1-oxopropyl, 1-oxobutyl, and 1-oxopentyl.

10

The term "alkylcarbonylalkyl", as used herein, alone or in any combination, refers to an alkylcarbonyl group appended to the parent molecular moiety through an alkyl group. Representative examples of alkylcarbonylalkyl include, but are not limited to, 2-oxopropyl, 3,3-dimethyl-2-oxopropyl, 3-oxobutyl, and 3-oxopentyl.

15

The term "alkylcarbonyloxy", as used herein, alone or in any combination, refers to an alkylcarbonyl group appended to the parent molecular moiety through an oxygen bridge. Representative examples of alkylcarbonyloxy include, but are not limited to, acetyloxy, ethylcarbonyloxy, and *tert*-butylcarbonyloxy.

20

The term "alkylsulfinyl", as used herein, alone or in any combination, refers to an alkyl group appended to the parent molecular moiety through a sulfinyl group. Representative examples of alkylsulfinyl include, but are not limited to, methylsulfinyl and ethylsulfinyl.

25

The term "alkylsulfinylalkyl", as used herein, alone or in any combination, refers to an alkylsulfinyl group appended to the parent molecular moiety through an alkyl group. Representative examples of alkylsulfinylalkyl include, but are not limited to, methylsulfinylmethyl and ethylsulfinylmethyl.

30

The term "alkylsulfonyl", as used herein, alone or in any combination, refers to an alkyl group appended to the parent molecular moiety through a sulfonyl group. Representative examples of alkylsulfonyl include, but are not limited to, methylsulfonyl and ethylsulfonyl.

The term "alkylsulfonylalkyl", as used herein, alone or in any combination, refers to an alkylsulfonyl group appended to the parent molecular moiety through an alkyl group. Representative examples of alkylsulfonylalkyl include, but are not limited to, methylsulfonylmethyl and ethylsulfonylmethyl.

5

The term "alkylthio" (synonym "alkylsulfanyl"), as used herein, alone or in any combination, refers to an alkyl group appended to the parent molecular moiety through a thio group. Representative examples of alkylthio include, but are not limited to, methylthio, ethylthio, *tert*-butylthio, and hexylthio.

10

The term "alkylthioalkyl" (synonym "alkylsulfanylalkyl"), as used herein, alone or in any combination, refers to an alkylthio group appended to the parent molecular moiety through an alkyl group. Representative examples of alkylthioalkyl include, but are not limited to, methylthiomethyl and 2-(ethylthio)ethyl.

15

The term "amino", as used herein, alone or in any combination, refers to a $-NR_eR_f$ group, wherein R_e and R_f are substituents, each individually and independently selected from hydrogen, alkyl, aryl, arylalkyl, acyl, alkylcarbonyl, arylcarbonyl, carbamoyl, ureido, formyl, alkylsulfonyl, arylsulfonyl, and the like. Representative examples of amino include, but are not limited to, dimethylamino, ethylamino, and benzyl-(methyl)amino.

20

The term "aminoalkyl", as used herein, alone or in any combination, refers to an amino group appended to the parent molecular moiety through an alkyl group. Representative examples of aminoalkyl include, but are not limited to, aminomethyl, 2-(amino)ethyl, benzyl-(methyl)aminomethyl, and dimethylaminomethyl.

25

The term "aminocarbonyl" or "carbamoyl", as used herein, alone or in any combination, refers to an amino group appended to the parent molecular moiety through a carbonyl group. Representative examples of aminocarbonyl include, but are not limited to, dimethylaminocarbonyl, benzyl-aminocarbonyl, and ethylaminocarbonyl.

30

The term "aminocarbonylalkyl", as used herein, alone or in any combination, refers to an aminocarbonyl group appended to the parent molecular moiety through an alkyl group.

Representative examples of aminocarbonylalkyl include, but are not limited to, 2-amino-2-oxoethyl, 2-(benzylamino)-2-oxoethyl, 2-(methylamino)-2-oxoethyl, 4-amino-4-oxobutyl, and 4-(dimethylamino)-4-oxobutyl.

- 5 The term "aryl", as used herein, alone or in any combination, refers to a carbocyclic group having at least one aromatic ring, *e.g.* phenyl or biphenyl, especially phenyl, or multiple condensed ring systems, in which at least one ring is aromatic, *e.g.* 1,2,3,4-tetrahydronaphthyl, naphthyl, anthryl, phenanthryl, and fluorenyl, especially 1,2,3,4-tetrahydronaphthyl, naphthyl or indanyl. The aryl group may be optionally substituted with
- 10 one or more functional groups individually and independently selected from alkenyl, alkoxy, alkoxyalkyl, alkoxycarbonyl, alkoxycarbonylalkyl, alkyl, alkylcarbonyl, alkylcarbonylalkyl, alkylcarbonyloxy, alkylendioxy (especially methylenedioxy), alkylsulfinyl, alkylsulfinylalkyl, alkylsulfonyl, alkylsulfonylalkyl, alkylthio, alkylthioalkyl, alkynyl, amino, aminoalkyl, aminocarbonyl, aminocarbonylalkyl, aryl, arylalkenyl,
- 15 arylalkyloxy, arylalkyl, aryloxy, aryloxycarbonyl, aryloxycarbonylalkyl, arylsulfinyl, arylsulfinylalkyl, arylsulfonyl, arylsulfonylalkyl, arylthio, arylthioalkyl, carboxy, carboxyalkyl, cyano, cyanoalkyl, formyl, formylalkyl, halogen, haloalkoxy, haloalkyl, heterocyclyl, hydroxy, hydroxyalkyl, mercapto, nitro, and the like, such as also arylcarbonyl like especially phenylcarbonyl.
- 20 If R⁷ and/or R⁸ represents "aryl", this term preferably represents a phenyl or naphthyl, preferably a phenyl, radical, said radicals being optionally mono-substituted by halogen, alkyl, alkoxy, -CF₃, -OCF₃, phenyl-alkyl or phenyl-carbonyl; or di-substituted with a substituent independently selected from halogen, alkoxy and phenyl; and most preferably represents phenyl optionally mono-substituted by halogen, alkyl, alkoxy, -CF₃, -OCF₃,
- 25 phenyl-alkyl or phenyl-carbonyl; phenyl, di-substituted with a substituent independently selected from halogen, alkoxy and phenyl; or naphthyl.

The term "arylalkenyl", as used herein, alone or in any combination, refers to an aryl group appended to the parent molecular moiety through an alkenyl group. The aryl group may be

30 unsubstituted or substituted, especially as defined hereinabove for the aryl group. Representative examples of arylalkenyl include, but are not limited to, 2-phenylethenyl, 3-phenylpropen-2-yl, and 2-naphth-2-ylethenyl.

The term "aryl-alkoxy", as used herein, alone or in any combination, refers to an aryl group appended to the parent molecular moiety through an alkoxy group. The aryl group may be unsubstituted or substituted, especially as defined hereinabove for the aryl group.

Representative examples of aryl-alkoxy include, but are not limited to, 2-phenylethoxy, 5-phenylpentoxy, and 3-naphth-2-ylpropoxy. If R^6 is "aryl-alkoxy", this term preferably represents benzyloxy or 1-phenyl-ethoxy.

The term "arylalkyl", as used herein, alone or in any combination, refers to an aryl group appended to the parent molecular moiety through an alkyl group. The aryl group may be unsubstituted or substituted, especially as defined hereinabove for the aryl group.

Representative examples of arylalkyl include, but are not limited to, benzyl, 2-phenylethyl, 3-phenylpropyl, and 2-naphth-2-ylethyl, such as especially benzyl, phenylethyl, phenylpropyl, 1-phenyl-ethyl, 2-phenyl-ethyl, diphenylmethyl, 4-phenyl-benzyl, 3,4-dichloro-benzyl, 2,4-dichloro-benzyl, difluoromethoxy-benzyl, 2-chloro-benzyl, 4-chloro-benzyl, 2-methylsulfanyl-benzyl, 2-fluoro-benzyl, 3-fluoro-benzyl, 4-fluoro-benzyl, 2-trifluoromethyl-benzyl, 3-trifluoromethyl-benzyl, 2,4-difluoro-benzyl, 2,5-difluoro-benzyl, 2,6-difluoro-benzyl, 3,5-difluoro-benzyl, 4-chloro-2-fluoro-benzyl, (2-fluoro-phenyl)-ethyl, (3-fluoro-phenyl)-ethyl, (4-fluoro-phenyl)-ethyl, (4-chloro-phenyl)-ethyl, (2,6-dichloro-phenyl)-ethyl, naphthalene-1-ylmethyl, and 2,2-diphenyl-ethyl.

If R^7 and/or R^8 represents "aryl-alkyl", this term preferably represents phenyl-alkyl optionally substituted in the alkyl moiety by phenyl or optionally substituted in the phenyl ring by methylenedioxy; phenyl-alkyl which is di-substituted by halogen or mono-substituted by halogen, $-CF_3$, $-OCHF_2$, alkyl or alkylsulfanyl; or naphthyl-alkyl.

The term "aryloxy", as used herein, alone or in any combination, refers to an aryl group appended to the parent molecular moiety through an oxygen bridge. The aryl group can be unsubstituted or substituted, especially as defined hereinabove for the aryl group.

Representative examples of aryloxy include, but are not limited to, phenoxy, naphthyloxy, 3-bromophenoxy, 4-chlorophenoxy, 4-methylphenoxy, and 3,4-dimethoxyphenoxy.

The term "carbamoyl", as used herein, alone or in any combination, refers to a $-C(O)NR_eR_f$ group. R_e and R_f are substituents, each individually and independently selected from hydrogen, alkyl, arylalkyl, and the like.

Similarly, the term "thiocarbamoyl", as used herein, alone or in any combination, refers to a -C(S)NR₆R_f group.

- 5 The term "carbonyl", as used herein, alone or in any combination, refers to a -C(O)- group.

The term "carboxy", as used herein, alone or in any combination, refers to a -CO₂H group.

- 10 The term "carboxyalkyl", as used herein, alone or in any combination, refers to a carboxy group appended to the parent molecular moiety through an alkyl group. Representative examples of carboxyalkyl include, but are not limited to, carboxymethyl, 2-carboxyethyl, and 3-carboxypropyl.

- 15 The term "cyano", as used herein, alone or in any combination, refers to a -C≡N group.

- The term "cyanoalkyl", as used herein, alone or in any combination, refers to a cyano group appended to the parent molecular moiety through an alkyl group. Representative examples of cyanoalkyl include, but are not limited to, cyanomethyl, 2-cyanoethyl, and 3-cyanopropyl, especially 2-cyanoethyl.

- 20 The term "cycloalkyl", as used herein, alone or in any combination, refers to a saturated cyclic hydrocarbon moiety containing 3-10 carbon atoms, optionally substituted with one or more groups, each individually and independently selected from alkenyl, alkoxy, alkoxyalkyl, alkoxycarbonyl, alkoxycarbonylalkyl, alkyl, alkylcarbonyl, alkylcarbonylalkyl, alkylcarbonyloxy, alkylendioxy, alkylsulfinyl, alkylsulfinylalkyl, alkylsulfonyl, alkylsulfonylalkyl, alkylthio, alkylthioalkyl, alkynyl, amino, aminoalkyl, aminocarbonyl, aminocarbonylalkyl, aryl, arylalkenyl, arylalkyloxy, arylalkyl, aryloxy, aryloxycarbonyl, aryloxycarbonylalkyl, arylsulfinyl, arylsulfinylalkyl, arylsulfonyl, arylsulfonylalkyl, arylthio, arylthioalkyl, carboxy, carboxyalkyl, cyano, cyanoalkyl, formyl, formylalkyl, halogen, haloalkoxy, haloalkyl, heterocyclyl, hydroxy, hydroxyalkyl, mercapto, nitro, and the like. Representative examples of cycloalkyl include, but are not limited to, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, and cyclooctyl, especially cyclopentyl and
- 30

cyclohexyl. In polycyclic cycloalkyl groups one of the distal rings may be aromatic, *e.g.*, 1-indanyl, 2-indanyl, tetrahydronaphthyl, bicyclo[4.2.0]octa-1,3,5-trien-7-yl, and the like. If R⁷ and/or R⁸ represents "cycloalkyl", this term preferably represents a cyclopentyl or cyclohexyl radical, said radicals, especially the cyclohexyl radical, being optionally substituted with an annellated benzene ring.

The terms "cycloalkenyl" and "cycloalkynyl", as used herein, alone or in any combination, refer to unsaturated cyclic hydrocarbon moieties containing at least one carbon-carbon double or carbon-carbon triple bond, respectively. Such moieties may optionally be substituted with one or more groups as discussed hereinabove for the cycloalkyl groups.

The term "formyl", as used herein, alone or in any combination, refers to a -C(O)H group.

The term "formylalkyl", as used herein, alone or in any combination, refers to a formyl group, appended to the parent molecular moiety through an alkyl group. Representative examples of formylalkyl include, but are not limited to, formylmethyl and 2-formylethyl.

The term "halo" or "halogen", as used herein, alone or in any combination, refers to fluorine, bromine, chlorine, and iodine.

The term "haloalkyl", as used herein, alone or in any combination, refers to an alkyl group having at least one hydrogen atom replaced with a halogen atom. Representative examples of haloalkyl include, but are not limited to, chloromethyl, 2-fluoroethyl, trifluoromethyl, pentafluoroethyl, and 2-chloro-3-fluoropentyl.

The term "haloalkoxy", as used herein, alone or in any combination, refers to an alkoxy group having at least one hydrogen atom replaced with a halogen atom. Representative examples of haloalkoxy include, but are not limited to, chloromethoxy, 2-fluoroethoxy, trifluoromethoxy, and pentafluoroethoxy.

The term "heterocyclyl", as used herein, alone or in any combination, refers to a monocyclic, bicyclic or polycyclic ring system containing up to 15 ring atoms, at least one of these being a heteroatom, preferably one to three heteroatoms, independently selected

from nitrogen, oxygen or sulfur. The ring system may be saturated, partially unsaturated, unsaturated or aromatic, mono- or bicyclic. Representative examples of heterocyclyl include, but are not limited to, furyl, imidazolyl, imidazoliny, imidazolidinyl, isothiazolyl, isoxazolyl, morpholinyl, oxadiazolyl, oxazolyl, oxazoliny, oxazolidinyl, piperazinyl, 5 piperidinyl, pyranal, pyrazinyl, pyrazolyl, pyridyl, pyrimidinyl, pyridazinyl, pyrrolyl, pyrrolinyl, pyrrolidinyl, tetrahydrofuranal, tetrahydrothienyl, thiadiazolyl, thiazolyl, thiazolinyl, thiazolidinyl, thienyl, thiomorpholinyl, 1,1-dioxothiomorpholinyl, benzimidazolyl, phthalazinyl, benzothiazolyl, benzothienyl, benzoxazolyl, benzofuranal, indolyl, indolinyl, indazolyl, isobenzofuranal, isobenzothienyl, isoindolyl, isoindolinyl, 10 isoquinolinyl, quinolinyl, and quinazolinyl. Defined heterocyclyl moieties may be optionally substituted with one or more groups, each individually and independently selected from alkenyl, alkoxy, alkoxyalkyl, alkoxycarbonyl, alkoxycarbonylalkyl, alkyl, alkylcarbonyl, alkylcarbonylalkyl, alkylcarbonyloxy, alkylendioxy, alkylsulfinyl, alkylsulfinylalkyl, alkylsulfonyl, alkylsulfonylalkyl, alkylthio, alkylthioalkyl, alkynyl, 15 amino, aminoalkyl, aminocarbonyl, aminocarbonylalkyl, aryl, arylalkenyl, arylalkyloxy, arylalkyl, aryloxy, arylcarbonyl, arylalkylcarbonyl, (diaryl)alkylcarbonyl, aryloxy carbonyl, aryloxy carbonylalkyl, arylsulfinyl, arylsulfinylalkyl, arylsulfonyl, arylsulfonylalkyl, arylthio, arylthioalkyl, carboxy, carboxyalkyl, cyano, cyanoalkyl, cycloalkyl, cycloalkylcarbonyl, cycloalkylalkylcarbonyl, formyl, formylalkyl, halogen, haloalkoxy, 20 haloalkyl, heterocyclyl, heterocyclylcarbonyl, hydroxy, hydroxyalkyl, mercapto, nitro, and the like.

If R^7 and R^8 , together with the nitrogen atom to which they are attached, form a heterocyclic 5-, 6-, 7-, or 8-membered ring system, this ring system may be saturated, partially saturated or unsaturated, preferably saturated or partially saturated (partially 25 saturated preferably meaning that 1 double bond is present), and preferably contains one or two ring heteroatoms selected from nitrogen, and which ring system is optionally substituted as defined for Formula I. Preferably the ring system is optionally substituted with (i) one or two annellated benzene rings, which benzene rings are unsubstituted or substituted with one or two substituents independently selected from alkoxy and $-CF_3$; (ii) a mono-substituted 30 phenyl ring substituted with halogen; or (iii) phenyl-alkyl, wherein the alkyl moiety is substituted with phenyl.

The term "heteroaryl", as used herein, alone or in any combination, is a special case of

heterocyclyl and refers to a mono- or bicyclic or polycyclic aromatic ring system, in which at least one heterocyclic ring is aromatic.

5 The term "saturated heterocyclyl" is another special case of "heterocyclyl" and refers to saturated rings as defined above for "heterocyclyl".

The term "heterocyclylalkenyl", as used herein, alone or in any combination, refers to a heterocyclyl group appended to the parent molecular moiety through an alkenyl group. Representative examples of heterocyclylalkenyl include, but are not limited to, 2-pyrid-3-
10 ylethenyl, 3-quinolin-3-ylpropen-2-yl, and 5-pyrid-4-ylpenten-4-yl.

The term "heterocyclylalkoxy", as used herein, alone or in any combination, refers to a heterocyclyl group appended to the parent molecular moiety through an alkoxy group. Representative examples of heterocyclylalkoxy include, but are not limited to, 2-pyrid-3-
15 ylethoxy, 3-quinolin-3-ylpropoxy, and 5-pyrid-4-ylpentyloxy.

The term "heterocyclylalkyl", as used herein, alone or in any combination, refers to a heterocyclyl group appended to the parent molecular moiety through an alkyl group. Representative examples of heterocyclylalkyl include, but are not limited to, 2-pyrid-3-
20 ylmethyl and 2-pyrimidin-2-ylpropyl.

The term "heterocycliloxy", as used herein, alone or in any combination, refers to a heterocyclyl group appended to the parent molecular moiety through an oxy group. Representative examples of heterocycliloxy include, but are not limited to, pyrid-3-yloxy
25 and quinolin-3-yloxy.

The term "hydroxy" or "hydroxyl" as used herein, alone or in any combination, refers to an -OH group

30 The term "hydroxyalkyl", as used herein, alone or in any combination, refers to an alkyl group having at least one hydrogen atom replaced by a hydroxy group. Representative examples of hydroxyalkyl include, but are not limited to, hydroxymethyl, 2-hydroxyethyl, 3-hydroxypropyl, and 2-ethyl-4-hydroxyheptyl.

The term "nitro", as used herein, alone or in any combination, refers to a -NO₂ group.

The term "oxo", as used herein, alone or in any combination, refers to an =O group.

5

The term "oxy", as used herein, alone or in any combination, refers to an -O- group.

The terms "mercapto" and "thiol", as used herein, alone or in any combination, refer to an -SH group.

10

The terms "thio"(synonym "sulfanyl"), "sulfinyl" and "sulfonyl", as used herein, alone or in any combination, refer to a -S(O)_n group with n= 0, 1 and 2, respectively.

Within the scope of the present invention, unless indicated otherwise, compounds of
15 Formula I or pharmaceutically acceptable salts thereof are included, that may exist in, and be isolated in, isomeric forms, including *cis*- or *trans*-isomers or mixtures thereof, and tautomers. Other compounds of this invention may contain one or more stereogenic or asymmetric centers, such as one or more asymmetric carbon atoms, and thus may give rise to optically pure enantiomers, mixtures of enantiomers, racemates, enantiomer-pure
20 diastereomers, mixtures of diastereomers, epimers, and other stereoisomeric forms that may be defined, in terms of absolute stereochemistry, as (*R*)-, (*S*)- or (*R,S*)-configured, preferably in the (*R*)- or (*S*)-configuration. Such isomers can be obtained by methods within the knowledge of one skilled in the art, *e.g.* by stereochemically controlled synthesis using chiral synthons or chiral reagents, or by means of classical separation techniques, such as
25 chromatographic or crystallization methods, or by other methods known in the art, such as through formation of diastereomeric salts, for example by salt formation with an enantiomerically pure chiral acid, or with an enantiomerically pure chiral base, or by means of chromatography, for example by using chromatographic materials modified with chiral ligands. Furthermore, the present invention refers to compounds containing centers of any
30 geometric asymmetry, like, for example, unsymmetrically substituted olefinic double bond, including *E* or *Z* geometric isomers and mixtures thereof. Generally, pure isomers of compounds of Formula I are preferred over isomeric mixtures.

In the present invention, the compounds of Formula I may be used in the form of pharmaceutically acceptable salts. The term "pharmaceutically acceptable salts" refers to relatively nontoxic, inorganic or organic acid and base addition salts, which retain the biological effectiveness and properties of the parent compound, and which are not
5 biologically or otherwise undesirable (see, *e.g.* Berge *et al.*, *J. Pharm. Sci.* 1977, 66, 1-19).

Certain compounds of the present invention can contain one or more basic functional groups, such as amino, alkylamino, or arylamino, and, thus, be capable of forming pharmaceutically acceptable acid addition salts. These acid addition salts may be prepared
10 by standard procedures in a suitable solvent from the parent compound of Formula I, with an appropriate amount of an inorganic acid, including, but not limited to, for example, hydrochloric acid, hydrobromic acid, sulfuric acid, or phosphoric acid; or of an organic acid, including, but not limited to, acetic acid, propionic acid, octanoic acid, decanoic acid, glycolic acid, pyruvic acid, oxalic acid, maleic acid, malonic acid, succinic acid, fumaric
15 acid, tartaric acid, citric acid, ascorbic acid, amino acids, such as glutamic acid or aspartic acid, benzoic acid, cinnamic acid, salicylic acid, mandelic acid, methanesulfonic acid, ethanesulfonic acid, *p*-toluenesulfonic acid, or other acidic organic compounds.

Certain compounds of the present invention may, on the other hand, contain one or more
20 acidic functional groups and, thus, be capable of forming pharmaceutically acceptable base addition salts. These salts can be prepared by addition of an appropriate amount, usually in stoichiometric ratio, of an alkaline reagent, such as hydroxide, carbonate or alkoxide, containing the appropriate cation, to the free acid in a suitable solvent. Preferred inorganic salts include, but are not limited to, ammonium, sodium, potassium, calcium or magnesium,
25 also zinc salts and the like. Preferred salts derived from organic bases include, but are not limited to, salts of primary, secondary, and tertiary amines, substituted amines, cyclic amines, and basic ion exchange resins, such as isopropylamine, trimethylamine, diethylamine, triethylamine, tripropylamine, ethanolamine, lysine, arginine, *N*-ethylpiperidine, piperidine, polyamine resins, and the like.

30 Compounds of the present invention containing both acidic and basic groups can also form internal salts (zwitter ions).

For isolation or purification purposes, it is also possible to use pharmaceutically unacceptable salts, for example perchlorates, picolimates, picrates, or the like. For therapeutic use, only pharmaceutically acceptable salts or free compounds are employed, where applicable in the form of pharmaceutical preparations, and these are therefore
5 preferred.

Certain compounds of Formula I, including their salts, may exist in solvated as well as unsolvated forms, such as, for example, hydrated forms, or their crystals may, for example, include the solvent used for crystallization. Different crystalline forms may be present. The
10 present invention encompasses all such solvated and unsolvated forms.

The present invention also relates to prodrug derivatives of the parent compounds of Formula I. The term "prodrug" refers to pharmacologically inactive precursors of a drug that may be converted into its therapeutically active form under physiological conditions *in vivo*,
15 for example, when they undergo solvolysis, or enzymatic degradation in blood, or in cells, (Bundgard H., in *Design of Prodrugs*, pp. 7-9, 21-24, Elsevier, Amsterdam (1985); Silverman R. B., in *The Organic Chemistry of Drug Design and Drug Action*, pp. 352-401, Academic Press, San Diego, CA (1992); Higuchi T. *et al.*, "Pro-drug as Novel Delivery Systems", A.C.S. Symposium Series, Vol. 14). The term "prodrug" also includes any
20 covalently bonded carriers, which release the active parent compound *in vivo* when administered to a mammal. Prodrug modifications of a compound often offer advantages of solubility, bioavailability, absorption, tissue compatibility, tissue distribution, or delayed release in the mammalian organism. Prodrugs are variations or derivatives of the compounds of Formula I, which have groups cleavable under metabolic conditions, for
25 example, pharmaceutically acceptable esters, or amides. Such groups can be cleaved enzymatically or non-enzymatically, or hydrolytically to the free hydroxy, carboxy, or amino group of the active parent compound. In another embodiment, the prodrug is a reduced form, which is oxidized *in vivo* to the therapeutic compound, for example, a thiol, which is oxidized to a sulfonate or sulfate, or an alcohol which is oxidized to a carboxylic
30 acid.

Further included within the scope of the present invention are pharmaceutically acceptable esters of the compounds of Formula I. The term "pharmaceutically acceptable esters" refers

to relatively non-toxic, esterified products of the parent compound. These esters can be prepared *in situ* during the final isolation and purification of the compounds, or by separately reacting the purified compounds in its free acid or hydroxyl form with a suitable esterifying agent. Carboxylic acids can be converted into esters *via* treatment with an alcohol in the presence of a catalyst. Hydroxyl containing derivatives can be converted into esters *via* treatment with an esterifying agent such as alkanoyl halides. The term further includes lower hydrocarbon groups capable of being solvated under physiological conditions, for example, alkyl esters, preferred methyl, ethyl, and propyl esters, methoxymethyl ester, methylthiomethyl ester, pivaloyloxymethyl ester and the like (see, *e.g.* Berge *et al.*, *J. Pharm. Sci.* 1977, 66, 1-19).

The compounds of the present invention have useful, in particular pharmacologically useful, properties. They are able to specifically antagonize the effect of endogenous PGD₂ on the CRTH2 receptor, and may be used for the prevention and/or treatment of chronic and acute allergic immune disorders comprising allergic asthma, rhinitis, chronic obstructive pulmonary disease (COPD), dermatitis, inflammatory bowel disease, rheumatoid arthritis, allergic nephritis, conjunctivitis, atopic dermatitis, bronchial asthma, food allergy, systemic mast cell disorders, anaphylactic shock, urticaria, eczema, itching, inflammation, ischemia-reperfusion injury, cerebrovascular disorders, pleuritis, ulcerative colitis, eosinophil-related diseases comprising Churg-Strauss syndrome and sinusitis, and basophil-related diseases, comprising basophilic leukemia and basophilic leukocytosis, in humans and other mammals.

A compound or a pharmaceutical composition of the invention may be used as a drug (medicine) or therapeutic agent for prevention and/or treatment of both chronic and acute allergic/immune disorders such as allergic asthma, rhinitis, COPD, dermatitis, inflammatory bowel disease, rheumatoid arthritis.

In another aspect, the compounds of Formula I may be used as standard or reference compounds in tests or assays involving the inhibition of the CRTH2 receptor. Such compounds could be made commercially available for use as a reference, quality standard or control, for example in pharmaceutical research when developing new assays or protocols related to CRTH2 activity.

As mentioned earlier, compounds of Formula I, or salts, or prodrugs thereof, antagonize the PGD₂ activation of the CRTH2 receptor. The biological effect of such compounds may be tested in a variety of *in vitro*, *ex vivo* and *in vivo* assays.

5

The ability of the compounds of Formula I to bind to the CRTH2 receptor may be measured by methods similar to those described in Sawyer N. *et al.*, *Br. J. Pharmacol.*, 2002, 137, 1163-1172 and by the method described below in the experimental part.

10 With this type of assay, IC₅₀ values (*i.e.* the concentrations where half-maximal inhibition of the interaction is found) in the range of 0.001 to 10 μM, preferably values below 1 μM, in particular values below 0.05 μM, are found with test compounds of Formula I. Exemplary IC₅₀ values determined in this test are given below in Table 4.

15 A functional assay with cells expressing the human CRTH2 receptor may be used to detect changes in the levels of intracellular calcium concentration following compound treatment. After addition of the compound the cells are challenged with PGD₂. In a Fluorescent Imaging Plate Reader (FLIPR™, Molecular Devices, Sunnyvale, California) fluorescence emission is recorded during both additions, emission peak values above base level after
20 PGD₂ addition were exported, normalized to low controls (no PGD₂) and high controls (no active compound). The relative values of the remaining activity were used to determine IC₅₀ values by curve fitting the data to a single site to a four-parameter logistic sigmoid dose response curve of the equation $A + \frac{(B-A)}{1 + ((C/x)^D)}$.

25 The ability of the compounds to inhibit PGD₂ induced change of intracellular calcium levels *via* CRTH2 activation may be measured by methods known to one skilled in the art or by the method described below in the experimental part.

With this assay, IC₅₀ values (*i.e.* the concentration of a compound at which the remaining
30 activity is 50%) in the range of 0.001 and 10 μM, preferably below 0.5 μM, are obtained with test compounds of Formula I. Exemplary IC₅₀ values determined in this test are given below in Table 5.

The results of these assays clearly demonstrate, that the present invention provides functional antagonists of the PGD₂ receptor.

On the basis of the biological studies discussed hereinabove, a compound of Formula I
5 according to the invention may show therapeutic efficacy against chronic and acute allergic/immune disorders such as allergic asthma, rhinitis, chronic obstructive pulmonary disease (COPD), dermatitis, inflammatory bowel disease, and rheumatoid arthritis.

A compound of Formula I, a pharmaceutically acceptable salt or a prodrug thereof, can be
10 administered alone in pure form or in combination with one or more other therapeutic agents, possible combination therapy taking the form of fixed combinations or the administration of a compound of the invention and one or more other therapeutic agents being staggered or given independently of one another, or the combined administration of fixed combinations and one or more other therapeutic agents. A compound of Formula I can
15 besides or in addition be administered especially for prevention and/or treatment of both chronic and acute allergic disorders or immune disorders in combination with other inflammatory diseases. Long-term therapy is equally possible as is adjuvant therapy in the context of other treatment strategies, as described above. Other possible treatments are preventive therapies, for example in patients at risk.

20 The invention relates also to pharmaceutical compositions comprising compounds of Formula I, to their use in therapeutic, in a broader aspect of the invention also prophylactic treatment or a method of treatment of the diseases mentioned above, to the compounds for said use and to the preparation of pharmaceutical formulations (medicines).

25 The pharmaceutically acceptable compounds of the present invention may be used, for example, for the preparation of pharmaceutical compositions that comprise an effective amount of the active ingredient together or in admixture with a significant amount of one or more inorganic, organic, solid or liquid, pharmaceutically acceptable carriers.

30 The invention relates also to a pharmaceutical composition that is suitable for administration to a warm-blooded animal, especially a human (or to cells or cell lines derived from a warm-blooded animal, especially a human, for the treatment or, in a broader aspect of the

invention, prevention of (*i.e.* prophylaxis against) a disease that responds to blockade of the interaction of the CRTH2 receptor with PGD₂, comprising an amount of a compound of Formula I or a pharmaceutically acceptable salt or a prodrug thereof, which is effective for said inhibition, together with at least one pharmaceutically acceptable carrier.

5

The pharmaceutical compositions according to the invention are those for enteral administration, such as nasal, buccal, rectal, dermal or, especially oral administration, and for parenteral administration, such as intramuscular, intravenous or subcutaneous, intrasternal, intravitreal, injection or infusion, to warm-blooded animals, especially humans.

10 Such compositions comprise an effective dose of the pharmaceutically active ingredient, alone or together with a significant amount of a pharmaceutically acceptable carrier. The dosage of the active ingredient depends on the species of warm-blooded animal, the body weight, the age and the individual conditions, individual pharmacokinetic data, the disease to be treated and the mode of administration.

15

The invention relates also to a process or a method for the treatment of a pathological condition mentioned hereinabove, especially a disease, which responds to blockade of the interaction of the CRTH2 receptor with PGD₂, especially allergic asthma, rhinitis, chronic obstructive pulmonary disease (COPD), dermatitis, inflammatory bowel disease, and
20 rheumatoid arthritis. The compounds of Formula I or salts or prodrugs thereof can be administered as such or especially in the form of pharmaceutical compositions.

The dose to be administered to warm-blooded animals, for example humans of approximately 70 kg body weight, is preferably from approximately 3 mg to
25 approximately 3000 mg, more preferably from approximately 10 mg to approximately 1000 mg per person per day, divided preferably into 1 to 3 single doses which may, for example, be of the same size. The amount of the compound actually administered will typically be determined by a physician, in the light of the relevant circumstances, including the condition to be treated, the chosen route of administration, the actual compound
30 administered, the age, the weight, and response of the individual patient, the severity of the patient's symptoms, and the like, for example, children usually receive half of the adults dose.

The pharmaceutical compositions comprise from approximately 1% to approximately 95%, preferably from approximately 20% to approximately 90%, active ingredient.

Pharmaceutical compositions according to the invention may be, for example, in unit dosage forms such as coated and uncoated tablets, pills, ampoules, vials, suppositories, dragées, or capsules. Further dosage forms are, for example, ointments, creams, pastes, emulsions, foams, chewable gums, tinctures, lip-sticks, drops, sprays or aerosols, syrups or elixirs, dispersions, transdermal patches or pads, or *via* an intravitreal device that releases the compound in a sustained capacity, and the like. Examples are capsules containing from about 0.05 g to about 1.0 g active ingredient.

The pharmaceutical compositions of the present invention are prepared in a manner known, *per se*, for example by means of conventional mixing, granulating, coating, dissolving, lyophilizing or confectioning processes.

Solutions of the active ingredient, and also suspensions, and especially isotonic aqueous solutions or suspensions, are preferably used, it being possible, for example in the case of lyophilized compositions, that comprise the active ingredient alone or together with a carrier, for example mannitol, for such solutions or suspensions to be produced prior to use. The pharmaceutical compositions may be sterilized and/or may comprise excipients, for example preservatives, stabilizers, wetting agents and/or emulsifiers, solubilizers, salts for regulating osmotic pressure and/or buffers and are prepared in a manner known *per se*, for example by means of conventional dissolving or lyophilizing processes. The said solutions or suspensions may comprise viscosity-increasing substances, such as sodium carboxymethylcellulose, carboxymethylcellulose, dextran, polyvinylpyrrolidone or gelatin.

Suspensions in oil comprise as the oil component the vegetable, synthetic or semi-synthetic oils customary for injection purposes. There may be mentioned as such especially liquid fatty acid esters that contain as the acid component a long-chain fatty acid having from 8 to 22, especially from 12 to 22, carbon atoms, for example lauric acid, tridecylic acid, myristic acid, pentadecylic acid, palmitic acid, margaric acid, stearic acid, arachidic acid, behenic acid or corresponding unsaturated acids, for example oleic acid, elaidic acid, erucic acid, brasidic acid or linoleic acid, if desired with the addition of antioxidants, for example vitamin E, β -carotene or 3,5-di-*tert*-butyl-4-hydroxytoluene. The alcohol component of

- those fatty acid esters has a maximum of 6 carbon atoms and is mono- or poly-hydroxy, for example a mono-, di- or trihydroxy, alcohol, for example methanol, ethanol, propanol, butanol, or pentanol or the isomers thereof, but especially glycol and glycerol. The following examples of fatty acid esters are therefore to be mentioned: ethyl oleate, isopropyl
- 5 myristate, isopropyl palmitate, "Labrafil M2375" (polyoxyethylene glycerol trioleate, Gattefossé, Paris), "Miglyol 812" (triglyceride of saturated fatty acids with chain lengths of C8 to C12, Hüls AG, Germany), but especially vegetable oils, such as cottonseed oil, almond oil, olive oil, castor oil, sesame oil, soybean oil and more especially groundnut oil.
- 10 The injection or infusion compositions are prepared in customary manner under sterile conditions; the same applies also to introducing the compositions into ampoules or vials and sealing the containers.

Pharmaceutical compositions for oral administration can be obtained by combining the

15 active ingredient with solid carriers, if desired granulating a resulting mixture, and processing the mixture, if desired or necessary, after the addition of appropriate excipients, into tablets, dragée cores or capsules. It is also possible for them to be incorporated into plastics carriers that allow the active ingredients to diffuse or be released in measured amounts.

- 20 Suitable carriers are especially fillers, such as sugars, for example lactose, saccharose, mannitol or sorbitol, cellulose preparations and/or calcium phosphates, for example tricalcium phosphate or calcium hydrogen phosphate, and binders, such as starch pastes using for example corn, wheat, rice, or potato starch, gelatin, tragacanth, methylcellulose,
- 25 hydroxypropylmethylcellulose, sodium carboxymethylcellulose and/or polyvinylpyrrolidone, and/or, if desired, disintegrators, such as the above-mentioned starches, and/or carboxymethyl starch, crosslinked polyvinylpyrrolidone, agar, alginic acid or a salt thereof, such as sodium alginate. Excipients are especially flow conditioners and lubricants, for example silicic acid, talc, stearic acid or salts thereof, such as magnesium or
- 30 calcium stearate, and/or polyethylene glycol. Dragée cores are provided with suitable, optionally enteric, coatings, there being used, inter alia, concentrated sugar solutions which may comprise gum Arabic, talc, polyvinylpyrrolidone, polyethylene glycol, and/or titanium dioxide, or coating solutions in suitable organic solvents, or, for the preparation of enteric

coatings, solutions of suitable cellulose preparations, such as ethylcellulose phthalate or hydroxypropylmethylcellulose phthalate. Capsules are dry-filled capsules made of gelatin and of soft sealed capsules made of gelatine and a plasticiser, such as glycerol or sorbitol. The dry-filled capsules may comprise the active ingredient in the form of granules, for
5 example with fillers, such as lactose, binders, such as starches, and/or glidants, such as talc or magnesium stearate, and if desired with stabilizers. In soft capsules the active ingredient is preferably dissolved or suspended in suitable oil excipients, such as fatty oils, paraffin oil or liquid polyethylene glycols, it being possible also for stabilizers and/or antibacterial
10 agents to be added. Dyes or pigments may be added to the tablets or dragée coatings or the capsule casings, for example for identification purposes or to indicate different doses of active ingredient.

For parenteral administration, aqueous solutions of an active ingredient in water-soluble form, for example of a water-soluble salt, or aqueous injection suspensions that contain
15 viscosity-increasing substances and stabilizers, are especially suitable. The active ingredient, optionally together with excipients, can also be in the form of a lyophilizate and be made into a solution before parenteral administration by the addition of solvents.

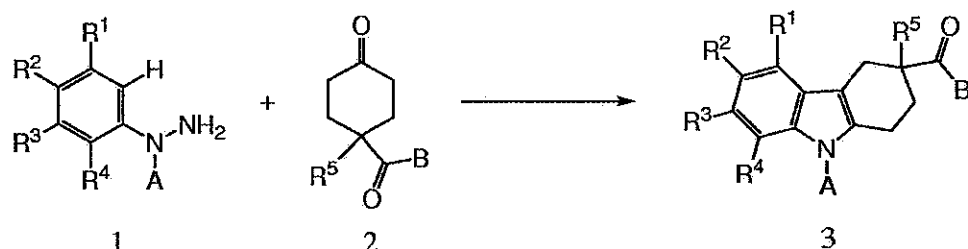
A further object of the invention is a process for preparing 2,3,4,9-tetrahydro-1*H*-carbazole
20 compounds of Formula I. Compounds according to Formula I of the present invention can be prepared according to the general sequence of reactions outlined in Scheme 1 below. The compounds obtained may also be converted into a pharmaceutically acceptable salt thereof in a manner known *per se*.

25 Compounds of the invention may be manufactured by the application or adaptation of known methods, by which is meant methods used heretofore or described in the literature, for example those described by Larock R. C. in "*Comprehensive organic transformations: a guide to functional group preparations*", VCH publishers, 1999.

In the reactions described hereinafter, it may be necessary to protect reactive functional
30 groups, for example hydroxy, amino, imino, thio or carboxy groups, where these are desired in the final product, to avoid their unwanted participation in the reactions. Conventional protecting groups may be used in accordance with standard practice, for example see

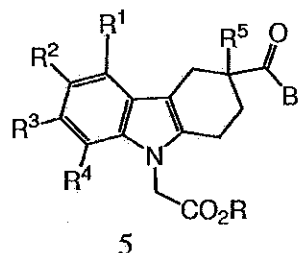
Greene T. W. and Wuts P. G. M. in "*Protective groups in organic synthesis*" Wiley-Interscience (1999).

Generally, the key step to build up a 2,3,4,9-tetrahydro-1*H*-carbazole compound of Formula 3 comprises a Fischer indole reaction known to a skilled person (e.g. M. H. Block *et al.*, *J. Med. Chem.* 2002, 45, 3509-3523). In such a reaction, a phenylhydrazine of Formula 1, wherein R¹, R², R³, and R⁴ are as defined for Formula I, and A represents hydrogen, CH₂COOH or CH₂COOalkyl, is condensed with a cyclohexanone derivative of Formula 2, wherein B represents hydroxy or alkoxy, and R⁵ is as defined for Formula I, in the presence of an acid, such as hydrochloric acid, in a solvent such as ethanol, etc., as outlined in Scheme 1.

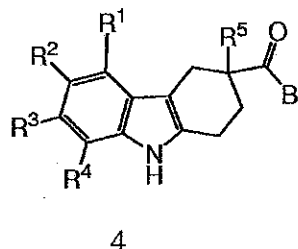


Scheme 1

A compound of Formula 5



is obtained in an alkylation reaction from an intermediate of Formula 4



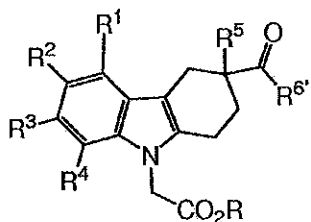
with an alkylating reagent of Formula L-CH₂CO₂R, wherein R represents an alkyl group, preferably ethyl or *tert.*-butyl, and L is a leaving group, in the presence of a base, such as cesium carbonate, sodium hydride, potassium *tert.*-butoxide or the like, in a suitable solvent,

such as acetone, tetrahydrofuran, dioxane or *N,N*-dimethylformamide.

Suitable leaving groups L are such as halo, in particular bromo or chloro, mesyloxy or tosyloxy. Preferably, the compound of Formula L-CH₂CO₂R is ethyl bromo-acetate.

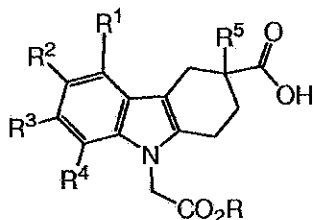
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A precursor of Formula 6



6

wherein R^{6'} represents -NR⁷R⁸ as defined hereinabove for Formula I, and R represents alkyl, is obtained by condensing an intermediate of Formula 7

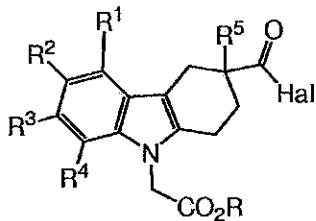


7

10

with a primary or secondary amine of Formula HNR⁷R⁸ by means of standard coupling reagents such as 1,3-dicyclohexylcarbodiimide, 1,3-diisopropylcarbodiimide, *O*-(7-azabenzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate and the like, in the presence of a base; or

15 *via* the corresponding acid halide of Formula 8



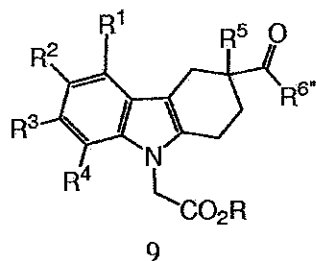
8

wherein Hal represents halogen, preferably chloro or bromo, in the presence of a base. The acid halide is obtained by reacting the corresponding acid with a halogenating reagent under conditions known to a skilled person, preferably by means of oxalyl chloride in the presence of a catalytic amount of *N,N*-dimethylformamide, phosphorous oxychloride or

20

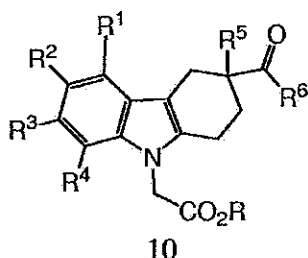
bis(trichloromethyl)carbonate.

A precursor of Formula 9



- 5 wherein $R^{6''}$ represents alkoxy or aryl-alkoxy, and R represents alkyl, is obtained by condensing the corresponding carboxylic acid with a reagent of Formula $X-R^{6''}$, wherein X is a leaving group such as halogen, in particular chloro or bromo; or alkyl- or arylsulfonate, such as mesylate or tosylate; in the presence of a base.

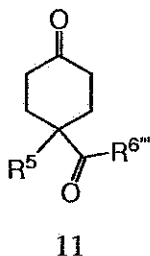
- 10 Hydrolysis of the CO_2R group in Formula 10



is carried out using routine procedures, for example by treatment with aqueous sodium hydroxide or lithium hydroxide; or trifluoroacetic acid or hydrochloric acid, to give a compound of Formula I.

15

A precursor of Formula 11



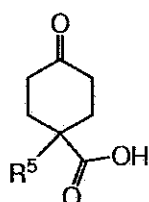
- wherein $R^{6'''}$ represents $-NR^7R^8$ and R^7 and R^8 are as defined hereinabove for Formula I, is obtained by condensing the corresponding 4-oxo-cyclohexanecarboxylic acid with a
 20 primary or secondary amine of Formula HNR^7R^8 by means of a standard coupling reagent such as 1,3-dicyclohexylcarbodiimide, 1,3-diisopropylcarbodiimide, *O*-(7-azabenzotriazol-

1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate and the like, in the presence of a base; or

via the corresponding 4-oxo-cyclohexanecarbonyl halide, preferably chloride or bromide, in the presence of a base. The acid halide is obtained by reacting the corresponding acid with a halogenating reagent under conditions known to a skilled person, preferably by means of oxalyl chloride, or phosphorous oxychloride, or bis(trichloromethyl)carbonate.

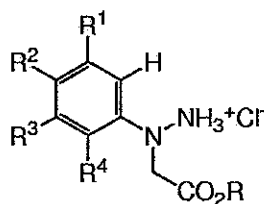
A precursor of above Formula 11, wherein $R^{6''}$ represents alkoxy, or aryl-alkoxy, and R represents alkyl, is obtained by condensing the corresponding carboxylic acid with a reagent of Formula $X-R^{6''}$, wherein X is a leaving group such as halogen, in particular chloro or bromo; or alkyl- or arylsulfonate, such as mesylate or tosylate, respectively, in the presence of a base.

4-Oxo-cyclohexanecarboxylic acid of Formula 12



wherein R^5 is alkyl (other than hydrogen), is easily accessed by means of known methods (e.g. *J. Am. Chem. Soc.* 1946, 68, 338-340).

A phenyl hydrazine of Formula 13



is prepared in three steps according to known methods starting from the corresponding aniline:

(i) alkylation with an alkylating reagent of Formula $L-CH_2CO_2R$, wherein R represents an alkyl group, preferably ethyl or *tert.*-butyl, and L is a leaving group as defined above in the reaction with a compound of Formula 4, in the presence of a base, such as ethyl-diisopropyl-

amine, cesium carbonate, sodium hydride, potassium *tert*.-butoxide or the like, in a suitable solvent, such as acetone, acetonitrile, tetrahydrofuran, dioxane or *N,N*-dimethylformamide. Suitable L is a leaving group such as halo, in particular bromo or chloro; or mesyloxy or tosyloxy (preferably, the compound of Formula L-CH₂CO₂R is ethyl bromo-acetate);

- 5 (ii) nitrosation by means of sodium nitrite, or amyl nitrite, in hydrochloric acid; (iii) and subsequent reduction with zinc metal in acetic acid.

Suitable are organic solvents, which are inert under the chosen reaction conditions.

- Preferred solvents are ethers, such as diethyl ether, diisopropyl ether, 2-methoxy-2-methyl-
10 propane, [1,4]dioxane, tetrahydrofuran or 1,2-dimethoxy-ethane; or alcohols, such as methanol, ethanol, propanol, isopropanol, butanol, isobutanol or *tert*-butanol; or hydrocarbons, such as benzene, toluene, xylene, hexane, cyclohexane or petroleum derived hydrocarbons, or halogenated hydrocarbons, such as dichloromethane, trichloro-methane, tetrachloromethane, dichloroethane, trichloroethane or chlorobenzene; or esters, such as
15 ethyl acetate; or amines, such as triethylamine, 4-methyl-morpholine, pyridine, and the like; as well as dimethylsulfoxide, *N,N*-dimethyl-formamide, *N*-methyl-pyrrolidone, *N,N*-dimethyl-acetamide, hexamethyl-phosphoric triamide, acetonitrile, acetone or nitromethane. Likewise, mixtures of above-mentioned solvents can be employed.

- 20 Suitable bases used in the processes described herein can be of inorganic or organic nature. Preferred are alkali metal hydroxide, for example lithium, sodium or potassium hydroxide, earth alkali metal hydroxide, for example barium hydroxide, alkali metal carbonate such as sodium or potassium carbonate, earth alkali metal carbonate, such as calcium carbonate, or alkali metal or earth alkali metal alkoxide, such as sodium or potassium methoxide,
25 ethoxide, or *tert*-butoxide; or organic amines, for example trialkyl-(C₁-C₆)-amine, such as triethylamine, ethyl-diisopropyl-amine, or heterocyclic amines, such as 1,4-diazabicyclo[2.2.2]octane (DABCO), 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU), pyridine, 2,6-lutidine, 4-dimethylamino-pyridine, 1-methyl-piperidine or 4-methyl-morpholine. Furthermore, alkali metals, such as sodium, or hydrides thereof, such as sodium hydride,
30 can be employed. The above-mentioned bases can be used as antacid agents.

Suitable coupling reagents act as dehydrating reagents, for example carbodiimides, such as diisopropylcarbodiimide, dicyclohexylcarbodiimide, or *N*-(3-dimethylaminopropyl)-*N'*-

ethylcarbodiimide hydrochloric acid salt; or carbonyl compounds, like carbonyldiimidazole, or 1,2-oxazolium compounds, like 2-ethyl-5-phenyl-isoxazolium-3-sulfonate, as well as 2,4,6-tripropyl-1,3,5,2,4,6-trioxatriphosphorinane-2,4,6-trioxide (PPACA; propanephosphonic acid cyclic anhydride), or alkyl chloroformates, such as iso-butyl
 5 chloroformate; or (benzotriazol-1-yloxy)-tris-(dimethylamino)-phosphonium-hexafluorophosphate (BOP), *O*-(benzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium-tetrafluoroborat (TBTU), *N',N',N',N'*-tetramethyl-*O*-(7-azabenzotriazole-1-yl)-uronium-hexafluorophosphate (HATU) and the like; or bis(trichloromethyl)carbonate, or diphenylphosphoramidate, or methanesulfonyl chloride; if indicated in the presence of a
 10 base such as triethylamine, or 4-ethyl-morpholine, or 4-methyl-piperidine, or ethyl-diisopropyl-amine.

Particular embodiments of the invention are described in the following Examples, which serve to illustrate the invention in more detail without limiting its scope in any way.

15

Experimental part:

Abbreviations (as used herein):

	AcOH	acetic acid
20	BSA	Bovine Serum Albumine
	calcd	calculated
	CH ₂ Cl ₂	dichloromethane
	CDCl ₃	deuterated chloroform
	conc.	concentrated
25	DIEA	ethyl-diisopropyl-amine
	DMAP	4- <i>N,N</i> -Dimethylaminopyridine
	DMF	<i>N,N</i> -dimethylformamide
	DMSO	dimethylsulfoxide
	EDTA	ethylenediaminetetraacetic acid
30	ESI-MS	electrospray ionization - mass spectrometry
	Et	ethyl
	EtOAc	ethyl acetate
	EtOH	ethanol

Ex.	Example
FLIPR	Fluorescent Imaging Plate Reader
g	gram(s)
h	hour(s)
5 HBSS	Hank's Balanced Salt Solution
HEPES	4-(2-hydroxyethyl)-piperazine-1-ethanesulfonic acid
H ₂ O	water
HCl	hydrochloric acid
HPLC	High-Performance Liquid Chromatography
10 H ₂ SO ₄	sulfuric acid
k	kilo
K ₂ CO ₃	potassium carbonate
KH ₂ PO ₄	potassium dihydrogenphosphate
KOtBu	potassium <i>tert</i> -butoxide
15 l	liter(s)
LC	liquid chromatography
μ	micro
m	milli
M	molar
20 MeOH	methanol
min	minute(s)
mol	mole(s)
MS	Mass Spectroscopy
MW	molecular weight
25 N	normality of solution
NaH	sodium hydride
NaCl	sodium chloride
NaHCO ₃	sodium hydrogencarbonate
Na ₂ CO ₃	sodium carbonate
30 NaNO ₂	sodium nitrite
NaOAc	sodium acetate
NaOH	sodium hydroxide
Na ₂ SO ₄	sodium sulfate

t_R	retention time
rt	room temperature
THF	tetrahydrofuran
TBME	<i>tert</i> -butylmethylether
5 TFA	trifluoroacetic acid

Chemistry

General remarks:

10 All solvents and reagents are used as obtained from commercial sources unless otherwise indicated. The starting materials are obtained from commercial sources or synthesized using standard literature procedures.

Temperatures are indicated in degrees Celsius (°C). Unless otherwise indicated, the reactions take place at room temperature (rt).

15 In mixtures, relations of parts of solvent or eluent or reagent mixtures in liquid form are given as volume relations (v/v), unless indicated otherwise.

Preparative chiral separations are performed on a HPLC instrument, equipped with a Dinox P580 binary pump and a Photodiode Array Detector Dionex PDA-100, using a Daicel Chiralcel OD column (20 x 250 mm, 10 µm) under isocratic elution conditions (% B) employing EtOH (eluent A) and hexane (eluent B), run time 30 min; flow rate 10 ml/min, 20 detection at 220 nm. The compounds are dissolved in EtOH before loading.

HPLC/MS analyses (LC-1 to LC-4) are performed on a Waters 2795 Alliance HPLC instrument, equipped with a Waters 996 Photodiode Array Detector and a Micromass ZQTM Waters mass spectrometer (electron spray ionization), detection at 200-400 nm, or on a 25 Agilent 1100 system, equipped with a Dionex P580 binary pump, a Dionex PDA-100 Photodiode Array Detector and a Finnigan AQA mass spectrometer (LC-5).

- LC-1: Analytical HPLC on a Waters XterraTM MS C₁₈ column (2.1 x 50mm, 5 µm). Linear gradient of water/0.06% formic acid (A) and acetonitrile/0.06% formic acid (B) from 5% to 95% B over 2 min; flow rate 0.75 ml/min.

30 - LC-2: Analytical HPLC on a Waters XterraTM MS C₁₈ column (4.6 x 50mm, 5 µm). Linear gradient of water/0.06% formic acid (A) and acetonitrile/0.06% formic acid (B) from 5% to 95% B over 1 min; flow rate 3 ml/min.

- LC-3: Analytical HPLC on a GromSil MS C₁₈ column (2.1 x 50 mm, 5 µm, Waters).
Linear gradient of water/0.06% formic acid (A) and acetonitrile/0.06% formic acid (B) from 5% to 95% B over 6 min; flow rate 0.25 ml/min.
- LC-4: Analytical HPLC on a Zorbax SB-AQ column (4.6 x 50 mm, 5 µm): Linear gradient
5 of water/0.06% formic acid (A) and acetonitrile/0.06% formic acid (B) from 5% to 95% B
over 1 min; flow rate 3 ml/min.
- LC-5: Analytical HPLC on a Zorbax SB-AQ column (50 x 4.6 mm, 5 µm): Linear gradient
of water/0.04% trifluoroacetic acid (A) and acetonitrile (B) from 5% to 95% B over 1 min;
flow rate 4.5 ml/min, detection at 210, 220, 230, 254 and 280 nm.
- 10 - LC-6: Analytical chiral HPLC performed on a HPLC instrument, equipped with a Dinoex
P580 binary pump, a Dionex PDA-100 Photodiode Array Detector and a Jasco OR-1590
Chiral Detector, using a Daicel Chiralcel OD column (4.6 x 250 mm, 10 µm) under isocratic
elution conditions (75% B) employing EtOH/0.1% TFA (eluent A) and hexane (eluent B),
run time 30 min; flow rate 0.8 ml/min, detection at 210 nm.
- 15 - LC-7: Analytical chiral HPLC performed on a HPLC instrument, equipped with a Dinoex
P580 binary pump, a Dionex PDA-100 Photodiode Array Detector and a Jasco OR-1590
Chiral Detector, using a Daicel ChiralPak IA column (4.6 x 250 mm, 5 µm) under isocratic
elution conditions (95%B) employing EtOH/0.1% TFA (eluent A) and hexane (eluent B),
run time 40 min; flow rate 1.0 ml/min, detection at 220 nm.
- 20 - LC-8: Analytical chiral HPLC performed on a HPLC instrument, equipped with a Dinoex
P580 binary pump, a Photodiode Array Detector Dionex PDA-100, a Jasco OR-1590 chiral
detector and a polarimeter, using a Daicel ChiralPak IA column (4.6 x 250 mm, 5 mm)
under isocratic elution conditions (70% B) employing EtOH + 1% TFA (eluent A) and
hexane (eluent B), run time 20 min; flow rate 0.8 ml/min, detection at 210 nm.
- 25 - LC-9: Analytical chiral HPLC performed on a HPLC instrument, equipped with a Dinoex
P580 binary pump, a Photodiode Array Detector Dionex PDA-100 and a Jasco OR-1590
chiral detector and a polarimeter, using a Daicel ChiralPak IA column (4.6 x 250 mm,
5 mm) under isocratic elution conditions (80% B) employing EtOH + 1% TFA (eluent A)
and hexane (eluent B), run time 20 min; flow rate 0.8 ml/min, detection at 210 nm.

Optical rotation of chiral compounds are measured on a Jasco P-1030 polarimeter in ethanolic solution at 20°C.

¹H NMR spectra are recorded on a Varian Mercury 300VX FT-NMR spectrometer.

- 5 Chemical shifts (δ) are reported in parts per million (ppm) relative to proton resonances resulting from incomplete deuteration of the NMR solvent, *e.g.* for dimethylsulfoxide δ (H) 2.49 ppm, for chloroform δ (H) 7.24 ppm.

Intermediate 1.1:

- 10 9-Ethoxycarbonylmethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid

Method A

a) 2,3,4,9-Tetrahydro-1*H*-carbazole-3-carboxylic acid

- A mixture of 4-oxo-cyclohexanecarboxylic acid (4.05 g, 28.5 mmol) and phenylhydrazine
15 hydrochloride (4.12 g, 28.5 mmol) in EtOH (200 ml) is stirred at reflux for 3 h. Then, the solvent is evaporated and the residue is dissolved in boiling toluene. Precipitating ammonium chloride is filtered off and the clear filtrate is cooled to rt. The resulting precipitate is filtered and dried under high vacuum to give pure subtitle compound as a white solid (5.16 g) in 84% yield. t_R (LC-3) 2.21 min; ESI-MS (positive ion): m/z 216.30 $[M+H]^+$
20 (calcd 215.09 for C₁₃H₁₃NO₂). ¹H-NMR (CDCl₃): 2.01 (m, 1H); 2.28 (m, 1H); 2.83 (m, 4H); 3.04 (dd, $J=14.8$ Hz, 4.7 Hz, 1H); 7.02 (m, 2H); 7.19 (d, $J=7.2$ Hz, 1H); 7.38 (d, $J=7.0$ Hz, 1H); 7.63 (s br, 1H).

4-Oxo-cyclohexanecarboxylic acid

- 25 Following the method described by D. Wustrow *et al.* (*J. Med. Chem.* 1998, 41, 760-771), subtitle compound is obtained from ethyl 4-oxo-cyclohexanecarboxylate as a white solid in 74% yield. ¹H-NMR (CDCl₃): 1.72 (m, 2H); 2.02 (m, 2H); 2.17 (m, 2H); 2.32 (m, 2H); 2.64 (m, 1H); 12.2 (s, 1H).

- 30 b) 9-Ethoxycarbonylmethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid

To a stirred solution of 2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid (0.40 g, 1.86 mmol) in dry DMF (4 ml) is added NaH (0.16 g, 4.1 mmol, 60% in mineral oil) in one portion. After gas evolution has ceased, the reaction mixture is kept stirring at rt for 1 h.

Then, ethyl bromoacetate (0.21 ml, 1.9 mmol) is added and stirring is continued overnight. Saturated KH_2PO_4 solution is added and the mixture is extracted with dichloromethane. The organic layers are combined and washed with H_2O and brine, dried over Na_2SO_4 , filtered and concentrated. The residue is purified by flash chromatography on silica gel (hexane/
5 EtOAc 3:1) to afford pure title compound as an off-white solid (0.34 mg) in 61% yield. t_{R} (LC-3) 2.05 min; ESI-MS (positive ion): m/z 324.15 $[\text{M}+\text{Na}]^+$ (calcd 301.13 for $\text{C}_{17}\text{H}_{19}\text{NO}_4$). ^1H -NMR ($\text{DMSO}-d_6$): 1.13 (t, $J=7.0$ Hz, 3H); 1.79 (m, 1H); 2.14 (m, 1H); 2.64 (m, 4H); 2.86 (dd, $J=14.3$ Hz, 4.8 Hz, 1H); 4.05 (q, $J=7.0$ Hz, 2H); 4.88 (s, 2H); 6.90 (dd, $J=7.6$ Hz, 7.0 Hz, 1H); 6.97 (dd, $J=7.6$ Hz, 7.0 Hz, 1H); 7.23 (d, $J=8.1$ Hz, 1H); 7.3 (d, $J=6.8$ Hz, 1H); 12.08 (s br, 1H).

Method B

A mixture of ethyl (*N*-phenyl-hydrazino)-acetate hydrochloric acid salt (2.3 g, 10 mmol) and 4-oxo-cyclohexanecarboxylic acid (1.42 g, 10 mmol) in glacial AcOH (20 ml) is stirred
15 at 70 °C for 1 h. The suspension is allowed to cool to rt, poured onto H_2O and stirred for 1 h. The resulting precipitate is filtered, washed with H_2O and dried under high vacuum to give pure 9-ethoxycarbonylmethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid as a white solid (2.85 g) in 93% yield.

20 Intermediate 1.2:

9-Ethoxycarbonylmethyl-6-fluoro-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid
Intermediate 1.2 is prepared analogous to the procedures described for Intermediate 1.1, using (4-fluoro-phenyl)-hydrazine in place of phenylhydrazine (Method A), or ethyl [*N*-(4-fluoro-phenyl)-hydrazino]-acetate hydrochloric acid salt in place of ethyl (*N*-phenyl-
25 hydrazino)-acetate hydrochloric acid salt (Method B).

Racemic 9-ethoxycarbonylmethyl-6-fluoro-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid is resolved into enantiomerically pure Intermediate 1.2.1 and 1.2.2, respectively, by applying chiral crystallization techniques by means of a chiral base forming the respective diastereomeric salts.

30

Intermediate 1.2.1:

(+)-9-Ethoxycarbonylmethyl-6-fluoro-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid

Racemic 9-ethoxycarbonylmethyl-6-fluoro-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid (15.0 g, 47.0 mmol) in EtOAc (250 ml) is heated to reflux and treated with *S*(-)- α -methylbenzylamine (2.99 ml, 23.5 mmol). The clear solution is cooled to rt and the crystals formed are collected by filtration. Recrystallization is repeated four times from EtOAc (2x 250 ml, 200 ml and 175 ml). The final crystal fraction is dissolved in EtOAc, extracted with 1N HCl, and washed with water. The organic layer is dried over Na₂SO₄ and the solvent evaporated to deliver title compound as a beige crystalline solid with an enantiomeric excess (ee) >96%. *t_R* (LC-7) 23.65 min; $\alpha_D = +33.5232^\circ$, $\pm 0.0215^\circ$ (20 °C in ethanol).

10 Intermediate 1.2.2:

(-)-9-Ethoxycarbonylmethyl-6-fluoro-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid

The title compound is prepared analogous to the crystallization procedure described for Intermediate 1.2.1, using *R*(+)- α -methylbenzylamine, with ee>96%: *t_R* (LC-7) 21.33 min.

15

Intermediate 1.3:

9-Ethoxycarbonylmethyl-8-chloro-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid

Intermediate 1.3 can be prepared analogous to the procedures described for Intermediate 1.1, using (2-chloro-phenyl)-hydrazine in place of phenylhydrazine (Method A), or ethyl [N-(2-chloro-phenyl)-hydrazino]-acetate hydrochloric acid salt in place of ethyl (N-phenyl-hydrazino)-acetate hydrochloric acid salt (Method B), or with Method C (described below).

Method C

a) 8-Chloro-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid

25 A mixture of 4-oxo-cyclohexanecarboxylic acid (3.1 g, 21.7 mmol) and 2-chlorophenylhydrazine hydrochloride (4.0 g, 21.7 mmol) in glacial acetic acid (740 ml) is stirred at reflux for 3 h. The solvent is evaporated and water is added to the residue. The resulting precipitate is filtered off and dried under high vacuum to give the subtitle compound as a brownish solid (4.5 g) in 83% yield. *t_R* (LC-5) 0.88 min; ESI-MS (positive ion): *m/z* 250.14 [M+H]⁺ (calcd 249.70 for C₁₃H₁₂NO₂Cl). ¹H-NMR (DMSO-*d*₆): 1.75 (m, 1H); 2.14 (m, 1H); 2.62 (m, 5H); 6.90 (t, *J*=7.9 Hz, 1H); 7.02 (d, *J*=7.3 Hz, 1H); 7.30 (d, *J*=7.6 Hz, 1H); 7.92 (br s, 1H); 10.9 (br s, 1H).

30

b) 8-Chloro-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid benzyl ester

A stirred mixture of crude 8-chloro-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid (4.45 g, 17.8 mmol) and cesium carbonate (2.90 g, 8.92 mmol) in ethanol (150 ml) and water (25 ml) is refluxed until a clear solution is obtained (15 min). Then the solvents are
5 evaporated and the dried residue is dissolved in dry DMF. This solution is treated with benzylbromide (2.27 ml, 18.7 mmol) and stirred at 45 °C for 15 min. The resulting precipitate is filtered off and the filtrate is concentrated. The residue is purified by flash chromatography on silica gel (heptane/ EtOAc 3:1) to deliver pure title compound as a yellowish solid (4.2 g) in 69% yield. t_R (LC-5) 1.09 min; ESI-MS (positive ion): m/z 340.19
10 $[M+H]^+$ (calcd 339.82 for $C_{20}H_{18}NO_2Cl$). 1H -NMR (DMSO- d_6): 1.90 (m, 1H); 2.19 (m, 1H); 2.86 (m, 5H); 5.13 (s, 2H); 6.91 (t, $J=7.6$ Hz, 1H); 7.04 (d, $J=7.6$ Hz, 1H); 7.32 (m, 6H); 11.0 (br s, 1H).

c) 8-Chloro-9-ethoxycarbonylmethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid benzyl ester

Cesium carbonate (6.32 g, 19.2 mmol) and ethyl bromoacetate (1.16 ml, 10.2 mmol) are added to a stirred solution of 8-chloro-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid benzyl ester (2.17 g, 6.4 mmol) in dry DMF (20 ml). The reaction mixture is stirred at 60 °C for 3 h. The precipitate is filtered off and rinsed thoroughly with acetone. The filtrate is
20 concentrated and the residue is purified by flash chromatography on silica gel (heptane/ EtOAc 4:1) to afford pure title compound as a yellow solid (2.54 mg) in 93% yield. t_R (LC-5) 1.14 min; ESI-MS (positive ion): m/z 426.16 $[M+H]^+$ (calcd 425.91 for $C_{24}H_{24}NO_4Cl$). 1H -NMR (CDCl₃): 1.25 (t, $J=7.0$ Hz, 3H); 2.07 (m, 1H); 2.37 (m, 1H); 2.69 (m, 2H); 2.89 (m, 2H); 3.07 (dd, $J=14.4, 4.4$ Hz, 1H); 4.22 (q, $J=7.3$ Hz, 2H); 5.07 (d, $J=18.2$ Hz, 1H);
25 5.18 (s, 2H); 5.19 (d, $J=18.5$ Hz, 1H); 6.97 (t, $J=7.9$ Hz, 1H); 7.08 (dd, $J=7.6, 0.9$ Hz, 1H); 7.34 (m, 6H).

d) 9-Ethoxycarbonylmethyl-8-chloro-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid

A mixture of 8-chloro-9-ethoxycarbonylmethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid benzyl ester (2.5 g, 5.9 mmol) and 10% palladium on activated charcoal (480 mg) in ethanol (200 ml) is stirred under a hydrogen atmosphere for 1 h. The catalyst is filtered off and washed with ethanol. The filtrate is concentrated to dryness to deliver Intermediate 1.3. t_R (LC-5) 0.96 min; ESI-MS (positive ion): m/z 336.23 $[M+H]^+$ (calcd

335.79 for $C_{17}H_{18}NO_4Cl$). 1H -NMR (DMSO- d_6): 1.18 (t, $J=7.0$ Hz, 3H); 1.85 (m, 1H); 2.19 (m, 1H); 2.66 (m, 4H); 2.90 (m, 1H); 4.14 (q, $J=7.0$ Hz, 2H); 5.17 (s, 2H); 6.96 (t, $J=7.6$ Hz, 1H); 7.04 (d, $J=7.3$ Hz, 1H); 7.38 (d, $J=7.6$ Hz, 1H); 12.3 (br s, 1H).

5 Intermediate 2.1:

9-Ethoxycarbonylmethyl-3-methyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid

Intermediate 2.1 is prepared analogous to the procedures described for Intermediate 1.1, using 1-methyl-4-oxo-cyclohexanecarboxylic acid in place of 4-oxo-cyclohexanecarboxylic acid.

10

1-Methyl-4-oxo-cyclohexanecarboxylic acid

Applying reported methods (*e.g.* M. Rubin, H. Wishinsky, *J. Amer. Chem. Soc.* 1946, 68, 338-340; N. B. Chapman, S. Sotheeswaran, K. J. Toyne, *J. Org. Chem.* 1970, 35, 917-923; H. A. Bruson, T. W. Riener, *J. Amer. Chem. Soc.* 1942, 64, 2850-2858) provided subtitle

15 compound as beige waxy solid.

1H -NMR ($CDCl_3$): 1.38 (s, 3H); 1.73 (m, 2H); 2.43 (m, 6H).

Intermediate 2.2:

9-Ethoxycarbonylmethyl-6-fluoro-3-methyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-

20 carboxylic acid

Intermediate 2.2 is prepared analogous to the procedures described for Intermediate 1.2, using 1-methyl-4-oxo-cyclohexanecarboxylic acid in place of and 4-oxo-cyclohexanecarboxylic acid.

25 Intermediate 3.1:

9-Ethoxycarbonylmethyl-3-ethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid

Intermediate 3.1 is prepared analogous to the procedures described for Intermediate 1.1, using 1-ethyl-4-oxo-cyclohexanecarboxylic acid in place of 4-oxo-cyclohexanecarboxylic acid.

30

1-Ethyl-4-oxo-cyclohexanecarboxylic acid

Applying reported methods (*e.g.* M. Rubin, H. Wishinsky, *J. Amer. Chem. Soc.* 1946, 68, 338-340; N. B. Chapman, S. Sotheeswaran, K. J. Toyne, *J. Org. Chem.* 1970, 35, 917-923;

H. A. Bruson, T. W. Riener, *J. Amer. Chem. Soc.* 1942, 64, 2850-2858) provided subtitle compound as viscous oil.

¹H-NMR (CDCl₃): 0.95 (t, *J*=7.2 Hz, 3H); 1.70 (m, 4H); 2.43 (m, 6H).

5 Intermediate 3.2:

9-Ethoxycarbonylmethyl-3-ethyl-6-fluoro-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid

Intermediate 3.2 is prepared analogous to the procedures described for Intermediate 1.2, using 1-ethyl-4-oxo-cyclohexanecarboxylic acid in place of 4-oxo-cyclohexanecarboxylic acid.

10

Intermediate 4.1:

9-Ethoxycarbonylmethyl-3-propyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid

Intermediate 4.1 is prepared analogous to the procedures described for Intermediate 1.1, using 4-oxo-1-propyl-cyclohexanecarboxylic acid in place of 4-oxo-cyclohexanecarboxylic acid.

15

4-Oxo-1-propyl-cyclohexanecarboxylic acid

Applying reported methods (*e.g.* M. Rubin, H. Wishinsky, *J. Amer. Chem. Soc.* 1946, 68, 338-340; N. B. Chapman, S. Sotheeswaran, K. J. Toyne, *J. Org. Chem.* 1970, 35, 917-923; H. A. Bruson, T. W. Riener, *J. Amer. Chem. Soc.* 1942, 64, 2850-2858) provided subtitle compound as viscous oil.

¹H-NMR (CDCl₃): 0.93 (t, *J*=7.0 Hz, 3H); 1.35 (m, 2H); 1.67 (m, 4H); 2.43 (m, 6H).

25 Intermediate 4.2:

9-Ethoxycarbonylmethyl-6-fluoro-3-propyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid

Intermediate 4.2 is prepared analogous to the procedures described for Intermediate 1.2, using 4-oxo-1-propyl-cyclohexanecarboxylic acid in place of 4-oxo-cyclohexanecarboxylic acid.

30

Intermediate 5.1

3-cyano-2,3,4,9-tetrahydro-1*H*-carbazole

A mixture of 4-cyano-cyclohexanone (0.48 g, 3.9 mmol) and phenylhydrazine hydrochloride (0.42 g, 3.9 mmol) in glacial acetic acid (5 ml) is stirred at reflux for 1 h. The solvent is evaporated and the residue chromatographed on silica gel (heptane / EtOAc 2:1) to deliver the title compound as a brownish solid (158 mg) in 50% yield. t_R (LC-5) 0.91 min; ESI-MS (positive ion): m/z 197.10 $[M+H]^+$ (calcd 196.25 for $C_{13}H_{12}N_2$). 1H -NMR ($CDCl_3$): 2.25 (m, 2H); 2.96 (m, 5H); 7.14 (m, 2H); 7.30 (d, $J=7.7$ Hz, 1H); 7.45 (d, $J=7.1$ Hz, 1H); 7.79 (br s, 1H).

4-Cyano-cyclohexanone

10 Applying reported methods (e.g. M. Rubin, H. Wishinsky, *J. Amer. Chem. Soc.* 1946, 68, 338-340; N. B. Chapman, S. Sotheeswaran, K. J. Toyne, *J. Org. Chem.* 1970, 35, 917-923; H. A. Bruson, T. W. Riener, *J. Amer. Chem. Soc.* 1942, 64, 2850-2858) provided subtitle compound as slightly yellow oil.

15 Ethyl (*N*-phenyl-hydrazino)-acetate hydrochloric acid salt

a) Ethyl phenylamino-acetate

According to Kotake *et al.* (*Chem. Pharm. Bull.* 1995, 43, 829-841), a stirred solution of aniline (15.3 g, 0.16 mol) and DIEA (56 ml, 0.33 mol) in acetonitrile (120 ml) is heated to 60 °C and ethyl bromoacetate (18.2 ml, 0.16 mmol) is added dropwise within 2 h. The reaction mixture is kept stirring at this temperature for another 3 h and is then evaporated to dryness. Water is added to the residue, the solid is filtered and thoroughly rinsed with H_2O . Crystallization from toluene provided pure subtitle compound as beige crystals (25.5 g) in 87% yield. t_R (LC-3) 2.33 min; ESI-MS (positive ion): m/z 180.27 $[M+H]^+$ (calcd 179.22 for $C_{10}H_{13}NO_2$). 1H -NMR ($CDCl_3$): 1.29 (t, $J=7.0$ Hz, 3H); 3.90 (s, 2H); 4.25 (m, 3H); 6.61 (d, $J=8.2$ Hz, 2H); 6.75 (t, $J=7.1$ Hz, 1H); 7.21 (m, 2H).

b) Ethyl *N*-nitroso-phenylamino-acetate

30 A solution of $NaNO_2$ (9.5 g, 0.14 mol) in H_2O (27 ml) is added to an ice-cold stirred solution of ethyl phenylamino-acetate (25 g, 0.14 mol) in conc. HCl (20 ml) and H_2O (27 ml) within 2 h. The reaction mixture is kept stirring for 1 h and then extracted twice with CH_2Cl_2 . The combined organic layers are washed with H_2O and brine, dried over

Na₂SO₄ and filtered. The solvent is evaporated and the crude subtitle compound thus obtained as orange oil is used without further purification. *t_R* (LC-4) 1.02 min.

c) Ethyl (*N*-phenyl-hydrazino)-acetate hydrochloric acid salt

- 5 According to Arcari *et al.* (*Farmaco* 1992, 47, 405-425), crude ethyl *N*-nitroso-phenylamino-acetate (0.14 mol) is dissolved in glacial AcOH (50 ml) and added to an ice-cold stirred suspension of zinc (32 g, 0.49 mol) in H₂O (50 ml) within 1 h. The reaction mixture is kept stirring at rt for 2 h, then the resulting precipitate is filtered and thoroughly washed with MeOH. The filtrate is acidified to pH 1 by the addition of conc. HCl and
- 10 washed with TBME, then basified to pH 10 by the addition of aqueous NH₄OH and extracted with CH₂Cl₂. The combined organic layers are dried over Na₂SO₄ and evaporated to give desired hydrazine as an oil. The compound is converted in its hydrochloride salt by treatment of the free hydrazine in diethyl ether with HCl (2N in Et₂O) to give title compound (20 g) in 63% yield (over two steps). *t_R* (LC-3) 2.33 min; ESI-MS (positive ion):
- 15 *m/z* 180.27 [M+H]⁺ (calcd 179.22 for C₁₀H₁₃NO₂). ¹H-NMR (DMSO-*d*₆): 1.15 (t, *J*=7.0 Hz, 3H); 3.59 (br s, 1H); 4.11 (q, *J*=7.0 Hz, 2H); 4.65 (s, 2H); 7.00 (m, 1H); 7.07 (m, 2H); 7.33 (m, 2H); 10.6 (br s, 1H).

Preparation of amines of formula HNR⁷R⁸:

20

The following amines are prepared according to the references listed below.

N-Phenyl-phenethylamine

- a) M. Beller, C. Breindl, T. H. Riermeier, M. Eichberger, H. Trauthwein, *Angew. Chme. Int. Ed.* 1998, 37, 3389-3391; b) F. Y. Kwang, A. Klapars, S. L. Buchwald, *Org. Lett.* 2002, 4, 581-584; c) L. Guy, C. Schaeffer, *Heterocycles* 1998, 48, 171-174.
- 25

5,6,11,12-Tetrahydro-dibenzo[*b,f*]azocine

A. M. Monro, R. M. Quinton, T. I. Wrigley, *J. Med. Chem.* 1963, 6, 255-261.

30

6,11-Dihydro-5*H*-dibenzo[*b,e*]azepine

a) E. J. Warawa, B. M. Migler, C. J. Ohnmacht, A. L. Needles, G. C. Gatos, F. M. McLaren, C. L. Nelson, K. M. Kirkland, *J. Med. Chem.* 2001, 44, 372-389; b) L. H. Werner, S. Ricca, E. Mohacsi, A. Rossi, V. P. Arys, *J. Med. Chem.* 1965, 8, 74-80.

5 5,11-Dihydrodibenzo[*b,e*] [1,4]oxazepine

B. J. Margolis, J. J. Swidorski, B. N. Rogers, *J. Org. Chem.* 2003, 68, 644-647.

Diphenethylamine

W. S. Bryant, I. A. Guzei, A. L. Rheingold, J. S. Merola, H. W. Gibson, *J. Org. Chem.*
10 1998, 63, 7634-7639.

[2-(4-Fluoro-phenyl)-ethyl]-phenyl-amine

J. D. Albright, V. G. DeVries, E. E. Largis, T. G. Miner, M. F. Reich, S. A. Schaffer, R. G. Shepherd, J. Upešlacis, *J. Med. Chem.* 1983, 26, 1378-1393.

15 [2-(4-chloro-phenyl)-ethyl]-phenyl-amine, phenyl-(3-phenyl-propyl)-amine, phenyl-(2-thiophen-3-yl-ethyl)-amine and (2,2-diphenyl-ethyl)-phenyl-amine are prepared applying the same procedure.

20 Example A-01

(±)-(3-Dibenzylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid

a): Ethyl (±)-(3-dibenzylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetate

A solution of Intermediate 1.1 (30 mg, 0.1 mmol), *N,N,N',N'*-tetramethyl-*O*-(7-azabenzotriazole-1-yl)-uronium-hexafluorophosphate (38 mg, 0.1 mmol) and DIEA (51 µl, 0.3 mmol) in THF/ DMF 4:1 (1 ml) is stirred at rt for 10 min and then is treated with dibenzylamine (20 mg, 0.1 mmol). After further stirring overnight at rt, the reaction mixture is diluted with H₂O (3 ml) and extracted three times with diethyl ether. The combined organic layers are evaporated and the obtained crude subtitle compound is used for further
25 synthesis without purification. *t_R* (LC-3) 3.03 min; MS (positive ion) *m/z* 481.29 [M+H]⁺
30 (calcd 480.62 for C₃₁H₃₂N₂O₃).

b) (±)-(3-Dibenzylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid

A stirred solution of crude ethyl ($\pm$)-(3-dibenzylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetate (0.1 mmol) in THF (1 ml) is treated with 0.2 N aqueous NaOH (0.5 ml, 0.1 mmol) added at rt for 15 min. Then, the reaction mixture is extracted twice with diethyl ether (2 ml), acidified to pH 1 by addition of 1M HCl and extracted with CH₂Cl₂. The combined organic layers are dried (Na₂SO₄), filtered and evaporated. The resulting solid is recrystallized from diethyl ether to yield title compound as a pale yellow solid. t_R (LC-2) 2.74 min; MS (positive ion) m/z 453.26 [M+H]⁺ (calcd 452.55 for C₂₉H₂₈N₂O₃). ¹H-NMR (DMSO-*d*₆): 1.96 (m, 2H); 2.68 (m, 4H); 2.95 (m, 1H); 4.05 (m, 2H); 4.66 (m, 2H); 4.81 (s, 2H); 6.98 (m, 2H); 7.26 (m, 12H); 12.8 (s, 1H).

Example A-02

{3-[(4-Chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid

a) Ethyl ($\pm$)-{3-[(4-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetate and its enantiomers

At -10 °C, bis(trichloromethyl)carbonate (66 mg, 0.22 mmol) and collidine (0.29 ml, 2.2 mmol) are added to a stirred solution of Intermediate 1.1 (0.20 g, 0.66 mmol) in dry THF (3 ml). The resulting suspension is stirred at this temperature for 1 min and then treated with 4-chloro-*N*-methylanilin (94 mg, 0.66 mmol). After stirring for 30 min the reaction is complete, the mixture poured onto 1M HCl (50 ml) and extracted three times with diethyl ether. The combined organic layers are dried over Na₂SO₄ and evaporated to dryness to leave the crude subtitle compound (0.20 g) in 71% yield as a pale yellow glassy solid. This material is used for further synthesis without purification. t_R (LC-1) 1.24 min; MS (positive ion) m/z 446.96 [M+Na]⁺ (calcd 424.92 for C₂₄H₂₅N₂O₃Cl). Retention times of the two enantiomers are respectively 10.1 min and 13.1 min (LC-6, 55% B), the more polar enantiomer displaying a rotation of the plane of the polarized light in a clockwise direction, the less polar enantiomer displaying a rotation in an anticlockwise direction, respectively, resulting a positive (+) and a negative (-) signal of the polarimetric plot, as co-monitored with an interfaced polarimeter.

Resolution into the enantiomers is carried out on a preparative chiral HPLC column (Chiralcel OD, 20 x 250 mm, 10 μ m) under isocratic elution conditions (EtOH/ hexane 45:55), runtime 30 min; flow rate 10 ml/min, detection at 220 nm. The (+) enantiomer of

ethyl -{3-[(4-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetate is eluting after 15.0 min, the (–) enantiomer after 18.5 min. Both enantiomers are obtained in >99% ee, as verified with LC-6 (55% B).

- 5 b) (±)-{3-[(4-Chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid

The title compound is obtained using conditions for the hydrolysis of crude ethyl (±)-{3-[(4-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetate analogous to Example 1: t_R (LC-1) 1.09 min; ESI-MS (positive ion): m/z 396.92 $[M]^+$ (calcd 396.12 for $C_{22}H_{21}N_2O_3Cl$). 1H -NMR ($CDCl_3$): 2.05 (m, 2H); 2.42 (m, 1H); 2.70 (m, 3H); 2.92 (m, 1H); 3.28 (s, 3H); 4.65 (s, 2H); 7.11 (m, 5H); 7.37 (m, 3H). Retention times of the two enantiomers are respectively 12.4 min and 15.5 min (LC-6, 75% B), the more polar enantiomer displaying a rotation of the plane of the polarized light in a clockwise direction, the less polar enantiomer displaying a rotation in an anticlockwise direction, respectively, resulting a positive (+) and a negative (–) signal of the polarimetric plot, as co-monitored with an interfaced polarimeter.

- b1) (+)-{3-[(4-Chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid

20 The title compound is obtained using conditions for the hydrolysis of ethyl (+)-{3-[(4-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetate analogous to Example 1: t_R (LC-1) 1.09 min; ESI-MS (positive ion): m/z 396.92 $[M]^+$; t_R (LC-6, 75% B) 12.4 min, positive signal of the polarimetric plot; ee >99%.

- 25 b2) (–)-{3-[(4-Chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid

The title compound is obtained using conditions for the hydrolysis of ethyl (–)-{3-[(4-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetate analogous to Example 1: t_R (LC-1) 1.09 min; ESI-MS (positive ion): m/z 396.92 $[M]^+$; t_R (LC-6, 75% B) 15.5 min, negative signal of the polarimetric plot; ee >99%.

Table 1. Examples A-03 to A-61 are prepared using procedures analogous to those described for Example A-01 or A-02.

Ex.	Compound Name	Formula MW	LC-MS	
			t_R (Method)	ESI m/z
A-03	(±)-[3-(3,4-Dihydro-1 <i>H</i> -isoquinoline-2-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₄ N ₂ O ₃ 388.46	2.28 (LC-3)	389.18
A-04	(±)-(3-Phenylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid	C ₂₁ H ₂₀ N ₂ O ₃ 348.401	2.08 (LC-3)	371.15 [M+Na] ⁺
A-05	(±)-{3-[Benzyl-((S)-1-phenyl-ethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₃₀ N ₂ O ₃ 466.579	2.55 (LC-3)	467.21
A-06	(±)-{3-[Benzyl-((R)-1-phenyl-ethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₃₀ N ₂ O ₃ 466.579	2.53 (LC-3)	467.21
A-07	(±)-[3-(Methyl-phenethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₆ N ₂ O ₃ 390.481	2.22 (LC-3)	391.21
A-08	(±)-[3-(Benzyl-phenethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₃₀ H ₃₀ N ₂ O ₃ 466.579	2.48 (LC-3)	467.28
A-09	(±)-(3-Diphenethylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid	C ₃₁ H ₃₂ N ₂ O ₃ 480.606	2.56 (LC-3)	481.23

A-10	(±)-[3-(6,11-Dihydro-dibenzo[b,e]azepine-5-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₉ H ₂₆ N ₂ O ₃ 450.536	1.14 (LC-4)	450.98
A-11	(±)-{3-[Phenyl-(3-phenyl-propyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₃₀ N ₂ O ₃ 466.579	1.19 (LC-4)	467.04
A-12	(±)-[3-(Phenyl-thiophen-3-ylmethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₆ H ₂₄ N ₂ O ₃ S 444.554	1.13 (LC-4)	444.96
A-13	(±)-[3-(Isopropyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₆ N ₂ O ₃ 390.481	1.11 (LC-4)	391.04
A-14	(±)-[3-(11,12-Dihydro-6H-dibenzo[b,f]azocine-5-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₃₀ H ₂₈ N ₂ O ₃ 464.563	2.46 (LC-3)	465.17
A-15	(±)-[3-(Butyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₅ H ₂₈ N ₂ O ₃ 404.508	2.47 (LC-3)	405.23
A-16	(±)-[3-(3,4-Dihydro-2H-quinoline-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₄ N ₂ O ₃ 388.466	2.22 (LC-0)	389.31
A-17	(±)-[3-(Benzyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₈ H ₂₆ N ₂ O ₃ 438.525	2.32 (LC-3)	439.26

A-18	(±)-[3-(2-Methoxy-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₂ N ₂ O ₄ 378.427	2.12 (LC-3)	379.05
A-19	(±)-[3-(4-Methoxy-biphenyl-3-ylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₈ H ₂₆ N ₂ O ₄ 454.524	2.41 (LC-3)	455.37
A-20	(±)-[3-(Ethyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₄ N ₂ O ₃ 376.455	2.13 (LC-3)	377.23
A-21	(±)-[3-(Methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₂ N ₂ O ₃ 362.428	2.03 (LC-3)	363.26
A-22	(±)-[3-(3-Benzoyl-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₈ H ₂₄ N ₂ O ₄ 452.509	2.28 (LC-3)	453.21
A-23	(±)-[3-(5-Chloro-2-methoxy-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₁ N ₂ O ₄ Cl 412.872	2.33 (LC-3)	435.30 [M+Na] ⁺
A-24	(±)-[3-(2-Benzyl-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₈ H ₂₆ N ₂ O ₃ 438.525	2.27 (LC-3)	439.25
A-25	(±)-[3-(Dibenzo[b,f]azepine-5-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₉ H ₂₄ N ₂ O ₃ 448.521	2.29 (LC-3)	449.28

A-26	(±)-[3-(6,7-Dimethoxy-3,4-dihydro-1 <i>H</i> -isoquinoline-2-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₆ H ₂₈ N ₂ O ₅ 448.517	1.98 (LC-3)	449.18
A-27	(±)-{3-[(4-Fluoro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ F 380.418	2.10 (LC-0)	381.28
A-28	(±)-[3-(Ethyl-naphthalen-1-yl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₇ H ₂₆ N ₂ O ₃ 426.514	1.08 (LC-4)	426.98
A-29	(±)-[3-(Cyclohexyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₇ H ₃₀ N ₂ O ₃ 430.546	1.14 (LC-4)	430.97
A-30	(±)-[3-(Phenethyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₉ H ₂₈ N ₂ O ₃ 452.552	1.14 (LC-4)	453.05
A-31	(±)-[3-(Methyl-o-tolyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₄ N ₂ O ₃ 376.455	1.06 (LC-4)	376.99
A-32	(±)-[3-(Allyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₄ N ₂ O ₃ 388.466	1.09 (LC-4)	388.96
A-33	(±)-[3-(7-Trifluoromethyl-3,4-dihydro-2 <i>H</i> -quinoline-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₅ H ₂₃ N ₂ O ₃ F ₃ 456.463	1.17 (LC-4)	457.14

A-34	(±)-[3-(Benzyl-isopropyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₅ H ₂₈ N ₂ O ₃ 404.508	1.13 (LC-1)	405.02
A-35	(±)-{3-[(3-Chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ Cl 396.873	1.08 (LC-1)	396.92
A-36	(±)-[3-(Benzhydryl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₈ H ₂₆ N ₂ O ₃ 438.525	1.12 (LC-1)	439.07
A-37	(±)-{3-[Methyl-(2-trifluoromethyl-phenyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₃ H ₂₁ N ₂ O ₃ F ₃ 430.425	1.08 (LC-1)	430.89
A-38	(±)-{3-[(3,4-Dichloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₂ H ₂₀ N ₂ O ₃ Cl ₂ 431.318	1.14 (LC-1)	452.77 [M+Na] ⁺
A-39	(±)-[3-(2,3-Dihydro-indole-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₂ N ₂ O ₃ 374.439	1.09 (LC-1)	374.92
A-40	(±)-{3-[(3-Fluoro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ F 380.418	1.03 (LC-1)	380.94
A-41	(±)-{3-[Ethyl-(4-trifluoromethoxy-phenyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₄ H ₂₃ N ₂ O ₄ F ₃ 460.451	1.16 (LC-1)	461.02

A-42	(±)-[3-(Benzhydryl-methyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₉ H ₂₈ N ₂ O ₃ 452.552	1.18 (LC-1)	453.05
A-43	(±)-{3-[Benzyl-(2-cyano-ethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₅ H ₂₅ N ₃ O ₃ 415.492	1.02 (LC-1)	416.02
A-44	(±)-{3-[Methyl-(2-pyridin-2-yl-ethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₃ H ₂₅ N ₃ O ₃ 391.47	0.69 (LC-1)	392.01
A-45	(±)-{3-[(Benzo[1,3]dioxol-5-ylmethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₃ H ₂₂ N ₂ O ₅ 406.437	0.99 (LC-1)	406.87
A-46	(±)-{3-[4-(4-Fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₅ H ₂₆ N ₃ O ₃ F 435.497	1.07 (LC-1)	436.03
A-47	(±)-[3-(2-Fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₁ H ₁₉ N ₂ O ₃ F 366.391	1.04 (LC-4)	389.06 [M+Na]
A-48	(±)-[3-(3-Fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₁ H ₁₉ N ₂ O ₃ F 366.391	1.05 (LC-4)	367.03
A-49	(±)-[3-(4-Fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₁ H ₁₉ N ₂ O ₃ F 366.391	1.03 (LC-4)	367.1

A-50	(±)-{3-[Benzyl-(3-fluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₉ H ₂₇ N ₂ O ₃ F 470.542	1.14 (LC-4)	471.13
A-51	(±)-{3-[Benzyl-(4-fluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₉ H ₂₇ N ₂ O ₃ F 470.542	1.13 (LC-4)	469.19 [M-H]
A-52	(±)-{3-[Benzyl-(2-trifluoromethyl-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₂₇ N ₂ O ₃ F ₃ 520.549	1.15 (LC-4)	521.07
A-53	(±)-{3-[Benzyl-(3-trifluoromethyl-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₂₇ N ₂ O ₃ F ₃ 520.549	1.17 (LC-4)	521.21
A-54	(±)-{3-[Benzyl-(2,5-difluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₉ H ₂₆ N ₂ O ₃ F ₂ 488.532	1.14 (LC-4)	489.14
A-55	(±)-{3-[Benzyl-(3,5-difluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₉ H ₂₆ N ₂ O ₃ F ₂ 488.532	1.14 (LC-4)	489.21
A-56	(±)-{3-[Benzyl-(4-chloro-2-fluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₉ H ₂₆ N ₂ O ₃ ClF 504.987	1.16 (LC-4)	505.14
A-57	(±)-{3-[Benzyl-[2-(2-fluoro-phenyl)-ethyl]-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₂₉ N ₂ O ₃ F 484.569	1.14 (LC-4)	485.19

A-58	(±)-(3-{Benzyl-[2-(3-fluoro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid	C ₃₀ H ₂₉ N ₂ O ₃ F 484.569	1.14 (LC-4)	485.19
A-59	(±)-(3-{Benzyl-[2-(4-fluoro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid	C ₃₀ H ₂₉ N ₂ O ₃ F 484.569	1.14 (LC-4)	485.19
A-60	(±)-(3-{Benzyl-[2-(4-chloro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid	C ₃₀ H ₂₉ N ₂ O ₃ Cl 501.024	1.17 (LC-4)	501.12
A-61	(±)-(3-{Benzyl-[2-(2,6-dichloro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid	C ₃₀ H ₂₈ N ₂ O ₃ Cl ₂ 535.469	1.19 (LC-4)	535.13

Table 2. Examples B-01 to B-71 are prepared using procedures analogous to those described for Example A-01 or A-02.

Ex.	Compound Name	Formula MW	LC-MS	
			t_R (Method)	ESI m/z
B-01	(±)-[3-Methyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₄ N ₂ O ₃ 376.455	0.96 (LC-5)	377.32
B-02	(±)-[3-(2,3-Dihydro-indole-1-carbonyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₄ N ₂ O ₃ 388.466	1.14 (LC-1)	388.96
B-03	(±)-[3-(6,11-Dihydro-dibenzo[b,e]azepine-5-carbonyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₃₀ H ₂₈ N ₂ O ₃ 464.563	1.24 (LC-1)	465.03
B-04	(±)-[3-(Benzyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₉ H ₂₈ N ₂ O ₃ 452.552	1.23 (LC-1)	453.05
B-05	(±)-{3-Methyl-3-[phenyl(3-phenyl-propyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₁ H ₃₂ N ₂ O ₃ 480.606	1.29 (LC-2)	481.09
B-06	(±)-[3-(Benzhydryl-methyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₃₀ H ₃₀ N ₂ O ₃ 466.579	1.24 (LC-2)	488.98 [M+Na] ⁺

B-07	(±)-[3-Methyl-3-(phenyl-thiophen-3-ylmethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₇ H ₂₆ N ₂ O ₃ S 458.58	1.22 (LC-1)	458.94
B-08	(±)-[3-(3,4-Dihydro-2 <i>H</i> -quinoline-1-carbonyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₅ H ₂₆ N ₂ O ₃ 402.492	1.15 (LC-1)	402.94
B-09	(±)-[3-(Cyclohexyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₈ H ₃₂ N ₂ O ₃ 444.573	1.31 (LC-1)	467.04 [M+Na] ⁺
B-10	(±)-{3-[(3-Chloro-phenyl)-methyl-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₃ H ₂₃ N ₂ O ₃ Cl 410.9	1.15 (LC-1)	432.91 [M+Na] ⁺
B-11	(±)-[3-Methyl-3-(phenethyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₃₀ H ₃₀ N ₂ O ₃ 466.579	1.26 (LC-2)	466.93
B-12	(±)-{3-[(4-Fluoro-phenyl)-methyl-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₃ H ₂₃ N ₂ O ₃ F 394.445	1.10 (LC-1)	416.92 [M+Na] ⁺
B-13	(±)-{3-[(3,4-Dichloro-phenyl)-methyl-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₃ H ₂₂ N ₂ O ₃ Cl ₂ 445.345	1.21 (LC-1)	466.9 [M+Na] ⁺
B-14	(±)-[3-Methyl-3-(methyl- <i>o</i> -tolyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₆ N ₂ O ₃ 390.481	1.13 (LC-1)	390.97

B-15	(±)-[3-(Isopropyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₅ H ₂₈ N ₂ O ₃ 404.508	1.19 (LC-1)	427.03 [M+Na] ⁺
B-16	(±)-[3-(Ethyl-naphthalen-1-yl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₈ H ₂₈ N ₂ O ₃ 440.541	1.03 (LC-5)	441.34
B-17	(±)-[3-(Allyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₅ H ₂₆ N ₂ O ₃ 402.492	1.00 (LC-5)	403.34
B-18	(±)-{3-[Benzyl-((R)-1-phenyl-ethyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₁ H ₃₂ N ₂ O ₃ 480.606	1.25 (LC-1)	503.04
B-19	(±)-[3-(2-Benzyl-phenylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₉ H ₂₈ N ₂ O ₃ 452.552	1.18 (LC-1)	453.05
B-20	(±)-[3-(Benzyl-phenethyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₃₁ H ₃₂ N ₂ O ₃ 480.606	1.07 (LC-5)	481.43
B-21	(±)-[3-(2,3-Dihydro-indole-1-carbonyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₆ H ₂₈ N ₂ O ₃ 416.519	1.14 (LC-4)	438.93
B-22	(±)-[3-(2,3-Dihydro-indole-1-carbonyl)-3-ethyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₅ H ₂₆ N ₂ O ₃ 402.492	1.11 (LC-4)	402.78

B-23	(±)-(3-Ethyl-3-phenylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid	C ₂₃ H ₂₄ N ₂ O ₃ 376.455	1.07 (LC-4)	377.04
B-24	(±)-[3-Ethyl-3-(4-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₃ N ₂ O ₃ F 394.445	1.07 (LC-4)	416.99 [M+Na]
B-25	(±)-(3-Phenylcarbamoyl-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid	C ₂₄ H ₂₆ N ₂ O ₃ 390.481	1.08 (LC-4)	413.04 [M+Na]
B-26	(±)-(3-Methyl-3-phenylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid	C ₂₂ H ₂₂ N ₂ O ₃ 362.428	1.05 (LC-4)	361.11 [M-H]
B-27	(±)-[3-(2-Fluoro-phenylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ F 380.418	1.04 (LC-4)	403.00 [M+Na]
B-28	(±)-[3-(3-Fluoro-phenylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ F 380.418	1.07 (LC-4)	403.00 [M+Na]
B-29	(±)-[3-(4-Fluoro-phenylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ F 380.418	1.05 (LC-4)	379.04 [M-H]
B-30	(±)-[3-(Methyl-phenyl-carbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₅ H ₂₈ N ₂ O ₃ 404.508	1.12 (LC-4)	427.03 [M+Na]

B-31	(±)-[3-(4-Fluoro-phenylcarbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₅ N ₂ O ₃ F 408.472	1.11 (LC-4)	431.05 [M+Na]
B-32	(±)-[3-(3-Fluoro-phenylcarbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₅ N ₂ O ₃ F 408.472	1.12 (LC-4)	431.05 [M+Na]
B-33	(±)-[3-(2-Fluoro-phenylcarbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₅ N ₂ O ₃ F 408.472	1.10 (LC-4)	430.89 [M+Na]
B-34	(±)-[3-(Benzhydryl-carbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₃₁ H ₃₂ N ₂ O ₃ 480.606	1.16 (LC-4)	503.14 [M+Na]
B-35	(±)-[3-Ethyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₆ N ₂ O ₃ 390.481	1.09 (LC-4)	413.04 [M+Na]
B-36	(±)-[3-(Benzhydryl-carbamoyl)-3-ethyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₃₀ H ₃₀ N ₂ O ₃ 466.579	1.13 (LC-4)	489.08 [M+Na]
B-37	(±){3-[Benzyl-(3-fluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₂₉ N ₂ O ₃ F 484.569	1.17 (LC-4)	485.19
B-38	(±){3-[Benzyl-(4-fluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₂₉ N ₂ O ₃ F 484.569	1.17 (LC-4)	485.19

B-39	(±)-{3-[Benzyl-(2-trifluoromethyl-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₁ H ₂₉ N ₂ O ₃ F ₃ 534.576	1.20 (LC-4)	557.23 [M+Na]
B-40	(±)-{3-[Benzyl-(3-trifluoromethyl-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₁ H ₂₉ N ₂ O ₃ F ₃ 534.576	1.20 (LC-4)	557.16 [M+Na]
B-41	(±)-{3-[Benzyl-(2,5-difluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₂₈ N ₂ O ₃ F ₂ 502.559	1.17 (LC-4)	503.2
B-42	(±)-{3-[Benzyl-(3,5-difluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₂₈ N ₂ O ₃ F ₂ 502.559	1.17 (LC-4)	503.2
B-43	(±)-{3-[Benzyl-(4-chloro-2-fluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₂₈ N ₂ O ₃ ClF 519.014	1.20 (LC-4)	541.16 [M+Na]
B-44	(±)-{3-[Benzyl-[2-(2-fluoro-phenyl)-ethyl]-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₁ H ₃₁ N ₂ O ₃ F 498.596	1.17 (LC-4)	499.18
B-45	(±)-{3-[Benzyl-[2-(3-fluoro-phenyl)-ethyl]-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₁ H ₃₁ N ₂ O ₃ F 498.596	1.17 (LC-4)	499.18
B-46	(±)-{3-[Benzyl-[2-(4-fluoro-phenyl)-ethyl]-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₁ H ₃₁ N ₂ O ₃ F 498.596	1.17 (LC-4)	499.18

B-47	(±)-(3-{Benzyl-[2-(4-chloro-phenyl)-ethyl]-carbamoyl}-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid	C ₃₁ H ₃₁ N ₂ O ₃ Cl 515.051	1.20 (LC-4)	515.11
B-48	(±)-(3-{Benzyl-[2-(2,6-dichloro-phenyl)-ethyl]-carbamoyl}-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid	C ₃₁ H ₃₀ N ₂ O ₃ Cl ₂ 549.496	1.23 (LC-4)	549.13
B-49	(±)-[3-Ethyl-3-(2-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₃ N ₂ O ₃ F 394.445	1.06 (LC-4)	395.15
B-50	(±)-[3-Ethyl-3-(3-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₃ N ₂ O ₃ F 394.445	1.08 (LC-4)	395.15
B-51	(±)-[3-(Benzhydryl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₉ H ₂₈ N ₂ O ₃ 452.552	1.14 (LC-4)	453.19
B-52	(±)-[3-(Benzyl-methyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₆ N ₂ O ₃ 390.481	1.07 (LC-4)	391.13
B-53	(±)-{3-Methyl-3-[(R)-(1,2,3,4-tetrahydro-naphthalen-1-yl)carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₆ H ₂₈ N ₂ O ₃ 416.519	1.08 (LC-4)	439.13 [M+Na]
B-54	(±)-{3-Methyl-3-[(naphthalen-1-ylmethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₇ H ₂₆ N ₂ O ₃ 426.514	1.08 (LC-4)	427.21

B-55	(±)-[3-(Benzyl-butyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₇ H ₃₂ N ₂ O ₃ 432.562	1.16 (LC-4)	453.19 [M+Na]
B-56	(±)-{3-[Benzyl-(2-cyano-ethyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₆ H ₂₇ N ₃ O ₃ 429.518	1.06 (LC-4)	452.15 [M+Na]
B-57	(±)-[3-(Benzyl-ethyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₅ H ₂₈ N ₂ O ₃ 404.508	1.10 (LC-4)	405.19
B-58	(±)-[3-(Benzyl-isopropyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₆ H ₃₀ N ₂ O ₃ 418.535	1.12 (LC-4)	417.38 [M-H]
B-59	(±)-[3-(1 <i>H</i> ,3 <i>H</i> -Benzo[<i>d,e</i>]isoquinoline-2-carbonyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₈ H ₂₆ N ₂ O ₃ 438.525	1.11 (LC-4)	461.08 [M+Na]
B-60	(±)-[3-Methyl-3-(4-pentyl-benzylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₈ H ₃₄ N ₂ O ₃ 446.589	1.17 (LC-4)	447.23
B-61	(±)-[3-(4-Fluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₃ N ₂ O ₃ F 394.445	1.03 (LC-4)	395.15
B-62	(±)-[3-(2-Fluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₃ N ₂ O ₃ F 394.445	1.04 (LC-4)	395.15

B-63	(±)-[3-Methyl-3-(2-methylsulfonyl-benzylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₆ N ₂ O ₃ S 422.547	1.07 (LC-4)	423.13
B-64	(±)-[3-(4-Chloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₃ N ₂ O ₃ Cl 410.9	1.06 (LC-4)	411.08
B-65	(±)-[3-(2-Chloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₃ N ₂ O ₃ Cl 410.9	1.06 (LC-4)	411.08
B-66	(±)-[3-(2-Difluoromethoxy-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₄ N ₂ O ₄ F ₂ 442.461	1.07 (LC-4)	443.14
B-67	(±)-[3-(2,4-Dichloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₂ N ₂ O ₃ Cl ₂ 445.345	1.10 (LC-4)	445.01
B-68	(±)-[3-(3,5-Difluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₂ N ₂ O ₃ F ₂ 412.435	1.04 (LC-4)	413.09
B-69	(±)-[3-(3,4-Dichloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₂ N ₂ O ₃ Cl ₂ 445.345	1.09 (LC-4)	446.19
B-70	(±)-[3-(2,4-Difluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₂ N ₂ O ₃ F ₂ 412.435	1.05 (LC-4)	411.15 [M-H]

B-71	(±)-[3-(2,6-Difluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₂ N ₂ O ₃ F ₂ 412.435	1.04 (LC-4)	413.16
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Example C-01

(±)-[6-Cyano-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid

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a) (±)-6-Cyano-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid methyl-phenyl-amide

A stirred solution of 4-oxo-cyclohexanecarboxylic acid methyl-phenyl-amide (50 mg, 0.22 mmol) and (4-cyanophenyl)-hydrazine hydrochloric acid salt (36.5 mg, 0.22 mmol) in glacial AcOH (1 ml) is heated to reflux for 1 h. At rt, two drops of conc. H₂SO₄ are added and the reaction mixture is again heated to reflux for another 1 h, then cooled to rt and poured into 5% aqueous NaOAc. The resulting precipitate is filtered, rinsed several times with H₂O and dried under high vacuum providing pure subtitle compound (35 mg) in 49% yield as a white solid. *t_R* (LC-4) 1.06 min; ESI-MS (positive ion): *m/z* 329.96 [M]⁺ (calcd 329.40 for C₂₁H₁₉N₃O). ¹H-NMR (CDCl₃): 2.12 (m, 2H); 2.53 (m, 1H); 2.73 (m, 3H); 2.93 (m, 1H); 3.34 (s, 3H); 7.25 (m, 3H); 7.32 (m, 2H); 7.41 (m, 2H), 7.71 (s, 1H); 8.04 (br s, 1H).

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a1) 4-Oxo-cyclohexanecarboxylic acid methyl-phenyl-amide

Oxalyl chloride (0.89 g, 7.0 mmol) and a few drops of dry DMF are added to a stirred solution of 4-oxo-cyclohexanecarboxylic acid (0.50 g, 3.5 mmol) in CH₂Cl₂ (7 ml) and the reaction is kept stirring at rt overnight. The volatiles are removed under reduced pressure, the residue is azeotroped with toluene and dried under high vacuum yielding crude acid chloride. This material is reacted without further purification with *N*-methylaniline (0.16 g, 1.5 mmol) in CH₂Cl₂ (7 ml) in the presence of DIEA (0.58 g, 4.5 mmol). After stirring overnight at rt, saturated aqueous NaHCO₃ is added and the mixture extracted three times with CH₂Cl₂. The combined organic layers are dried over Na₂SO₄ and evaporated. The residue is column chromatographed on silica gel (hexane/ EtOAc 2:1) and the title

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compound is obtained as a pale yellow solid (0.47 g) in 57% yield. t_R (LC-4) 0.89 min; ESI-MS (positive ion): m/z 231.98 $[M]^+$ (calcd 231.13 for $C_{14}H_{17}NO_2$).

1H -NMR ($CDCl_3$): 2.04 (m, 6H); 2.45 (m, 2H); 2.65 (m, 1H); 3.28 (s, 3H); 7.22 (m, 2H); 7.44 (m, 3H).

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b) Ethyl ($\pm$)-[6-cyano-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetate

To a stirred solution of ($\pm$)-6-cyano-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid methyl-phenyl-amide (0.14 g, 0.43 mmol) in dry DMF (1.5 ml) is added KOtBu (58 mg, 0.52 mmol) and ethyl bromoacetate (79 mg, 0.47 mmol). The reaction mixture is stirred at rt overnight, then poured into saturated aqueous KH_2PO_4 and extracted with CH_2Cl_2 . The combined organic layers are washed with H_2O and brine, dried over Na_2SO_4 , filtered and evaporated. The residue is purified by column chromatography on silica gel (hexane/ EtOAc 1:1) to provide subtitle compound (91 mg) as a beige solid in 51% yield. t_R (LC-3) 1.15 min; ESI-MS (positive ion): m/z 416.09 $[M+H]^+$ (calcd 415.19 for $C_{25}H_{25}N_3O_3$). 1H -NMR ($CDCl_3$): 1.13 (t, $J=7.0$ Hz, 3H); 1.79 (m, 1H); 2.05 (m, 1H); 2.25 (m, 1H); 2.65 (m, 1H); 2.72 (m, 2H); 3.19 (s, 3H); 3.24 (m, 1H); 4.07 (q, $J=7.0$ Hz, 2H); 5.00 (d, $J=3.3$ Hz, 2H); 7.33 (m, 1H); 7.41 (m, 5H); 7.50 (d, $J=8.0$ Hz, 1H); 7.87 (s, 1H).

20 c) ($\pm$)-[6-Cyano-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid
Ethyl [6-cyano-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetate is saponified analogous to the conditions described for Example 1 yielding the title compound: t_R (LC-4) 1.05 min; ESI-MS (positive ion): m/z 388.06 $[M+H]^+$ (calcd 387.44 for $C_{23}H_{21}N_3O_3$). 1H -NMR ($CDCl_3$): 1.76 (m, 1H); 1.99 (m, 1H); 2.20 (m, 1H); 2.64 (m, 3H); 3.15 (s, 3H); 3.26 (m, 1H); 4.83 (s, 2H); 7.36 (m, 7H); 7.81 (s, 1H).

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Table 3. Examples C-02 to C-38 are prepared using a procedure analogous to that described for Example C-01, Example A-01 or A-02.

Ex.	Compound Name	Formula MW	LC-MS	
			t _R (Method)	ESI m/z
C-02	(±)-[6-Methyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₄ N ₂ O ₃ 376.455	2.28 (LC-3)	377.2
C-03	(±)-[6-Fluoro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ F 380.418	2.19 (LC-3)	381.15
C-04	(±)-[6-Chloro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ Cl 396.873	2.31 (LC-3)	397.14
C-05	(±)-[6-Iodo-3-(methyl-phenyl-carbamoyl)-carbazol-9-yl]-1,2,3,4-tetrahydro]-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ I 488.32	2.44 (LC-3)	489.00
C-06	(±)-[6-Bromo-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ Br 441.324	2.38 (LC-3)	443.00
C-07	(±)-[6-Methoxy-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₄ N ₂ O ₄ 392.454	2.17 (LC-3)	393.13
C-08	(±)-[7-Chloro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ Cl 396.873	2.34 (LC-3)	397.07

C-09	(±)-[6-Isopropyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₅ H ₂₈ N ₂ O ₃ 404.508	2.22 (LC-3)	405.16
C-10	(±)-[8-Chloro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ Cl 396.873	2.09 (LC-3)	397.07
C-11	(±)-[3-(Methyl-phenyl-carbamoyl)-6-nitro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₁ N ₃ O ₅ 407.425	1.08 (LC-4)	408.06
C-12	(±)-[3-(Methyl-phenyl-carbamoyl)-6-trifluoromethyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₁ N ₂ O ₃ F ₃ 430.425	1.13 (LC-4)	431.04
C-13	(±)-[7-Methyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₄ N ₂ O ₃ 376.455	1.03 (LC-2)	377.04
C-14	(±)-[8-Chloro-3-(indan-2-ylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₄ H ₂₃ N ₂ O ₃ Cl 422.911	0.97 (LC-5)	423.23
C-15	(±)-[8-Chloro-3-(2,2-diphenyl-ethylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₉ H ₂₇ N ₂ O ₃ Cl 486.997	1.02 (LC-5)	487.29
C-16	(±)-[8-Chloro-3-(2,3-dihydro-indole-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₁ N ₂ O ₃ Cl 408.884	0.99 (LC-5)	409.2

C-17	(±)-{8-Chloro-3-[4-(4-fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₅ H ₂₅ N ₃ O ₃ ClF 469.943	0.97 (LC-5)	470.27
C-18	(±)-[3-(Benzhydryl-methyl-carbamoyl)-8-chloro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₉ H ₂₇ N ₂ O ₃ Cl 486.997	1.06 (LC-5)	487.31
C-19	(±)-[3-(Benzhydryl-carbamoyl)-8-chloro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₈ H ₂₅ N ₂ O ₃ Cl 472.971	1.01 (LC-5)	473.3
C-20	(±)-[3-(4-Benzhydryl-piperazine-1-carbonyl)-8-chloro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₃₂ H ₃₂ N ₃ O ₃ Cl 542.077	0.87 (LC-5)	542.35
C-21	(+)-[3-(2,3-Dihydro-indole-1-carbonyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₁ N ₂ O ₃ F 392.429	13.8 (LC-8)	393.14
C-22	(+)-[6-Fluoro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₁ N ₂ O ₃ F 380.418	6.8 (LC-8)	381.09
C-23	(+)-[3-(Benzhydryl-methyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₉ H ₂₇ N ₂ O ₃ F 470.542	1.14 (LC-4)	471.13
C-24	(+)-{6-Fluoro-3-[4-(4-fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₅ H ₂₅ N ₃ O ₃ F ₂ 453.488	1.08 (LC-4)	454.23

C-25	(+)-[3-(Benzyl-isopropyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₅ H ₂₇ N ₂ O ₃ F 422.498	7.2 (LC-8)	423.16
C-26	(+)-[3-(Benzyl-phenethyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₃₀ H ₂₉ N ₂ O ₃ F 484.569	10.0 (LC-8)	485.26
C-27	(+)-[3-(Azocane-1-carbonyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₇ N ₂ O ₃ F 386.465	6.5 (LC-8)	387.25
C-28	(+)-{3-[Benzyl-((S)-1-phenyl-ethyl)-carbamoyl]-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₂₉ N ₂ O ₃ F 484.569	9.0 (LC-8)	485.19
C-29	(+)-{3-[Benzyl-((R)-1-phenyl-ethyl)-carbamoyl]-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₂₉ N ₂ O ₃ F 484.569	7.4 (LC-8)	485.33
C-30	(+)-[6-Fluoro-3-((R)-1-phenyl-ethylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₃ N ₂ O ₃ F 394.445	10.6 (LC-9)	395.15
C-31	(-)-[3-(Benzhydryl-methyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₉ H ₂₇ N ₂ O ₃ F 470.542	1.21 (LC-4)	471.47
C-32	(-)-{6-Fluoro-3-[4-(4-fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₂₅ H ₂₅ N ₃ O ₃ F ₂ 453.488	1.09 (LC-2)	454.23

C-33	(-)-[3-(Benzyl-isopropyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₅ H ₂₇ N ₂ O ₃ F 422.498	6.5 (LC-8)	423.2
C-34	(-)-[3-(Benzyl-phenethyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₃₀ H ₂₉ N ₂ O ₃ F 484.569	10.3 (LC-8)	485.19
C-35	(-)-[3-(Azocane-1-carbonyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₂ H ₂₇ N ₂ O ₃ F 386.465	7.8 (LC-8)	387.19
C-36	(-)-{3-[Benzyl-((S)-1-phenyl-ethyl)-carbamoyl]-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₂₉ N ₂ O ₃ F 484.569	9.0 (LC-8)	485.46
C-37	(-)-{3-[Benzyl-((R)-1-phenyl-ethyl)-carbamoyl]-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	C ₃₀ H ₂₉ N ₂ O ₃ F 484.569	8.0 (LC-8)	485.4
C-38	(-)-[6-Fluoro-3-((R)-1-phenyl-ethylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	C ₂₃ H ₂₃ N ₂ O ₃ F 394.445	7.5 (LC-9)	395.08

Example D-01

Benzyl (±)-9-carboxymethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylate

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a) Benzyl (±)-9-ethoxycarbonylmethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylate

Cesium carbonate (0.32 g, 1.0 mmol) and benzylbromide (60 mg, 0.35 mmol) are added to a stirred solution of Intermediate 1.1 (100 mg, 0.33 mmol) in dry DMF (1 ml). The resulting suspension is stirred at rt for 2 h, poured into saturated aqueous KH₂PO₄ solution and

extracted three times with diethyl ether. The combined organic layers are washed with H₂O and brine, dried over Na₂SO₄ and evaporated. The residue is crystallized from diisopropyl ether to give pure subtitle compound (95 mg) as a white solid in 73% yield. *t_R* (LC-4) 1.21 min; MS (positive ion) *m/z* 392.94 [M+H]⁺ (calcd 391.18 for C₂₄H₂₅NO₄).

- 5 ¹H-NMR (CDCl₃): 1.06 (m, 3H), 1.89 (m, 1H); 2.17 (m, 1H); 2.56 (m, 2H); 2.73 (m, 2H); 2.90 (m, 1H); 3.98 (m, 2H); 4.51 (d, *J*=12.8 Hz, 2H); 4.98 (d, *J*=12.8 Hz, 2H); 6.94 (m, 3H); 7.12 (m, 5H); 7.27 (m, 1H).

b) Benzyl (±)-9-carboxymethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylate

- 10 Benzyl (±)-9-ethoxycarbonylmethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylate is saponified analogous to the conditions described for Example A-01 yielding pure title compound: *t_R* (LC-4) 1.09 min; ESI-MS (positive ion): *m/z* 364.96 [M]⁺ (calcd 363.41 for C₂₂H₂₁NO₄).

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Example D-02

(*R*)-1-Phenyl-ethyl (±)-9-carboxymethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylate

- 20 a) Ethyl (±)-(3-chlorocarbonyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetate

Oxalyl chloride (1.26 g, 10 mmol) is dropwise added to a solution of Intermediate 1.1 (1.5 g, 5 mmol) in dry CH₂Cl₂ and the resulting orange solution is stirred overnight at rt. The volatiles are removed under reduced pressure and the residue is azeotroped twice with dry toluene and then dried under high vacuum to leave the crude subtitle compound as brownish oil, which is used without further purification.

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b): (*R*)-1-Phenyl-ethyl (±)-9-ethoxycarbonylmethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylate

- 30 Crude ethyl (±)-(3-chlorocarbonyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetate (5 mmol) is dissolved in dry CH₂Cl₂ and treated with (*R*)-1-phenyl-ethanol (0.55 g, 5.4 mmol), DIEA (1.94 g, 15 mmol) and DMAP (61 mg, 0.5 mmol). The resulting yellow solution is stirred at rt overnight, diluted with EtOAc and successively washed with 0.1 N HCl, saturated aqueous NaHCO₃, H₂O and brine. The organic layer is dried over Na₂SO₄, filtered and

evaporated. The residue is purified by column chromatography on silica gel (hexane/ EtOAc 5:1) to provide the subtitle compound as yellow oil (1.36 g) in 76% yield.

t_R (LC-4) 1.23 min; ESI-MS (positive ion): m/z 406.14 $[M+H]^+$ (calcd 405.49 for $C_{25}H_{27}NO_4$).

- 5 1H -NMR ($CDCl_3$): 1.24 (t, $J=7.1$ Hz, 3H); 1.56 (d, $J=6.5$ Hz, 3H); 2.05 (m, 1H); 2.34 (m, 1H); 2.87 (m, 4H); 3.12 (m, 1H); 4.18 (q, $J=6.5$ Hz, 2H); 4.71 (s, 2H); 5.95 (q, $J=6.5$ Hz, 1H); 7.14 (m, 3H); 7.34 (m, 5H); 7.48 (m, 1H).

c) (*R*)-1-Phenyl-ethyl ($\pm$)-9-carboxymethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylate

- 10 A stirred solution of (*R*)-1-phenyl-ethyl ($\pm$)-9-ethoxycarbonylmethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylate (50 mg, 0.123 mmol) in THF (1.1 ml) is treated with 0.2 N aqueous NaOH (0.555 ml, 0.111 mmol) at rt for 1 h. The basic reaction mixture is washed three times with diethyl ether and the aqueous phase is evaporated to dryness. The resulting solid is recrystallized from diisopropyl ether to give the pure sodium salt of the title compound as
15 a white solid (30 mg) in 61% yield.

t_R (LC-4) 1.16 min; ESI-MS (positive ion): m/z 399.97 $[M]^+$ (calcd 399.41 for $C_{23}H_{22}NNaO_4$).

- 1H -NMR ($DMSO-d_6$): 1.48 (d, $J=5.8$ Hz, 3H); 1.83 (m, 1H); 2.21 (m, 1H); 2.74 (m, 4H); 2.95 (m, 1H); 4.23 (s, 2H); 5.84 (q, $J=5.8$ Hz, 1H); 6.88 (t, $J=7.2$ Hz, 1H), 6.95 (t, $J=7.2$ Hz,
20 1H); 7.15 (d, $J=8.1$ Hz, 1H); 7.32 (m, 6H).

Biological assays:

Preparation of CRTH2 membranes and radioligand binding assay:

- Preparation of the membranes and radioligand binding assays are performed according to
25 known procedures, *e.g.* Sawyer N. *et al.* (*Br. J. Pharmacol.*, 2002, 137, 1163-1172). A clonal HEK 293 cell line, expressing high level of recombinant hCRTH2 receptor, is selected for the preparation of membranes. Cells are detached from culture plates in 5 ml buffer A (5 mM Tris, 1 mM $MgCl_2 \cdot 6H_2O$, 0.1 mM PMSF, 0.1 mM phenanthroline) per plate using a police rubber and transferred into centrifugation tubes and frozen at $-80^\circ C$.
30 After thawing, the cells are centrifuged at 500 g for 5 min and then resuspended in buffer A. Cells are then fragmented by homogenization with a Polytron homogenizer for 30 s. The membrane fragments are centrifuged at 3000 g for 40 min and resuspended in membranes in

buffer B (50 mM Tris, 25 mM MgCl₂, 250 mM saccharose, pH 7.4) and aliquots are stored frozen.

Binding assay is performed in a total volume of 250 μ l. In each well, 75 μ l buffer C (50 mM Tris, 100 mM NaCl, 1 mM EDTA, 0.1% BSA (protease free), 0.01 % NaN₃, pH 7.4) is
 5 mixed with 50 μ l {³H}-PGD₂ (at 2.5 nM (220.000 dpm per well) from Amersham, TRK734), 100 μ l CRTH2 membranes to give 80 μ g per well and 25 μ l of test compound in buffer C containing 1% DMSO. For unspecific binding, PGD₂ is added to the reaction
 mixture at 1 μ M final concentration. This binding assay mix is incubated at rt for 90 min and then filtered through a GF/C filter plate. The filter is washed three times with ice cold
 10 binding buffer. Then, 40 μ l per well Microscint-40 (Packard) are added and the bound radioactivity is quantified by means of Topcount (Packard).

Test for antagonist binding to the CRTH2 receptor:

Compounds of Formula I displayed IC₅₀ values of less than 10 μ M, as exemplified in the
 15 following Table 4.

Table 4

Compound Name	hCRTH2 BDG IC ₅₀ [μ M]
($\pm$)-{3-[(3-Chloro-phenyl)-methyl-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	0.001
($\pm$)-[3-(2,3-Dihydro-indole-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.002
($\pm$)-[3-Methyl-3-(phenyl-thiophen-3-ylmethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.002
($\pm$)-[3-Methyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.003
($\pm$)-[3-(Benzyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.003

(+)-[3-(Benzyl-isopropyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.003
(±)-[3-(3,5-Difluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.019
(±)-[3-(3-Fluoro-phenylcarbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.022
(-)-[3-(Benzyl-isopropyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.067

Intracellular calcium mobilization assay (FLIPR):

Cells (HEK-293), stably expressing the hCRTH₂ receptor under the control of the
 5 cytomegalovirus promotor from a single insertion of the expression vector pcDNA5
 (Invitrogen), are grown to confluency in DMEM (low glucose, Gibco) medium
 supplemented with 10% fetal calf serum (both Bioconcept, Switzerland) under standard
 mammalian cell culture conditions (37°C in a humidified atmosphere of 5% CO₂). Cells are
 detached from culture dishes using a dissociation buffer (0.02% EDTA in PBS, Gibco) for 1
 10 min, and collected by centrifugation at 200 g at rt for 5 min in assay buffer (equal parts of
 Hank's BSS (HBSS, Bioconcept) and DMEM (low glucose, without phenol red, Gibco)).
 After incubation for 45 min (37°C and 5% CO₂) in the presence of 1 µM Fluo-4 and 0.04%
 Pluronic F-127 (both Molecular Probes), 20 mM HEPES (Gibco) in assay buffer, the cells
 are washed with and resuspended in assay buffer, then seeded onto 384-well FLIPR assay
 15 plates (Greiner) at 50,000 cells in 66 µl per well, and sedimented by centrifugation.

Stock solutions of test compounds are made up at a concentration of 10 mM in DMSO, and
 serially diluted in assay buffer to concentrations required for inhibition dose response
 curves. Prostaglandin D₂ (Biomol, Plymouth Meeting, PA) is used as an agonist.

A FLIPR384 instrument (Molecular Devices) is operated according to the manufacturer's
 20 standard instructions, adding 4 µl of test compound dissolved at 10 mM in DMSO and
 diluted prior to the experiment in assay buffer to obtain the desired final concentration. 10 µl
 of 80 nM prostaglandin D₂ (Biomol, Plymouth Meeting, PA) in assay buffer, supplemented
 with 0.8% bovine serum albumin (fatty acid content <0.02%, Sigma), is then added to

obtain a final concentration of 10nM and 0.1%, respectively. Changes in fluorescence are monitored before and after the addition of test compounds at $\lambda_{\text{ex}}=488$ nm and $\lambda_{\text{em}}=540$ nm. Emission peak values above base level after prostaglandin D₂ addition are exported after base line subtraction. Values are normalized to high-level control (no test compound added) after subtraction of base line value (no prostaglandin D₂ added). The program XLfit 3.0 (IDBS) is used to fit the data to a single site dose response curve of the equation $(A+((B-A)/(1+((C/x)^D))))$ and to calculate the IC₅₀ values.

Antagonist analysis:

Compounds of Formula I antagonize prostaglandin D₂ mediated hCRTH2 receptor activity with an IC₅₀ less than 10 μ M as exemplified in the following Table 5.

Table 5

Compound Name	hCRTH2 FLIPR IC ₅₀ [μ M]
($\pm$)-[3-(Methyl-phenyl-carbamoyl)-6-nitro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.011
($\pm$)-[3-Methyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.013
($\pm$)-[3-(Benzhydryl-methyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.017
($\pm$)-{3-[(3-Chloro-phenyl)-methyl-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	0.020
($\pm$)-{3-[Methyl-(2-trifluoromethyl-phenyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	0.045
(+)-[3-(2,3-Dihydro-indole-1-carbonyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.028
(+)-[3-(Azocane-1-carbonyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.046
($\pm$)-{3-[Benzyl-(2-cyano-ethyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	0.036

[3-(2,3-Dihydro-indole-1-carbonyl)-3-ethyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid	0.067
(-)-{6-Fluoro-3-[4-(4-fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid	0.599

The compounds of the invention can be formulated as the active ingredient according to methods known *per se* to give pharmaceutical preparations of the following compositions:

5

Example E-01: Gelatin solution

A sterile-filtered aqueous solution, with 20% cyclodextrins as solubilisers, of one of the compounds of Formula I mentioned in the preceding Examples (*e.g.* Example A-02 as active ingredient), is so mixed under aseptic conditions, with heating, with a sterile gelatin solution containing phenol as preservative, that 1.0 ml of solution has the following composition:

10

active ingredient	3 mg
gelatin	150.0 mg
15 phenol	4.7 mg
dist. water with 20% cyclodextrins	
as solubilisers	1.0 ml

Example E-02: Sterile dry substance for injection

20 5 mg of one of a compound of Formula I mentioned in the preceding Examples (*e.g.* Example B-02) as active ingredient are dissolved in 1 ml of an aqueous solution with 20 mg of mannitol and 20% cyclodextrins as solubilisers. The solution is sterile-filtered and introduced under aseptic conditions into a 2 ml ampoule, deep-frozen and lyophilized. Before use, the lyophilisate is dissolved in 1 ml of distilled water or 1 ml of physiological saline solution. The solution is administered intramuscularly or intravenously. This formulation can also be introduced into a twin-chambered injection ampoule.

25

Example E-03: Film-coated tablets

The following ingredients are used for the preparation of 10,000 tablets each containing 100 mg of the active ingredient:

	active ingredient	1000 g
5	corn starch	680 g
	colloidal silica	200 g
	magnesium stearate	20 g
	stearic acid	50 g
	sodium carboxymethyl starch	250 g
10	water	quantum satis

A mixture of one of the compounds of Formula I mentioned in the preceding Examples (*e.g.* Example B-10) as active ingredient, 50 g of corn starch and the colloidal silica is processed with a starch paste, made from 250 g of corn starch and 2.2 kg of demineralised water, to form a moist mass. This is forced through a sieve having a mesh size of 3 mm and dried at 45°C for 30 min in a fluidized bed drier. The dry granulates are pressed through a sieve having a mesh size of 1 mm, mixed with a pre-sieved mixture (1 mm sieve) of 330 g of corn starch, the magnesium stearate, the stearic acid and the sodium carboxymethyl starch, and compressed to form slightly biconvex tablets.

20

Example E-04: Soft capsules

5000 soft gelatin capsules, each comprising as active ingredient 0.05 g of one of the compounds of Formula I mentioned in the preceding Examples (*e.g.* C-03) are prepared as follows:

25

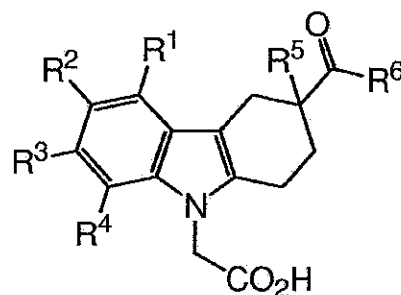
active ingredient	250 g
lauroglycol®	2 liters

The pulverized active ingredient is suspended in Lauroglykol® (propylene glycol laureate, Glattefossé S. A., Saint Priest, France) and ground in a wet pulverizer to produce a particle size of about 1 to 3 µm. 0.42 g portions of the mixture are then introduced into soft gelatin capsules using a capsule-filling machine.

30

Claims:

1. A compound selected from the group consisting of 2,3,4,9-tetrahydro-1*H*-carbazoles
 5 of the Formula I:



I

wherein

- R^1 , R^2 , R^3 and R^4 independently represent hydrogen, alkyl, alkoxy, halogen, nitro, cyano,
 10 trifluoromethyl, or formyl;
 R^5 represents hydrogen, alkyl, or $-CF_3$;
 R^6 represents alkoxy, aryl-alkoxy, or $-NR^7R^8$;
 R^7 and R^8 independently represent hydrogen, alkyl, cyano-alkyl, alkenyl, aryl, aryl-alkyl,
 phenylcarbonyl, cycloalkyl, pyridyl-alkyl, thienyl-alkyl, furanyl-alkyl, or imidazolyl-alkyl;
 15 or
 R^7 and R^8 , together with the nitrogen atom to which they are attached, form a heterocyclic
 5-, 6-, 7-, or 8-membered ring system with one to three heteroatoms which are selected from
 nitrogen, oxygen and sulfur and which ring system is optionally substituted with (i) one or
 two annellated benzene rings, which benzene rings are unsubstituted or substituted with one
 20 or two substituents independently selected from C1-C4 alkyl, C1-C4 alkoxy, halogen, $-CF_3$,
 and $-OCF_3$; (ii) an unsubstituted phenyl ring; (iii) a mono- or di-substituted phenyl ring,
 wherein the substituents are independently selected from halogen, C1-C4 alkyl, C1-C4
 alkoxy, $-CF_3$, and $-OCF_3$; or (iv) phenyl-alkyl wherein the alkyl moiety is substituted by
 phenyl;
 25 and optically pure enantiomers, mixtures of enantiomers such as racemates, optically pure
 diastereomers, mixtures of diastereomers, mixtures of enantiomers and diastereomers such as

diastereomeric racemates, meso forms, and geometric isomers;
prodrugs of such compounds in which a prodrug forming group is present, as well as
solvates and morphological forms, and pharmaceutically acceptable salts thereof.

- 5 2. A compound according to claim 1, wherein
R¹, R², R³ and R⁴ independently represent hydrogen, alkyl, alkoxy, halogen, nitro, cyano,
trifluoromethyl, or formyl;
R⁵ represents hydrogen, alkyl, or -CF₃;
R⁶ represents alkoxy; aryl-alkoxy; mono-, di-, or tri-substituted aryl-alkoxy substituted
10 independently in the aryl moiety with halogen, C1-C4 alkyl, C1-C4 alkoxy, -CF₃, -OCF₃; or
-NR⁷R⁸;
R⁷ and R⁸ independently represent hydrogen; alkyl; cyano-alkyl; alkenyl; aryl; mono-, di-,
or tri- substituted aryl substituted independently in the aryl moiety with halogen, C1-C4
alkyl, C1-C4 alkoxy, -CF₃, -OCF₃; aryl-alkyl; mono-, di-, or tri- substituted aryl-alkyl
15 substituted independently in the aryl moiety with halogen, C1-C4 alkyl, C1-C4 alkoxy,
-CF₃, -OCF₃; phenylcarbonyl; phenyl-C1-C4 alkyl optionally substituted with a
methylenedioxy group at the phenyl ring; cycloalkyl; pyridyl-alkyl; thienyl-alkyl; furanyl-
alkyl; or imidazolyl-alkyl; or
R⁷ and R⁸, together with the nitrogen atom to which they are attached, form a heterocyclic
20 5-, 6-, 7-, or 8-membered ring system with one to three heteroatoms which are selected from
nitrogen, oxygen and sulfur and which ring system is optionally substituted with (i) one or
two annellated benzene rings, which benzene rings are unsubstituted or substituted with one
or two substituents selected from C1-C4 alkyl, C1-C4 alkoxy, halogen, -CF₃, -OCF₃; (ii) an
unsubstituted phenyl ring; or (iii) a mono- or di-substituted phenyl ring substituted with
25 halogen, C1-C4 alkyl, C1-C4 alkoxy, -CF₃, or -OCF₃.
3. A compound according to claim 1 or 2, wherein R¹, R², R³ and R⁴ represent C₁-C₄ alkyl,
C₁-C₄ alkoxy, halogen, nitro, cyano, or trifluoromethyl.
- 30 4. A compound according to claim 1 or 2, wherein R¹, R², R³ and R⁴ represent methyl,
trifluoromethyl, methoxy, fluoro, chloro, bromo, or iodo.

5. A compound according to any one of claims 1 to 4, wherein R⁷ represents alkenyl, alkyl, aryl-C1-C4 alkyl, cycloalkyl, thienyl-C1-C4 alkyl, furanyl-C1-C4 alkyl, pyridyl-C1-C4 alkyl, or imidazolyl-C1-C4 alkyl; and R⁸ represents hydrogen, aryl, aryl-C1-C4 alkyl, furanyl-C1-C4 alkyl, pyridyl-C1-C4 alkyl, or thienyl-C1-C4 alkyl.

5

6. A compound according to any one of claims 1 to 4, wherein R⁷ represents hydrogen, allyl, 2-cyano-ethyl, methyl, butyl, ethyl, isopropyl, benzyl, 1-phenyl-ethyl, 2-phenyl-ethyl, phenyl-propyl, cyclohexyl, or thiophen-3-ylmethyl; and R⁸ represents phenyl, 2-benzyl-phenyl, 2-methoxy-phenyl, 2-methyl-phenyl, 2-trifluoromethyl-phenyl, 3,4-dichloro-phenyl, 10 3-benzoyl-phenyl, 3-chloro-phenyl, 2-fluoro-phenyl, 3-fluoro-phenyl, 4-chloro-phenyl, 4-fluoro-phenyl, 4-methoxy-biphenyl-3-yl, 4-trifluoromethoxy-phenyl, 5-chloro-2-methoxy-phenyl, naphthalen-1-yl, benzo[1,3]dioxol-5-ylmethyl, benzyl, diphenylmethyl, 1-phenyl-ethyl, 2-phenyl-ethyl, 2-pyridin-2-yl-ethyl, 4-phenyl-benzyl, 3,4-dichloro-benzyl, 2,4-dichloro-benzyl, difluoromethoxy-benzyl, 2-chloro-benzyl, 4-chloro-benzyl, 2-

15 methylsulfanyl-benzyl, 2-fluoro-benzyl, 3-fluoro-benzyl, 4-fluoro-benzyl, 2-trifluoromethyl-benzyl, 3-trifluoromethyl-benzyl, 2,4-difluoro-benzyl, 2,5-difluoro-benzyl, 2,6-difluoro-benzyl, 3,5-difluoro-benzyl, 4-chloro-2-fluoro-benzyl, (2-fluoro-phenyl)-ethyl, (3-fluoro-phenyl)-ethyl, (4-fluoro-phenyl)-ethyl, (4-chloro-phenyl)-ethyl, (2,6-dichloro-phenyl)-ethyl, naphthalene-1-ylmethyl, 1,2,3,4-tetrahydro-naphthalen-1-yl, indan-2-yl, or 20 2,2-diphenyl-ethyl.

7. A compound according to any one of claims 1 to 4, wherein R⁷ and R⁸, together with the nitrogen atom to which they are attached, form a dihydro-dibenzo[b,f]azocine, dihydro-indole, dihydroisoquinoline, dihydroquinoline, or dibenzazepine ring.

25

8. A compound according to any one of claims 1 to 4, wherein R⁷ and R⁸, together with the nitrogen atom to which they are attached, form a heterocyclic ring system selected from the group consisting of 11,12-dihydro-6H-dibenzo[b,f]azocine-5-yl, 2,3-dihydro-indole-1-yl, 3,4-dihydro-1H-isoquinoline-2-yl, 3,4-dihydro-2H-quinoline-1-yl, 4-(4-fluoro-phenyl)-piperazine-1-yl, 6,11-dihydro-dibenzo[b,e]azepine-5-yl, 6,7-dimethoxy-3,4-dihydro-1H-isoquinoline-2-yl, 7-trifluoromethyl-3,4-dihydro-2H-quinoline-1-yl, dibenzo[b,f]azepine-5-yl, 1H,3H-benzo[d,e]isoquinoline-2-yl, 4-benzhydryl-piperazine-1-yl, and azocane-1-yl.

30

9. A compound according to any one of claims 1 to 4, wherein R⁶ represents (*R*)-1-phenyl-ethyloxy or benzyloxy.
10. A compound according to claim 1, wherein
- 5 R¹, R², R³ and R⁴ independently represent hydrogen, alkyl, alkoxy, halogen, nitro, cyano, or trifluoromethyl;
- R⁵ represents hydrogen or alkyl;
- R⁶ represents phenyl-alkoxy or -NR⁷R⁸;
- R⁷ and R⁸ independently represent hydrogen; alkyl; cyano-alkyl; alkenyl; phenyl optionally
- 10 mono-substituted by halogen, alkyl, alkoxy, -CF₃, -OCF₃, phenyl-alkyl or phenyl-carbonyl; phenyl, di-substituted with a substituent independently selected from halogen, alkoxy and phenyl; phenyl-alkyl optionally substituted in the alkyl moiety by phenyl or optionally substituted in the phenyl ring by methylenedioxy; phenyl-alkyl which is di-substituted by halogen or mono-substituted by halogen, -CF₃, -OCHF₂, alkyl or alkylsulfanyl; naphthyl;
- 15 naphthyl-alkyl; cycloalkyl which is optionally substituted with an annellated benzene ring; pyridyl-alkyl; or thienyl-alkyl; or
- R⁷ and R⁸, together with the nitrogen atom to which they are attached, form a heterocyclic 5-, 6-, 7-, or 8-membered ring system with one or two heteroatoms selected from nitrogen and which ring system is optionally substituted with (i) one or two annellated benzene rings,
- 20 which benzene rings are unsubstituted or substituted with one or two substituents independently selected from alkoxy and -CF₃; (ii) a mono-substituted phenyl ring substituted with halogen; or (iii) phenyl-alkyl, wherein the alkyl moiety is substituted with phenyl.
- 25 11. A compound according to claim 1 selected from the group consisting of:
- [3-methyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[(3-chloro-phenyl)-methyl-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(2,3-dihydro-indole-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 30 [3-methyl-3-(phenyl-thiophen-3-ylmethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(benzyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

- {3-[(4-fluoro-phenyl)-methyl-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(6,11-dihydro-dibenzo[*b,e*]azepine-5-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 5 [3-(benzyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[phenyl-(3-phenyl-propyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(2,3-dihydro-indole-1-carbonyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[(3,4-dichloro-phenyl)-methyl-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 10 [3-(allyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[benzyl-((*S*)-1-phenyl-ethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- {3-[methyl-(2-trifluoromethyl-phenyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(methyl-phenyl-carbamoyl)-6-trifluoromethyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 15 acid;
- [3-methyl-3-(methyl-*o*-tolyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(2-benzyl-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(dibenzo[*b,f*]azepine-5-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(ethyl-naphthalen-1-yl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 20 [3-(benzhydryl-methyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-methyl-3-[phenyl-(3-phenyl-propyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(ethyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(phenyl-thiophen-3-ylmethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 25 [3-(benzyl-isopropyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[4-(4-fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(benzyl-phenethyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(3,4-dihydro-2*H*-quinoline-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 30 [3-(cyclohexyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[(3-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-methyl-3-(phenethyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(2-methoxy-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

- [3-(allyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(benzyl-phenethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(phenethyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(methyl-*o*-tolyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
5 {3-[benzyl-((*R*)-1-phenyl-ethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
[3-(methyl-phenyl-carbamoyl)-6-nitro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(7-trifluoromethyl-3,4-dihydro-2*H*-quinoline-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(3,4-dihydro-2*H*-quinoline-1-carbonyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic
10 acid;
{3-[(4-fluoro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
[3-(4-methoxy-biphenyl-3-ylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[6-fluoro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[7-chloro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
15 [6-chloro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
{3-[methyl-(2-pyridin-2-yl-ethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
{3-[(4-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
{(*R*)-3-[(4-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
[3-(isopropyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
20 {3-[(3,4-dichloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
[3-(benzhydryl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[6-cyano-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
{3-[benzyl-((*R*)-1-phenyl-ethyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-
acetic acid;
25 [3-(2-benzyl-phenylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[6-isopropyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[6-methyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(11,12-dihydro-6*H*-dibenzo[*b,f*]azocine-5-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-
acetic acid;
30 [3-(methyl-phenethyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[6-bromo-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(3-benzoyl-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
(3-dibenzylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;

- [3-(ethyl-naphthalen-1-yl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[(3-fluoro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- {3-[benzyl-(2-cyano-ethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 5 (3-phenylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
- [3-(isopropyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- (3-diphenethylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
- [3-(benzhydryl-methyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(5-chloro-2-methoxy-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 10 [3-(butyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[ethyl-(4-trifluoromethoxy-phenyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [6-iodo-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 15 [6-methoxy-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 9-carboxymethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid (*R*)-1-phenyl-ethyl ester;
- {{(*S*)-3-[(4-chloro-phenyl)-methyl-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(6,11-dihydro-dibenzo[*b,e*]azepine-5-carbonyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-
- 20 yl]-acetic acid;
- [8-chloro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 9-carboxymethyl-2,3,4,9-tetrahydro-1*H*-carbazole-3-carboxylic acid benzyl ester;
- [3-(3,4-dihydro-1*H*-isoquinoline-2-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(6,7-dimethoxy-3,4-dihydro-1*H*-isoquinoline-2-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-
- 25 yl]-acetic acid;
- {3-[(benzo[1,3]dioxol-5-ylmethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid; and
- [3-(cyclohexyl-phenyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid.
- 30 12. A compound according to claim 1 selected from the group consisting of:
- {3-[benzyl-((*R*)-1-phenyl-ethyl)-carbamoyl]-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(benzhydryl-methyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

- {3-[benzyl-((S)-1-phenyl-ethyl)-carbamoyl]-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(benzyl-phenethyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(2,3-dihydro-indole-1-carbonyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 5 [3-(benzyl-isopropyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(azocane-1-carbonyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- (3-phenylcarbamoyl-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
- [3-(methyl-phenyl-carbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [6-fluoro-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 10 {6-fluoro-3-[4-(4-fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(4-fluoro-phenylcarbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {6-fluoro-3-[4-(4-fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 15 [3-(benzhydryl-methyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- (3-{benzyl-[2-(2,6-dichloro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
- {3-[benzyl-((S)-1-phenyl-ethyl)-carbamoyl]-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 20 [3-(3-fluoro-phenylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(3,5-difluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(3-fluoro-phenylcarbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(3-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(2,3-dihydro-indole-1-carbonyl)-3-ethyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 25 [3-ethyl-3-(3-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-ethyl-3-(4-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(2-chloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(benzyl-phenethyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-methyl-3-(2-methylsulfanyl-benzylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 30 acid;
- (3-ethyl-3-phenylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
- {3-[benzyl-((R)-1-phenyl-ethyl)-carbamoyl]-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;

- [3-ethyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(2-fluoro-phenylcarbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(2-fluoro-phenylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
{3-methyl-3-[(naphthalen-1-ylmethyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic
5 acid;
{3-[benzyl-(2-cyano-ethyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic
acid;
[3-(benzyl-butyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(4-fluoro-phenylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
10 [3-(4-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(2,4-dichloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
{3-[benzyl-(4-fluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
[3-(benzyl-isopropyl-carbamoyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(2-difluoromethoxy-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic
15 acid;
[3-(2-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(benzyl-ethyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
(3-methyl-3-phenylcarbamoyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
[3-ethyl-3-(2-fluoro-phenylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
20 [3-(4-chloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
(3-{benzyl-[2-(4-fluoro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic
acid;
[3-methyl-3-(4-pentyl-benzylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
{3-[benzyl-(4-chloro-2-fluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic
25 acid;
[3-(azocane-1-carbonyl)-6-fluoro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(2-fluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(3,4-dichloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
(3-{benzyl-[2-(4-chloro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic
30 acid;
[3-(2,4-difluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
[3-(benzhydryl-carbamoyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;

- {8-chloro-3-[4-(4-fluoro-phenyl)-piperazine-1-carbonyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [8-chloro-3-(2,3-dihydro-indole-1-carbonyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(2,6-difluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 5 {3-[benzyl-(3-fluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(benzhydryl-methyl-carbamoyl)-8-chloro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(benzhydryl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[benzyl-(3-trifluoromethyl-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 10 [3-(benzhydryl-carbamoyl)-3-ethyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(4-fluoro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- (3-{benzyl-[2-(3-fluoro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
- [3-(benzyl-isopropyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 15 {3-[benzyl-(3,5-difluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- (3-{benzyl-[2-(2-fluoro-phenyl)-ethyl]-carbamoyl}-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
- {3-[benzyl-(2-trifluoromethyl-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 20 {3-[benzyl-(4-fluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- [3-(1*H*,3*H*-benzo[*de*]isoquinoline-2-carbonyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[benzyl-(2,5-difluoro-benzyl)-carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 25 {3-[benzyl-(3-fluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- {3-methyl-3-[(*R*)-(1,2,3,4-tetrahydro-naphthalen-1-yl)carbamoyl]-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- {3-[benzyl-(3,5-difluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 30 acetic acid;
- [3-(benzyl-methyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- (3-{benzyl-[2-(4-chloro-phenyl)-ethyl]-carbamoyl}-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;

- (3-{benzyl-[2-(4-fluoro-phenyl)-ethyl]-carbamoyl}-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
- [8-chloro-3-(2,2-diphenyl-ethylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[benzyl-(3-trifluoromethyl-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 5 [3-(4-benzhydryl-piperazine-1-carbonyl)-8-chloro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- {3-[benzyl-(4-chloro-2-fluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 10 [3-(benzhydryl-carbamoyl)-8-chloro-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(2,3-dihydro-indole-1-carbonyl)-3-propyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- (3-{benzyl-[2-(3-fluoro-phenyl)-ethyl]-carbamoyl}-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
- (3-{benzyl-[2-(2,6-dichloro-phenyl)-ethyl]-carbamoyl}-3-methyl-1,2,3,4-tetrahydro-
- 15 carbazol-9-yl)-acetic acid;
- {3-[benzyl-(2-trifluoromethyl-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- {3-[benzyl-(2,5-difluoro-benzyl)-carbamoyl]-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl}-acetic acid;
- 20 [8-chloro-3-(indan-2-yl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- (3-{benzyl-[2-(2-fluoro-phenyl)-ethyl]-carbamoyl}-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl)-acetic acid;
- [6-fluoro-3-((*R*)-1-phenyl-ethylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [6-fluoro-3-((*R*)-1-phenyl-ethylcarbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- 25 [3-(benzyl-cyanomethyl-carbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- [3-(2,3-dichloro-benzylcarbamoyl)-3-methyl-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid;
- and
- [7-methyl-3-(methyl-phenyl-carbamoyl)-1,2,3,4-tetrahydro-carbazol-9-yl]-acetic acid.
- 30 13. A pharmaceutical composition comprising a compound according to any one of claims 1 to 12 and a pharmaceutically acceptable carrier.

14. A compound according to any one of claims 1 to 12, or composition according to claim 13, for use as a medicament.

15. Use of a compound according to any one of claims 1 to 12 for the preparation of a
5 pharmaceutical composition for the prevention and/or treatment of diseases selected from
the group consisting of both chronic and acute allergic diseases or immune disorders
comprising allergic asthma, rhinitis, chronic obstructive pulmonary disease, dermatitis,
inflammatory bowel disease, rheumatoid arthritis, allergic nephritis, conjunctivitis, atopic
dermatitis, bronchial asthma, food allergy, systemic mast cell disorders, anaphylactic shock,
10 urticaria, eczema, itching, inflammation, ischemia-reperfusion injury, cerebrovascular
disorders, pleuritis, ulcerative colitis, and eosinophil-related diseases, such as Churg-Strauss
syndrome and sinusitis, and basophil-related diseases, such as basophilic leukemia and
basophilic leukocytosis.