Title: INDOL-S-CARBONYL-SPIRO-PIPERIDINE DERIVATIVES AS VIA RECEPTOR ANTAGONISTS

Abstract: This invention relates to indol-3-yl-carbonyl-spiro-piperidine derivatives which act as Via receptor antagonists and which are represented by Formula (I); wherein the spiro-piperidine head group A and the residues R' , R'' and R''' are as defined herein. The invention further relates to pharmaceutical compositions containing such compounds, their use in medications against dysmenorrhea, hypertension, chronic heart failure, inappropriate secretion of vasopressin, liver cirrhosis, nephrotic syndrome, obsessive compulsive disorder, anxious and depressive disorders, and methods of preparation thereof.
The present invention relates to compounds of the general formula (I)

wherein

A is selected from the following groups (a), (b), (c), (d), (e), (f), (g) and (h):
wherein in (a) the dotted line is either nil or a double bond;

\[ R^1 \text{ is } \begin{cases} H, \\ \text{or is } \text{Ci}_{6}-\text{alkyl optionally substituted by CN,} \\ \text{or is aryl, 5 or 6 membered heteroaryl or sulfonylaryl which are optionally substituted by one or more B,} \\ \text{or is } -(\text{CH}_2)_m-R^a \text{ wherein } R^a \text{ is:} \\ \text{CN,} \\
\text{OR}, \\
\text{NR\text{R}_v}, \\
\text{C}_{3-6}-\text{cycloalkyl, 3 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,} \\
\text{or is } -(\text{CH}_2)_n-(\text{CO})-R^b \text{ or } -(\text{CH}_2)_n-(\text{SO}_2)-R^b \text{ wherein } R^b \text{ is:} \\ \text{Ci}_{6}-\text{alkyl,} \\
\text{Ci}_{6}-\text{alkoxy,} \\
\text{C}_{3-6}-\text{cycloalkyl,} \\
-(\text{CH}_2)_m\text{NR}\text{R}^v, \\
\text{NR\text{R}_v}, \\
\text{C}_{3-6}-\text{cycloalkyl, 4 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,} \\
\text{or } R^1 \text{ and } R^3 \text{ together with the indole ring to which they are attached form a 5 or 6 membered heterocycloalkyl which can be substituted by } =0, \text{ C(O)O-C}_{1-6}-\text{alkyl or } \text{C}_{1-6}-\text{alkyl;} \\
\text{R}^2 \text{ is } \begin{cases} \text{one or more of H, OH, halo, CN, nitro, C}_{1-6}-\text{alkyl optionally substituted by } -\text{NR}\text{R}^v, \text{Ci}_{6}-\text{alkoxy, } \text{O-CH}_2\text{-C}_{2-6}-\text{alkenyl, benzyloxy,} \\ \text{or two } R^2 \text{ may form an oxo or dioxo bridge together with the indole ring to which they are attached;} \\ \text{R}^3 \text{ is } \begin{cases} H, \\ \text{or is halo,} \\ \text{or is } -(\text{CO})-R^c \text{ wherein } R^c \text{ is:} \\
\text{C}_{1-6}-\text{alkyl,} \\
-(\text{CH}_2)_m\text{VNR\text{R}_v}, \\
-(\text{CH}_2)_n\text{NR}\text{R}^v, \\
\text{5 or 6 membered heterocycloalkyl optionally substituted by } \text{C}_{1-6}-\text{alkyl,} \\
\text{or is } \text{Ci}_{6}-\text{alkyl or aryl, which are optionally substituted by} \\
\text{halo,} \\
\text{-O(CO)-C}_{1-6}-\text{alkyl,} \end{cases} \end{cases} \]
or by \(-\text{NH(CO})R\) \(^d\), wherein \(R^d\) is \(C^\wedge\text{-alky!}\) optionally substituted by halo or nitro, or \(R^d\) is aryl or a 5 or 6 membered heteroaryl, which are optionally substituted by halo, nitro, \(C^\wedge\text{-alky!}\) or \(C_1\text{-}C_6\)-haloalkyl;

\(R^4\) is one or more of \(H\), halo, \(Q\)-o-alkyl or \(C_1\text{-}C_6\)-alcohol optionally substituted by \(OH\), or two \(R^4\) may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;

\(R^5\) is \(H\), \(C_{1\text{-}6}\text{-alkyl or aryl};\)

\(R^6\) is \(H\) or \(C_1\text{-}\beta\text{-alkyl};\)

\(R^7\) is \(H\) or \(-\text{SO}_2\) \(R^5\) wherein \(R^5\) is \(C_{1\text{-}6}\text{-alkyl or aryl};\)

\(R^8\) is \(H\) or \(C_{1\text{-}6}\text{-alkyl};\)

\(X\) is \(\text{CH}_2\) or \(C=O;\)

\(B\) is halo, \(CN\), \(N\text{R}R^w\), \(C_{1\text{-}6}\text{-alkyl optionally substituted by CN, halo or }C_{1\text{-}6}\text{-alkoxy, }C_{1\text{-}6}\text{-alkoxy, }C_1\text{-}C_6\text{-haloalkoxy, }C_{3\text{-}6}\text{-cycloalkyl, }-\text{C(O)O-C}_{1\text{-}6}\text{-alkyl, }-\text{C(O) NR R}^w\text{-, }-\text{C(O)-C}_{1\text{-}6}\text{-alkyl, }-\text{S(O)}_2\text{-C}_{1\text{-}6}\text{-alkyl, }-\text{S(O)}_2\text{-}N\text{R}R^w\text{-, (C R}^m\text{R}^w\text{)}_n\text{-phenyl, or }-\text{S(O)}_2\text{-}C_{1\text{-}6}\text{-alkyl optionally substituted by one or more substituent(s) selected from the group consisting of:}\)

\(\text{halo, CN, N\text{R}R^w}, C_{1\text{-}6}\text{-alkyl optionally substituted by CN, halo or }C_{1\text{-}6}\text{-alkoxy, }C_1\text{-}C_6\text{-alkoxy, }C_{3\text{-}6}\text{-cycloalkyl, }-\text{C(O)O-C}_{1\text{-}6}\text{-alkyl, }-\text{C(O)-NR R}^w\text{-, }-\text{C(O)-C}_{1\text{-}6}\text{-alkyl, }-\text{S(O)}_2\text{-C}_{1\text{-}6}\text{-alkyl, }-\text{S(O)}_2\text{-}N\text{R}R^w\text{-;};\)

\(R^1\) and \(R^w\) are \(H\), \(C_{1\text{-}6}\text{-alkyl, }C_{1\text{-}6}\text{-alkyl-NR}^m\text{R}^w\text{-, }-\text{(C(O)O-C}_{1\text{-}6}\text{-alkyl, }-\text{C(O)-NR}^m\text{R}^w\text{-, }-\text{C(O)-C}_{1\text{-}6}\text{-alkyl, }-\text{S(O)}_2\text{-C}_{1\text{-}6}\text{-alkyl, }-\text{S(O)}_2\text{-}N\text{R}^m\text{R}^w\text{ or }OH;\)

\(R^w\) and \(R^w\) are \(H\) or \(C_{1\text{-}6}\text{-alkyl;}\)

\(m\) is 1 to 6;

\(n\) is \(O\) to 4;

as well as pharmaceutically acceptable salts thereof.

The compounds of formula (I) may contain some asymmetric carbon atoms. Accordingly, the present invention includes all stereoisomeric forms of the compounds of formula (I), including each of the individual enantiomers and mixtures thereof.

It has been found that the compounds of formula (I) have a good activity on the Via receptor. Therefore, the invention provides the use of a compound of formula (I) or a pharmaceutically acceptable salt thereof in the manufacture of medicaments for the treatment of dysmenorrhea, hypertension, chronic heart failure, inappropriate secretion of vasopressin, liver cirrhosis, nephrotic syndrome, obsessive compulsive disorder, anxious and depressive disorders.
Vasopressin is a 9 amino acid peptide mainly produced by the paraventricular nucleus of the hypothalamus. Three vasopressin receptors, all belonging to the class I G-protein coupled receptors, are known. The Via receptor is expressed in the brain, liver, vascular smooth muscle, lung, uterus and testis, the VIb or V3 receptor is expressed in the brain and pituitary gland, the V2 receptor is expressed in the kidney where it regulates water excretion and mediates the antidiuretic effects of vasopressin.


Thus vasopressin receptor antagonists are useful as therapeutics in the conditions of dysmenorrhea, hypertension, chronic heart failure, inappropriate secretion of
vasopressin, liver cirrhosis, nephrotic syndrome, obsessive compulsive disorder, anxious and depressive disorders.

The preferred indications with regard to the present invention are the treatment of anxious and depressive disorders.

As used herein, the term "aryl" means a monovalent cyclic aromatic hydrocarbon moiety consisting of a mono-, bi- or tricyclic aromatic ring. Examples of aryl moieties include, but are not limited to, optionally substituted phenyl, naphthyl, phenanthryl, fluorenyl, indenyl, pentalenyl, azulenyl, oxydiphenyl, biphenyl, methylenediphenyl, aminodiphenyl, diphenylsulfidyl, diphenylsulfonyl, diphenylisopropylidenyl, benzodioxanyl, benzofuranyl, benzodioxylyl, benzopyranyl, benzoxazinyl, benzoxazinonyl, benzopiperadinyl, benzopiperazinyl, benzopyrrolidinyl, benzomorpholinyl, methylenedioxyphenyl, ethylenedioxyphenyl, as well as those specifically illustrated by the examples herein below. Substituents for aryl include but are not limited to halogen, C1-6-alkyl, C1-6-alkoxy as well as those specifically listed and illustrated by the description and examples herein below. Preferred aryl are phenyl and naphthyl and still preferably phenyl.

The term "C1-6-alkyl" denotes a saturated straight- or branched-chain group containing from 1 to 6 carbon atoms, for example, methyl, ethyl, propyl, isopropyl, n-butyl, i-butyl, 2-butyl, t-butyl and the like. Preferred C1-6-alkyl groups are C1-4-groups, i.e. with 1-4 carbon atoms.

The term "C1-6-alkoxy" denotes a group wherein the alkyl residues are as defined above, and which is attached via an oxygen atom. Preferred C1-6-alkoxy groups are methoxy and ethoxy as well as those specifically illustrated by the examples herein below.

The term "C2-6-alkenyl" denotes a carbon chain of 2 to 6 carbon atoms comprising a double bond in its chain. C2-6-alkenyl groups include ethenyl, propen-1-yl, propen-2-yl, buten-1-yl, buten-3-yl, penten-1-yl, penten-2-yl, penten-3-yl, penten-4-yl, hexen-1-yl, hexen-2-yl, hexen-3-yl, hexen-4-yl and hexen-5-yl, as well as those specifically illustrated by the examples herein below.

The term "benzyloxy" denotes a benzyl group attached via an oxygen atom.

The term "halogen" or "halo" denotes chlorine (Cl), iodine (I), fluorine (F) and bromine (Br).
The term "Ci-6-haloalkyl" denotes a Ci-6-alkyl group as defined above which is substituted by one or more halogen. Examples of Ci-6-haloalkyl include but are not limited to methyl, ethyl, propyl, isopropyl, isobutyl, sec-butyl, tert-butyl, pentyl or n-hexyl substituted by one or more Cl, F, Br or I atom(s) as well as those groups specifically illustrated by the examples herein below. Preferred Ci-6-haloalkyl are difluoro- or trifluoro-methyl or ethyl.

"Ci-6-haloalkoxy" denotes a C1-6-alkoxy group as defined above which is substituted by one or more halogen. Examples of C1-6-haloalkoxy include but are not limited to methoxy or ethoxy, substituted by one or more Cl, F, Br or I atom(s) as well as those groups specifically illustrated by the examples herein below. Preferred C1-6-haloalkoxy are difluoro- or trifluoro-methoxy or ethoxy.

The term "C3-6-cycloalkyl" denotes a monovalent or divalent saturated carbocyclic moiety consisting of a monocyclic ring. Cycloalkyl can optionally be substituted with one, two, three or four substituents, wherein each substituent is independently hydroxy, Ci-6-alkyl, C1-6-alkoxy, halogen, amino, unless otherwise specifically indicated. Examples of cycloalkyl moieties include optionally substituted cyclopropyl, optionally substituted cyclobutyl, optionally substituted cyclopentyl and optionally substituted cyclohexyl as well as those specifically illustrated by the examples herein below.

The term "3 to 7 membered heterocycloalkyl" means a monovalent saturated moiety, consisting of one ring of 3 to 7 atoms as ring members, including one, two, or three heteroatoms chosen from nitrogen, oxygen or sulfur, the rest being carbon atoms. 3 to 7 membered heterocycloalkyl can optionally be substituted with one, two, three or four substituents, wherein each substituent is independently hydroxy, C1-6-alkyl, C1-6-alkoxy, Ci-6-thioalkyl, halo, C1-6-haloalkyl, Ci-6-hydroxyalkyl, alkoxy, Ci-6-carbalkylamino, di(C1-6)alkylamino, aminocarbonyl, or carbamoyl, unless otherwise specifically indicated. Examples of heterocyclic moieties include, but are not limited to, oxirane, optionally substituted oxetane, optionally substituted tetrahydro-furan, optionally substituted piperidinyl, optionally substituted pyrrolidinyl, optionally substituted morpholinyl, optionally substituted piperazinyl, optionally substituted azepane or homopiperazine, and the like or those which are specifically exemplified herein. Substituents can be selected from Ci-6-alkyl, Ci-6-alkoxy, Ci-6-haloalkyl, halo, CN, OH, NH₂, as well as those substituents which are are specifically illustrated in the examples hereinafter.
The term "5 or 6 membered heteroaryl" means an aromatic ring of 5 or 6 ring atoms as ring members containing one, two, or three ring heteroatoms selected from N, O, or S, the rest being carbon atoms. 5 or 6 heteroaryl can optionally be substituted with one, two, three or four substituents, wherein each substituent is independently hydroxy, C<sub>1-6</sub>-alkyl, C<sub>1-6</sub>-alkoxy, Ci-6-thioalkyl, halo, Ci-6-haloalkyl, Ci-6-hydroxyalkyl, alkoxy carbonyl, amino, Ci-6-alkylamino, di(C<sub>1-6</sub>)alkylamino, aminocarbonyl, or carbamylamino, unless otherwise specifically indicated. Examples of heteroaryl moieties include, but are not limited to, optionally substituted imidazolyl, optionally substituted oxazolyl, optionally substituted thiazolyl, optionally substituted pyrazinyl, optionally substituted pyrolyl, optionally substituted pyrazinyl, optionally substituted pyridinyl, optionally substituted pyrimidinyl, optionally substituted furanyl, and those which are specifically exemplified herein.

The term "sulfonylaryl" denotes an aryl group as defined hereinabove which is attached via a sulfonyl group.

The expression "two R<sub>2</sub> may form an oxo or dioxo bridge together with the indole ring to which they are attached notes an oxo or dioxo bridge of the following formulae:

![O]

or

![O]

which bind two adjacent carbon atoms of the phenyl or indole ring of the compound of formula (I) to which either R<sub>2</sub> is binding.

Examples of group illustrating the expression "R<sub>1</sub> and R<sub>3</sub> together with the indole ring to which they are attached form a 5 or 6 membered heterocycloalkyl which can be substituted by =0. C(O)O-C<sub>1-6</sub>-alkyl or Q-e-alkyl" are:

![N]

or

![N]

as well as those specifically illustrated by the examples.

The term "pharmaceutically acceptable acid addition salts" embraces salts with inorganic and organic acids, such as hydrochloric acid, nitric acid, sulfuric acid, phosphoric acid, citric acid, formic acid, fumaric acid, maleic acid, acetic acid, succinic
acid, tartaric acid, methane-sulfonic acid, p-toluenesulfonic acid, as well as those specifically illustrated by the examples herein below.

In a certain embodiment, the compounds of formula (I) are those compounds wherein:

$R^1$ is $H$,

or is C1-C6-alkyl optionally substituted by CN,

or is aryl, 5 or 6 membered heteroaryl or sulfonylaryl which are optionally substituted by one or more B,

or is -(CH$_2$)$_m$-$R^a$ wherein $R^a$ is:

OR$_1$,

CN,

NR$_1$R$_2$,

C$_3$-$C_6$-cycloalkyl, 3 to 7 membered-heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,

or is -(CH$_2$)$_n$-(CO)-R$_b$ or -(CH$_2$)$_n$-(SO$_2$)-R$_b$, wherein $R_b$ is:

C$_1$-$C_6$-alkoxy,

NR$_3$R$_4$,

4 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,

or $R^1$ and $R^3$ together with the indole ring to which they are attached form a 5 or 6 membered heterocycloalkyl which can be substituted by =O, C(O)O-C$_1$-$C_6$-alkyl or C$_1$-$C_6$-alkyl;

$R^2$ is one or more of H, OH, halo, C$_1$-$C_6$-alkyl optionally substituted by -NR$_m$R$_b$, C$_1$-$C_6$-alkoxy;

$R^3$ is $H$,

or is halo

or is -(CO)-R$_c$ wherein $R_c$ is:

C$_1$-$C_6$-alkyl

-(CH$_2$)$_m$VNR$_1$R$_1$,

-(CH$_2$)$_n$NR$_m$R$_b$,

5 or 6 membered heterocycloalkyl optionally substituted by C$_1$-$C_6$-alkyl, or is C$_1$-$C_6$-alkyl or aryl, which are optionally substituted by halo,
R^4 is one or more of H, halo, or C_i_6^-alkoxy optionally substituted by OH, or two R^4 may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;

R^5 is H or aryl;

R^6 is H;

R^7 is H or -SO_2^-R^6 wherein R^6 is C_i_6^-alkyl or aryl;

R^8 is H or C_1-β-alkyl;

X is CH_2 or C=O;

B is halo, CN, NH_2, C_i-6^-alkyl optionally substituted by CN or C_i-6^-alkoxy, C_1-6^-alkoxy, C_i-6^-cycloalkyl, -C(O)O-C_1-6^-alkyl, -(C=O)O-C_1-6^-alkyl, -(C(O)O)_{n-phenyl}, wherein the phenyl is optionally substituted by one or more substituents selected from the group consisting of:

halo, C_i-6^-alkyl optionally substituted by CN or halo, C_i-6^-alkoxy;

R^1 and R^u are H, C_1-6^-alkyl, C_i-6^-alkyl-NR^mR^l, -(CO)O-C_1-6^-alkyl, -(C(O)O)_{n-phenyl}, -(C(O)-NR^mR^l, -C(O)-C_1-6^-alkyl, -S(O)_{2-C_i-6^-alkyl} or -S(O)_{2-NR^mR^l} or OH;

R^u and R^v are H or C_1-6^-alkyl;

m is 1 to 6;

n is 0 to 4;

as well as pharmaceutically acceptable salts thereof.

In another embodiment, the compounds of formula (I) are those compounds wherein:

A is selected from (a), (b), (c), (d) or (e), and wherein

R^1 is H,

or is C_i-6^-alkyl optionally substituted by CN,

or is aryl, 5 or 6 membered heteroaryl or sulfanylaryl which are optionally substituted by one or more B,

or is -(CH_2)_{m-R^a} wherein R^a is:

CN,

OR^1,

NR^mR^l,

C_3-6^-cycloalkyl, 4 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,

or is -(CH_2)_{n-(CO)-R^b} or -(CH_2)_{n-(SO_2)-R^b}, wherein R^b is:

C_i-β-alkyl,

C_i-6^-alkoxy,
C₅₋₆-cycloalkyl,
- (CH₂)ₘ-NRₘRᵣ⁻,
NRᵣ⁻,
C₅₋₆-cycloalkyl, 4 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
or R¹ and R³ together with the indole ring to which they are attached form a 5 or 6 membered heterocycloalkyl which can be substituted by (CO);
R² is one or more of H, OH, halo, CN, nitro, C₁₋₆-alkyl optionally substituted by -NRᵐRᵣ⁻, C₁₋₆-alkoxy, -O-CH₂-C₂₋₆-alkenyl, benzyloxy,
or two R² may form an oxo or dioxo bridge together with the indole ring to which they are attached;
R³ is H,
or is halo,
or is -(CO)-Rₖ wherein Rₖ is:
C₁₋₆-alkyl,
- (CH₂)ₙ-NRᵣ⁻,
- (CH₂)ₙ-NRᵐRᵣ⁻,
5 or 6 membered heterocycloalkyl optionally substituted by C₁₋₆-alkyl,
or is C₁₋₆-alkyl or aryl, which are optionally substituted by
halo,
-O(CO)-C₁₋₆-alkyl,
or by-NH(CO)Rᵈ wherein Rᵈ is C₁₋₆-alkyl optionally substituted by halo or nitro, or Rᵈ is aryl or a 5 or 6 membered heteroaryl, which are optionally substituted by halo, nitro, C₁₋₆-alkyl or C₁₋₆-haloalkyl;
R⁴ is one or more of H, halo, C₁₋₆-alkyl or C₁₋₆-alkoxy or two R⁴ may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;
R⁵ is H, C₁₋₆-alkyl or aryl;
R⁶ is H or C₁₋₆-alkyl;
B is
halo, CN, NRᵐRᵣ⁻, C₁₋₆-alkyl optionally substituted by CN, halo or C₁₋₆-alkoxy, C₁₋₆-alkoxy, C₁₋₆-haloalkoxy, C₅₋₆-cycloalkyl, -C(O)-C₁₋₆-alkyl, -C(O) NRᵐRᵣ⁻, -C(O)-C₁₋₆-alkyl, -S(O)₂-C₁₋₆-alkyl, -S(O)₂ NRᵣ⁻, (CRᵐRᵣ⁻)ₙ-phenyl, or (CRᵐRᵣ⁻)ₙ-5 or 6 membered heteroaryl wherein the phenyl or 5 or 6 membered heteroaryl moiety is optionally substituted by one or more substituent(s)
selected from the group consisting of:
halo, CN, NR\textsuperscript{R}, Ci\textsubscript{6}-alkyl optionally substituted by CN or Ci\textsubscript{6}-alkoxy,
Ci\textsubscript{6}-alkoxy, Q-e-haloalkoxy, C\textsubscript{3-6}-cycloalkyl, -C(O)-O-C\textsubscript{1-6}-alkyl, -C(O)-
NR\textsuperscript{R}, -C(O)-Ci\textsubscript{6}-alkyl, -S(O)\textsubscript{2}Ci\textsubscript{6}-alkyl, -S(O)\textsubscript{2}-NR\textsuperscript{R};
\text{R\textsuperscript{1}} and \text{R\textsuperscript{1}} are H, C\textsubscript{1-6}-alkyl, Ci\textsubscript{6}-alkyl-NR\textsubscript{m}RV,
-C(O)O-C\textsubscript{1-6}-alkyl, -C(O)-NR\textsubscript{m}RV, -C(O)-Ci\textsubscript{6}-alkyl,
-S(O)\textsubscript{2}-Ci\textsubscript{6}-alkyl or -S(O)\textsubscript{2}-NR\textsuperscript{R}R; 
\text{R\textsuperscript{v}} and \text{R\textsuperscript{v}} are H or C\textsubscript{1-6}-alkyl;
\text{m} is 1 to 6;
\text{n} is O to 4;
as well as pharmaceutically acceptable salts thereof.

In another embodiment, the compounds of formula (I) are those compounds wherein:
\text{A} is selected from (a), (b), (c), (d) or (e), and wherein
\text{R\textsuperscript{1}} is H,
or is Ci\textsubscript{6}-alkyl,
or is aryl, 5 or 6 membered heteroaryl or sulfonylaryl which are optionally
substituted by one or more B,
or is -(CH\textsubscript{2})\textsuperscript{m}-R\textsuperscript{3} wherein \text{R\textsuperscript{3}} is:
CN,
NR\textsuperscript{R},
C\textsubscript{3-6}-cycloalkyl, 4 to 7 membered-heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally
substituted by one or more B,
or is -(CH\textsubscript{2})\textsuperscript{n}-(CO)-R\textsuperscript{5} or -(CH\textsubscript{2})\textsuperscript{n}-(SO\textsubscript{2})-R\textsuperscript{5}, wherein \text{R\textsuperscript{5}} is:
C\textsubscript{1-6}-alkoxy,
NR\textsuperscript{R},
4 to 7 membered-heterocycloalkyl, aryl, which are optionally substituted by
one or more B,
or \text{R\textsuperscript{1}} and \text{R\textsuperscript{3}} together with the indole ring to which they are attached form a 5 or
6 membered heterocycloalkyl which can be substituted by (CO);
\text{R\textsuperscript{2}} is one or more of H, OH, halo, C\textsubscript{1-6}-alkyl optionally substituted by -NR\textsuperscript{m}RV, C\textsubscript{1-6}-
alkoxy;
\text{R\textsuperscript{3}} is H,
or is -(CO)-R\textsuperscript{C}, wherein \text{R\textsuperscript{C}} is:
-(CH\textsubscript{2})\textsuperscript{m}VNR\textsuperscript{R},
-(CH\textsubscript{2})\textsuperscript{n}NR\textsuperscript{m}R,
5 or 6 membered heterocycloalkyl optionally substituted by C\textsubscript{1-6}-alkyl,
or is C1-6-alkyl or aryl, which are optionally substituted by halo,

R4, R5 and R6 are H;

R7 is H or -SO2-R6 wherein R6 is C1-6-alkyl or aryl;

B is halo, NH2, Q-o-alkyl optionally substituted by CN or C1-6-alkoxy, C1-6-alkoxy,

5 Ci-e-haloalkoxy, C3-6-cycloalkyl, -C(O)O-C1-6-alkyl, -(CRmRn)-phenyl, wherein

the phenyl is optionally substituted by one or more substituent(s) selected from

the group consisting of:

halo, C1-6-alkyl optionally substituted by CN or halo, C1-6-alkoxy;

R1 and R6 are H, Ci-1-alkyl, Ci-6-alkyl-NRmRn, -(CO)O-Ci-6-alkyl, -(C(O)O)NRmRn, -(C(O)-

10 Ci-6-alkyl, -S(O)2-Ci-6-alkyl or -S(O)2-NRmRn;

Rm and Rn are H or Ci-6-alkyl;

m is 1 to 6;

n is O to 4;

as well as pharmaceutically acceptable salts thereof.

In another embodiment, the compound of formula (I) are those compounds

wherein:

A is selected from (a), (b), (c), (d) or (e), and wherein

R1 is H or,

Ci-6-alkyl optionally substituted by CN or,

20 Ci-6-alkoxy or,

aryl or,

5 or 6 membered heteroaryl or,

sulfonylaryl or,

-(CH2)m-R6 wherein R6 is C3-6-cycloalkyl, 5 or 6 membered-heterocycloalkyl,

25 aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or

more substituents selected from the group consisting of:

halo, CN, Ci-6-alkyl, Ci-6-alkoxy, C1-6-haloalkoxy, -(C(O)O)-C1-6-alkyl

and phenyl optionally substituted by halo, Q-o-alkyl, Ci-6-haloalkyl or

C1-0-alkoxy,

30 -(CH2)n-NR-R6 or,

-(CH2)n-(CO)-Rb wherein Rb is aryl or 5 or 6 membered-heterocycloalkyl;

R2 is one or more of H, halo, CN, nitro, C1-6-alkyl, C1-6-alkoxy, -O-CH2-C2-6-alkenyl,

35 benzyl, or two R2 may form an oxo or dioxo bridge together with the indole

ing to which they are attached;

R3 is H or,
halo or, 
-(CO)-R C, wherein R C is C 1-6 -alkyl, 5 or 6 membered heterocycloalkyl optionally substituted by C 1-6 -alkyl, or R C is -(CH 2 ) VR 1 or, 
Ci-6-alkyl or aryl, which are optionally substituted by: 
-O(CO)-C 1-6 -alkyl, 
or by-NH(CO)R d, wherein R d is C^-alkyl optionally substituted by halo or nitro, or R d is aryl or a 5 or 6 membered heteroaryl, which are optionally substituted by halo, nitro, C^-alkyl or C 176-haloalkyl; 
R 4 is one or more of H, halo, Ci-6-alkyl or C 176-alkoxy or two R 4 may form an oxo or dioxo bridge together with the phenyl ring to which they are attached; 
R 5 is H, Ci-6-alkyl or aryl; 
R 6 is H or C 1-β-alkyl; 
R 7 is H or -SO 2 -R e wherein R e is Ci-6-alkyl or aryl; 
R 1 and R v are independently selected from H, C^-alkyl! or -(CO)O-C 1-6 -alkyl; 
m is 1 to 6; 
n is 0 to 4; 
as well as pharmaceutically acceptable salts thereof. 

As it can be seen from the definition of A in the compounds of formula (I), said compounds of formula (I) encompass the compounds of formulae (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) and (I-h) as follows:
wherein R\textsuperscript{1} to R\textsuperscript{8} and X are as defined hereinabove in connection with formula (I).

In a certain embodiment the compounds of the invention are those compounds of formula (I-a):

\[
\text{(I-a)}
\]

wherein the dotted line is either nil or a double bond;

R\textsuperscript{1} is H,

or is Ci-6-alkyl optionally substituted by CN,

or is sulfonylaryl,

or is -(CH\textsubscript{2})\textsuperscript{m}-R\textsuperscript{a} wherein R\textsuperscript{a} is:

OR\textsuperscript{1},

CN,

NR\textsuperscript{1}R'\textsuperscript{1},

C\textsubscript{5-6}-cycloalkyl, 3 to 6 membered-heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,

or is -(CH\textsubscript{2})\textsuperscript{n}-(CO)-R\textsuperscript{b}, wherein R\textsuperscript{b} is:

Ci-g-alkoxy,
NR\textsuperscript{\textregistered},
6-membered-heterocycloalkyl, aryl, or 5 or 6-membered heteroaryl which are optionally substituted by one or more B;
R\textsuperscript{2} is one or more of H, halo, C\textsubscript{1-6}-alkyl;
R\textsuperscript{3} is H,
or is Ci\textsubscript{6}-alkyl,
or is -(CO)-R\textsuperscript{c}, wherein R\textsuperscript{c} is:
Ci\textsubscript{6}-alkyl
-(CH\textsubscript{z})\textsubscript{n}-NR\textsuperscript{R}\textsuperscript{v},
10 R\textsuperscript{4} is one or more of H, halo, or C\textsubscript{1-6}-alkoxy optionally substituted by OH, or two R\textsuperscript{4} may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;
R\textsuperscript{5} is H;
R\textsuperscript{6} is H;
15 B is halo, CN, Q-0-alkyl optionally substituted by CN or C\textsubscript{1-6}-alkoxy, C\textsubscript{1-6}-alkoxy, C\textsubscript{1-6}-haloalkoxy, C\textsubscript{3-6}-cycloalkyl, -C(O)O-C\textsubscript{1-6}-alkyl, -(CR\textsuperscript{m}R\textsuperscript{iv})\textsubscript{n}-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:
halo, C\textsubscript{1-6}-alkyl optionally substituted by CN or halo, C\textsubscript{1-6}-alkoxy;
20 R\textsuperscript{1} and R\textsuperscript{iv} are H, C\textsubscript{1-6}-alkyl, Ci\textsubscript{6}-alkyl-NR\textsuperscript{R}\textsuperscript{v}, -C(O)-C\textsubscript{1-6}-alkyl, -S(O)\textsubscript{2}-Ci\textsubscript{6}-alkyl or OH;
R\textsuperscript{iv} and R\textsuperscript{v} are H or C\textsubscript{1-6}-alkyl;
m is 1 to 6;
n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

Further, in a certain embodiment the compounds of the invention are those compounds of formula (I-a) as described above, wherein
R\textsuperscript{1} is H or,
-(CH\textsubscript{2})\textsubscript{m}-R\textsuperscript{a} wherein R\textsuperscript{a} is aryl which is optionally substituted by one or more substituents selected from the group consisting of:
halo, CN, C\textsubscript{1-6}-alkyl, Ci\textsubscript{6}-alkoxy, Ci-e-haloalkoxy, -C(O)O-C\textsubscript{1-6}-alkyl
and phenyl optionally substituted by halo, C\textsubscript{1-6}-alkyl, Ci\textsubscript{6}-haloalkyl or C\textsubscript{1-0}-alkoxy;
R\textsuperscript{2} is H or halo;
R\textsuperscript{3} is H or Q-6-alkyl; and
35 R\textsuperscript{4}, R\textsuperscript{5} and R\textsuperscript{6} are H;
m is 1 to 6; as well as pharmaceutically acceptable salts thereof.

The following compounds are examples according to the invention:

l′-[1-benzyl-2-methyl-lH-indol-3-yl]carbonyl]spiro[indene-l,4′-piperidine];

1′-[(1-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-2,3-dihydrospiro[indene-1,4′-piperidine] ;

l′-[1-benzyl-lH-indol-3-yl]carbonyl]spiro[indene-l,4′-piperidine];
r-[(2-methyl-lH-indol-3-yl)carbonyl]spiro[indene-l,4′-piperidine];
r-[(6-chloro-lH-indol-3-yl)carbonyl]spiro[indene-l,4′-piperidine]; and

r-[(6-chloro-lH-indol-3-yl)carbonyl]-2,3-dihydrospiro[indene-1,4′-piperidine].

In a certain embodiment the compounds of the invention are those compounds of formula (I-b):

![Chemical Structure](image)

R\(^1\) is H,

or is C\(_1-6\)-alkyl optionally substituted by CN,

or is sulfonylaryl,

or is -(CH\(_2\))\(_m\)-R\(^a\) wherein R\(^a\) is:

OR\(^1\),

CN,

NR\(^1\)R\(^2\),

C\(_3-6\)-cycloalkyl, 3 to 6 membered-heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,

or is -(CH\(_2\))\(_n\)-(CO) R\(^b\), wherein R\(^b\) is:

C\(_1-6\)-alkoxy,

NR\(^1\)R\(^2\),
5 or 6 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B;

R² is one or more of H, halo, C₁₋₆-alkyl;
R³ is H,
5 or is C₁₋₆-alkyl,
or is -(CO)-R⁶, wherein R⁶ is:
C₁₋₆-alkyl
-(CH₂)ₙ-NR'R'',
R⁴ is is one or more of H, halo, or C₁₋₆-alkoxy optionally substituted by OH, or two R⁴
10 may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;
R⁶ is H;
R⁷ is H or -SO₂-R⁶ wherein R⁶ is C₁₋₆-alkyl or aryl;
B is halo, CN, C₁₋₆-alkyl optionally substituted by CN or C₁₋₆-alkoxy, C₁₋₆-alkoxy, C₁₋₆-haloalkoxy, C₁₋₆-cycloalkyl, -(CO)O-C₁₋₆-alkyl, -(CR₃R⁴)ₙ-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:
halo, C₁₋₆-alkyl optionally substituted by CN or halo, C₁₋₆-alkoxy;
R¹ and R'' are H, C₁₋₆-alkyl, C₁₋₆-alkyl-NR³R⁴, -(CO)-C₁₋₆-alkyl, -S(O)₂-C₁₋₆-alkyl or OH;
20 R'' and R'''' are H or C₁₋₆-alkyl;
m is 1 to 6;
n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

Further, in a certain embodiment the compounds of the invention are those compounds of formula (I-b) as described above, wherein
R¹ is H,
or is -(CH₂)ₘ-Rₐ wherein Rₐ is aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more substituents selected from the group
30 consisting of:
halo, CN, C₁₋₆-alkyl, C₁₋₆-alkoxy, C₁₋₆-haloalkoxy, -(CO)O-C₁₋₆-alkyl
and phenyl optionally substituted by halo, C₁₋₆-haloalkyl or C₁₋₆-alkoxy,
or is -(CH₂)ₙ-NR'R'',
or is -(CH₂)ₙ-(CO)-Rₐ, wherein Rₐ is aryl or 5 or 6 membered-heterocycloalkyl
which are optionally substituted by one or more B.
or is \((\text{CH}_2)_n\)(CO)\(\text{R}^b\) wherein \(\text{R}^b\) is \(\text{N}\text{R}^1\text{R}^1\), \(\text{R}^2\) is H or halo;
\(\text{R}^3\) is H or \(\text{C}\_1\_6\)-alkyl;
\(\text{R}^4\) is H or halo;
\(\text{R}^5\) is H;
\(\text{R}^6\) is H or \(-\text{SO}_2\text{R}^c\) wherein \(\text{R}^c\) is \(\text{Ci}\_1\_6\)-alkyl or aryl;
\(\text{B}^\text{is}\) halo, \(\text{NH}_2\), Q-\(\text{o}\)-alkyl optionally substituted by \(\text{CN}\) or \(\text{C}\_1\_6\)-alkoxy, \(\text{C}\_1\_6\)-alkoxy, \(\text{Ci}\_1\_6\)-haloalkoxy, \(\text{C}\_3\_6\)-cycloalkyl, \(-\text{C}(\text{O})\text{O}-\text{C}\_1\_6\)-alkyl, \(-\text{(CR}^m\text{R}^n\}_\text{n}\)-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:
\[
\text{halo, Ci}\_1\_6\text{-alkyl optionally substituted by CN or halo, C}\_1\_6\text{-alkoxy;}
\]
\(\text{R}^1\), \(\text{R}^1\) are independently selected from H or \(\text{C}\_1\_6\)-alkyl;
\(\text{m}\) is 1 to 6;
\(\text{n}\) is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

In another embodiment thereof,
\(\text{R}^1\) is H or,
\(-\text{(CH}_2\}_\text{m}\)-\(\text{R}^a\) wherein \(\text{R}^a\) is aryl which is optionally substituted by one or more substituents selected from the group consisting of:
\[
\text{halo, CN, Q-6-alkyl, Ci}\_6\text{-alkoxy, Ci-e-haloalkoxy, -C(O)O-C}\_1\_6\text{-alkyl and phenyl optionally substituted by halo, Ci}\_1\_6\text{-haloalkyl or C}\_1\_6\text{-alkoxy;}
\]
\(\text{R}^2\) is H or halo;
\(\text{R}^3\) is H or \(\text{C}\_1\_6\)-alkyl;
\(\text{R}^4\) is H or halo;
\(\text{R}^6\) is H;
\(\text{R}^7\) is H or \(-\text{SO}_2\text{R}^c\) wherein \(\text{R}^c\) is \(\text{C}\_1\_6\)-alkyl;
\(\text{m}\) is 1 to 6;
as well as pharmaceutically acceptable salts thereof.
The following compounds are examples according to the invention:

\(\text{l'-(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]\ -l-(methylsulfonyl)-1,2-dihydrospiro[indole-3,4'-piperidine];\)
\(\text{r'-(6-chloro-1-(3-fluorobenzoyl)-1H-indol-3-yl)carbonyl]}-1,2\text{-dihydrospiro[indole-3,4'-piperidine];}\)
Preferred are the following compounds:

1'-[(6-chloro-lH-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4'-piperidine];
1'-[(l-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4'-piperidine];
r-lfo-chloro-l-CS^-difluorobenzoyO-lH-indol-S-ylcarbonylJ-l^-dihydropirofindole-
3,4'-piperidine;  
2-[6-chloro-3-(l,2-dihydro-l H -spiro[indole-3,4'-piperidin]-r-ylcarbonyl)-lH-indol-l-
yl]-l-(3,5-difluorophenyl)ethanone;  
2-[6-chloro-3-(l,2-dihydro-l H -spiro[indole-3,4'-piperidin]-r-ylcarbonyl)-lH-indol-l-
yl]-l-(3,4-difluorophenyl)ethanone;  
2-[6-chloro-3-(l,2-dihydro-l H -spiro[indole-3,4'-piperidin]-r-ylcarbonyl)-lH-indol-l-
yl]-l-(2-fluorophenyl)ethanone;  
2-[6-chloro-3-(l,2-dihydro-l H -spiro[indole-3,4'-piperidin]-r-ylcarbonyl]-lH-indol-l-
yl]-N,N-diethylethanamine; and

r-{[6-chloro-l-(pyridin-2-ylmethyl)-lH-indol-3-yl]carbonyl]-1,2-dihydrospiro[indole-
3,4'-piperidine].
In a certain embodiment the compounds of the invention are those compounds of formula (I-c):

\[
\begin{align*}
\text{R}^1 & \text{is } H, \\
& \text{or is } C_{1-6}-\text{alkyl optionally substituted by CN,} \\
& \text{or is aryl, 5 or 6 membered heteroaryl or sulfonylaryl which are optionally substituted by one or more B,} \\
& \text{or is } -(\text{CH}_2)_n-R^a \text{ wherein } R^a \text{ is:} \\
& \quad \text{OR}, \\
& \quad \text{CN}, \\
& \quad \text{NR}^mR^n, \\
& \quad C_{3-6}\text{-cycloalkyl, 4 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,} \\
& \text{or is } -(\text{CH}_2)_n-(\text{CO})-R^b \text{ wherein } R^b \text{ is:} \\
& \quad \text{Ci}_{6}\text{-alkoxy,} \\
& \quad \text{NR}^mR^n, \\
& \quad 4 \text{ to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,} \\
& \text{or } R^1 \text{ and } R^3 \text{ together with the indole ring to which they are attached form a 5 or 6 membered heterocycloalkyl which can be substituted by } =0; \\
\text{R}^2 & \text{is one or more of H, halo, C\text{-alkyl optionally substituted by } NR^mR^n, C_{1-6}\text{-alkoxy;}} \\
\text{R}^3 & \text{is } H, \\
& \text{or is } C_{1-6}\text{-alkyl,} \\
& \text{or is halo,} \\
& \text{or is } -(\text{CO})-R^c \text{ wherein } R^c \text{ is:} \\
& \quad \text{C}_{1-6}\text{-alkyl,}
\end{align*}
\]
-(CH₂)ₙ-NR₁R₂,
-(CH₂)ₙ-NR₃R₄,
5 or 6 membered heterocycloalkyl optionally substituted by C₁₋₆-alkyl;
R⁴ is one or more of H, halo, or C₁₋₆-alkoxy optionally substituted by OH, or two R⁴ may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;
B is halo, NH₂, Q-o-alkyl optionally substituted by CN or C₁₋₆-alkoxy, C₁₋₆-alkoxy, Ci-e-haloalkoxy, C₃₋₆-cycloalkyl, -C(O)O-C₁₋₆-alkyl, -(CR₃R⁴)ₙ-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:
halo, C₁₋₆-alkyl optionally substituted by CN or halo, C₁₋₆-alkoxy;
R¹ and R⁵ are H, C₁₋₆-alkyl, Ci₋₆-alkyl-NR₃R⁴, -(CO)O-C₁₋₆-alkyl, -(CO)O-C₁₋₆-NR₃R⁴, -C(O)-Ci₋₆-alkyl, -S(O)₂Ci₋₆-alkyl, -S(O)₂R₁₋₆-NR₃R⁴;
R⁵ and R⁶ are H or C₁₋₆-alkyl;
m is 1 to 6;
n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

In a further embodiment thereof,
R¹ is H or,
C₁₋₆-alkyl optionally substituted by CN or,
-(CH₂)ₙ-R₃ wherein R₃ is aryl which is optionally substituted by one or more substituents selected from the group consisting of:
halo, CN, C₁₋₆-alkyl, C₁₋₆-alkoxy, C₁₋₆-haloalkoxy, -C(O)O-C₁₋₆-alkyl and phenyl optionally substituted by halo, Ci₋₆-haloalkyl or C₁₋₆-alkoxy;
R² is H or halo;
R³ is H, C₁₋₆-alkyl or,
-(CO)-R₄, wherein R₄ is 5 or 6 membered heterocycloalkyl optionally substituted by Q-6-alkyl, or R₄ is -(CH₂)ₙVNRᵣ;
R⁴ is H, halo, Ci₋₆-alkoxy or two R⁴ may form a dioxo bridge together with the phenyl ring to which they are attached;
R¹ and R⁵ are independently selected from H, Ci₋₆-alkyl and -(CO)O-C₁₋₆-alkyl;
m is 1 to 6;
n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.
The following compounds are examples according to the invention:
r-[(1-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
4- {6-chloro-3- [(3-oxo-1'H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-lH-indol-1-yl}butanenitrile;
5- {6-chloro-3-[(3-oxo-1'H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-lH-indol-1-yl}propanenitrile;
1'-{(1-2-(3,4-dimethoxyphenyl)ethyl]-5-methoxy-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-{(1-(4-ethoxyphenyl)-5-methoxy-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
5-bromo-r-{{6-chloro-1-(2-fluorobenzoyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
5-bromo-l'-{{6-chloro-1-(2,3-difluorobenzoyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
S-bromo-l'-do-chloro-l-fCS^-difluoropheny^sulfonyy-lH-indol-S-ylJcarbony^-SH-spiro[2-benzofuran-1,4'-piperidin]-3-one; and
l'-[(1-biphenyl-3-yl-6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one.

Preferred are the following compounds:
l'-[(6-chloro-lH-indol-3-yl)carbonyl]-6-methoxy-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
6-chloro-r-{{6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
r-{{6-chloro-2-[(4-methylpiperidin-1-yl)carbonyl]-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
6-chloro-3-[(5-fluoro-3-oxo-1'H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-y]carbonyl]-1H-indole-2-carboxamide;
6-chloro-N-2-(dimethylamino)ethyl]-3-[(5-fluoro-3-oxo-1'H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-y]carbonyl]-1H-indole-2-carboxamide;
l'-[(6-chloro-2-(piperazin-1-ylcarbonyl)-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
r-{{6-chloro-2-(morpholin-4-ylcarbonyl)-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
r-{{6-chloro-2-[(4-methylpiperazin-1-yl)carbonyl]-lH-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]-5-bromo-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
S-bromo-r-fC 6-chloro-lH-indol-S-y^carbonyy-SH-spirofl-benzofuran-l^'-piperidin]-
3-one;
1'-{[6-chloro-l-(3-fluorobenzoyl)-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
5-bromo-r-{[6-chloro-l-(3-fluorobenzoyl)-1H-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
1'-(r-{6-chloro-l-[3-fluoro-5-(3,5-difluorobenzoyl)]-1H-indol-3-yl}carbonyl)-5-fluoro-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one;
1'-(r-{6-chloro-l-(2-fluorobenzoyl)-1H-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-
1,4'-piperidin]-3-one;
r-((r-{6-chloro-l-(3,5-difluorobenzoyl)]-1H-indol-3-yl}carbonyl)-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one;
1'-(r-{6-chloro-l-(2-fluorobenzoyl)-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one;
r-((r-{6-chloro-l-(3,5-difluorobenzoyl)]-1H-indol-3-yl}carbonyl)-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one;
1'-(r-{6-chloro-l-(2-fluorobenzoyl)-1H-indol-3-yl]carbonyl]-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one;
r-((r-{6-chloro-l-(3,5-difluorobenzoyl)]-1H-indol-3-yl}carbonyl)-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one;
5-bromo-r-([6-chloro-l-(3,5-difluorobenzoyl)]-1H-indol-3-yl]carbonyl]-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one;
1'-(r-{[4-amino-2-(methoxymethyl)pyrimidin-5-yl]methyl}-6-chloro-lH-indol-3-
yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-(r-{(1-[4-amino-2-methylpyrimidin-5-yl]methyl}-6-chloro-lH-indol-3-yl]carbonyl]-5-
fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
5-bromo-r-{[6-chloro-l-(3,5-difluorobenzoyl)]-1H-indol-3-yl]carbonyl]-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one;
1'-(r-{[4-amino-2-(methoxymethyl)pyrimidin-5-yl]methyl}-6-chloro-lH-indol-3-
yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
r-((r-{1-[4-amino-2-methylpyrimidin-5-yl]methyl}-6-chloro-lH-indol-3-yl]carbonyl]-5-
fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
r-tCT-chloro-l-oxo-l^'^^'-tetrahydropyrazinofl^-aJindol-I0-y^carbonyy-S-fluoro-
3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
Particularly preferred are the following compounds:

r-[(l-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-6-chloro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-[(l-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-4-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r'-[(l-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-lH-indol-l-yl]-N-[2-(dimethylamino)ethyl] acetamide; and

r-[(6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
3. \([\text{6-chloro-3-}\{(3-OXO-1H,3H-}\text{spiro[2-benzofuran-1,4'-piperidin]}-1'-yl}\text{carbonyl}\}-1H-\text{indol-1-yl}]\text{acetonitrile;}

5. \([\text{6-chloro-3-}\{(3-}\text{o xo-1H,3H-}\text{spiro[2-benzofuran-1,4'-piperidin]}-r-yl]\text{carbonyl}]\)-1H-indol-1-yl]acetonitrile;

10. \([\text{6-chloro-1-}\{(3,5-}\text{difluorobenzyl}-1H-\text{indol-3-yl}]\text{carbonyl]}\)-1H-indol-1-yl]acetonitrile;

15. \([\text{6-chloro-1-}\{(3,5-}\text{difluorobenzyl}-1H-\text{indol-3-yl}]\text{carbonyl]}\)-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

20. \([\text{6-chloro-1-}\{(3-}\text{fluorobenzyl}-1H-\text{indol-3-yl}]\text{carbonyl]}\)-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

25. \([\text{6-chloro-1-}\{(3-}\text{fluorobenzyl}-1H-\text{indol-3-yl}]\text{carbonyl]}\)-1H-indol-1-yl]acetonitrile;

30. 5-bromo-\([\text{6-chloro-1-}\{(3-}\text{fluorobenzyl}-1H-\text{indol-3-yl}]\text{carbonyl]}\)-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

35. \([\text{6-chloro-1-}\{(6-}\text{chboro-1-}\{(3-}\text{fluorobenzyl}-1H-\text{indol-3-yl}]\text{carbonyl]}\)-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-(6-chloro-1-[2-(3,4-difluorophenyl)-2-oxoethyl]-1H-indol-3-yl)carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
5-r-lf 6-chloro-l-CS-fluoropheny^-lH-indol-S-yllcarbonylJ-SH-spirofl-benzofuran-l,.^-
piperidin]-3-one;
5-r-([6-chloro-l-(2-oxo-2-piperidin-l-ylethyl)-1H-indol-3-yl]carbonyl)-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one;
r-([6-chloro-l-(2-morpholin-4-yl-2-oxoethyl)-1H-indol-3-yl]carbonyl)-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one;
5-([3-OXO-l'1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl]carbonyl]-1H-indol-1-yl]-N,N-dimethylacetamide;
5-[3-OXO-l'1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl]carbonyl]-IH-indol-1-yl]-N,N-diethylacetamide;
5-[(6-chloro-l-pyridin-2-yl-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
5-([6-chloro-l-(2-methylpyridin-4-yl)methyl]-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
5-([6-chloro-l-(6-chloropyridin-3-yl)methyl]-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
5-([6-chloro-l-(3-chloro-6-methylpyridazin-4-yl)methyl]-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
5-([6-chloro-l-(pyridin-4-ylmethyl)-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
5-([6-chloro-l-(pyridin-4-ylmethyl)-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
5-([6-chloro-l-(pyridin-4-ylmethyl)-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
5-([6-chloro-l-(pyridin-4-ylmethyl)-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
5-([6-chloro-l-(pyridin-4-ylmethyl)-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
5-([6-chloro-l-(pyridin-4-ylmethyl)-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-
piperidin]-3-one;
r-({6-chloro-l-(pyridin-2-ylmethyl)-lH-indol-3-yl}carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
2-l6-chloro-S-fCS-fluoro-S-oxo-lH ^4H-spirofl-benzofuran-l'4'-piperidin]-l'-yl)carbonyl]-lH-indol- 1-yl]-N,N-dimethylacetamide;
5 r-((6-chloro-l-[2-(dimethylamino)ethyl]-lH-indol-3-yl)carbonyl)-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;
r-([6-chloro-l-(pyridin-3-ylmethyl)-lH-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;
r-([6-chloro-l-(pyrimidin-5-ylmethyl)-lH-indol-3-yl]carbonyl)-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;
3-6-chloro-3-[(5-fluoro-3-oxo-lH ,3H-spiro[2-benzofuran-l,4'-piperidin]-l'-yl)carbonyl]-lH-indol- 1-yl]propanenitrile;
tert-butyl (6-chloro-3-[(5-fluoro-3-oxo-lH ,3H-spiro[2-benzofuran-l,4'-piperidin]-l'-yl]carbonyl]-lH-indol- 1-yl)acetate;
l'-(6-chloro-l-(2-morpholin-4-yl-2-oxoethyl)-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;
r-((1-[4-benzylmorpholin-2-yl)methyl]-6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;
r-([6-chloro-l-(pyridin-2-ylmethyl)-lH-indol-3-yl]carbonyl)-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;
r-([6-chloro-l-(pyridin-3-ylmethyl)-lH-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;
l'-(6-chloro-1-[2-(4-methylpiperazin-l-yl)-2-oxoethyl]-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;
r-([6-chloro-l-(pyridin-2-ylmethyl)-lH-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;
l'-(6-chloro-1-[5-chloro-1H-imidazol-5-yl)methyl]-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;
l'-(6-chloro-1-[(3,5-dimethylisoxazol-4-yl)methyl]-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;
l'-(6-chloro-l-[(3,5-dimethylisoxazol-4-yl)methyl]-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;
r-({6-chloro-1-[(2,5-dimethyl-1,3-oxazol-4-yl)methyl]-1H-indol-3-yl}carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-({6-chloro-1-[(3-fluoroacetan-3-yl)methyl]-1H-indol-3-yl}carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
5
r-({6-chloro-1-[(3-fluoroacetan-3-yl)methyl]-1H-indol-3-yl}carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-({6-chloro-1-[(1-methoxymethyl)cyclopropyl]methyl}-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1-[(6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-1H-indol-3-yl]methyl)cyclopropyl] acetonitrile;
1'-[(6-chloro-1-[(1-methoxymethyl)-cyclopropyl]methyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
[1-[(6-chloro-3-3-[(5-fluoro-3-oxo-1H ,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-1H-indol-1-yl)methyl]cyclopropyl] acetonitrile;
15
r-({6-chloro-1-[2-ethoxy-2H-pyran-4-yl]ethyl}-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one hydrochloride;
1'-[(6-chloro-1-[2-ethoxy-2H-pyran-4-yl]ethyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
tert-butyl 2-{6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-1H-indol-3-yl]methyl)morpholine-4-carboxylate;
tert-butyl 2-{6-chloro-3-[(3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-1H-indol-3-yl]methyl)morpholine-4-carboxylate;
r-{{6-chloro-l-(morpholin-2-ylmethyl)-1H-indol-3-yl]carbonyl}-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one dihydrochloride;
25
{6-chloro-l-(morpholin-2-ylmethyl)-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one hydrochloride;
2-{6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-1H-indol-3-yl]acetamide;
2-{6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-1H-indol-3-yl]N-methylacetamide;
30
1'-[(6-chloro-l-(2-oxo-2-piperazin-1-ylethyl)-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
r-{{1-[(3,5-difluorobenzyl)-1H-indol-3-yl]carbonyl}]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
35
r-{{1-[(3,5-difluorobenzyl)-1H-indol-3-yl]carbonyl}]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
N,N-diethyl-2-{3-[(3-oxo-1H,3H-spiro[2-benzofuran-1,4’-piperidin]-r-yl)carbonyl]-1H-indol-1-yl}acetamide; and
2-{6-chloro-5-methyl-3-[(3-oxo-1H,3H-spiro[2-benzomran-1,4’-piperidin]-r-yl)carbonyl]-1H-indol-1-yl}-N,N-dimethylacetamide.

In a certain embodiment the compounds of the invention are those compounds of formula (I-d):

![Chemical Structure](image)

wherein

10 R^1 is H,
or is C\textsubscript{i}-C\textsubscript{6}-alkyl optionally substituted by CN,
or is aryl, 5 or 6 membered heteroaryl or sulfonylaryl which are optionally substituted by one or more B,
or is -(CH\textsubscript{2})\textsuperscript{m}-R\textsuperscript{a} wherein R\textsuperscript{a} is:
15 OR\textsuperscript{1},
CN,
NR\textsuperscript{1}R\textsuperscript{1},
C\textsubscript{3}-C\textsubscript{6}-cycloalkyl, 3 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
or is -(CH\textsubscript{2})\textsuperscript{n}-(CO)-R\textsuperscript{b} wherein R\textsuperscript{b} is:
20 C\textsubscript{i}-C\textsubscript{6}-alkoxy,
NR\textsuperscript{1}R\textsuperscript{1},
4 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
or R\textsuperscript{1} and R\textsuperscript{3} together with the indole ring to which they are attached form a 5 or 6 membered heterocycloalkyl which can be substituted by C(O)O-C\textsubscript{i}-C\textsubscript{6}-alkyl or C\textsuperscript{i}-C\textsubscript{6}-alkyl;
R^2 is one or more of H, halo, C_{1-6}-alkyl, C_{1-6}-alkoxy;
R^3 is H,
or is C_{1-6}-alkyl,
or is -(C(O)-R C, wherein R^c is C_{1-6}-alkyl, or -(CH_2)_n^m-NR^v R^w;

5 R^4 is one or more of H, halo, or C_{1-6}-alkoxy optionally substituted by OH, or two R^4 may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;
R^5 is H or aryl;
B is halo, NH_2, C_{1-6}-alkyl optionally substituted by CN or C_{1-6}-alkoxy, C_{1-6}-alkoxy,

10 Ci-e-haloalkoxy, C_{3-6}-cycloalkyl, -(C(O)-C_{1-6}-alkyl, -(CR^m R^v)_n-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:
halo, C_{1-6}-alkyl optionally substituted by CN or halo, C_{1-6}-alkoxy;
R^1 and R^u are H, C_{1-6}-alkyl, C_{1-6}-alkyl-NR^v, -(CO)-C_{1-6}-alkyl, -(C(O)-NR^v, -(C(O)-

15 C_{1-6}-alkyl, -(S(O)_{2}-C_{1-6}-alkyl, -(S(O)_{2}-NR^m R^v or OH;
R^w and R^v are H or C_{1-6}-alkyl;
m is 1 to 6;
n is Oto 4;
as well as pharmaceutically acceptable salts thereof.

20
In a further embodiment thereof,
R^1 is H or,
Ci-e-alkyl optionally substituted by CN or,
-(CH_2)_m^m-R^4 wherein R^4 is C_{3-6}-cycloalkyl, aryl, or 5 or 6 membered heteroaryI

25 which are optionally substituted by one or more substituents selected from the group consisting of:
halo, CN, C_{1-6}-alkyl, C_{1-6}-alkoxy, -(C(O)-C_{1-6}-alkyl and phenyl
optionally substituted by halo, C_{1-6}-alkyl Ci-e-haloalkyl or C_{1-6}-alkoxy,
-(CH_2)_m^m-NR R^w or,

30 -(CH_2)_n^m-(C(O)-R b, wherein R^b is aryl;
R^2 is H, halo or C_{1-6}-alkoxy;
R^3 is H or,
-(CO)-R C, wherein R^c is Ci-e-alkyl or,
C_{1-6}-alkyl;

35 R^4 is H or halo;
R^5 is H or aryl;
R\textsuperscript{1} and R\textsuperscript{11} are C\textsubscript{1-6}-alkyl;
m is 1 to 6;
n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

The following compounds are examples according to the invention:
\begin{itemize}
\item r-[(l-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-3-phenyl-3H-spiro[2-benzofuran-1,4'-piperidine];
\item l'-[(l-benzyl-5-methoxy-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
\item r-[(l-\{[2-(4-chlorophenyl)-l,3-thiazol-4-yl]methyl\}-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
\item r-[(l-\{[2-(4-chlorophenyl)-l,3-thiazol-4-yl]methyl\}-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
\item l'-\{[2-methyl-l-(5-methyl-2-[3-(trifluoromethyl)phenyl]-l,3-oxazol-4-yl]methyl\}-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
\item r-[(l-\{[2-(2-fluorophenyl)-5-methyl-l,3-oxazol-4-yl]methyl\}-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
\item l'-\{[2-(4-isopropylphenyl)-5-methyl-l,3-oxazol-4-yl]methyl\}-2-methyl-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
\item r-[(5-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
\item r-[(l-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
\item r-(lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
\item 2-[6-chloro-3-(l H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-1-yl]propanenitrile;
\item 3-[6-chloro-3-(l H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-1-yl]propanenitrile;
\end{itemize}
4-[6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4′-piperidin]-yl)carbonyl]-1H-indol-1-yl]butanenitrile; 
\( \text{r-d} \) 6-chloro-l-fCBl^d-difluoropheny^sulfonyy-lH-indol-S-ylJcarbony^SH-spirofl- 
benzofuran-1,4′-piperidine] ; 

1′-lfl- (biphenyl-S-ylcarbony^6-chloro-lH-indol-S-ylcarbonyljSH-spirofl- 
benzofuran-1,4′-piperidine] ; 
1′-[(l-biphenyl-2-yl-6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4′- 
piperidine] ; 
1′-[(l-biphenyl-2-yl-6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4′- 
piperidine] ; 
\( \text{r-d} \) 6-chloro-l-fCl^d-dibenzylpiperazin^y^methyy-lH-indol-S-ylJcarbony^SH- 
spiro[2-benzofuran-1,4′-piperidine] ; 
tert-butyl 10-(rH,3H-spiro[2-benzofuran-1,4′-piperidin]-l'-ylcarbonyl)-3,4- 
dihydropyrazino[1,2-a]indole-2(1H)-carboxylate; and 

The following compounds are preferred examples according to the invention: 
1′- [(1-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran- 
1,4′-piperidine] ; 
1′-[(1-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-6-chloro-3H-spiro[2-benzofuran- 
1,4′-piperidine] ; 
1′-[(2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4′-piperidine] ; 
1′-[(1-benzoyl-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran- 
1,4′-piperidine] ; 
1′-[(2-methyl-l-(phenylsulfonyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran- 
1,4′-piperidine] ; 
1′-[(l-(cyclohexylmethyl)-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran- 
1,4′- piperidine] ; 
r-[(2-methyl-l-(3-fluorobenzyl)-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran- 
1,4′-piperidine] ; 
r-[(2-methyl-l-[2-(trifluoromethoxy)benzyl]-lH-indol-3-yl)carbonyl]-3H-spiro[2- 
benzofuran-1,4′-piperidine] ; 
1′-[(l-(3,5-dimethylbenzyl)-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran- 
1,4′-piperidine] ;
methyl 4·[[2-methyl-3-([l'H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-ylcarbonyl)-IH-indol-1-yl]methyl]benzoate;
4·[[2-methyl-3-([l'H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-ylcarbonyl)-IH-indol-1-yl]methyl]benzonitrile;
1'·[[1-(3,5-difluorobenzyl)-2-methyl-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
1'·[[1-(2-chlorobenzyl)-2-methyl-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
1'·[[1-(2-methoxybenzyl)-2-methyl-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
1'·[[1-(4-methoxybenzyl)-2-methyl-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
r'·[[1-([2-(2-methoxyphenyl)-5-methyl-1,3-oxazol-4-yl]methyl)-2-methyl-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
r'·[[1-(benzyl-IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
r'·[[6-chloro-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
N,N-dimethyl-2-[3-([l'H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl]-IH-indol-1-yl]ethan amine;
2-methyl-1-[3-([l'H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl]-IH-indol-2-yl]butan-l-one;
[6-chloro-3-([l'H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-ylcarbonyl]-IH-indol-1-yl]acetonitrile;
1'·[[6-chloro-l-(3-fluorobenzoyl)-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
1'·[[6-chloro-l-(2-fluorobenzoyl)-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
1'·[[6-chloro-l-(3,5-difluorobenzoyl)-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
1'·[[6-chloro-l-(2,3-difluorobenzoyl)-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
1'·[[6-chloro-l-(3,5-difluorobenzyl)-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
1'·[[6-chloro-l-(3-fluorobenzyl)-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
1'·[[6-chloro-l-(2-oxo-2-piperidin-l-ylethyl)-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
1'·[[6-chloro-l-(2-oxo-2-piperidin-l-ylethyl)-IH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
1'-\{6-chloro-l-(2-morpholin-4-yl-2-oxoethyl)-lH-indol-3-yl\}carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine] ;
2-[6-chloro-3-(l H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-1-yl]-N,N-diethylacetamide;
2-[6-chloro-3-(l H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-1-yl]-N,N-dimethylacetamide;
2-[6-chloro-3-(l H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-1-yl]-N,N-diethylacetamide;
2-[6-chloro-3-(l H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-1-yl]-N,N-dimethylacetamide;
2-[6-chloro-3-(l H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-1-yl]-N,N-diethylacetamide;
2-[6-chloro-3-(l H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-1-yl]-N,N-diethylacetamide;
2-[6-chloro-3-(l H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-1-yl]-N,N-dimethylacetamide;
tert-butyl [6-chloro-3-(l H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-1-yl] acetate;
1'-\{6-chloro-l-(3,5-difluorophenyl)-lH-indol-3-yl\}carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine] ;
1'-\{6-chloro-l-(3-fluorophenyl)-lH-indol-3-yl\}carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine] ;
2-[6-chloro-3-(l H ,3H-spiro[2-benzofuran-1,4'-piperidin]-l'-ylcarbonyl)-lH-indol-1-yl]-l-(2-fluorophenylethanone;
r-\{6-chloro-l-pyridin-2-yl-lH-indol-3-yl\}carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine] ;
r-\{6-chloro-l-pyridin-4-ylmethyl\}-lH-indol-3-yl\}carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine] ;
2-[6-chloro-3-(l H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-1-yl]-l-pyridin-2-ytlethanone;
r-\{6-chloro-l-(pyridin-3-ylmethyl)-lH-indol-3-yl\}carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine] ;
r-\{6-chloro-l-(pyrimidin-5-ylmethyl)-lH-indol-3-yl\}carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine] ;
r-\{6-chloro-l-(pyridin-2-ylmethyl)-lH-indol-3-yl\}carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine] ;
r-\{6-chloro-l-(pyrazin-2-ylmethyl)-lH-indol-3-yl\}carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine] ;
2-[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4'-piperidin]-l'-ylcarbonyl)-lH-indol-1-yl]-N,N-dimethyllethan eamine;
r-\{6-chloro-l-(2-oxo-2-piperazin-1-ylethyl)-lH-indol-3-yl\}carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine] ;
r-\{6-chloro-l-(2-morpholin-4-yl-2-oxoethyl)-lH-indol-3-yl\}carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine] ;
r-({6-chloro-1-[(5-methylisoxazol-3-yl)methyl]-1H-indol-3-yl}carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidine];
l'-({6-chloro-1-[2-(4-methylpiperazin-1-yl)-2-oxoethyl]-1H-indol-3-yl}carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidine];
4-[(1-{[2-cyclopropyl-4-methylcyclopenta-1,4-dien-1-yl}methyl]-6-methyl-1H-inden-3-yl}vinyl)-2',3'-dihydropiro[cyclohexane-1,1'-indene];
rd'-chloro-1-CCl-methyl-1H-imidazol-S-ylmethyl-1H-indol-S-yl]carbonyl-3H-spiro[2-benzofuran-1,4'-piperidine];
r-({6-chloro-1-[(3-methylisoxazol-5-yl)methyl]-1H-indol-3-yl}carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidine];
l'-({6-chloro-1-[(3,5-dimethylisoxazol-4-yl)methyl]-1H-indol-3-yl}carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidine];
r-({6-chloro-1-[(2,5-dimethyl-1,3-oxazol-4-yl)methyl]-1H-indol-3-yl}carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidine];
l'-({6-chloro-1-[(2-tetrahydro-2H-pyran-4-yl)ethyl]-1H-indol-3-yl}carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidine];
tert-butyl 2-{[6-chloro-3-(1 H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl]methyl}morpholine-4-carboxylate;
l'-({6-chloro-1-[(morpholin-2-ylmethyl)-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidine];
2-[6-chloro-3-(1 H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl]-N-[2-(dimethylamino)ethyl]acetamide;
2-[6-chloro-5-methyl-3-(1 H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl]-N,N-dimethylacetamide;
2-[6-chloro-3-(1 H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl]acetamide;
2-[6-chloro-3-(1 H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl]-N-[2-(methylamino)ethyl]acetamide;
N-(2-aminoethyl)-2-[6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidin]-ylcarbonyl)-1H-indol-1-yl] acetamide;
2-[6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidin]-ylcarbonyl)-1H-indol-1-yljethan amine;
2-[6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidin]-ylcarbonyl)-1H-indol-1-yl]-N-methylethan amine;
2-[6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidin]-ylcarbonyl)-1H-indol-1-yl]-N-methylacetamide;
l’-{[6-chloro-l-(2-morpholin-4-ylethyl)-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4’-piperidine];
l’-{[6-chloro-l-(3-morpholin-4-ylpropyl)-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4’-piperidine];
r’-{[6-chloro-l-(oxiran-2-ylmethyl)-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4’-piperidine];
r’-[{6-chloro-l-(2-methylpyridin-4-yl)methyl]-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4’-piperidine];
r’-[{6-chloro-l-(3S)-piperidin-3-ylmethyl]-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4’-piperidine];
r’-[{6-chloro-l-(tetrahydro-2H-pyran-4-ylmethyl)-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4’-piperidine];
r’-[{6-chloro-l-(l-methylpyrrolidin-3-yl)methyl]-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4’-piperidine];
r’-[{6-chloro-l-(3S)-l-methylpiperidin-3-yl)methyl]-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4’-piperidine];
r’-[{6-chloro-l-(pyrrolidin-3-ylmethyl)-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4’-piperidine];
r’-[{6-chloro-l-(2S)-pyrrolidin-2-ylmethyl)-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4’-piperidine];
r’-[{6-chloro-l-(3S)-pyrrolidin-3-ylmethyl)-1H-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4’-piperidine];
r’-{[6-chloro-l-(2-methyl-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];
r’-(1,2,3,4-tetrahydropyrazino[1,2-a]indol-10-ylcarbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] hydrochloride;
In a certain embodiment the compounds of the invention are those compounds of formula (I-e):

\[
\begin{align*}
&\text{wherein} \\
&R^1 \text{ is } H, \\
&\text{or is } \text{Ci-alkyl optionally substituted by } \text{CN}, \\
&\text{or is sulfonaryl}, \\
&\text{or is } -(\text{CH}_2)_m-R^a \text{ wherein } R^a \text{ is:} \\
&\text{OR}, \\
&\text{CN},
\end{align*}
\]
NR\textsuperscript{1},
C\textsubscript{3-6}-cycloalkyl, 3 to 6 membered-heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B.

or is -(CH\textsubscript{2})\textsubscript{n}-(CO)-R\textsuperscript{b}, wherein R\textsuperscript{b} is:

5  Ci\textsubscript{1-6}-alkoxy,

NR\textsuperscript{1},
5 or 6 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B;

R\textsuperscript{2} is one or more of H, halo, C\textsubscript{1-6}-alkyl;
10  R\textsuperscript{3} is H,
or is Ci\textsubscript{1-6}-alkyl,
or is -(CO)-R\textsuperscript{C}, wherein R\textsuperscript{c} is:

Ci\textsubscript{1-6}-alkyl

-(CHz)\textsubscript{n}-NR\textsuperscript{2},

15  R\textsuperscript{4} is one or more of H, halo, or C\textsubscript{1-6}-alkoxy optionally substituted by OH, or two R\textsuperscript{4} may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;

R\textsuperscript{6} is H;
B is halo, CN, Q-o-alkyl optionally substituted by CN or C\textsubscript{1-6}-alkoxy, C\textsubscript{1-6}-alkoxy or,

6-haloalkoxy, C\textsubscript{3-6}-cycloalkyl, -C(O)O-C\textsubscript{1-6}-alkyl, -(CR\textsuperscript{m}R\textsuperscript{v})\textsubscript{n}-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:

halo, Ci\textsubscript{1-6}-alkyl optionally substituted by CN or halo, C\textsubscript{1-6}-alkoxy;

R\textsuperscript{1} and R\textsuperscript{v} are H, C\textsubscript{1-6}-alkyl, Ci\textsubscript{1-6}-alkyl-NR\textsuperscript{m}R\textsuperscript{v}, -C(O)-C\textsubscript{1-6}-alkyl, -S(O)\textsubscript{2}-Ci\textsubscript{1-6}-alkyl or OH;
25  R\textsuperscript{m} and R\textsuperscript{v} are H or C\textsubscript{1-6}-alkyl;

m is 1 to 6;

n is 0 to 4;

as well as pharmaceutically acceptable salts thereof.

Further, in a certain embodiment the compounds of the invention are those compounds of formula (I-e), wherein

R\textsuperscript{1} is H or,

-(CH\textsubscript{2})\textsubscript{m}-R\textsuperscript{a} wherein R\textsuperscript{a} is 5 or 6 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl or,

-(CH\textsubscript{2})\textsubscript{n}-NR\textsuperscript{v} or,

35  -(CH\textsubscript{2})\textsubscript{n}-(CO)-R \textsuperscript{b}, wherein R\textsuperscript{b} is 5 or 6 membered-heterocycloalkyl;
R² is one or more of H or halo;
R³ is H, or is C₁₋₆-alkyl;
R⁴ is one or more of H or halo;
R⁶ is H;
R¹ and R⁺ are C₁₋₆-alkyl;
m is 1 to 6;
n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

The following compounds are examples according to the invention:

r-[(6-chloro-lH-indol-5-ylcarbonyl)spiro[1-benzofuran-3,4'-piperidine];
r-fCl-benzyl-l-methyl-lH-indol-5-ylcarbonyl]spiro[1-benzofuran-3,4'-piperidine];
r-[(IH-indol-3-ylcarbonyl)spiro[1-benzofuran-3,4'-piperidine];
r-[(6-chloro-5-fluoro-lH-indol-3-yl)carbonyl]spiro[1-benzofuran-3,4'-piperidine];

2-[6-chloro-3-(1H-spiro[1-benzofuran-3,4'-piperidin]-l'-ylcarbonyl)-1H-indol-1-yl]-N,N-dimethylethan amine;
r-[(6-chloro-l-(2-pyrrolidin-1-ylethyl)-lH-indol-3-yl)carbonyl]spiro[1-benzofuran-3,4'-piperidine];
3-[6-chloro-3-(1H-spiro[1-benzofuran-3,4'-piperidin]-l'-ylcarbonyl)-1H-indol-1-yl]-N,N-dimethylpropan-1-amine;
r-[(6-chloro-l-(2-morpholin-4-ylethyl)-lH-indol-3-yl)carbonyl]spiro[1-benzofuran-3,4'-piperidine];
2-[6-chloro-3-(1H-spiro[1-benzofuran-3,4'-piperidin]-l'-ylcarbonyl)-1H-indol-1-yl]-N,N-diethylethan amine;

l'-(6-chloro-l-[2-(lH-pyrrol-1-yl)ethyl]-lH-indol-3-yl)carbonyl]spiro[1-benzofuran-3,4'-piperidine]; and
l'-(6-chloro-l-(2-oxo-2-piperidin-1-ylethyl)-lH-indol-3-yl)carbonyl]spiro[1-benzofuran-3,4'-piperidine].

In a certain embodiment the compounds of the invention are those compounds of formula (I-f):
wherein

R
1
is H,
or is C
1-6-
alkyl optionally substituted by CN,
or is sulfonylaryl,
or is -(CH
2
)m-R
a
wherein R
a
is:

OR
1
,
CN,
NR
R
1
R
1
,
C
3-6-
cycloalkyl,
3 to 6 membered-heterocycloalkyl,
aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
or is -(CH
2
)n-(CO)-R
b
wherein R
b
is:

C
1-6-
alkoxy,
NR
R
c
R
c
,

C
3-6-
cycloalkyl,
3 to 6 membered-heterocycloalkyl,
aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B;

R
2
is one or more of H, halo, C
1-6-
alkyl;
R
3
is H,
or is C
1-6-
alkyl,
or is -(CO)-R
c
, wherein R
c
is:

C
1-6-
alkyl
-(CH
2
)VNR
R
v
,

R
4
is one or more of H, halo, or C
1-6-
alkoxy optionally substituted by OH, or two R
4
may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;
B is halo, CN, C
1-6-
alkyl optionally substituted by CN or C
1-6-
alkoxy, C
1-6-
alkoxy, C
1-6-
haloalkoxy, C
3-6-
cycloalkyl, -C(O)O-C
1-6-
alkyl, -(CR
m
R
v
)n-phenyl, wherein the
phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:
halo, C_{1-6}-alkyl optionally substituted by CN or halo, C_{1-6}-alkoxy;
R^1 and R' are H, C_{1-6}-alkyl, C_{1-6}-alkyl-NR_{m}R_{l}, -C(O)-C_{1-6}-alkyl, -S(O)_{2}C_{1-6}-alkyl or OH;
R'' and R''' are H or C_{1-6}-alkyl;
m is 1 to 6;
n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

Further, in a certain embodiment the compounds of the invention are those compounds of formula (I-f), wherein
R^1 is H;
R^2 is one or more of H or halo;
R^3 is H;
R^4 is one or more of H or halo;
as well as pharmaceutically acceptable salts thereof.

The following compounds are examples according to the invention:
5-bromo-r-(IH-indol-3-ylcarbonyl)spiro[indole-3,4'-piperidin]-2(IH)-one; and
5-bromo-r-[(6-chloro-IH-indol-3-yl)carbonyl]spiro[indole-3,4'-piperidin]-2(IH)-one.

In a certain embodiment the compounds of the invention are those compounds of formula (I-g):

![Diagram](image)

wherein
R^1 is H,
or is C_{1-6}-alkyl optionally substituted by CN,
or is sulfonylaryl,

or is -(CH\(_2\))\(_m\)-Ra wherein Ra is:

OR\(_1\),

CN,

NRR\(_n\),

C\(_3\)\(_6\)-cycloalkyl, 3 to 6 membered-heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,

or is -(CH\(_2\))\(_n\)-(CO)-R\(_b\), wherein R\(_b\) is:

Ci-6'-alkoxy,

NRR\(_n\),

6 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B;

R\(_2\) is one or more of H, halo, C\(_1\)\(_6\)-alkyl;

R\(_3\) is H,

or is Ci-6'-alkyl,

or is -(CO)-R\(_c\), wherein R\(_c\) is:

Ci-6'-alkyl

-(CH\(_2\))\(_n\)-NR\(_m\)R\(_l\),

R\(_4\) is one or more of H, halo, or C\(_1\)\(_6\)-alkoxy optionally substituted by OH, or two R\(_4\) may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;

B is halo, CN, Ci-6'-alkyl optionally substituted by CN or C\(_1\)\(_6\)-alkoxy, C\(_1\)\(_6\)-alkoxy, C\(_1\)\(_6\)-haloalkoxy, C\(_3\)\(_6\)-cycloalkyl, -C(O)O-C\(_1\)\(_6\)-alkyl, -(CR\(_m\)R\(_n\))\(_n\)-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:

halo, Ci-6'-alkyl optionally substituted by CN or halo, C\(_1\)\(_6\)-alkoxy;

R\(_1\) and R\(_n\) are H, C\(_1\)\(_6\)-alkyl, Ci-6'-alkyl-NR\(_m\)R\(_l\), -C(O)-C\(_1\)\(_6\)-alkyl, -S(O)\(_2\)-Ci-6'-alkyl or OH;

R\(_n\) and R\(_n\) are H or C\(_1\)\(_6\)-alkyl;

m is 1 to 6;

n is 0 to 4;

as well as pharmaceutically acceptable salts thereof.

Further, in a certain embodiment the compounds of the invention are those compounds of formula (I-g), wherein

R\(_1\) is H or,
-(CH₂)ₘ-Rᵃ wherein Rᵃ is aryl;
R² is one or more of H or halo;
R³ is H,
   or is Ci₆-alkyl;
5  R⁴ is H;
R⁶ is H;
as well as pharmaceutically acceptable salts thereof.

The following compounds are examples according to the invention:
(SS,RR)-r-[(l-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-3',5'-dimethyl-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one;
(RS,SR)-l'-[(6-chloro-lH-indol-3-yl)carbonyl]-3',5'-dimethyl-3H-spiro[2-benzofuran-
1,4'-piperidin]-3-one;
(RS,SR)-r-[(l-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-3',5'-dimethyl-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one; and
(RR,3'R,5'S)-r-[(l-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-3',5'-dimethyl-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one.

In a certain embodiment the compounds of the invention are those compounds of
formula (I-h):

\[ \text{[Image of chemical structure]} \]

wherein
R¹ is H,
   or is Ci₆-alkyl optionally substituted by CN,
   or is sulfonaryl,
   or is -(CH₂)ₘ-Rᵃ wherein Rᵃ is:
   \[
   \text{OR¹,}
   \]
   \[
   \text{OR²,}
   \]
   \[
   \text{OR³,}
   \]
   \[
   \text{OR⁴,}
   \]
   \[
   \text{OR⁶.}
   \]

\[ \text{(I-h)} \]
CN,
NR\textsubscript{1}\textsubscript{1},
C\textsubscript{3}-\textsubscript{6}-cycloalkyl, 3 to 6 membered-heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
or is -(CH\textsubscript{2})\textsubscript{n}-(CO)-R\textsubscript{b}, wherein R\textsubscript{b} is:
Ci-\textsubscript{6}-alkoxy,
NR\textsubscript{1}\textsubscript{1},
6 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B;

\begin{align*}
R^2 & \text{ is one or more of H, halo, C}_1^6\text{-alkyl;} \\
R^3 & \text{ is H, or is Ci-6-alkyl,} \\
& \text{ or is -(CO)-R}^\text{C}, \text{ wherein R}^\text{C} \text{ is:} \\
& \text{ Ci-6-alkyl} \\
& \text{ -(CH}_2\text{VNR} \textsubscript{1}\text{R} \textsubscript{1},} \\
R^4 & \text{ is one or more of H, halo, or C}_1^6\text{-alkoxy optionally substituted by OH, or two R}^4 \text{ may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;} \\
R^8 & \text{ is H or Ci-e-alkyl;} \\
X & \text{ is CH}_2 \text{ or C=O;} \\
B & \text{ is halo, CN, C}^\text{N}-\text{alkyl! optionally substituted by CN or C}_1^6\text{-alkoxy, C}_1^6\text{-alkoxy, C}_1^6\text{ haloalkoxy, C}_3^6\text{-cycloalkyl, } \text{-C}(O)O-C}_1^6\text{-alkyl, -(CR}_m\text{R}^\text{b})_n\text{-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:} \\
& \text{ halo, Ci-6-alkyl optionally substituted by CN or halo, C}_1^6\text{-alkoxy;} \\
R^1\text{ and R}^\text{w} & \text{ are H, Q-e-alkyl, Q-e-alkyl-NR}_\text{m}\text{R}^\text{w}, \text{-C}(O)O-C}_1^6\text{-alkyl, -S(O)}_2\text{-C}_1^6\text{-alkyl or OH;} \\
R^\text{w} & \text{ and R}^\text{v} \text{ are H or Ci-6-alkyl;} \\
m & \text{ is 1 to 6;} \\
n & \text{ is 0 to 4;} \\
\text{ as well as pharmaceutically acceptable salts thereof.}
\end{align*}

Further, in a certain embodiment the compounds of the invention are those compounds of formula (I-h), wherein
\begin{align*}
R^1 & \text{ is H,} \\
or is -(CH}_2\text{V (CO)-R}^\text{b}, \text{ wherein R}^\text{b} \text{ is } NR\textsubscript{1}\textsubscript{1};
\end{align*}
**R**\(^2\) is one or more of H or halo;
\(R^3\) is H;
\(R^4\) is H;
\(R^8\) is H or \(C_{1-6}\)-alkyl;
\(X\) is \(CH_2\) or C=O;
\(R^1\) and \(R^6\) are H or \(Cl_{1-6}\)-alkyl;
as well as pharmaceutically acceptable salts thereof.

The following compounds are examples according to the invention:

l\(^'\)-[(6-chloro-lH-indol-3-yl)carbonyl]spiro[iso indole-1,4'-piperidin]-3(2H)-one;

1\(^'\)-[(6-chloro-1H-indol-3-yl)carbonyl]-2-methyl-2,3-dihydrospiro[isoindole-1,4'-piperidine] ;

r\(^'\)-[(6-chloro-1H-indol-S-yl]carbonyl\(^{-}\)dihydrospiroisoindole-1\(^'\)-piperidinel]

and


The invention also encompasses the compounds of formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h) for a use in the prevention or treatment of dysmenorrhea, hypertension, chronic heart failure, inappropriate secretion of vasopressin, liver cirrhosis, nephrotic syndrome, obsessive compulsive disorder, anxious and depressive disorders.

The invention also encompasses a pharmaceutical composition comprising a compound of formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h), which pharmaceutical composition is useful against dysmenorrhea, hypertension, chronic heart failure, inappropriate secretion of vasopressin, liver cirrhosis, nephrotic syndrome, obsessive compulsive disorder, anxious and depressive disorders.

The invention further encompasses the use of a compound of formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h) for the preparation of a medicament which is useful against dysmenorrhea, hypertension, chronic heart failure, inappropriate secretion of vasopressin, liver cirrhosis, nephrotic syndrome, obsessive compulsive disorder, anxious and depressive disorders.

In a certain embodiment, the compounds of the invention can be manufactured according to a process comprising the step of reacting a compound of formula (II):
with a compound of formula A-H in order to obtain the compound of formula (I),
wherein A, R₁, R₂ and R₃ are as defined hereinabove.

In another embodiment, the compounds of the invention can be manufactured
according to a process comprising the step of reacting a compound of formula (III):

with an electrophile compound of formula R¹-Y in order to obtain the compound
of formula (I), wherein R¹, R² and R₃ are as defined hereinabove and Y is halo, preferably
Br or Cl.

In still another embodiment, the compounds of the invention can be manufactured
according to a process comprising the step of reacting a compound of formula (IV):

with an amine of formula HNR₁R¹₁ in order to obtain the compound of formula (I),
wherein R¹, R², R₃, Rᵢ and Rᵠ are as defined hereinabove.

The preparation of the compounds of the invention of formulae (I), (I-a), (I-b), (I-c),
(I-d), (I-e), (I-f), (I-g) or (I-h) is described more in details with the following general
schemes A, B and C, wherein R¹, R², R₃ and A are as defined hereinabove:
Compounds of formula (I) can be prepared via an amide coupling between an indole 3-carboxylic acid (II) and a spiro-piperidine (A-H). Indole 3-carboxylic acids (II) are either commercially available or readily prepared using a procedure described in J. Med. Chem. 1991, 34, 140. Alternatively, they can be prepared following the general scheme D as described hereinafter. The spiro-piperidine derivatives A-H are either commercially available or can be prepared using commercially available starting materials and conventional methods. Spiropiperidines A-H of group (h) can be prepared as described in general scheme E. General scheme A is hereinafter further illustrated with general scheme and procedure I to which, e.g., examples 1 to 24, 34, 39-43, 66-70, 77-80 and 123-126 refer.

Compounds of formula (I) with R¹ different from H can be prepared by N-deprotonation of an indole derivative (III) (a compound of formula (I) wherein R¹ is H) followed by treatment with an electrophilic reactant R¹-Y (wherein Y is a leaving group) which is either commercially available or easily prepared according to methods well known in the art and commercially available starting materials. Alternatively, compounds (I) can be prepared by coupling of an indole derivative (III) with a boronic acid R¹.
B(OH)$_2$ using a transition metal catalyst such as Cu(OAc)$_2$ in the presence of pyridine, molecular sieves and air in dichloromethane. Derivatives (III) are prepared using the method described in the general scheme A. General scheme B is hereinafter further illustrated with general schemes and procedures II and III to which, e.g., examples 46 to 65, 81 to 87, 93-103, 133-136, 142-144, 150-154, 161-168, 170, 172-175, 180, 182, 183, 188-193, 195-197, 201-203, 220, 224, 228-247, 253-260, 263, 266, and 267 refer, as well as general schemes and procedures V (e.g., examples 104-115, 117, 204-212), VI (e.g., examples 155-157, 222, 223, 225, 226), VII (e.g., examples 88-92, 128-132, 137-141, 145-149, 158, 214-218, 221) and VIII.

General scheme C

Compounds of formula (I) wherein $R^3$ is an amide (-CONH$R^1$) can be prepared via an amide coupling between an indole 2-carboxylic acid (IV) and an amine NH$R^1$R$^2$. The indole 2-carboxylic acid derivatives (IV) are readily prepared using commercially available starting products and conventional methods. General scheme C is hereinafter further illustrated with general scheme and procedure IV to which, e.g., examples 24 to 33 refer.
The treatment of an indole derivative (Va) with trifluoroacetic anhydride in DMF affords intermediate (VI) which can be hydrolysed with an aqueous sodium hydroxide solution to give the 3-carboxylic acid indole derivative (Ha). Alternatively, (VI) can react with an electrophilic reactant $R^1\cdot Y$ to give (VII), which is then converted to the corresponding carboxylic acid derivative (lib) with NaH/H$_2$O in DMF (see J. Org Chem., 1993, 10, 2862). Intermediate (VII) can alternatively be obtained by treatment of an indole derivative (Vb) with trifluoroacetic anhydride in DMF, dichloromethane or 1,2-dichloroethane. Addition of a suitable base may be advantageous.
Cyclization of a bromophenylacetonitrile derivative (VIII) affords a 4-bromoaryl-2-cyanopiperidine derivative (IX). The cyano group is transformed to an azidocarbonyl group using standard functional group transformations to give a compound of formula (X). Upon heating in toluene an azide (X) undergoes a Curtius rearrangement. The crude intermediate isocyanate is trapped to form a lactam of formula (XI) after bromine-lithium exchange using tert-butyl lithium at -100 °C. N-Deprotection affords spiropiperidine (XII). Alternatively, lactam (XI) can be N-alkylated to give lactam (XV).

Both, lactam (XI) and (XV), are reduced using standard conditions to afford spiropiperidines (XIV) and (XVI) after N-deprotection, respectively.
The following general procedures I to VIII are meant to give examples of the preparation of the compounds of the invention according to general schemes A to C.

The following general procedure I is an example of the preparation of the compounds of the invention according to general scheme A

General procedure I

General procedure I - amide coupling:

To a stirred solution of an indole-3-carboxylic acid derivative (1 mmol) in 10 ml CH₂Cl₂ was added (1.3 mmol) EDC, (1.3 mmol) HOBt, (1.3 mmol) Et₃N and (1 mmol) of the amine derivative. The mixture was stirred overnight at RT and then poured onto water and extracted with CH₂Cl₂. The combined organic phases were dried over Na₂SO₄ and concentrated in vacuo. Flash chromatography or preparative HPLC afforded the title compound.

The following general procedures II and III are examples of the preparation of the compounds of the invention according to general scheme B:

General procedure II

General procedure II:

To a stirred solution of 30 mg (0.09 mmol) of l'-(2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidine] (the preparation of which have
been described in example 43) in 3 ml DMF was added 4 mg (0.10 mmol) NaH (60% in oil). The mixture was stirred at RT for 30 min. and then the electrophilic reactant $R^1-Y$ (0.15 mmol) was added. The mixture was stirred an additional 18 hours and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na$_2$SO$_4$ and concentrated in vacuo. Purification by preparative HPLC afforded the corresponding derivatives.

General procedure III:

To a stirred solution of 20 mg (0.054 mmol) of 1'-[6-chloro-1H-indol-3-yl)carbonyl]spiro[1-benzofuran-3,4'-piperidine] (the preparation of which have been described in example 77) in 3 ml DMF was added 8.8 mg (0.11 mmol) NaH (60% in oil). The mixture was stirred at RT for 30 min. and then the electrophilic reactant $R^1-Y$ (QOS mmol) was added. The mixture was stirred an additional 18 hours at 60°C and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na$_2$SO$_4$ and concentrated in vacuo. Purification by preparative HPLC afforded the corresponding derivatives.

The following general procedure IV is an example of preparation of the compounds of the invention according to general scheme C:
General procedure IV:

To a stirred solution of 6-chloro-3-[(5-fluoro-3-oxo-1'H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-1'H-indole-2-carboxylic acid (1 mmol) (the preparation of which have been described in example 24d) in 10 ml CH₂Cl₂ was added (1.3 mmol) EDC, (1.3 mmol) HOBt, (1.3 mmol) Et₃N and (1 mmol) of the amine derivative (wherein R¹ and R'' are as defined hereinabove). The mixture was stirred overnight at RT and then poured onto water and extracted with CH₂Cl₂. The combined organic phases were dried over Na₂SO₄ and concentrated in vacuo. Flash chromatography or preparative HPLC afforded the title compound.

The following general procedures V to VIII are examples of the preparation of the compounds of the invention according to general scheme B:

General procedure V
General procedure V:

To a stirred solution of an indole derivative (III) (0.079 mmol) in 2 ml DMF was added NaH (0.10 mmol, 60% in oil). The mixture was stirred at room temperature for 30 min. and then the electrophile R1-Y (0.15 mmol) was added. The mixture was stirred for an additional 18 hours and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na₂SO₄ and concentrated in vacuo. Purification by preparative HPLC afforded the corresponding derivatives.

General procedure VI

General procedure VI - aryl coupling:

To a solution of the indole in dichloromethane was added anhydrous Cu(OAc)₂ (2 eq), the boronic acid (3 eq) and pyridine (4 eq), and the reaction mixture stirred 16h at room temperature in the presence of 0.4 nm molecular sieve under an open to air atmosphere. Filtration through decalite, washing with dichloromethane and concentration gave the crude product which was purified by silica gel chromatography (hexane/ethyl acetate) to give the desired product.

General procedures VII and VIII

General procedure VII - acylation:
A solution of the indole in dry DMF was treated with sodium hydride (1.05 eq) and stirred for 15 min at room temperature, then treated with an acid chloride (1.1 eq) and stirred at room temperature for 2h. Purification by preparative HPLC yielded the desired product.

General Procedure VIII - sulphonylation :

A solution of the indole in dry DMF was treated with sodium hydride (1.05 eq) and stirred for 15 min at room temperature, then treated with a sulphonyl chloride (1.1 eq) and stirred at room temperature for 2h. Purification by preparative HPLC yielded the desired product.

Results - Via activity

Material & Method:

The human Via receptor was cloned by RT-PCR from total human liver RNA. The coding sequence was subcloned in an expression vector after sequencing to confirm the identity of the amplified sequence. To demonstrate the affinity of the compounds from the present invention to the human Via receptor binding studies were performed. Cell membranes were prepared from HEK293 cells transiently transfected with the expression vector and grown in 20 liter fermenters with the following protocol.

50g of cells are resuspended in 30ml freshly prepared ice cold Lysis buffer (50mM HEPES, 1mM EDTA, 10mM MgCl2 adjusted to pH= 7.4 + complete cocktail of protease inhibitor (Roche Diagnostics)). Homogenized with Polytron for 1min and sonicated on ice for 2x 2 minutes at 80% intensity (Vibracell sonicator). The preparation is centrifuged 20 min at 500 g at 4°C, the pellet is discarded and the supernatant centrifuged 1hour at 43'000g at 4°C (19'000rpm). The pellet is resuspended in 12.5 mL Lysis buffer+ 12.5ml Sucrose 20% and homogenized using a Polytron for 1-2 min. The protein concentration is determined by the Bradford method and aliquots are stored at -80°C until use. For binding studies 60mg Yttrium silicate SPA beads (Amersham) are mixed with an aliquot of membrane in binding buffer (50 mM Tris, 120mM NaCl, 5 mM KCl, 2 mM CaCl2, 10 mM MgCl2) for 15 minutes with mixing. 50ul of bead/membrane mixture is then added to each well of a 96 well plate, followed by 50ul of 4 nM 3H-Vasopressin (American Radiolabeled Chemicals). For total binding measurement 100ul of binding buffer are added to the respective wells, for non-specific binding 100ul of 8.4mM cold vasopressin and for compound testing 100ul of a serial dilution of each compound in 2% DMSO. The plate is incubated 1h at room temperature, centrifuged 1 min at 1000g and counted on a Packard
Top-Count. Non-specific binding counts are subtracted from each well and data is normalized to the maximum specific binding set at 100%. To calculate an IC 50 the curve is fitted using a non-linear regression model (XLfit) and the Ki is calculated using the Cheng-Pru ssoff equation.

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The compounds of formula (I) as well as their pharmaceutically usable acid addition salts can be used as medicaments, e.g. in the form of pharmaceutical preparations. The pharmaceutical preparations can be administered orally, e.g. in the form of tablets, coated tablets, dragees, hard and soft gelatine capsules, solutions, emulsions or suspensions. The administration can, however, also be effected rectally, e.g. in the form of suppositories, or parenterally, e.g. in the form of injection solutions.

The compounds of formula (I) and their pharmaceutically usable acid addition salts can be processed with pharmaceutically inert, inorganic or organic excipients for the production of tablets, coated tablets, dragees and hard gelatine capsules. Lactose, corn starch or derivatives thereof, talc, stearic acid or its salts etc can be used as such excipients e.g. for tablets, dragees and hard gelatine capsules.

Suitable excipients for soft gelatine capsules are e.g. vegetable oils, waxes, fats, semisolid and liquid polyols etc.

Suitable excipients for the manufacture of solutions and syrups are e.g. water, polyols, saccharose, invert sugar, glucose etc.

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<td>6.2</td>
<td>202</td>
<td>4.3</td>
<td>280</td>
<td>67.6</td>
<td></td>
</tr>
<tr>
<td>109</td>
<td>1.1</td>
<td>203</td>
<td>11.7</td>
<td>281</td>
<td>19.1</td>
<td></td>
</tr>
<tr>
<td>110</td>
<td>0.8</td>
<td>204</td>
<td>1.1</td>
<td>283</td>
<td>63.1</td>
<td></td>
</tr>
<tr>
<td>111</td>
<td>2.4</td>
<td>205</td>
<td>1.0</td>
<td>284</td>
<td>138.0</td>
<td></td>
</tr>
<tr>
<td>112</td>
<td>2.0</td>
<td>206</td>
<td>0.5</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Suitable excipients for injection solutions are e.g. water, alcohols, polyols, glycerol, vegetable oils etc.

Suitable excipients for suppositories are e.g. natural or hardened oils, waxes, fats, semi-liquid or liquid polyols etc.

Moreover, the pharmaceutical preparations can contain preservatives, solubilizers, stabilizers, wetting agents, emulsifiers, sweeteners, colorants, flavorants, salts for varying the osmotic pressure, buffers, masking agents or antioxidants. They can also contain still other therapeutically valuable substances.

The dosage can vary within wide limits and will, of course, be fitted to the individual requirements in each particular case. In general, in the case of oral administration a daily dosage of about 10 to 1000 mg per person of a compound of general formula (I) should be appropriate, although the above upper limit can also be exceeded when necessary.

The following Examples illustrate the present invention without limiting it. All temperatures are given in degrees Celsius.

**Example A**

Tablets of the following composition are manufactured in the usual manner:

<table>
<thead>
<tr>
<th>mg/tablet</th>
</tr>
</thead>
<tbody>
<tr>
<td>Active substance</td>
</tr>
<tr>
<td>Lactose</td>
</tr>
<tr>
<td>Corn starch</td>
</tr>
<tr>
<td>Microcrystalline cellulose</td>
</tr>
<tr>
<td>Magnesium stearate</td>
</tr>
<tr>
<td>Tablet weight</td>
</tr>
</tbody>
</table>

**Example B**

Capsules of the following composition are manufactured:

<table>
<thead>
<tr>
<th>mg/capsule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Active substance</td>
</tr>
<tr>
<td>Lactose</td>
</tr>
<tr>
<td>Corn starch</td>
</tr>
<tr>
<td>Microcrystalline cellulose</td>
</tr>
<tr>
<td>Magnesium stearate</td>
</tr>
<tr>
<td>Tablet weight</td>
</tr>
</tbody>
</table>
The active substance, lactose and corn starch are firstly mixed in a mixer and then in a comminuting machine. The mixture is returned to the mixer, the talc is added thereto and mixed thoroughly. The mixture is filled by machine into hard gelatine capsules.

Example C

Suppositories of the following composition are manufactured:

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>mg/supp.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Active substance</td>
<td>15</td>
</tr>
<tr>
<td>Suppository mass</td>
<td>1285</td>
</tr>
<tr>
<td>Total</td>
<td>1300</td>
</tr>
</tbody>
</table>

The suppository mass is melted in a glass or steel vessel, mixed thoroughly and cooled to 45°C. Thereupon, the finely powdered active substance is added thereto and stirred until it has dispersed completely. The mixture is poured into suppository moulds of suitable size, left to cool; the suppositories are then removed from the moulds and packed individually in wax paper or metal foil.

EXAMPLES

Examples of compounds of formula (I-a)

Example 1
l'-(1-benzyl-2-methyl-1'H-indol-3-yl)carbonyl|spiro|in dene- 1,4'-piperidine]
Amide coupling according to general procedure I described hereinabove:

- Amine: spiro[indene-1,4'-piperidine] prepared as described in J. Med. Chem. 1992, 35, 2033,

- Acid: 1-Benzyl-2-methyl-1H-indole-3-carboxylic acid (the preparation of which is described hereinafter).

ES-MS m/e (%): 433.5 (M+H+).

1-Benzyl-2-methyl-1H-indole-3-carboxylic acid

To a stirred solution of 0.50 g (3.10 mmol) of 2-methyl-1H-indole-3-carboxylic acid (described in J. Heterocyclic Chem. 1977, 14, 1123) in 5 ml DMF was added 0.27 g (6.75 mmol) of NaH (60% in oil). The mixture was stirred at RT for 30 min. and then 0.39 ml (3.28 mmol) of benzyl bromide was added. The mixture was stirred an additional hour and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na2SO4 and concentrated in vacuo. Crystallization in Et2O afforded 1-Benzyl-2-methyl-1H-indole-3-carboxylic acid.

Example 2

1'-(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]-2,3-dihydrospiro[indene-1,4'-piperidine]
- Amine: 2,3-dihydrospiro[indene-1,4'-piperidine] prepared as described in *J. Med. Chem.* 1992, 35, 2033,
- Acid: 1-Benzyl-2-methyl-IH-indole-3-carboxylic acid (described in example 1), ES-MS m/e (%): 435.5 (M+H+).

Example 3
r-[(1-benzyl-IH-indol-3-yl)carbonyl]spiro[indene-1,4'-piperidine]

Amide coupling according to general procedure I described hereinabove:
- Amine: spiro[indene-1,4'-piperidine] prepared as described in *J. Med. Chem.* 1992, 35, 2033,
- Acid: 1-benzyl-IH-indole-3-carboxylic acid (the preparation of which is described hereinafter),
ES-MS m/e (%): 419.4 (M+H+).

1-benzyl-IH-indole-3-carboxylic acid

To a stirred solution of 0.50 g (3.10 mmol) IH-indole-3-carboxylic acid in 5 ml DMF was added 0.27 g (6.75 mmol) of NaH (60% in oil). The mixture was stirred at RT for 30 min. and then 0.39 ml (3.28 mmol) of benzyl bromide was added. The mixture was stirred an additional hour and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na2SO4 and concentrated in vacuo.

Crystallization in Et2O afforded 0.61 g (78%) of 1-benzyl-IH-indole-3-carboxylic acid as a white solid. ES-MS m/e (%): 250 (M-H+).

Example 4
r-[(2-methyl-IH-indol-3-yl)carbonyl]spiro[indene-1,4'-piperidine]
Amide coupling according to general procedure I described hereinabove:
- Amine: spiro[indene-1,4'-piperidine] prepared as described in *J.Med.Chem.* 1992, 35, 2033,
- Acid: 2-Methyl-1H-indole-3-carboxylic acid prepared as described in *J.Heterocyclic Chem.* 1977, 14, 1123,

ES-MS m/e (%): 343.2 (M+H+).

Example 5
l'-(6-chloro-1H-indol-3-yl)carbonyl]spiro[indene-1,4'-piperidine]

Amide coupling according to general procedure I described hereinabove:
- Amine: spiro[indene-1,4'-piperidine] prepared as described in *J.Med.Chem.* 1992, 35, 2033,
- Acid: 6-chloro-1H-indole-3-carboxylic acid (the preparation of which is described hereinafter),

ES-MS m/e (%): 363.4 (M+H+).

6-chloro-1H-indole-3-carboxylic acid
Using a procedure described in *J. Med. Chem.* 1991, 34, 140, from 7.0 g (0.046 mmol) of 6-chloro-1H-indole was prepared 5.80 g (64%) of 6-chloro-1H-indole-S-carboxylic acid as a light brown solid. ES-MS m/e (%): 194 (M-H+).

Example 6

r-[(6-chloro-1H-indol-3-yl)carbonyl]-2,3-dihydrospiro[indene-1,4′-piperidine]

Amide coupling according to general procedure I described hereinafore:
- Amine: 2,3-dihydrospiro[indene-1,4′-piperidine] prepared as described in *J. Med. Chem.* 1992, 35, 2033,
- Acid: 6-chloro-1H-indole-S-carboxylic acid (described in example 5), ES-MS m/e (%): 365.4 (M+H+).

Examples of compounds of formula (I-b)

Example 7

r-[(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]-l-(methylsulfonyl)-l,2-dihydrospiro[indole-3,4′-piperidine]
Amide coupling according to general procedure I described hereinabove:
- Amine: 1-(methylsulfonyl)-1,2-dihydrospiro[indole-3,4’-piperidine] prepared as described in *Tetrahedron*, 1997, 53, 10983,
- Acid: 1-Benzyl-2-methyl-1H-indole-3-carboxylic acid (described in example 1),

ES-MS m/e (%): 514.6 (M+H+).

Example 8

r-[(6-chloro-1H-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4’-piperidine]

Amide coupling according to general procedure I described hereinabove:
- Amine: 1,2-dihydrospiro[indole-3,4’-piperidine] prepared as described in *Tetrahedron*, 2004, 60, 4875-4878,
- Acid: 6-chloro-1H-indole-5-carboxylic acid (described in example 5),

ES-MS m/e (%): 366.4 (M+H+).

Example 9

l’-[(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4’-piperidine]
Amide coupling according to general procedure I described hereinabove:
- Amine: 1,2-dihydrospiro[indole-3,4'-piperidine] prepared as described in Tetrahedron, 2004, 60, 4875-4878,
- Acid: l-Benzyl-l-methyl-lH-indole-3-carboxylic acid (described in example 1),
ES-MS m/e (%): 436.6 (M+H+).

Examples of compounds of formula (I-c)

Example 10

r-[(l-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-6-chloro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure I described hereinabove:
- Amine: 6-chloro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in
European patent application EP722941,
- Acid: l-Benzyl-2-methyl-lH-indole-3-carboxylic acid (described in example 1),
ES-MS m/e (%): 485.5 (M+H+).
Example 1

l'-(l-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-4-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one

Amide coupling according to general procedure I described hereinabove:
- Amine: 4-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one prepared as described in EP722941,
- Acid: l-Benzyl-2-methyl-lH-indole-3-carboxylic acid (described in example 1),
ES-MS m/e (%): 469.5 (M+H+).

Example 2

r'-(l-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-6-methoxy-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure I described hereinabove:
- Amine: 6-methoxy-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in EP722941,
- Acid: l-Benzyl-2-methyl-lH-indole-3-carboxylic acid (described in example 1),
Example 13

l’-[l-benzyl-2-methyl-lH-indol-3-yl]carbonyl]-5-methoxy-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure I described hereinabove:
- Amine: 5-methoxy-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in EP722941,
- Acid: l-Benzyl-2-methyl-lH-indole-3-carboxylic acid (described in example 1),

ES-MS m/e (%): 481.6 (M+H+).

Example 14

r-[l-benzyl-2-methyl-lH-indol-3-yl]carbonyl]-7-chloro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure I described hereinabove:
- Amine: 7-chloro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (the preparation of which is described hereinafter),
- Acid: l-Benzyl-2-methyl-lH-indole-3-carboxylic acid (described in example 1),

ES-MS m/e (%): 485.5 (M+H+).
7-chloro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure described in J.Org.Chem. 1976,41, 2628, from 2-bromo-3-chloro-benzoic acid (the preparation of which is described in J.Org.Chem. 203, 68, 2030) and 1-benzyl-piperidin-4-one was prepared 7-chloro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one, after debenzylation (Pd/C, H₂, MeOH/HCl), as a white solid. ES-MS m/e (%): 238.7 (M+H⁺).

Example of compounds of formula (I-g)

Example 15

(^,i?i?)-r-[(1-benzyl-2-methyl-1H-indol-3-yl)carnbonyl]-3',5'-dimethyl-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure I described hereinabove:
- Amine: (SS,RR)-3',5'-dimethyl-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in WO9929696,
- Acid: 1-Benzyl-2-methyl-1H-indole-3-carboxylic acid (described in example 1), ES-MS m/e (%): 479.6 (M+H⁺).

Examples of compounds of formula (I-c)

Example 16

r-[(6-chloro-1H-indol-3-yl)carnbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one
Amide coupling according to general procedure I described hereinabove:
- Amine: 3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in J. Org. Chem. 1976, 41, 2628,
- Acid: 6-chloro-lH-indole-S-carboxylic acid (described in example 5),
  ES-MS m/e (%): 381.4 (M+H+).

Example 17
l’-[(6-chloro-lH-indol-3-yl)carbonyl]-6-methoxy-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure I described hereinabove:
- Amine: 6-methoxy-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in EP722941,
- Acid: 6-chloro-lH-indole-S-carboxylic acid (described in example 5),
  ES-MS m/e (%): 411.4 (M+H+).

Example of compounds of formula (I-g)
Example 18
(iS,Si)-r-[(6-chloro-lH-indol-3-yl)carbonyl]-3',5'-dimethyl-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure I described hereinabove:

- Amine: (lr,3'R,5'S)-3',5'-dimethyl-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in WO9929696;
- Acid: 6-chloro-lH-indole-S-carboxylic acid (described in example 5), ES-MS m/e (%): 409.4 (M+H+).

Examples of compounds of formula (I-c)

Example 19
r-[(6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure I described hereinabove:
Example 20

6-chloro-\(\text{r-}[\{6\text{-chloro-lH-indol-3-yl}\text{carbonyl}\}3\text{-H-spiro[2-benzofuran-l,4'-piperidin]}\]-3-one

Amide coupling according to general procedure I described hereinabove:

- Amine: 6-chloro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in EP722941,
- Acid: 6-chloro-lH-indole-S-carboxylic acid (described in example 88), ES-MS m/e (%): 415.3 (M\(^+\)).
Amide coupling according to general procedure I described hereinabove:
- **Amine:** (lr,3'R,5'S)-3',5'-dimethyl-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one
  prepared as described in WO9929696,
- **Acid:** l-Benzyl^-methyl-lH-indole-S-carboxylic acid (described in example 1),

ES-MS m/e (%): 479.6 (M+H +).

**Examples of compounds of formula (I-c)**

**Example 22**
\[
\text{r-}[(l\text{-benzyl-2-methyl-lH-indol-3-yl})\text{carbonyl}]-3\text{H-spiro}[2\text{-benzofuran-l,4'-piperidin}]-3\text{-one}
\]

Amide coupling according to general procedure I described hereinabove:
- **Amine:** 3H-spiro[2-benzofuran-l,4'-piperidin]-3-one prepared as described in *J. Org. Chem.* 1976, 41, 2628,
- **Acid:** l-Benzyl-2-methyl-lH-indole-3-carboxylic acid (described in example 1),

ES-MS m/e (%): 451.6 (M+H +).

**Example 23**
\[
\text{r-}[(l\text{-benzyl-2-methyl-lH-indol-3-yl})\text{carbonyl}]-5\text{-fluoro-3H-spiro}[2\text{-benzofuran-l,4'-piperidin}]-3\text{-one}
\]
Amide coupling according to general procedure I described hereinabove:
- Amine: 5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in WO2001014376,
- Acid: 1-Benzyl-1-methyl-1H-indole-S-carboxylic acid (described in example 1),
ES-MS m/e (%): 469.6 (M+H+).

Example 24

\[ \text{r-} \left( \{6\text{-chloro-2-}\{4\text{-methylpiperidin-1-yl} \text{carbonyl}\}-\text{indol-3-yl} \text{carbonyl}\}-5\text{-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one} \right) \]

\[ \begin{align*}
\text{a) 6-Chloro-3-formyl-1H-indole-2-carboxylic acid ethyl ester} \\
\text{To a stirred solution of 2.36 g (0.011 mol) of 6-chloro-lH-indole-2-carboxylic acid ethyl ester in DMF (20 ml) was added phosphorus oxychloride (1.08 ml, 0.012 mol) at RT. The solution was heated at 60}^\circ\text{C for 2 hours, then cooled to RT, and poured into water. The pH was adjusted to 7 by careful addition of aq. NaOH 2N. The resulting brown precipitate was collected by filtration and dried over night in a vacuum oven (50}^\circ\text{C). The title compound was obtained as light brown solid, 2.05 g (77%).} 
\end{align*} \]
b) 6-Chloro-lH-indole^^-dicarboxylic acid 2-ethyl ester

To a solution of 0.104 g (0.413 mmol) of 6-Chloro-S-formyl-lH-indole-l-carboxylic acid ethyl ester in a mixture of tert-butanol (10 ml) and H₂O (5 ml), was added 2-methyl-2-butene (2 ml) followed by a solution OfNaClO₂ (0.344 g, 3.80 mmol) and NaH₂PO₄ (0.399 g, 2.90 mmol) in water (2 ml). The mixture was stirred overnight at RT. The organic solvents were removed, and then the aqueous solution diluted with water, and washed twice with hexane. The pH of the aqueous phase was adjusted to 3 by addition of aq. HCl IN, and the product extracted with EtOAc. The combined organic phases were dried over Na₂SO₄ and concentrated in vacuo to afford 15 mg (14%) of 6-chloro-lH-indole-2,3-dicarboxylic acid 2-ethyl ester as a light yellow solid.

c) lH-indole-2-carboxylic acid, 6-chloro-3-r(5-fluoro-3-oxospiroisobenzofuran-1(3H),4'-piperidin)-l'-yl-2-carbonylll -ethyl ester

Amide coupling according to general procedure I described hereinabove:

- Amine: 5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in WO2001014376,
- Acid: 6-chloro-lH-indole-2,3-dicarboxylic acid 2-ethyl Ester (the preparation of which is described hereinafter),

d) 6-chloro-3-r(5-fluoro-3-oxo-lH,3H-spiro[2-benzofuran-1,4'-piperidin]-r'-yl)-2-carbonylll -lH-indole-2-carboxylic acid

To a solution of 1.89 g (4.02 mmol) of lH-indole-2-carboxylic acid, 6-chloro-3-[5-fluoro-3-oxospiro(isobenzofuran-1(3H),4'-piperidin]-r-yl]carbonyl]-ethyl ester in EtOH (150 ml), was added an aqueous solution of LiOH (7.8 ml, 1M). The resulting white suspension was stirred at 80°C overnight, cooled to RT, and then poured in 500 ml aq. HCl (IN). The product was extracted with 3 times 500 ml OfCH₂Cl₂, and the combined organic phases were dried over Na₂SO₄ and concentrated in vacuo to afford 1.42 g (72%) of 6-chloro-3-[5-fluoro-3-oxo-lH,3H-spiro[2-benzofuran-1,4'-piperidin]-r'-yl]carbonyl]-lH-indole-2-carboxylic acid as a white solid.

e) r-(6-chloro-2-r(4-methylpiperidin-l-yl)carbonylll-lH-indol-3-yl]carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure IV described hereinabove:

- Amine: commercially available 4-methyl-piperidine,
- Acid: 6-chloro-3-[5-fluoro-3-oxo-lH,3H-spiro[2-benzofuran-1,4'-piperidin]-r'-yl]carbonyl]-lH-indole-2-carboxylic acid (described in step d hereinabove),
Example 25

6-chloro-3-[(5-fluoro-3-oxo-1'H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl]-1'H-indole-2-carboxamide

Amide coupling according to general procedure IV described hereinabove:
- Amine: commercial available ammonium hydroxide,
- Acid: 6-chloro-3-[(5-fluoro-3-oxo-rH,3H-spiro[2-benzofuran-1,4'-piperidin]-l'-yl)carbonyl]-1'H-indole-2-carboxylic acid (described in example 24),

ES-MS m/e (%): 525 (M+H+).

Example 26

6-chloro-N-[2-(dimethylamino)ethyl]-3-[(5-fluoro-3-oxo-rH,3H-spiro[2-benzofuran-1,4'-piperidin]-l'-yl)carbonyl]-1'H-indole-2-carboxamide

Amide coupling according to general procedure IV described hereinabove:
- Amine: commercially available N,N-dimethyl-ethane-1,2-diamine,
- Acid: 6-chloro-3-[(5-fluoro-3-oxo-rH,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl]-1'H-indole-2-carboxylic acid (described in example 24),
Example 27

1'-[(6-chloro-2-(piperazin-1-ylcarbonyl)-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure IV described hereinabove:
- Amine: commercially available piperazine,
- Acid: 6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl]-1H-indole-2-carboxylic acid (described in example 24),

ES-MS m/e (%): 513 (M+H+).

Example 28

r-[(6-chloro-2-(morpholin-4-ylcarbonyl)-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure IV described hereinabove:
- Amine: commercially available morpholine,
- Acid: 6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl]-1H-indole-2-carboxylic acid (described in example 24),
Example 29

6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-yl)carbonyl]-N,N-dimethyl-1H-indole-2-carboxamide

Amide coupling according to general procedure IV described hereinabove:
- Amine: commercially available dimethylamine,
- Acid: 6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-yl)carbonyl]-1H-indole-2-carboxylic acid (described in example 24),

ES-MS m/e (%): 512 (M+H+).

Example 30

tert-butyl {2-[(6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-yl)carbonyl]-1H-indol-2-yl]carbonyl}amino[ethyl]methylcarbamate

Amide coupling according to general procedure IV described hereinabove:
- Amine: commercially available (2-Amino-ethyl)-methyl-carbamic acid tert-butyl ester,
- Acid: 6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-yl)carbonyl]-1H-indole-2-carboxylic acid (described in example 24),

ES-MS m/e (%): 471 (M+H+).
ES-MS m/e (%): 599 (M+H+).

Example 31
6-chloro-N,N-diethyl-3-[(5-fluoro-3-oxo-l'H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl]-lH-indole-2-carboxamide

Amide coupling according to general procedure IV described hereinabove:
- Amine: commercially available diethyl-amine,
- Acid: 6-chloro-3-[(5-fluoro-3-oxo-l'H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl]-lH-indole-2-carboxylic acid (described in example 24),

ES-MS m/e (%): 498 (M+H+).

Example 32
r-[(6-chloro-2-[(4-methylpiperazin-1-yl)carbonyl]-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure IV described hereinabove:
- Amine: commercially available 1-methyl-piperazine,
- Acid: 6-chloro-3-[(5-fluoro-3-oxo-l'H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl]-lH-indole-2-carboxylic acid (described in example 24),
ES-MS m/e (%): 524 (M+H+).

Example 33
l'-[(6-chloro-2-(piperidin-1-ylcarbonyl)-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one

Amide coupling according to general procedure IV described hereinabove:
- Amine: commercially available piperidine,
- Acid: 6-chloro-3-[(5-fluoro-3-oxo-rH,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-1H-indole-2-carboxylic acid (described in example 24),

ES-MS m/e (%): 510 (M+H+).

Example 34
r-[(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]-7H-spiro[furo[3,4-
f][1,3]benzodioxole-5,4'-piperidin]-7-one

Amide coupling according to general procedure I described hereinabove:
- Amine: 7H-spiro[furo[3,4-f][1,3]benzodioxole-5,4'-piperidin]-7-one prepared as described in DE2458176A1,
- Acid: l-Benzyl-2-methyl-1H-indole-3-carboxylic acid (described in example 1),
ES-MS m/e (%): 495 (M+H+).

Example 35

3-{6-chloro-3-[3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl]carbonyl]-1H-indol-1-yl}propanenitrile

To a stirred solution of 100 mg (0.26 mmol) of 1'-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]-3-one (the preparation of which has been described in example 16) in 5 ml DMF was added 11.5 mg (0.28 mmol) NaH (60% in oil). The mixture was stirred at RT for 30 min. and then 35 mg (0.26 mmol) of 3-bromo-propionitrile was added. The mixture was stirred an additional hour and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na₂SO₄ and concentrated in vacuo. Flash chromatography (CH₂Cl₂/MeOH, 99:1; SiO₂) afforded 48 mg (29%) of 3-{6-chloro-3-[3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl]carbonyl]-1H-indol-1-yl}propanenitrile as a light yellow solid.

ES-MS m/e (%): 434(M+H+).

Example 36

4-{6-chloro-3-[3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1-yl]carbonyl]-1H-indol-1-yl}butanenitrile
To a stirred solution of 100 mg (0.26 mmol) of \((\text{6-chloro-lH-indol-3-yl})\text{carbonyl}\)-3H-spiro[2-benzofuran-1,4'-piperidine]-3-one (the preparation of which has been described in example 16) in 5 ml DMF was added 11.5 mg (0.28 mmol) NaH (60% in oil). The mixture was stirred at RT for 30 min. and then 37 mg (0.26 mmol) of 4-bromo- butyronitrile was added. The mixture was stirred overnight and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na$_2$SO$_4$ and concentrated in vacuo. Flash chromatography (CH$_2$Cl$_2$/MeOH, 99:1; SiO$_2$) afforded 69 mg (59%) of 4-\{(6-chloro-3-\{(3-oxo-lH,3H-spiro[2-benzofuran-1,4'-piperidin]-1-yl)\}carbonyl\}-lH-indol-1-yl\}butanenitrile as a light yellow solid.

ES-MS m/e (%): 448(M+H$^+$).

Example 37

\{6-chloro-3-\{(3-oxo-rH,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)\}carbonyl\}-lH-indol-1-yl\}acetonitrile

To a stirred solution of 100 mg (0.26 mmol) of \((\text{6-chloro-lH-indol-3-yl})\text{carbonyl}\)-3H-spiro[2-benzofuran-1,4'-piperidine]-3-one (the preparation of which has been described in example 16) in 5 ml DMF was added 11.5 mg (0.28 mmol) NaH (60% in oil). The mixture was stirred at RT for 30 min. and then 31 mg (0.26 mmol) of bromo-
acetonitrile was added. The mixture was stirred overnight and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na₂SO₄ and concentrated in vacuo. Flash chromatography (EtOAc/Hx, 1:6; SiO₂) afforded 58 mg (34%) of [6-chloro-3-[(3-oxo-l H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl]carbonyl]-lH-indol-1-yl]acetonitrile as a light yellow solid.

ES-MS m/e (%): 420(M+H⁺).

Example 38

2-[(6-chloro-3-[(3-oxo-l'H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl]carbonyl]-lH-indol-1-yl]propanenitrile

To a stirred solution of 100 mg (0.26 mmol) of l'-(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (the preparation of which has been described in example 16) in 5 ml DMF was added 11.5 mg (0.28 mmol) NaH (60% in oil). The mixture was stirred at RT for 30 min. and then 35 mg (0.26 mmol) of 2-bromo-propionitrile was added. The mixture was stirred overnight and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na₂SO₄ and concentrated in vacuo. Flash chromatography (EtOAc/Hx, 2:1; SiO₂) afforded 54 mg (34%) of 2-[(6-chloro-3-[(3-oxo-l'H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-lH-indol-1-yl]propanenitrile as a light yellow solid.

ES-MS m/e (%): 434(M+H⁺).

Examples of compounds of formula (I-d)

Example 39

r-[(1-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]
Amide coupling according to general procedure I described hereinabove:
- Amine: spiro[isobenzofuran-1(3H),4'-piperidine] prepared as described in *J. Org. Chem.* 1976, 41, 2628,
- Acid: l-Benzyl-2-methyl-lH-indole-3-carboxylic acid (described in example 1).
ES-MS m/e (%): 437.5 (M+H+).

**Example 40**

r-[(l-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-6-chloro-3H-spiro[2-benzofuran-1,4'-piperidine]

Amide coupling according to general procedure I described hereinabove:
- Amine: 6-chloro-3H-spiro[2-benzofuran-1,4'-piperidine] prepared as described in WO2004004714,
- Acid: l-Benzyl-2-methyl-lH-indole-3-carboxylic acid (described in example 1)
ES-MS m/e (%): 471.3 (M+H+).
Example 41

l'-[(1-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-3-phenyl-3H-spiro[2-benzofuran-1,4'-piperidine]

Amide coupling according to general procedure I described hereinabove:
- Amine: 3-phenyl-3H-spiro[2-benzofuran-1,4'-piperidine] prepared as described in J. Med. Chem. 1976, 19, 1315,
- Acid: l-Benzyl^-methyl-lH-indole-S-carboxylic acid (described in example 1)

ES-MS m/e (%): 513.6 (M+H^+).

Example 42

r-[(l-benzyl-5-methoxy-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]

Amide coupling according to general procedure I described hereinabove:
- Amine: spiro[isobenzofuran-l(3H),4'-piperidine] prepared as described in J. Org. Chem. 1976, 41, 2628,
- Acid: l-benzyl-S-methoxy-l-methyl-lH-indole-S-carboxylic acid (commercially available),
  ES-MS m/e (%): 467.4 (M+H +).

Example 43

\[
\text{r-}[\text{(2-methyl-lH-indol-3-yl)carbonyl}]-3H\text{-spiro}[2\text{-benzofuran-1,4'-piperidine}]
\]

Amide coupling according to general procedure I described hereinafore:
- Amine: spiro[isobenzofuran-l(3H),4'-piperidine] prepared as described in J. Org. Chem. 1976, 41, 2628,
- Acid: 2-Methyl-lH-indole-3-carboxylic acid (described in J. Heterocyclic Chem. 1977, 14, 1123),
  ES-MS m/e (%): 347.3 (M+H +).

Example 44

\[
\text{l'-[(l-benzoyl-2-methyl-lH-indol-3-yl)carbonyl}-3H\text{-spiro}[2\text{-benzofuran-1,4'-piperidine}]
\]

To a stirred solution of 40 mg (0.11 mmol) of l'-[(2-methyl-lH-indol-3-yl)carbonyl]-
3H-spiro[2-benzofuran-1,4'-piperidine] (the preparation of which has been described in example 43) in 3 ml DMF was added 5 mg (0.11 mmol) NaH (60% in oil). The mixture was stirred at RT for 30 min. and then 19 mg (0.13 mmol) of benzoyl chloride was added.
The mixture was stirred an additional hour and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na₂SO₄ and concentrated in vacuo. Re-crystallization in Et₂O afforded 51 mg (98%) of l’-[l-benzoyle-2-methyl-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] as white crystals.

ES-MS m/e (%): 451.3 (M+H+).

Example 45
r-[[2-methyl-l-(phenylsulfonyl)-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine]

To a stirred solution of 40 mg (0.11 mmol) of l’-[2-methyl-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (the preparation of which has been described in example 43) in 3 ml DMF was added 5 mg (0.11 mmol) NaH (60% in oil). The mixture was stirred at RT for 30 min. and then 24 mg (0.14 mmol) of benzenesulfonyl chloride was added. The mixture was stirred an additional hour and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na₂SO₄ and concentrated in vacuo. Re-crystallization in Et₂O afforded 45 mg (80%) of l’-[2-methyl-l-(phenylsulfonyl)-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] as white crystals. ES-MS m/e (%): 487.4 (M+H+).

Example 46
r-[[l-(cyclohexylmethyl)-2-methyl-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine]
Following the general procedure II described hereinabove above with bromomethyl-cyclohexane as electrophile, the title compound was obtained as white solid. ES-MS m/e (%): 443.5 (M+H^+).

Example 47

1'{-[l-(3-fluorobenzyl)-2-methyl-lH-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure II described hereinabove with 1-bromomethyl-3-fluorobenzene as electrophile; the title compound was obtained as white solid. ES-MS m/e (%): 455.4 (M+H^+).

Example 48

r-{(2-methyl-l-[2-(trifluoromethoxy)benzyl]-lH-indol-3-yl)carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidine]
Following the general procedure II described hereinabove with 1-bromomethyl-2-trifluoromethoxy-benzene as electrophile; the title compound was obtained as white solid.

ES-MS m/e (%): 521.4 (M+H+).

Example 49

1'-[[1-(3,5-dimethylbenzyl)-2-methyl-1H-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure II described hereinabove with 1-bromomethyl-3,5-dimethyl-benzene as electrophile, the title compound was obtained as white solid.

ES-MS m/e (%): 465.4 (M+H+).

Example 50

methyl 4-[[2-methyl-3-(1'H,3H-spiro[2-benzofuran-1,4'-piperidin]-l'-ylcarbonyl)-1H-indol-1-yl]methyl]benzoate
Following the general procedure II described hereinabove with 4-bromomethyl-benzoic acid methyl ester as electrophile; the title compound was obtained as white solid.
ES-MS m/e (%): 495.5 (M+H+).

Example 51
4-[[2-methyl-3-(rH,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl carbonyl)-lH-indol-1-yl]methyl]benzonitrile

Following the general procedure II described hereinabove with 4-bromomethyl-benzonitrile as electrophile; the title compound was obtained as white solid.
ES-MS m/e (%): 462.4 (M+H+).

Example 52
1'-(1-(3,5-difluorobenzyl)-2-methyl-lH-indol-3-yl[carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]
Following the general procedure II described hereinabove with 1-bromomethyl-3,5-difluoro-benzene as electrophile; the title compound was obtained as white solid.

ES-MS m/e (%): 473.4 (M+H⁺).

Example 53

l'-(l-(2-chlorobenzyl)-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure II described hereinabove with 1-bromomethyl-2-chloro-benzene as electrophile; the title compound was obtained as white solid.

ES-MS m/e (%): 471.3 (M+H⁺).

Example 54

l'-(l-(2-methoxybenzyl)-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]
Following the general procedure II described hereinabove with 1-chloromethyl-2-methoxy-benzene as electrophile; the title compound was obtained as white solid.

ES-MS m/e (%): 467.4 (M+H+).

Example 55

1'-%{[l-(4-methoxybenzyl)-2-methyl-lH-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure II described hereinabove with 1-chloromethyl-4-methoxy-benzene as electrophile; the title compound was obtained as white solid.

ES-MS m/e (%): 467.4 (M+H+).

Example 56

1'-%{[l-{[2-(4-chlorophenyl)-l,3-thiazol-4-yl]methyl}-2-methyl-lH-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidine]
Following the general procedure II described hereinabove with 4-chloromethyl-2-(4-chloro-phenyl)-thiazole (commercially available) as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 554.3 (M+H+).

Example 57

l'-{[1-[[2-(4-chlorophenyl)-1,3-thiazol-4-yl]methyl]-2-methyl-lH-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure II described hereinabove with 4-bromomethyl-2-(4-chloro-phenyl)-5-methyl-thiazole (the preparation of which has been described in WO2004020420) as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 568.3 (M+H+).

Example 58

l'-{[1-[[2-(2-methoxyphenyl)-5-methyl-1,3-oxazol-4-yl]methyl]-2-methyl-lH-indol-3-yl]carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidine]
Following the general procedure II described hereinabove with 4-chloromethyl-2-(2-methoxy-phenyl)-5-methyl-oxazole (the preparation of which has been described in WO2002092084 & WO2004031162) as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 548.5 (M+H+).

Example 59

r-[[2-methyl-l-{{5-methyl-2-[3-(trifluoromethyl)phenyl]-1,3-oxazol-4-yl}methyl}-1H-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure II described hereinabove with 4-chloromethyl-5-methyl-2-(3-trifluoromethyl-phenyl)-oxazole (the preparation of which has been described in WO2004031162) as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 586.3 (M+H+).

Example 60

r-[[l-{{2-(2-fluorophenyl)-5-methyl-1,3-oxazol-4-yl}methyl}-2-methyl-1H-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]
Following the general procedure II described hereinabove with 4-chloromethyl-2-(2-fluoro-phenyl)-5-methyl-oxazole (commercially available) as electrophile, the title compound was obtained as a white solid.

5 ES-MS m/e (%): 536.4 (M+H^+).

Example 61

r'-(1-{{2-(4-isopropylphenyl)-5-methyl-1,3-oxazol-4-yl}methyl}-2-methyl-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure II described hereinabove with 4-chloromethyl-2-(4-isopropyl-phenyl)-5-methyl-oxazole (the preparation of which has been described in WO2002092084 & WO2004031162) as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 560.4 (M+H^+).

Example 62

r-[(1-{{2-(4-ethylphenyl)-5-methyl-1,3-oxazol-4-yl}methyl}-2-methyl-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]
Following the general procedure II described hereinabove with 4-chloromethyl-2-(4-ethyl-phenyl)-5-methyl-oxazole (the preparation of which has been described in WO2002092084) as electrophile, the title compound was obtained as a white solid.

Example 63

r-[2-methyl-l-{{5-methyl-2-(2-methylphenyl)-1,3-oxazol-4-yl}methyl}-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidine]

Following the general procedure II described hereinabove with 4-chloromethyl-5-methyl-2-o-tolyl-oxazole (the preparation of which has been described in WO2004031162) as electrophile, the title compound was obtained as a white solid.

Example 64

r-[2-methyl-l-{{5-methyl-2-[4-(trifluoromethyl)phenyl]-1,3-oxazol-4-yl}methyl}-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidine]
Following the general procedure II described hereinabove with 4-chloromethyl-5-methyl-2-(4-trifluoromethyl-phenyl)-oxazole (the preparation of which has been described in *J. Med. Chem.* 2000, 43, 995-1010) as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 586.3 (M+H+).

**Example 65**

r-ICl-methyl-l-dS-methyl-l-tl-Ctrifluoromethyl^{phenyll-|}oxazoM-y1methyl|-!!|-indol-3-y1|carbonyl|]-3H-spio[2-benzofuran-l,4'-piperidine]

Following the general procedure II described hereinabove with 4-chloromethyl-5-methyl-2-(2-trifluoromethyl-phenyl)-oxazole (the preparation of which has been described in WO2004031162) as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 586.5 (M+H+).

**Example 66**

r-[(5-chloro-lH-indol-3-yl)carbonyl]-3H-spio[2-benzofuran-l,4'-piperidine]
Amide coupling according to general procedure I described hereinabove:
- Amine: spiro[isobenzofuran-l(3H),4'-piperidine] prepared as described in J.Org.Chem. 1976, 41, 2628,
- Acid: S-Chloro-lH-indole-S-carboxylic acid (commercially available), ES-MS m/e (%): 367.1 (M+H+).

Example 6

r-[(l-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidine]

Amide coupling according to general procedure I described hereinabove:
- Amine: spiro[isobenzofuran-l(3H),4'-piperidine] prepared as described in J.Org.Chem. 1976, 41, 2628,
- Acid: l-Methyl-lH-indole-S-carboxylic acid (commercially available), ES-MS m/e (%): 347.5 (M+H+).

Example 6

r-[(l-benzyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidine]
Amide coupling according to general procedure I described hereinabove:
- Acid: l-benzyl-lH-indole-3-carboxylic acid (described in example 3), ES-MS m/e (%): 423.6 (M+H⁺).

**Example 69**

r-[[(6-chloro-lH-indol-3-y1)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]

Amide coupling according to general procedure I described hereinabove:
- Acid: 6-chloro-lH-indole-3-carboxylic acid (the preparation of which is described hereinafter), ES-MS m/e (%): 367.2 (M+H⁺).

6-chloro-lH-indole-3-carboxylic acid

Using a procedure described in *J.Med. Chem.* 1991, 34, 140, from 7.0 g (0.046 mmol) of 6-chloro-lH-indole was prepared 5.80 g (64%) of 6-chloro-lH-indole-3-carboxylic acid as a light brown solid.

ES-MS m/e (%): 194 (M-H⁻).
Example 70

l'-(lH-indol-3-ylcarbonyl)-3H-spiro[2-benzofuran-1,4'-piperidine] Amide coupling according to general procedure I described hereinabove:

- Amine: spiro[isobenzofuran-l(3H),4'-piperidine] prepared as described in J. Org. Chem. 1976, 41, 2628,
- Acid: lH-Indole-3-carboxylic acid (commercially available), ES-MS m/e (%): 333.3 (M+H+).

Example 71

N,N-dimethyl-2-[3-(rH,3H-spiro[2-benzofuran-1,4'-piperidine]-r-ylcarbonyl)-lH-indol-1-yl]ethan amine

To a stirred solution of 20 mg (0.060 mmol) of l'-(lH-indol-3-ylcarbonyl)-3H-spiro[2-benzofuran-1,4'-piperidine] (the preparation of which has been described in example 70) in DMF (3 ml) at RT, was added 2.9 mg (0.072 mmol) of NaH (60% in oil). The mixture was stirred 20 min. and then 13 mg (0.072 mmol) of (2-chloro-ethyl)-dimethyl-amine in 1 ml of DMF was added. The mixture was stirred an additional 5 hours at 50°C and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na2SO4 and concentrated in vacuo. Flash chromatography (CH2Cl2/MeOH 8/2) afforded 11 mg (48%) of N,N-dimethyl-2-[3-(lH,3H-spiro[2-benzofuran-1,4'-
piperidin]-r-ylcarbonyl)-lH-indol-l-yl]ethanamine as a viscous oil. ES-MS m/e (%): 404.3 (M+H+).

Example 72
1-methyl-l-tS-CrH^H-spiroCl-benzofuran-l'-piperidinj-r-ylcarbony^-lH-indol-l-
yl]butan-l-one

Amide coupling according to general procedure I described hereinafore:
- Amine: spiro[isobenzofuran-l(3H),4'-piperidine] prepared as described in J. Org. Chem. 1976, 41, 2628,
- Acid: 2-(2-Methyl-butyryl)-lH-indole-3-carboxylic acid (commercially available), ES-MS m/e (%): 417 (M+H+).

Example 73
2-[6-chloro-3-(rH,3H-spiro[2-benzofuran-l,4'-piperidin]-l'-ylcarbonyl]-lH-indol-l-
yl]propanenitrile

To a stirred solution of 100 mg (0.26 mmol) of l'-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidine] (the preparation of which has been described in example 69) in 5 ml DMF was added 11.5 mg (0.28 mmol) NaH (60% in oil). The mixture was stirred at RT for 30 min. and then 35 mg (0.26 mmol) of 2-bromo-propionitrile was added. The mixture was stirred overnight and then poured onto water
and extracted with ethyl acetate. The combined organic phases were dried over Na₂SO₄
and concentrated in vacuo. Flash chromatography (EtOAc/Hx, 1:1; SiO₂) afforded 93 mg
(91%) of 2-[6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl]-IH-
indol-1-yl] propanenitrile as a light yellow solid.

ES-MS m/e (%): 420(M+H⁺).

Example 74
3-[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4'-piperidin]-l'-ylcarbonyl]-IH-indol-l-
yl]propanenitrile

![Chemical structure]

To a stirred solution of 100 mg (0.26 mmol) of 1'-(6-chloro-IH-indol-3-yl)carbonyl]-
3H-spiro[2-benzofuran-1,4'-piperidine] (the preparation of which has been described in
example 69) in 5 ml DMF was added 11.5 mg (0.28 mmol) NaH (60% in oil). The
mixture was stirred at RT for 30 min. and then 35 mg (0.26 mmol) of 3-bromo-
propionitrile was added. The mixture was stirred for two days at RT and then poured
onto water and extracted with ethyl acetate. The combined organic phases were dried
over Na₂SO₄ and concentrated in vacuo. Flash chromatography (EtOAc/Hx, 2:1; SiO₂)
afforded 105 mg (96%) of 3-[6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-l'-
ylcarbonyl]-IH-indol-1-yl] propanenitrile as a white solid.

ES-MS m/e (%): 420(M+H⁺).

Example 75
4-[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl]-IH-indol-l-
yl]butanenitrile
To a stirred solution of 100 mg (0.26 mmol) of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (the preparation of which has been described in example 69) in 5 ml DMF was added 11.5 mg (0.28 mmol) NaH (60% in oil). The mixture was stirred at RT for 30 min. and then 35 mg (0.26 mmol) of 4-bromo-acetonitrile was added. The mixture was stirred for two days at RT and then poured onto water and extracted with ethyl acetate. The combined organic phases were dried over Na₂SO₄ and concentrated in vacuo. Flash chromatography (EtOAc/Hx, 2:1; SiO₂) afforded 99 mg (88%) of 4-[6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidin]-r-ylcarbonyl)-1H-indol-1-yl]butanenitrile as a white solid.

ES-MS m/e (%): 434(M+H⁺).

Example 76

[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4’-piperidin]-r-ylcarbonyl)-IH-indol-1-yljacetonitrile

To a stirred solution of 100 mg (0.26 mmol) of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (the preparation of which has been described in example 69) in 5 ml DMF was added 11.5 mg (0.28 mmol) NaH (60% in oil). The mixture was stirred at RT for 30 min. and then 35 mg (0.26 mmol) of bromo-acetonitrile was added. The mixture was stirred overnight at RT and then poured onto water and
extracted with ethyl acetate. The combined organic phases were dried over Na₂SO₄ and concentrated in vacuo. Flash chromatography (EtOAc/Hx, 2:1; SiO₂) afforded 43 mg (41%) of [6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl] acetonitrile as a white solid.

ES-MS m/e (%): 406(M+H⁺).

**Examples of compounds of formula (I-e)**

**Example 77**

r-[(6-chloro-1H-indol-3-yl)carbonyl]spiro[2-benzofuran-1,4'-piperidine]

Amide coupling according to general procedure I described hereinabove:
- Amine: commercially available spiro[1-benzofuran-3,4'-piperidine],
- Acid: 6-chloro-1H-indole-3-carboxylic acid (described in example 5),
ES-MS m/e (%): 367.4 (M+H⁺).

**Example 78**

r-[(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]spiro[1-benzofuran-3,4'-piperidine]

Amide coupling according to general procedure I described hereinabove:
- Amine: commercially available spiro[1-benzofuran-3,4'-piperidine],
- Acid: 1-Benzyl-2-methyl-1H-indole-3-carboxylic acid (described in example 1),
ES-MS m/e (%): 437.6 (M+H⁺).
Example 79

`1'-(IH-indol-3-ylcarbonyl)spiro[l-benzofuran-3,4'-piperidine]`

Amide coupling according to general procedure I described hereinabove:
- Amine: commercially available spiro[l-benzofuran-3,4'-piperidine],
- Acid: IH-Indole-3-carboxylic acid (commercially available),
ES-MS m/e (%): 333.4 (M+H+).

Example 80

- [(6-chloro-S-fluoro-IH-indol-S-ylcarbonyl)spiro[l-benzofuran-3,4'-piperidine]`

Amide coupling according to general procedure I described hereinabove:
- Amine: commercially available spiro[l-benzofuran-3,4'-piperidine],
- Acid: 6-chloro-S-fluoro-IH-indole-S-carboxylic acid (the preparation of which is described hereinafter),
ES-MS m/e (%): 385.1 (M+H+).

a) 6-chloro-5-fluoro-IH-indole:
Following the procedure described in WO9747598, from 6-chloro-5-fluoro-IH-indole-2,3-dione was prepared 6-chloro-5-fluoro-IH-indole.

b) 6-chloro-5-fluoro-IH-indole-3-carboxylic acid:
Following a procedure described in *J. Med. Chem.* 1991, 34, 140, from 0.25 g (1.47 mmol) of 6-chloro-5-fluoro-1H-indole was prepared 0.35 g (90%) of 6-chloro-5-fluoro-1H-indole-3-carboxylic acid as a light brown solid. ES-MS m/e (%): 213 (M-H⁻).

**Example 81**

2-[6-chloro-3-[(1H-spiro[benzofuran-3,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl]-N,N-dimethylethan amino

Following the general procedure III as described hereinabove with commercially available (2-chloro-ethyl)-dimethyl-amine as electrophile, the title compound was obtained as colorless viscous oil.

ES-MS m/e (%): 438.1 (M+H⁺).

**Example 82**

r-[[6-chloro-1-(2-pyrrolidin-1-ylethyl)-1H-indol-3-yl]carbonyl]spiro[1-benzofuran-3,4'-piperidine]

Following the general procedure III as described hereinabove with commercially available 1-(2-chloro-ethyl)-pyrrolidine as electrophile, the title compound was obtained as a colorless viscous oil.

ES-MS m/e (%): 464.0 (M+H⁺).
Example 83

3-[6-chloro-3-(1H-spiro[1-benzofuran-3,4'-piperidin]-1'-ylcarbonyl)-1H-indol-1-yl]-N,N-dimethylpropan-1-amine

Following the general procedure III as described hereinabove with commercially available (3-chloro-propyl)-dimethyl-amine as electrophile, the title compound was obtained as colorless viscous oil.

ES-MS m/e (%): 452.0 (M+H+).

Example 84

1'-{[6-chloro-1-(2-morpholin-4-ylethyl)-1H-indol-3-yl]carbonyl}spiro[1-benzofuran-3,4'-piperidine]

Following the general procedure III as described hereinabove with commercially available 4-(2-chloro-ethyl)-morpholine as electrophile, the title compound was obtained as a colorless viscous oil.

ES-MS m/e (%): 480.1 (M+H+).

Example 85

2-[6-chloro-3-(1H-spiro[1-benzofuran-3,4'-piperidin]-1'-ylcarbonyl)-1H-indol-1-yl]-N,N-diethylethan amine
Following the general procedure III as described hereinabove with commercially available (2-bromo-ethyl)-diethyl-amine as electrophile, the title compound was obtained as colorless viscous oil.

5 ES-MS m/e (%): 466.2 (M+H+).

Example 86
l'-{(6-chloro-l-[2-(lH-pyrrol-l-yl)ethyl]-lH-indol-3-yl)carbonyl}spiro[l-benzofuran-3,4'-piperidine]

Following the general procedure III as described hereinabove with commercially available l-(2-chloro-ethyl)-lH-pyrrole as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 460.2 (M+H+).

Example 87
l'-{(6-chloro-l-(2-oxo-2-piperidin-l-ylethyl)-lH-indol-3-yl)carbonyl}spiro[l-benzofuran-3,4'-piperidine]
Following the general procedure III as described hereinabove with commercially available 2-chloro-1-piperidin-1-yl-ethanone as electrophile, the title compound was obtained as a white solid.

\[ \text{ES-MS } m/e \text{ (\%): 492.2 (M+H\textsuperscript{+}).} \]

**Examples of compounds of formula (I-b)**

**Example 88**

\[ \text{r-[(6-Chloro-l-(3-fluorobenzoyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine} \]

Following the general procedure VII as described above, the acylation of l'-[(6-chloro-lH-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4'-piperidine] (prepared according to example 8) with commercially available 3-fluorobenzoyl chloride gave the title compound.

\[ \text{ES-MS } m/e \text{ (\%): 488.5(M+H\textsuperscript{+}).} \]
Example 89

l’-[(6-Chloro-1-(2-fluorobenzoyl)-1H-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4’-piperidine

Following the general procedure VII as described above, the acylation of l’-[6-chloro-1H-indol-3-yl]carbonyl]-1,2-dihydrospiro[indole-3,4’-piperidine] (prepared according to example 8) with commercially available 2-fluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 488.4(M+H+).

Example 90

r-[(6-Chloro-1-(3,5-difluorobenzoyl)-1H-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4’-piperidine

Following the general procedure VII as described above, the acylation of l’-[6-chloro-1H-indol-3-yl]carbonyl]-1,2-dihydrospiro[indole-3,4’-piperidine] (prepared according
to example 8 above) with commercially available 3,5-difluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 506.4(M+H^+).

Example 91

Following the general procedure VII as described above, the acylation of l'-(6-chloro-lH-indol-3-yl)carbonyl)-1,2-dihydrospiro[indole-3,4'-piperidine] (prepared according to example 8) with commercially available 2,3-difluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 506.4(M+H^+).

Example 92
Following the general procedure VII as described above, the sulphonylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4’-piperidine] (prepared according to example 8) with commercially available 3,5-difluorobenzencesulfonyl chloride gave the title compound.

ES-MS m/e (%): 542.4(M-H+).

Example 93

Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4’-piperidine] (prepared according to example 8) with commercially available 3,5-difluorobenzyl chloride gave the title compound.

ES-MS m/e (%): 492.4(M+H+).
Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4'-piperidine] (prepared according to example 8) with commercially available 3-fluorobenzyl chloride gave the title compound.

ES-MS m/e (%): 474.4(M+H+).

Example 95

Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4'-piperidine] (prepared according to example 8) with commercially available 2-chloro-1-(3,5-difluorophenyl)-ethanone gave the title compound.
Example 96

2-[6-Chloro-3-(1,2-dihydro-1H-spiro[indole-3,4'-piperidin]-1'-ylcarbonyl)-1H-indol-1-yl]-1-(3,4-difluorophenyl)ethanone

Following the general procedure III as described above, the alkylation of l'-[(6-chloro-1H-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4'-piperidine] (prepared according to example 8) with commercially available 2-chloro-l-(3,4-difluoro-phenyl)-ethanone gave the title compound.

ES-MS m/e (%): 520.4(M+H+).

Example 97

2-[6-Chloro-3-(1,2-dihydro-rH-spiro[indole-3,4'-piperidin]-r-ylcarbonyl]-1H-indol-1-yl]-l-(2-fluorophenyl)ethanone

Following the general procedure III as described above, the alkylation of l'-[(6-chloro-1H-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4'-piperidine] (prepared according...
to example 8) with commercially available 2-chloro-l-(2-fluoro-phenyl)-ethanone gave the title compound.

ES-MS m/e (%): 502.5(M+H +).

Example 98

2-[6-Chloro-3-(1,2-dihydro-1H-spiro[indole-3,4'-piperidin]-l'-ylcarbonyl)-1H-indol-1-yl]-N,N-diethylethan amine

Following the general procedure III as described above, the alkylation of l'-[(6-chloro-1H-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4'-piperidine] (prepared according to example 8) with commercially available diethylaminoethyl bromide gave the title compound.

ES-MS m/e (%): 465.4(M+H +).

Example 99

2-[6-Chloro-3-(1,2-dihydro-1H-spiro[indole-3,4'-piperidin]-l'-ylcarbonyl)-1H-indol-1-yl]-N,N-diethylacetamide
Following the general procedure III as described above, the alkylation of \( l'-(6\text{-chloro-}lH\text{-indol-3-yl})\text{carbonyl}\)-l,2-dihydrospiro[indole-3,4'-piperidine] (prepared according to example 8) with commercially available 2-chloro-\( N,N\text{-diethyl-acetamide} \) gave the title compound.

ES-MS m/e (%): 479.5 (M+H\(^+\)).

Example 100

2-[6-Chloro-3-(1,2-dihydro-rH-spiro[indole-3,4'-piperidin]-l'-ylcarbonyl]-lH-indol-1-yl]-N,N-dimethylacetamide

Following the general procedure III as described above, the alkylation of \( l'-(6\text{-chloro-}lH\text{-indol-3-yl})\text{carbonyl}\)-l,2-dihydrospiro[indole-3,4'-piperidine] (prepared according to example 8) with commercially available 2-chloro-\( N,N\text{-diethyl-acetamide} \) gave the title compound.

ES-MS m/e (%): 451.5 (M+H\(^+\)).

Example 101
2-[6-Chloro-3-(1,2-dihydro-rH-spiro[indole-3,4'-piperidin]-r-ylcarbonyl)-lH-indol-1-yl]-l-pyridin-2-ylethanone

Following the general procedure III as described above, the alkylation of 1'-%[6-chloro-lH-indol-3-yl]carbonyl]-l,2-dihydrospiro[indole-3,4'-piperidine] (prepared according to example 8) with commercially available 2-chloro-l-pyridin-2-yl-ethanone gave the title compound.

ES-MS m/e (%): 485.4(M+H+).

Example 102

Following the general procedure III as described above, the alkylation of 1'-%[6-chloro-lH-indol-3-yl]carbonyl]-l,2-dihydrospiro[indole-3,4'-piperidine] (prepared according to example 8) with commercially available 3-bromomethyl-pyridine gave the title compound.

ES-MS m/e (%): 499.4(M+H+)
Following the general procedure III as described above, the alkylation of 1'-(6-chloro-1H-indol-3-yl)carbonyl]-1,2-dihydrospiro[indole-3,4'-piperidine] (prepared according to example 8) with methanesulfonic acid pyridin-2-ylmethyl ester (described in WO 9955318) gave the title compound.
Following the general procedure V as described hereinabove, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (the preparation of which has been described in example 16) with commercially available 2-chloro-1-piperidin-1-yl-ethanone as electrophile, the title compound was obtained as a white powder.

ES-MS m/e (%): 506.0 (M+H+).

Example 105

1’-[(6-Chloro-1-(2-morpholin-4-yl-2-oxoethyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure V as described hereinabove, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (the preparation of which has been described in example 16) with commercially available 2-
chloro-l-morpholin-4-yl-ethanone as electrophile, the title compound was obtained as a white powder.

ES-MS m/e (%): 508.1 (M+H+).

Example 106

5 2-{6-Chloro-3-[(3-oxo-l'H,3H-spiro[2-benzofuran-l,4'-piperidin]-r-yl)carbonyl]-lH-indol-l-yl]-N,N-dimethylacetamide

Following the general procedure V as described hereinabove, the alkylation of l’-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (the preparation of which has been described in example 16) with commercially available 2-chloro-N,N-dimethylacetamide as electrophile, the title compound was obtained as a white powder.

ES-MS m/e (%): 466.1 (M+H+).

Example 107

15 2-{6-Chloro-3-[(3-oxo-l'H,3H-spiro[2-benzofuran-l,4'-piperidin]-r-yl)carbonyl]-lH-indol-l-yl]-N,N-diethylacetamide
Following the general procedure V as described hereinabove, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (the preparation of which has been described in example 16) with commercially available 2-chloro-N,N-diethyl-acetamide as electrophile, the title compound was obtained as a white powder.

ES-MS m/e (%): 494.1 (M+H+).

Example 108

1’-[(6-Chloro-1-(piperidin-1-ylcarbonyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-
benzofuran-1,4’-piperidin]-3-one

Following the general procedure V as described hereinabove, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (the preparation of which has been described in example 16) with commercially available piperidine-1-carbonyl chloride as electrophile, the title compound was obtained as a white powder.

ES-MS m/e (%): 492.1 (M+H+).
Example 109
tert-Butyl {6-chloro-3-[(3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl]-1H-indol-1-yl}acetate

Following the general procedure V as described hereinabove, the alkylation of 1'-(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (the preparation of which has been described in example 16) with commercially available chloro-acetic acid tert-butyl ester as electrophile, the title compound was obtained as a white powder.

ES-MS m/e (%): 495.2 (M+H+).

Example 110

2-[(6-Chloro-3-[(5-fluoro-3-oxo-rH,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl]-1H-indol-1-yl]-N,N-dimethylacetamide

Following the general procedure V as described hereinabove, the alkylation of 1'-(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one
(the preparation of which has been described in example 19) with commercially available 2-chloro-N,N-dimethyl-acetamide as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 484.0 (M+H+).

Example 111

r-({6-Chloro-l-[2-(dimethylamino)ethyl]-lH-indol-3-yl}carbonyl)-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one

Following the general procedure V as described hereinabove, the alkylation of l'-{(6-chloro-lH-indol-3-yl)carbonyl}-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one (the preparation of which have been described in example 19) with commercially available (2-chloro-ethyl)-dimethyl-amine as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 470.1 (M+H+).

Example 112

tert-Butyl {6-chloro-3-{(5-fluoro-3-oxo-lH ,3H-spiro[2-benzofuran-l,4'-piperidin]-r-yl)carbonyl] -lH-indol- 1-yl}acetate
Following the general procedure V as described hereinabove, the alkylation of 1’-[(6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (the preparation of which have been described in example 19) with commercially available chloro-acetic acid tert-butyl ester as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 513.2 (M+H+).

Example 113

l’-[(6-Chloro-l-(2-morpholin-4-yl-2-oxoethyl)-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

Following the general procedure V as described hereinabove, the alkylation of 1’-[(6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (the preparation of which have been described in example 19) with commercially available 2-chloro-l-morpholin-4-yl-ethanone as electrophile, the title compound was obtained as a white solid.
Following the general procedure V as described hereinabove, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (the preparation of which have been described in example 19) with commercially available 2-chloro-l-(4-methyl-piperazin-l-yl)-ethanone as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 539.4 (M+H+).

Example 115

2-{6-Chloro-3-[5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4’-piperidin]-r-yl]carbonyl]-lH-indol-1-yl}acetamide
Following the general procedure V as described hereinabove, the alkylation of 1’-[(6-chloro-1H-indol-3-y1)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (the preparation of which have been described in example 19) with commercially available chloro-acetic acid methyl ester as electrophile, an ester intermediate methyl [6-chloro-3’-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4’-piperidin]-1’-yl)carbonyl]-1H-indol-1-yl]acetate was obtained as a white solid.

This intermediate compound, methyl [6-chloro-3’-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4’-piperidin]-1’-yl)carbonyl]-1H-indol-1-yl]acetate, was then hydrolysed using standard conditions (aq.NaOH IM, MeOH, room temperature) to give [6-chloro-3’-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4’-piperidin]-1’-yl)carbonyl]-1H-indol-1-yl]acetic acid as a white solid.

An amide coupling between [6-chloro-3’-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4’-piperidin]-1’-yl)carbonyl]-1H-indol-1-yl]acetic acid and NH₂OH (25% in water) afforded the title compound as a white solid.

ES-MS m/e (%): 456.4 (M+H⁺).

Example 116

2’-{6-Chloro-3’-[(5-fluoro-3-oxo-rH,3H-spiro[2-benzofuran-1,4’-piperidin]-r-yl)carbonyl] -1H-indol- 1-yl}-N-methylacetamide

An amide coupling between [6-chloro-3’-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4’-piperidin]-1’-yl)carbonyl]-1H-indol-1-yl]acetic acid (described herein above) and methyl amine afforded the title compound as a white solid.

ES-MS m/e (%): 470.3 (M+H⁺).
Example 117

l’-{[6-Chloro-l-(2-oxo-2-piperazin-1-ylethyl)-lH-indol-3-yl]carbonyl}-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

Following the general procedure V as described hereinabove, the alkylation of l’-{[6-chloro-lH-indol-3-yl]carbonyl}-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (the preparation of which have been described in example 19) with commercially available 4-(2-Chloro-acetyl)-piperazine-l-carboxylic acid tert-butyl ester as electrophile, the title compound was obtained as a white solid after removal of the Boc protecting group under standard conditions (TFA/ dichloromethane, room temperature).

ES-MS m/e (%): 525.1 (M+H+).

Example 118

2-{6-Chloro-3-[(5-fluoro-3-oxo-rH,3H-spiro[2-benzofuran-1,4’-piperidin]-r-yl]carbonyl]-lH-indol-1-yl]-N-[2-(dimethylamino)ethyl]acetamide
An amide coupling between {6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4′-piperidin]-l′-yl)carbonyl]-IH-indol-3-yl}acetic acid (described herein above) and N,N-dimethyl-ethane-1,2-diamine afforded the title compound as a white solid.

ES-MS m/e (%): 527.2 (M+H+).

Example 119

2-[(6-Chloro-3-[(3-oxo-lH,3H-spiro[2-benzofuran-1,4′-piperidin]-l′-yl)carbonyl]-IH-indol-3-yl]-N,N-dimethylacetamide

An amide coupling between {6-chloro-3-[(3-oxo-lH,3H-spiro[2-benzofuran-1,4′-piperidin]-l′-yl)carbonyl]-IH-indol-3-yl}acetic acid (described herein above) and N,N-dimethyl-ethane-1,2-diamine afforded the title compound as a white solid.

ES-MS m/e (%): 509.2 (M+H+).

Example 120

2-[(6-Chloro-5-methyl-3-[(3-oxo-lH,3H-spiro[2-benzofuran-1,4′-piperidin]-l′-yl)carbonyl]-IH-indol-3-yl]-N,N-dimethylacetamide

a) l-(6-Chloro-5-methyl-lH-indol-3-yl)-2,2,2-trifluoroo-ethanone
Using a procedure described in J. Med. Chem. 1991, 34, 140, from 0.250 g (0.002 mol) of 6-chloro-5-methyl-1H-indole were prepared 0.38 g (96%) of 1-(6-chloro-5-methyl-1H-indol-3-yl)-2,2,2-trifluoro-ethanone as a white solid.

b) 2-r6-Chloro-5-methyl-3-(2,2,2-trifluoro-acetyl)-indol-1-yl-N,N-dimethyl-acetamide

\[\text{Using a similar procedure as described in J. Med. Chem. 1991, 34, 140, from 0.280 g of 2-}\]

\[\text{[6-chloro-5-methyl-3-(2,2,2-trifluoro-acetyl)-indol-1-yl]-N,N-dimethyl-acetamide were prepared 0.18 g (76%) of the title compound as a white solid.}\]

c) 6-Chloro-1-dimethylcarbamoylmethyl-5-methyl-1H-indole-3-carboxylic acid

\[\text{Using a similar procedure as described in J. Med. Chem. 1991, 34, 140, from 0.280 g of 2-}\]

\[\text{[6-chloro-5-methyl-3-(2,2,2-trifluoro-acetyl)-indol-1-yl]-N,N-dimethyl-acetamide were prepared 0.18 g (76%) of the title compound as a white solid.}\]

d) 2-{6-chloro-5-methyl-3-r(3-oxo-1H,3H-spiro2-benzofuran-1,4'-piperidinl-r-vDcarbonyll-1H-indol-1-yl]-N,N-dimethylacetamide}
Amide coupling according to general procedure I described hereinabove:


- Acid: 6-Chloro-1-dimethylcarbamoylmethyl-S-methyl-S-methyl-lH-indole-S-carboxylic acid, ES-MS m/e (%): 480.3 (M+H+).

Example 121

1'-(1-Benzyl-2-methyl-lH-indol-3-yl)carbonyl]-5-bromo-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

a) 5-Bromo-r-methyl-3H-spiro2-benzofuran-1,4'-piperidinl-3-one

Butyllithium (97.2 ml of 1.47 M solution in hexane, 143 mmol) was added dropwise to a solution of 2,5-Dibromo-benzoic acid (20 g, 72 mmol) in dry THF (300 ml) at -78° C over a period of 3.5 h under a nitrogen atmosphere. The reaction mixture was stirred at -78 °C for 2 h. A solution of N-methyl piperidone (11.31 g, 99 mmol) in hexane (40 mL) was added dropwise during 30 min to the reaction mixture at -78 °C. The reaction mixture was allowed to come to room temperature and stirring was continued for overnight. The reaction mixture was added to a mixture of water (500 ml) and ether (300 mL). The aqueous layer was extracted with ether (5 X 150 mL) and acidified with
concentrated HCl (to pH 2-3) and extracted with ether (2 X 150 ml). The acidic solution was boiled for 1 h and then cooled to 0-5 °C and made alkaline (to pH 9-10) with aqueous NaOH. The cold solution was rapidly extracted with chloroform (5 X 300 mL). The combined chloroform extracts were washed with water (150 ml), dried over sodium sulfate and evaporated under reduced pressure. The residue was purified by silica gel (100-200) column chromatography eluting with methanol in dichloromethane (0.5% to 2.5%) to afford the desired product (4.2 g, 20%).

$^1$H-NMR (400MHz, CDCl$_3$): $\delta$ 1.71 (d, $J = 14.2$ Hz, 2H), 2.15-2.24 (m, 2H), 2.37 (s, 3H), 2.45-2.52 (m, 2H), 2.83-2.87 (m, 2H), 7.26 (d, $J = 8.25$ Hz, IH), 7.75 (dd, $J = 8.0$, 1.7 Hz, IH).

$^1$C-NMR (100MHz, CDCl$_3$): $\delta$ 35.95, 46.05, 51.42, 84.00, 122.54, 122.97, 127.52, 128.64, 137.06, 152.24, 167.77.

b) 5-Bromo-3-oxo-1 H,3H-spiror2-benzofuran-l,4'-piperidinel-r-carbonitrile

A solution of the N-methylpiperidine (3.0 g, 10 mmol) in chloroform (50 ml) was added dropwise to a stirred boiling solution of cyanogen bromide (12.16 g, 120 mmol) in chloroform (100 ml) under a nitrogen atmosphere and the resulting solution was refluxed for overnight. The reaction mixture was cooled and washed with 25 mL of 5% HCl and then with 20 ml of water. The organic layer was dried over sodium sulfate and evaporated under reduced pressure. The residue was purified by silica gel column chromatography (100-200) eluting with methanol in dichloromethane (0.5% to 1.0%) to get the pure product (1.6 g, 51%).

$^1$H-NMR (400MHz, CDCl$_3$): $\delta$ 1.72 (d, $J = 14.2$Hz, 2H), 2.24-2.32 (m, 2H), 3.37-3.59 (m, 4H), 7.32 (d, $J = 8.2$ Hz, IH), 7.83 (dd, $J = 8.0$, 1.7Hz, IH), 8.03 (d, $J = 1.7$Hz, IH).

c) 5-Bromo-3H-spiror2-benzofuran-l,4'-piperidinl-3-one
A mixture of cyanoamine (1.0 g, 3.2 mmol) and 20%HCl (12 ml) was heated under reflux under a nitrogen atmosphere for 6 h. The reaction mixture was cooled to 0-5 °C and pH was adjusted to 9-10 with aqueous NaOH solution and rapidly extracted with chloroform (3 X 50 ml). The combined extracts were washed with water, the organic layer was dried over sodium sulfate and evaporated under reduced pressure. The residue was washed with distilled hexane and dried under high vacuum to get the pure product (0.64 g, 70%).

IR (KBr) 3333.84, 290.53, 2835.25, 2811.07, 2749.38, 1756.04, 1470.28, 1415.14, 1271.03, 1196.28, 1083.84, 929.07, 831.50, 792.35, 734.78, 691.24, 548.46, 534.50 cm⁻¹. ¹H-NMR (400MHz, CDCl₃): δ 1.66-1.72 (m, 2H), 2.02-2.09 (m, 2H), 3.07-3.18 (m, 4H), 7.29 (d, J = 7.8 Hz, IH), 7.77 (dd, J = 7.8, 1.7 Hz, IH), 7.99 (d, J = 1.7Hz, IH). ¹³C-NMR (100MHz, CDCl₃): δ 6.33, 42.49, 85.23, 122.61, 122.93, 127.39, 128.64, 137.07, 152.44, 167.91. HA-MS: 282.1 and 284.1; C₁₂H₁₂⁷BrNO₂ [MH⁺] requires 282.1. mp: 162-163 °C.

d) r-r(l-Benzyl-2-methyl-lH-indol-3-yl)carbonyll-5-bromo-3H-spiror2-benzofuran-1,4'-piperidinl-3 -one

Following the general procedure I as described above, the acylation of 5-bromo-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one with l-benzyl-2-methyl-lH-indole-3-carboxylic acid (preparation described in example 1), gave the title compound.

ES-MS m/e (%): 531.5(M+H⁺).
Example 122

1’-{(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl}-6-(2-hydroxyethoxy)-3H-spiro[2-
benzofuran-1,4’-piperidin]-3-one

a) 6-fluoro-r-methyl-3H-spiro2-benzofuran-l,4'-piperidin-3-one

5

To a solution of the substituted 2-bromo-4-fluoro-benzoic acid (10.9 g, 50 mmol) in dry
THF (200 ml) at -78 °C n-butyllithium (1.6 M in hexanes) (100 mmol) was added drop
wise (3 h) and the resulting solution was stirred for an additional 2 h at the same
temperature. Freshly distilled N-methyl 4-piperidone (7.91 g, 70 mmol) in dry hexane
(25 ml) was added over 30 min at the same temperature. The mixture was then allowed to
stir at rt and was finally added to ether (200 ml) and water (300 ml). The basic (aqueous)
layer was extracted with ether (5 X 100 ml) and the aqueous layer was acidified with
concentrated hydrochloric acid (pH 2-3) and extracted with ether. The aqueous solution
was boiled for 1 h and was then cooled to 0-5 °C and made alkaline (pH 9-10) with cold
aqueous sodium hydroxide. The cold solution was rapidly extracted with chloroform (5 X
200 ml). The combined chloroform extracts were washed with water, dried, concentrated
to give light yellow solid which was purified over neutral alumina eluting with a gradient
of 30-50% ethyl acetate-hexane to obtain 1.75 g (15%) of the desired product as a white
solid.

1H-NMR (CDCl3, 400 MHz): δ 1.68-1.75 (m, 2H), 2.18-2.19 (m, IH), 2.38 (s, 3H), 2.44-
2.52 (m, 2H), 2.68-2.84 (m, 2H), 2.84-2.85 (m, IH), 7.02-7.05 (m, IH), 7.19-7.22 (m,
IH), 7.84-7.87 (m, IH); HA-MS: 236 (M + 1).

b) 6-Fluoro-3-oxo-l Η ,3H-spiro2-benzofuran-l,4'-piperidinel-r-carbonitrile
To a solution of the 6-fluoro-r-methyl-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (1.17 g, 5 mmol) in dry chloroform (10 ml) was added cyanogenbromide (60 mmol) and the resulting solution was refluxed for 36 h. The reaction mixture was extracted with 5% HCl (5 ml) and then with water (2.5 ml). The chloroform solution was dried (anhdyrous MgSO₄) and concentrated to give a pale yellow solid which was chromatographed over SiO₂ eluting with 1% MeOH-dichloromethane to give 858 mg (70%) of the desired product as a white solid.

1H-NMR (CDCl₃, 400 MHz): δ 1.72-1.76 (m, 2H), 2.22-2.30 (m, 1H), 3.48-3.60 (m, 4H), 7.09-7.11 (m, 1H), 7.11-7.28 (m, 1H), 7.89-7.92 (m, 1H); IR (KBr): 3492, 3043, 2216, 1760, 1602, 1478 cm⁻¹.

c) 5-(2-hydroxyethoxy)-3H-spiro2-benzofuran-1,4'-piperidin1-3-one

6-Fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-carbonitrile (1.23 g, 5 mmol) was heated with ethylene glycol (5 ml) and sodium hydroxide (0.82 g, 20.5 mmol) for 15-20 min at 130 °C. Most of the ethylene glycol was removed by distillation under high vacum. The residual reaction mixture was diluted with water and extracted repeatedly with chloroform. The combined organics was dried and concentrated to give a semi solid material which was purified over Al₂O₃ column upon elution with 5-7% MeOH/CH₂Cl₂ containing NH₃ (aqueous) to yield 789 mg (60%) of the desired product as a pale yellow solid.

1H-NMR (d₆-DMSO, 400 MHz): δ 1.47-1.50 (m, 2H), 2.03-2.10 (m, 2H), 2.79-2.85 (m, 2H), 2.95-2.97 (m, 2H), 3.73-3.76 (m, 2H), 4.12-4.14 (m, 2H), 7.09 (d, J = 8.4 Hz, 1H),
7.20 (s, IH), 7.69 (d, J = 8.4 Hz, IH); $^{13}$C-NMR ($d_6$-DMSO, 100 MHz) : $\delta$ 35.9, 42.3, 59.3, 70.4, 84.6, 106.4, 116.6, 117.0, 126.8, 156.9, 163.9, 168.5; HA-MS: 264.3 (M + 1).

d) r-rfl-Benzyl-2-methyl-lH-indol-3-yl)carbonyll-6-(2-hydroxyethoxy)-3H-spiro2-
benzofuran- 1,4'-piperidinl -3-one

Following the general procedure I as described above, the acylation of 5-(2-
hydroxyethoxy)-3H-spiro[2-benzofuran- 1,4'-piperidin]-3-one with 1-benzyl-2-methyl-
IH-indole-3-carboxylic acid (preparation described in example 1), gave the title 
compound.

ES-MS m/e (%): 511.6(M+H$^+$).

Example 123

1'-(1-[2-(3,4-Dimethoxyphenyl)ethyl]-5-methoxy-2-methyl-lH-indol-3-yl)carbonyl)-
3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure I as described above, the acylation of 3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one prepared as described in example 16 above, with
commercially available 1-[2-(3,4-dimethoxy-phenyl)-ethyl]-5-methoxy-2-methyl-l \( H \)-indole-3-carboxylic acid, gave the title compound.

ES-MS m/e (%): 555.3 (M+H+).

Example 124

5 r-\{l-(4-Ethoxyphenyl)-5-methoxy-2-methyl-lH-indol-3-yl]carbonyl\}-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one

Following the general procedure I as described above, the acylation of 3H-spiro[2-benzofuran-l,4'-piperidin]-3-one prepared as described in example 16, with commercially available 1-(4-ethoxyphenyl)-5-methoxy-2-methyl-l \( H \)-indole-3-carboxylic acid, gave the title compound.

ES-MS m/e (%): 511.5 (M+H+).

Example 125

5-Bromo-l'-\{(6-chloro-lH-indol-3-yl)carbonyl\}-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one
Following the general procedure I as described above, the acylation of 5-bromo-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one with lH-indole-3-carboxylic acid (commercially available), gave the title compound.

ES-MS m/e (%): 459.3(M+H+).

Example 126

l'-(l-Benzyl-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure I as described above, the acylation of 3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in example 16 above with 1-benzyl-2-methyl-lH-indole-3-carboxylic acid (preparation described in example 1), gave the title compound.

ES-MS m/e (%): 451.6(M+H+).
Following the general procedure VI as described above, the arylation of 1'-(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 16) with commercially available 3,5-difluorophenylboronic acid gave the title compound.

ES-MS m/e (%): 493.1(M+H+).

Example 128

Following the general procedure VII as described above, the acylation of 1'-(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared
according to example 16 above) with commercially available 3-fluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 503.4(M+H+).

Example 129

\[ \text{l'-[6-Chloro-l-(2-fluorobenzoyl)-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one} \]

Following the general procedure VII as described above, the acylation of \( \text{l'-(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one} \) (prepared according to example 16) with commercially available 2-fluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 503.4(M+H+).

Example 130

\[ \text{r-[6-Chloro-l-(3,5-difluorobenzoyl)-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one} \]
Following the general procedure VII as described above, the acylation of l'-[(6-chloro-
IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 16) with commercially available 3,5-difluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 521.4(M+H+).

**Example 131**

r-{{6-Chloro-l-(2,3-difluorobenzoyl)-IH-indol-3-yl}carbonyl]-3H-spiro[2-benzofuran-
1,4'-piperidin]-3-one

Following the general procedure VII as described above, the acylation of l'-[(6-chloro-
IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 16) with commercially available 2,3-difluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 521.4(M+H+).

**Example 132**

r-{{6-Chloro-l-[(3,5-difluorophenyl)sulfonyl]-IH-indol-3-yl}carbonyl]-3H-spiro[2-
benzofuran-1,4'-piperidin]-3-one
Following the general procedure VII as described above, the sulphonylation of 1′-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4′-piperidin]-3-one (prepared according to example 16 above) with commercially available 3,5-difluorobenzenesulfonyl chloride gave the title compound.

ES-MS m/e (%): 557.4(M+H+).

Example 133

r-[(6-Chloro-1-(3,5-difluorobenzyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4′-piperidin]-3-one

Following the general procedure III as described above, the acylation of 1′-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4′-piperidin]-3-one (prepared according to example 16 above) with commercially available 3,5-difluorobenzyl chloride gave the title compound.

ES-MS m/e (%): 507.4(M+H+).

Example 134
Following the general procedure III as described above, the acylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 16) with commercially available 3-fluorobenzyl chloride gave the title compound.

ES-MS m/e (%): 489.4(M+H+).

Example 135

Following the general procedure III as described above, the acylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 16) with commercially available 2-chloro-l-(3-fluoro-phenyl)ethanone gave the title compound.
Example 136

r-((6-Chloro-1-[2-(2,5-difluorophenyl)-2-oxoethyl]-1H-indol-3-yl)carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the acylation of 1'-(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 16) with commercially available 2-chloro-1-(2,5-difluoro-phenyl)ethanoneethanone gave the title compound.

Example 137

1'-(6-Chloro-1-(3-fluorobenzoyl)-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one
Following the general procedure VII as described above, the acylation of l’-[(6-chloro-
1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one
(prepared according to example 19) with commercially available 3-fluorobenzoyl
chloride gave the title compound.

ES-MS m/e (%): 521.4(M+H+).

Example 138

\[ r\{-[6-Chloro-l-(2-fluorobenzoyl)-lH-indol-3-yl]carbonyl\}-5-fluoro-3H-spiro[2-
benzofuran-1,4’-piperidin]-3-one \]

Following the general procedure VII as described above, the acylation of l’-[(6-chloro-
1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one
(prepared according to example 19) with commercially available 2-fluorobenzoyl
chloride gave the title compound.

ES-MS m/e (%): 521.4(M+H+).

Example 139

\[ r\{-[6-Chloro-l-(2,3-difluorobenzoyl)-lH-indol-3-yl]carbonyl\}-5-fluoro-3H-spiro[2-
benzofuran-1,4’-piperidin]-3-one \]
Following the general procedure VII as described above, the acylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 19) with commercially available 3,5-difluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 539.4(M+H+).

Example 140

r’-[(6-Chloro-1-(2,3-difluorobenzoyl)-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

Following the general procedure VII as described above, the acylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 19) with commercially available 2,3-difluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 539.3(M+H+).
Example 141

\[ r-(\{6\text{-Chloro-1-[(3,5-difluorophenyl)sulfonyl]-1H-indol-3-yl} \text{ carbonyl}\}-5\text{-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]}-3\text{-one} \]

Following the general procedure VII as described above, the sulphonylation of \( l'-(6\text{-chloro-1H-indol-3-yl} \text{ carbonyl})-5\text{-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]}-3\text{-one} \) (prepared according to example 19) with commercially available 3,5-difluorobenzensulfonyl chloride gave the title compound.

ES-MS m/e (%): 575.3(M+H+).

Example 142

\[ r\{6\text{-Chloro-1-(3,5-difluorobenzyl)-1H-indol-3-yl} \text{ carbonyl}\}-5\text{-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]}-3\text{-one} \]
Following the general procedure III as described above, the alkylation of l’-[(6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with commercially available 3,5-difluorobenzyl chloride gave the title compound.

**Example 143**

l’-[(6-Chloro-l-(3-fluorobenzyl)-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

ES-MS m/e (%): 525.4(M+H+).

Following the general procedure III as described above, the alkylation of l’-[(6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with commercially available 3-fluorobenzyl chloride gave the title compound.

ES-MS m/e (%): 507.4(M+H+).

**Example 144**

l’-[(6-Chloro-l-[2-(3-fluorophenyl)-2-oxoethyl]-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one
Following the general procedure III as described above, the alkylation of 1'-(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one (prepared according to example 19) with commercially available 2-chloro-l-(3-fluorophenyl)-ethanone gave the title compound.

ES-MS m/e (%): 535.4 (M+H^+).

Example 145

5-Bromo-l'-(6-chloro-l-(3-fluorobenzoyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one

Following the general procedure VII as described above, the acylation of 5-bromo-l'-(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one with commercially available 3-fluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 581.2(M+H^+).
Example 146

5-Bromo-l’-([6-chloro-l-(2-fluorobenzoyl)-lH-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

Following the general procedure VII as described above, the acylation of 5-bromo-l’-([6-chloro-lH-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one with commercially available 3-fluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 581.2(M+H+).

Example 147

5-Bromo-l’-([6-chloro-l-(3,5-difluorobenzoyl)-lH-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

Following the general procedure VII as described above, the acylation of 5-bromo-l’-([6-chloro-lH-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one with commercially available 3,5-difluorobenzoyl chloride gave the title compound.
ES-MS m/e (%): 599.2(M+H+).

Example 148

5-Bromo-l'-{(6-chloro-l-(2,3-difluorobenzoyl)-lH-indol-3-yl)carbonyl}-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one

Following the general procedure VII as described above, the acylation of 5-bromo-l'-(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one with commercially available 2,3-difluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 599.2(M+H+)

Example 149

S-Bromo-l'-do-chloro-l-fCS^-difluoropheny^sulfonyy-lH-indol-S-ylJcarbony^-SH-spiro[2-benzofuran-l,4'-piperidin]-3-one
Following the general procedure VII as described above, the sulphonylation of 5-bromo-1-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one with commercially available 3,5-difluorobenzenesulfonyl chloride gave the title compound.

5 ES-MS m/e (%): 635.2(M+H+).

Example 150

5-Bromo-1'-%[6-chloro-1-(3,5-difluorobenzyl)-1H-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of 5-bromo-1-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one with commercially available 3,5-difluorobenzyl chloride gave the title compound.

ES-MS m/e (%): 585.2(M+H+).

Example 151

5-Bromo-1-[(6-chloro-1-(3-fluorobenzyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one
Following the general procedure III as described above, the alkylation of 5-bromo-1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one with commercially available 3-fluorobenzyl chloride gave the title compound.

Example 152

5-Bromo-1’-([6-chloro-1-[2-(3-fluorophenyl)-2-oxoethyl]-1H-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

Following the general procedure III as described above, the alkylation of 5-bromo-1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one with commercially available 2-chloro-1-(3-fluoro-phenyl)ethanone gave the title compound.

ES-MS m/e (%): 595.3(M+H+).

Example 153
1'-((6-Chloro-1-[2-(2-fluorophenyl)-2-oxoethyl]-1H-indol-3-yl)carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of 3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in example 16 with commercially available 2-chloro-1-(3-fluoro-phenyl)-ethanone gave the title compound.

ES-MS m/e (%): 517.4(M+H+).

Example 154

r-((6-Chloro-1-[2-(3,4-difluorophenyl)-2-oxoethyl]-1H-indol-3-yl)carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of 3H-spiro[2-benzofuran-1,4'-piperidin]-3-one prepared as described in example 16 above with commercially available 2-chloro-1-(3, 4-difluoro-phenyl)-ethanone gave the title compound.
ES-MS m/e (%): 535.4(M+H+).

Example 155

l’-[(6-Chloro-l-(3-fluorophenyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

Following the general procedure VI as described above, the arylation of l’-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 16 above) with commercially available 3-fluorophenylboronic acid gave the title compound.

ES-MS m/e (%): 475.0(M+H+).

Example 156

l’-[l-Biphenyl-3-yl-6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one
Following the general procedure VI as described above, the arylation of 1’-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 16) with commercially available 3-biphenylboronic acid gave the title compound.

ES-MS m/e (%): 533.0(M+H+).

Example 157

1’-[(1-Biphenyl-2-yl-6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

Following the general procedure VI as described above, the arylation of 1’-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 16) with commercially available 2-biphenylboronic acid gave the title compound.

ES-MS m/e (%): 533.0(M+H+).
Example 158

l'-(l-(Biphenyl-3-ylcarbonyl)-6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure VII as described above, the acylation of l'-(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 16) with commercially available biphenyl-3-carbonyl chloride gave the title compound.

ES-MS m/e (%): 561.4(M+H+).

Example 159

r-(6-Chloro-1-pyridin-2-yl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

To a solution of r-(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 16 above) in dry DMF was added NaH
(1 eq) and the reaction mixture stirred at room temperature for 30 min and then treated with 2-fluoropyridine (1.5 eq) and heated at 140 °C under microwave irradiation for 15 min. Purification by preparative HPLC gave the desired product in 38% yield.

ES-MS m/e (%): 458.1(M+H+).

Example 160

t-{[6-Chloro-l-pyridin-2-yl-lH-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

To a solution of 1'-{[6-chloro-lH-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) in dry DMF was added NaH (1 eq) and the reaction mixture stirred at room temperature for 30 min and then treated with 2-fluoropyridine (1.5 eq) and heated at 140 °C under microwave irradiation for 15 min. Purification by preparative HPLC gave the desired product in 36% yield.

ES-MS m/e (%): 476.0(M+H+).

Example 161

r-{[6-Chloro-l-[(2-methylpyridin-4-yl)methyl]-lH-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one
Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with 4-chloromethyl-2-methyl-pyridine (described in WO 2006023707) gave the title compound.

ES-MS m/e (%): 504.2(M+H+).

Example 162

r-[l-[[4-Amino-2-(methoxymethyl)pyrimidin-5-yl]methyl]-6-chloro-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with 5-chloromethyl-2-methoxy-pyrimidin-4-ylamine gave the title compound.
Following the general procedure III as described above, the alkylation of 1'-(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19 above) with 5-chloromethyl-2-methyl-pyrimidin-4-ylamine (described in Pharmaceutical Chemistry Journal (Translation of Khimiko-Farmatsevticheskii Zhurnal) (1999), 33(2), 101-103) gave the title compound.

ES-MS m/e (%): 520.2(M+H+).
Following the general procedure III as described above, the alkylation of l’-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with commercially available 2-chloro-5-chloromethylpyridine gave the title compound.

ES-MS m/e (%): 524.1(M+H+).

Example 165

r-((6-Chloro-l-[(3-chloro-6-methylpyridazin-4-yl)methyl]-1H-indol-3-yl)carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of l’-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19 above) with 3-chloro-4-chloromethyl-6-methylpyridazine (prepared by US 3453277) gave the title compound.
ES-MS m/e (%): 539.1(M+H+).

Example 166

r-[{6-Chloro-l-(pyridin-4-ylmethyl)-lH-indol-3-yl}carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of l’-{(6-chloro-lH-indol-3-yl)carbonyl}-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with commercially available 4-bromomethylpyridine gave the title compound.

1H-NMR (300MHz, CDC13): δ 1.75 (m, 2H), 2.26-2.39 (m, 2H), 3.3 (m, 2H), 4.45 (m, 2H), 5.6 (s, 2H), 7.15 (m, 3H), 7.65 (m, 3H), 7.8 (d,lH), 7.92 (m,lH), 8.03 (s,lH), 8.52(d,2H)

Example 167

r-[{6-Chloro-l-(2-pyridin-4-ylethyl)-lH-indol-3-yl}carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one
Following the general procedure III as described above, the alkylation of 1'-[(6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with commercially available 4-(2-bromo-ethyl)-pyridine gave the title compound.

ES-MS m/e (%): 504.2(M+H+).

Example 168

r-[(6-Chloro-l-(pyridin-4-ylmethyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of 1'-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 16) with commercially available 4-bromomethyl-pyridine gave the title compound.

ES-MS m/e (%): 472.3(M+H+).
Example 169

r-\{[6-Chloro-l-(2-oxo-2-pyridin-2-yethyl)-lH-indol-3-yl]carbonyl\}-5-fluoro-3H-
spiro[2-benzofuran-l,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of l'-\{[6-chloro-
H-indol-3-yl]carbonyl\}-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one
(prepared according to example 19 above) with commercially available 2-bromo-l-
pyridin-2-yethylone gave the title compound.

ES-MS m/e (%): 518.4(M+H+).

Example 170

l'-(6-Chloro-l-[2-(5-methyl-2-phenyl-1,3-oxazol-4-yl)-2-oxoethyl]-lH-indol-3-
yl]carbonyl)-5-fluoro-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one
Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 19) with 2-bromo-1-(5-methyl-2-phenyl-oxazol-4-yl)-ethanone (described in Journal of Medicinal Chemistry (1992), 35(14), 2617-26) gave the title compound.

ES-MS m/e (%): 598.4(M+H+).

Example 171

r-[[6-Chloro-1-(pyridin-2-ylmethyl)-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

a) Methanesulfonic acid pyridin-3-ylmethyl ester

To a mixture of 2-(hydroxymethyl)pyridine, DMAP and NEt3 was slowly added MsCl at 0 °C and the reaction mixture was stirred at room temperature for 1h. The reaction mixture was extracted with water and dichloromethane. The organic phase was dried on Na2SO4, filtered and the solvent was evaporated. Silica gel column chromatography (dichloromethane / MeOH 99:1) gave the title compound in 52% yield.

ES-MS m/e (%): 188.1 (M+H+).

b) r-[[6-chloro-1-(pyridin-2-ylmethyl)-1H-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one
(prepared according to example 19) with methanesulfonic acid pyridin-2-ylmethyl ester (described in WO 9955318) gave the desired product in 29% yield;

ES-MS m/e (%): 490.0(M+H+).

Example 172

r-\{[6-Chloro-1-(pyridin-3-ylmethyl)-1H-indol-3-yl]carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of l'-\{[6-chloro-1H-indol-3-yl]carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 16) with commercially available 3-bromomethyl-pyridine gave the title compound.

ES-MS m/e (%): 472.1(M+H+).

Example 173

l'-\{[6-Chloro-1-(pyrazin-2-ylmethyl)-1H-indol-3-yl]carbonyl\}-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one
Following the general procedure III as described above, the alkylation of l’-{(6-chloro-lH-indol-3-yl)carbonyl}-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 19) with methanesulfonic acid pyrazin-2-ylmethyl ester (preparation described in WO 2002064574) gave the title compound.

ES-MS m/e (%): 491.0(M+H+).

Example 174

r-{[6-Chloro-l-(pyrimidin-5-ylmethyl)-lH-indol-3-yl]carbonyl}-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

a) Methanesulfonic acid pyrimidin-5-ylmethyl ester

To a mixture of pyrimidin-5-yl-methanol, DMAP and NEt₃ was slowly added MsCl at 0 °C and the reaction mixture was stirred at room temperature for 3h. The reaction mixture was extracted with water and dichloromethane. The organic phase was dried on Na₂SO₄, filtered and the solvent was evaporated. Silica gel column chromatography (Ethyl acetate/hexane 1:1) gave the title compound in 40% yield.

b) l’-ire-Chloro-l^pyrimidin-5-ylmethyD-lH-indol^-ylcarbonyll-S-fluoro^H- spiro2-benzofuran-l,4’-piperidinl-3 -one
Following the general procedure III as described above, the alkylation of \( l'-[(6\text{-chloro}-1H\text{-indol-3-yl})\text{carbonyl}]\text{-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]}\text{-3-one} \) (prepared according to example 19) with methanesulfonic acid pyrimidin-5-ylmethyl ester gave the title compound.

ES-MS m/e (%): 491.0(M+H+).

Example 175

3-\{6-Chloro-3-[(5-fluoro-3-oxo-rH,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl]-1H-indol-1-yl\}propanenitrile

Following the general procedure III as described above, the alkylation of \( l'-[(6\text{-chloro}-1H\text{-indol-3-yl})\text{carbonyl}]\text{-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]}\text{-3-one} \) (prepared according to example 19) with commercially available 3-bromo-propionitrile gave the title compound.
ES-MS m/e (%): 452.0 (M+H+).

Example 176

1’-[(7-Chloro-1-oxo-1,2,3,4-tetrahydropyrazino[1,2-a]indol-10-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

a) 6-Chloro-l H-indole-2,3-dicarboxylic acid 2-ethyl ester

To a solution of commercially available 6-chloro-3-formyl-1H-indole-2-carboxylic acid ethyl ester in t-BuOH and 2-methyl-2-butene (50 eq) was added 9.2 eq NaClO₂ (9 eq) and aq. NaH₂PO₄ (7 eq) and stirred 12h at room temperature. After concentration the mixture was dissolved in H₂O and adjusted to pH7 and then extracted with EtOAc; partial concentration of the organic phase precipitated the product in 52% yield.

b) Ethyl 6-chloro-3-r(5-fluoro-3-oxo-l H,3H-spiror2-benzofuran-l,4'-piperidinl-r- vDcarbonyll -lH-indole-2-carbox ylate

Following the general procedure I as described above, the acylation of 1’-[(6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with 6-chloro-l H-indole-2,3-dicarboxylic acid 2-ethyl ester gave the title compound.

ES-MS m/e (%): 471.0 (M+H+).
c) 6-Chloro-3\{(5-fluoro-3-oxo-1H,3H-spiro\[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl\}-1H-indole-2-carboxylic acid

A solution of the ethyl 6-chloro-3\{(5-fluoro-3-oxo-1H,3H-spiro\[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl\}-1H-indole-2-carboxylate in EtOH was treated with 2 eq. of aq. IN LiOH at 80 °C for 12h, then concentrated and dissolved in IN HCl and extracted with dichloromethane. Evaporation of the solvent gave the title compound in 72% yield.

d) 6-Chloro-N-f2-chloroethyl)-3-rf5-fluoro-3-oxo-lH,3H-spiro\[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl\}-1H-indole-2-carboxamide

To a stirred solution of 6-chloro-3\{(5-fluoro-3-oxo-1H,3H-spiro\[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl\}-1H-indole-2-carboxylic acid in dichloromethane were added EDC, HOBt, NEt₃ and 2-chloroethylamine hydrochloride. The reaction mixture was stirred at room temperature for 2 days. The reaction mixture was poured onto water and extracted with dichloromethane. The combined organic phases were dried over Na₂SO₄ and the solvent was evaporated. Column chromatography (ethyl acetate/hexane 1:1) gave the desired compound in 59% yield.

MS m/e (\%) : 504.0 (M+H⁺).
e) l'-[(7-Chloro-1-oxo-1,2,3,4-tetrahydropyrazino[1,2-alindol-10-vDcarbonyl]-5-fluoro-
3H-spiro[2-benzofuran-1,4'-piperidin]-1-3-one

A solution of the 6-chloro-N-(2-chloroethyl)-3-[(5-fluoro-3-oxo-l'H,3H-spiro[2-
5 benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-lH-indole-2-carboxamide in dry DMF was
treated with NaH (1 eq) and stirred for 1h at room temperature, then treated with a
further portion of NaH (1 eq) and stirred for 2h at room temperature. Dilution with
water and extraction with EtOAc followed by washing with aq. NH₄Cl gave the crude
product which was purified by silica gel chromatography (dichloromethane/MeOH 98:2)
to give 42% yield of the desired product.

ES-MS m/e (%): 468.3 (M+H⁺).

Example 177

l'-{(1-[(4-Benzylmorpholin-2-yl)methyl]-6-chloro-lH-indol-3-yl)carbonyl}-5-fluoro-
3H-spiro[2-benzofuran-1,4'-piperidin]-3-one
Following the general procedure III as described above, the alkylation of l’-[6-chloro-lH-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 19 above) with commercially available 4-benzyl-2-(chloromethyl)morpholine gave the title compound.

ES-MS m/e (%): 588.2(M+H+).

Example 178

l’-(6-Chloro-1-[l,4-dibenzylpiperazin-2-yl]methyl]-lH-indol-3-yl]carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

Following the general procedure III as described above, the alkylation of l’-[6-chloro-lH-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 19) with 1,4-dibenzyl-2-chloromethyl-piperazine (described in Journal of Medicinal Chemistry 1999, 42(9), 1587-1603) gave the title compound.

ES-MS m/e (%): 677.3(M+H+).

Example 179

l’-(6-Chloro-1-[5-methylisoxazol-3-yl]methyl]-lH-indol-3-yl]carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one
Following the general procedure III as described above, the alkylation of l'-(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with methanesulfonic acid 5-methyl-isoxazol-3-ylmethyl ester (described in WO 2004092172) gave the title compound.

ES-MS m/e (%): 494.1(M+H+).

Example 180

r-[(6-Chloro-l-(pyridin-2-ylmethyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of l'-(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 16 above) with methanesulfonic acid pyridin-2-ylmethyl ester (described in WO 9955318) gave the title compound.

ES-MS m/e (%): 472.1(M+H+).
Example 181

6-Chloro-3-[(5-fluoro-3-oxo-1'H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-N-methyl-1H-indole-2-carboxamide

A suspension of the 6-chloro-3-[(5-fluoro-3-oxo-1'H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-1H-indole-2-carboxylic acid in dichloromethane was treated with EDC (1.2 eq), HOBt (1.2 eq) and Et$_3$N (1.2 eq) and the solution stirred at room temperature for 15 min. Methylamine (1 eq) was then added and the reaction mixture stirred at room temperature for 16h. Purification by preparative HPLC yielded the desired product.

ES-MS m/e (%): 456.1(M+H$^+$).

Example 182

l'-{(6-Chloro-1-[(5-cyclopropyl-2-methyl-1,3-oxazol-4-yl)methyl]-1H-indol-3-yl)carbonyl}-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

a) 2-Acetylamino-2-cyclopropanecarbonyl-malonic acid diethyl ester

To a solution of 6 eq KOt-Bu in THF at room temperature was added 6 eq 2-acetylamino-malonic acid diethyl ester and after 5 min 1 eq cyclopropanecarbonylchloride. After 15min, the mixture was concentrated and partitioned between
EtOAc and H₂O. The organic layer was concentrated to give the desired product in 60% yield.

ES-MS m/e (%): 286.2 (M+H+).

b) 5-Cyclopropyl-2-methyl-oxazole-4-carboxylic acid ethyl ester

\[ \text{5-Cyclopropyl-2-methyl-oxazole-4-carboxylic acid ethyl ester} \]

2-Acetylamino-2-cyclopropanecarbonyl-malonic acid diethyl ester in DMSO was treated with 2 eq H₂O and stirred at room temperature for 6h then extracted with Et₂O to give after concentration the title compound in 34% yield.

c) (5-Cyclopropyl-2-methyl-oxazol-4-yl)-methanol

\[ \text{(5-Cyclopropyl-2-methyl-oxazol-4-yl)-methanol} \]

A solution of 5-cyclopropyl-2-methyl-oxazole-4-carboxylic acid ethyl ester in Et₂O at 0°C was treated sequentially with (2 x 2.3 eq) LiBH₄ and stirred for 1h. 10 eq of MeOH were then added and the solution stirred for 4h at room temperature. Sequential addition of aq NH₄Cl (H₂ evolution) and then Na₂CO₃/NaCl gave a mixture which was extracted with EtOAc to give after concentration the title compound in 16% yield.

ES-MS m/e (%): 154.1 (M+H+).

d) r-f(6-Chloro-1-ry-5-cyclopropyl-2-methyl-1,3-oxazol-4-yl)methyl-IH-indol-3-yl[carbonyl]-5-fluoro-3H-spiro2-benzofuran-1,4'-piperidin1-3-one
Following the general procedure III as described above, the alkylation of l'-(6-chloro-1H-indol-3-yl)carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with methanesulfonic acid 5-cyclopropyl-2-methyl-oxazol-4-ylmethyl ester (prepared by mesylation of (5-cyclopropyl-2-methyl-oxazol-4-yl)-methanol) gave the title compound.

ES-MS m/e (%): 534.2(M+H+).

Example 183

r-([6-Chloro-1-[(l-methyl-lH-imidazol-5-yl)methyl]-lH-indol-3-yl]carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of l'-(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with methanesulfonic acid 3-methyl-3H-imidazol-4-
ylmethyl ester (prepared by mesylation of the commercially available (3-methyl-3H-imidazo1-4-yl)-methanol) gave the title compound.

ES-MS m/e (%): 493.1(M+H+).

Example 184

r-(6-Chloro-l-(3-methylisoxazol-5-yl)methyl]-lH-indol-3-yl)carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of l’-[(6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with methanesulfonic acid 3-methyl-isoxazol-5-ylmethyl ester (described in Heterocycles, 23(3), 571-83; 1985) gave the title compound.

ES-MS m/e (%): 494.1(M+H+).

Example 185

r-(6-Chloro-l-[1,5-dimethyl-lH-pyrazol-3-yl)methyl]-lH-indol-3-yl)carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one
Following the general procedure III as described above, the alkylation of 1'-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with methanesulfonic acid 1,5-dimethyl-1H-pyrazol-3-ylmethyl ester (prepared by mesylation of commercially available methanesulfonic acid 1,5-dimethyl-1H-pyrazol-3-ylmethyl ester) gave the title compound.

**ES-MS m/e (%):** 507.2(M+H+).

**Example 186**

r-{(6-Chloro-1-[(3,5-dimethylisoxazol-4-yl)methyl]-1H-indol-3-yl)carbonyl}-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of 1'-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19 above) with methanesulfonic acid 3,5-dimethyl-
isoazol-4-ylmethyl ester (prepared by mesylation of the commercially available 1,5-dimethyl-\textit{H}-pyrazol-3-yl)-methanol) gave the title compound.

ES-MS m/e (%): 508.2(M+H+).

Example 187

1'-(6-Chloro-1-[(2,5-dimethyl-1,3-oxazol-4-yl)methyl]-\textit{H}-indol-3-yl)carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of 1'-(6-chloro-\textit{H}-indol-3-yl)carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19 above) with methanesulfonic acid 2,5-dimethyl-oxazol-4-ylmethyl ester (prepared by mesylation of (2,5-dimethyl-oxazol-4-yl)-methanol, described in Organic Letters (1999), 1(1), 87-90) gave the title compound.

ES-MS m/e (%): 508.1(M+H+).

Example 188

1'-(6-Chloro-1-[(3-fluorooxetan-3-yl)methyl]-\textit{H}-indol-3-yl)carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one
Following the general procedure III as described above, the alkylation of l'-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with 3-bromomethyl-3-fluoro-oxetane (described in US2005215599) gave the title compound.

ES-MS m/e (%): 487.1(M+H+).

Example 189

l'-[(6-Chloro-1-[(3-fluorooxetan-3-yl)methyl]-1H-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of l'-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 16 above) with 3-bromomethyl-3-fluoro-oxetane (described in US2005215599) gave the title compound.
Example 190

r-[(6-Chloro-l-[(l-(methoxymethyl)cyclopropyl)methyl]-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of l'-[(6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with 1-bromomethyl-l-methoxymethyl-cyclopropane (described in WO 2001032633) gave the title compound.

Example 191

[1-((6-Chloro-3-[(5-fluoro-3-oxo-lH,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl] -lH-indol- 1-yl )methyl)cyclopropyl]acetonitrile
Following the general procedure III as described above, the alkylation of l'-[(6-chloro-
IH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one
(prepared according to example 19) with 1-bromomethyl-cyclopropanecarbonitrile
(described in EP 148004) gave the title compound.

ES-MS m/e (%): 492.5(M+H+).

Example 192

l'-[(6-Chloro-l-{[l-(methoxymethyl)cyclopropyl]methyl}-lH-indol-3-yl)carbonyl]-3H-
spirow[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure III as described above, the alkylation of l'-[(6-chloro-
IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared
according to example 16 above) with 1-bromomethyl-l-methoxymethyl-cyclopropane
(prepared as described in WO 2001032633) gave the title compound.

ES-MS m/e (%): 479.5(M+H+).

Example 193

{l-[(6-Chloro-3-{[3-oxo-rH,3H-spiro[2-benzofuran-1,4'-piperidin]-l'-yl]carbonyl]-
IH-indol-1-ylJmethyJcyclopropyl]acetonitrile

[1-[(6-Chloro-3-[3-oxo-rH,3H-spiro[2-benzofuran-1,4'-piperidin]-l'-yl]carbonyl]-
IH-indol-1-ylJmethyJcyclopropyl]acetonitrile
Following the general procedure III as described above, the alkylation of \( \text{l'-(6-chloro-1H-indol-3-yl)carbonyl-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one} \) (prepared according to example 16) with 1-bromomethyl-l-methoxymethyl-cyclopropane gave the title compound.

ES-MS m/e (%): 474.5(M+H+).

Example 194

6-Chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-N-[2-(methylamino)ethyl]-1H-indole-2-carboxamide hydrochloride

A solution of tert-butyl \( \{2-[(6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-1H-indol-2-yl]carbonyl)-amino\} \)ethylJmethylcarbamate was treated with 4 eq of HCl in dioxane and stirred at room temperature for 3h then
treated with a further portion of 4 eq of HCl in dioxane. The solution was stirred at room temperature for 16h and evaporated to give the product in 58% yield.

ES-MS m/e (%): 511.2(M+H+).

Example 195

r-d 6-Chloro-l-Cl-Ctetrahydro-lH-pyran- 4-y^ethyy-lH-indol-S-ylcarbony^S-fluoro-3H-spiro[2-benzo furan- 1,4'-piperidin]-3-one hydrochloride

Following the general procedure III as described above, the alkylation of l'-[6-chloro-lH-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzo furan-1,4'-piperidin]-3-one (prepared according to example 19) with 4-(2-bromo-ethyl)-tetrahydro-pyran (described in US 2004220214) gave the title compound.

ES-MS m/e (%): 511.2 (M+H+).

Example 196

r-(6-Chloro-l-[2-(tetrahydro-2H-pyran-4-yl)ethyl]-lH-indol-3-yl]carbonyl)-3H-spiro[2-benzo furan-1,4'-piperidin]-3-one
Following the general procedure III as described above, the alkylation of l’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 16) with 4-(2-bromo-ethyl)-tetrahydro-pyran (described in US 2004220214) gave the title compound.

ES-MS m/e (%): 493.2 (M+H+).

Example 197

tert-Butyl 2-[(6-chloro-3-[(3-oxo-rH,3H-spiro[2-benzofuran-1,4’-piperidin]-l’-yl)carbonyl] -1H-indol- 1-yl]methyl)morpholine-4-carboxylate

Following the general procedure III as described above, the alkylation of l’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 16) with 2-chloromethyl-morpholine-4-carboxylic acid tert-butyl ester (described in WO 2006020415) gave the title compound.
Example 198

1'-([6-Chloro-l-(morpholin-2-ylmethyl)-lH-indol-3-yl]carbonyl)-3H-spiro[2-benzo furan-1,4'-piperidin]-3-one hydrochloride

Example 199

tert-Butyl 2-((6-chloro-3-[(3-oxo-l'H,3H-spiro[2-benzo furan-1,4'-piperidin]-1'-yl)carbonyl]-lH-indol-1-yl)methyl)morpholine-4-carboxylate was dissolved in a solution of HCl (5 eq) in dioxane and stirred at room temperature for 5h. A further portion of HCl (5 eq) in dioxane was added and the solution stirred for an additional 5h at room temperature. Evaporation gave the desired product in 96% yield. ES-MS m/e (%): 480.2 (M+H+).
Following the general procedure III as described above, the alkylation of l'-(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 19) with 2-chloromethyl-morpholine-4-carboxylic acid tert-butyl ester (described in WO 2006020415) gave the title compound:

ES-MS m/e (%): 598.2(M+H+).

Example 200

l'-(6-Chloro-l-(morpholin-2-ylmethyl)-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one dihydrochloride

tert-Butyl 2-((6-chloro-3-((5-fluoro-3-oxo-1 H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl]-1H-indol-l-yl)methyl)morpholine-4-carboxylate was dissolved in a solution of HCl (5 eq) in dioxane and stirred at room temperature for 5h. A further portion of HCl (5 eq) in dioxane was added and the solution stirred for an additional 5h
at room temperature. Evaporation gave the desired product in quantitative yield. ES-MS m/e (%): 498.1 (M+H+).

Example 201

l’-[[l-(3,5-Difluorobenzyl)-lH-indol-3-yl]carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one

Following the general procedure III as described above, the alkylation of l’-[(6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one (prepared according to example 19) with commercially available 3,5-difluorobenzyl chloride gave the title compound.

ES-MS m/e (%): 491.5(M+H+).

Example 202

r’-[[l-(3,5-Difluorobenzyl)-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one
Following the general procedure III as described above, the alkylation of 1’-{(6-chloro-1H-indol-3-yl)carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 16) with commercially available 3,5-difluorobenzyl chloride gave the title compound.

ES-MS m/e (%): 473.5(M+H+).

Example 203

N,N-Diethyl-2-{3-[(3-oxo-1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1-yl)carbonyl]-1H-indol-1-yl}acetamide

Following the general procedure III as described above, the alkylation of 1’-{(6-chloro-1H-indol-3-yl)carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (prepared according to example 16) with commercially available 2-chloro-N,N-diethyl-acetamide gave the title compound.
ES-MS m/e (%): 460.6 (M+H+).

Examples of compounds of formula (I-d)

Example 204

r-[[6-Chloro-1-(2-oxo-2-piperidin-1-ylethyl)-1H-indol-3-yl] carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure V as described hereinabove, the alkylation of l'-[[6-chloro-1H-indol-3-yl] carbonyl] -3H-spiro[2-benzofuran-1,4'-piperidine] (the preparation of which have been described in example 69) with commercially available 2-chloro-l-piperidin-1-yl-ethanone as electrophile, the title compound was obtained as a white powder.

ES-MS m/e (%): 492.1 (M+H+).

Example 205

l'-[[6-Chloro-1-(2-morpholin-4-yl-2-oxoethyl)-1H-indol-3-yl] carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]
Following the general procedure V as described hereinabove, the alkylation of \( \text{L'[-[(6-chloro-lH-indol-3-yl)carbonyl] -3H-spiro[2-benzofuran-1,4'-piperidine]} \) (the preparation of which have been described in example 69) with commercially available 2-Chloro-1 morpholin-4-yl-ethanone as electrophile, the title compound was obtained as a white powder.

ES-MS m/e (%): 494.1 (M+H\(^+\)).

Example 206

2-\([6\text{-Chloro-3-}-(1\text{ H ,3H-spiro[2-benzofuran-1,4'-piperidin]-ylcarbonyl}-lH\text{-indol-1-yl}]-N,N\text{-dimethylacetamide}\)

Following the general procedure V as described hereinabove, the alkylation of \( \text{L'[-[(6-chloro-lH-indol-3-yl)carbonyl] -3H-spiro[2-benzofuran-1,4'-piperidine]} \) (the preparation of which have been described in example 69) with commercially available 2-chloro-N,N-dimethyl-acetamide as electrophile, the title compound was obtained as a white powder.

ES-MS m/e (%): 452.0 (M+H\(^+\)).

Example 207

2-\([6\text{-Chloro-3-}-(1\text{ H ,3H-spiro[2-benzofuran-1,4'-piperidin]-ylcarbonyl}-lH\text{-indol-1-yl}]-N,N\text{-diethylacetamide}\)
Following the general procedure V as described hereinabove, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (the preparation of which have been described in example 69) with commercially available 2-chloro-N,N-diethyl-acetamide as electrophile, the title compound was obtained as a white powder.

ES-MS m/e (%): 480.1 (M+H+).

Example 208

l’-[(6-Chloro-l-(piperidin-l-ylcarbonyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4’-piperidine]

Following the general procedure V as described hereinabove, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (the preparation of which have been described in example 69) with commercially available piperidine-1-carbonyl chloride as electrophile, the title compound was obtained as a white powder.

ES-MS m/e (%): 478.0 (M+H+).

Example 209
tert-Butyl [6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-ylcarbonyl)-1H-indol-1-yl] acetate

Following the general procedure V as described hereinabove, the alkylation of 1′-[6-chloro-1H-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (the preparation of which have been described in example 69) with commercially available chloro-acetic acid tert-butyl ester as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 481.3 (M+H+)

Example 210

2-[6-Chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1-yl)carbonyl]-1H-indol-1-yl]-N,N-dimethylethan amine

Following the general procedure V as described hereinabove, the alkylation of 1′-[6-chloro-1H-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (the preparation of which have been described in example 69) with commercially available (2-chloro-ethyl)-dimethyl-amine as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 438.4 (M+H+)
Following the general procedure V as described hereinabove, the alkylation of l'-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] with commercially available 4-(2-chloroacetyl)-piperazine-1-carboxylic acid tert-butyl ester as electrophile, the title compound was obtained as a white solid after removal of the Boc protecting group under standard conditions (TFA/dichloromethane, room temperature).

ES-MS m/e (%): 493.1 (M+H+).

Example 212

l'-[(6-Chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]
Following the general procedure as described hereinabove, the alkylation of \( l'-(6\)-chloro-1H-indol-3-yl)carbonyl \) -3H-spiro[2-benzofuran-1,4'-piperidine] (the preparation of which have been described in example 69) with commercially available 2-chloro-l-(4-methyl-piperazin-1-yl)-ethanone as electrophile, the title compound was obtained as a white solid.

ES-MS m/e (%): 507.4 (M+H\(^+\)).

Example 213

\[ 2-[6-Chloro-5-methyl-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidin]-r-y1carbonyl)-1H-indol-1-y1]-N,N-dimethy1acetamide \]

Amide coupling according to general procedure I described hereinabove:

- Amine: Spiro[isobenzofuran-1(3H),4’-piperidine] prepared as described in *J. Org. Chem.* 1976, 41, 2628,

- Acid: 6-Chloro-l-dimethylcarbamoylmethyl-S-methyl-IH-indole-S-carboxylic acid,

ES-MS m/e (%): 466.3 (M+H\(^+\)).

Example 214

\[ r-[6-Chloro-l-(3-fluorobenzoyl)-1H-indol-3-y1]carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine \]
Following the general procedure VII as described above, the acylation of \( l'\)-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with commercially available 3-fluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 489.4(M+H⁺).

Example 215

\[
\text{r-\{[6-Chloro-1-(2-fluorobenzoyl)-lH-indol-3-yl]carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine}
\]

Following the general procedure VII as described above, the acylation of \( l'\)-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with commercially available 2-fluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 489.4(M+H⁺).

Example 216
Following the general procedure VII as described above, the acylation of \( \text{I'}-[\text{6-chloro-lH-indol-3-yl} \text{carbonyl}]-3\text{H-spiro}[2\text{-benzofuran-1,4'-piperidine}] \) (prepared according to example 69) with commercially available 3,5-difluorobenzoyl chloride gave the title compound.

ES-MS m/e (%): 507.4(M+H+).

Example 217

Following the general procedure VII as described above, the acylation of \( \text{I'}-[(6\text{-chloro-lH-indol-3-yl})\text{carbonyl}]-3\text{H-spiro}[2\text{-benzofuran-1,4'-piperidine}] \) (prepared according to example 69) with commercially available 2,3-difluorobenzoyl chloride gave the title compound.
Example 218

r-\{(6-Chloro-l-[(3,5-difluorophenyl)sulfonyl]-lH-indol-3-yl}carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine

Following the general procedure VII as described above, the sulphonylation of \textit{l}'-[(5-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] \textit{\textit{(prepared according to example 69)}} with commercially available 3,5-difluorobenzenesulfonyl chloride gave the title compound.

ES-MS m/e (%): 543.3(M+H+).

Example 219

r-\{[6-Chloro-l-(3,5-difluorobenzyl)-lH-indol-3-yl]carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine
Following the general procedure III as described above, the alkylation of l'-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidine] (prepared according to example 69) with commercially available 3,5-difluorobenzyl chloride gave the title compound.

ES-MS m/e (%): 493.4(M+H+).

Example 220

r-[(6-Chloro-l-(3-fluorobenzyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidine]

Following the general procedure III as described above, the alkylation of l'-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidine] (prepared according to example 69) with commercially available 3-fluorobenzyl chloride gave the title compound.

ES-MS m/e (%): 475.4(M+H+).
Example 221

\[
l'-\{[1-(Biphenyl-3-ylcarbonyl)-6-chloro-1H-indol-3-yl]carbonyl\}-3H-spiro[2-benzofuran-1,4'-piperidine
\]

Following the general procedure VII as described above, the acylation of \( l'-[(6\text{-chloro-}1H\text{-indol-3-yl})\text{carbonyl]}\)-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with commercially available biphenyl-3-carbonyl chloride gave the title compound.

ES-MS m/e (%): 547.4(M+H+).

Example 222

\[
r-\{[6\text{-Chloro-}l-(3,5\text{-difluorophenyl})\text{-}1H\text{-indol-3-yl}]\text{carbonyl]}\}-3H\text{-spirol[2-benzofuran-1,4'-piperidine
}\]

Following the general procedure VI as described above, the arylation of \( l'-[(6\text{-chloro-}1H\text{-indol-3-yl})\text{carbonyl]}\)-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to
example 69) with commercially available 3,5-difluorophenylboronic acid gave the title compound.

ES-MS m/e (%): 478.9(M+H+).

Following the general procedure VI as described above, the arylation of 1'-(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with commercially available 3-fluorophenylboronic acid gave the title compound.

ES-MS m/e (%): 461.1(M+H+).

Example 224

2-[6-Chloro-3-(rH,3H-spiro[2-benzofuran-1,4'-piperidin]-l'-ylcarbonyl]-1H-indol-1-yl]-l-(2-fluorophenyl)ethanone
Following the general procedure III as described above, the alkylation of l'-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with commercially available 2-bromo-l-(2-fluoro-phenyl)-ethanone gave the title compound.

ES-MS m/e (%): 503.4(M+H+).

Example 225

l'-[(l-Biphenyl-2-yl-6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure VI as described above, the arylation of l'-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with commercially available 2-biphenylboronic acid gave the title compound.

ES-MS m/e (%): 519.3(M+H+).
Example 226

1’-[(1-Biphenyl-2-yl-6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine

Following the general procedure VI as described above, the arylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (prepared according to example 69) with commercially available 3-biphenylboronic acid gave the title compound.

ES-MS m/e (%): 519.3(M+H+).

Example 227

r’-[(6-Chloro-1-pyridin-2-yl-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine

To a solution of r’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (prepared according to example 69) in dry DMF was added NaH (1 eq) and
the reaction mixture stirred at room temperature for 30 min and then treated with 2-fluoropyridine (1.5 eq) and heated at 140 °C under microwave irradiation for 15 min. Purification by preparative HPLC gave the desired product in 24% yield.

ES-MS m/e (%): 444.1(M+H+).

Example 228

r-[[6-Chloro-1-(pyridin-4-ylmethyl)-1H-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine

Following the general procedure III as described above, the alkylation of l'-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with commercially available 4-bromomethyl-pyridine gave the title compound.

ES-MS m/e (%): 458.4(M+H+).

Example 229

2-[6-Chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl]-1-pyridin-2-ylethanol
Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (prepared according to example 69 above) with commercially available 2-bromo-1-pyridin-2-ylenone gave the title compound.

ES-MS m/e (%): 486.4(M+H+).

Example 230

r-[(6-Chloro-l-(pyridin-3-ylmethyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine]

Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (prepared according to example 69) with commercially available 3-bromomethyl-pyridine gave the title compound.
ES-MS m/e (%): 458.4(M+H⁺).

Example 231

r-{{6-Chloro-l-(pyridin-2-ylmethyl)-lH-indol-3-yl}carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure III as described above, the alkylation of l'{{6-chloro-lH-indol-3-yl}carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with methanesulfonic acid pyridin-2-ylmethyl ester (described in WO 9955318) gave the title compound.

ES-MS m/e (%): 458.3(M+H⁺).

Example 232

r-{{l-[(4-Benzylmorpholin-2-yl)methyl]-6-chloro-lH-indol-3-yl}carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidine]
Following the general procedure III as described above, the alkylation of l’-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (prepared according to example 69) with commercially available 4-benzyl-2-(chloromethyl)morpholine gave the title compound.

ES-MS m/e (%): 556.3(M+H+).

Example 233

l’-[(6-Chloro-l-[(1,4-dibenzylpiperazin-2-yl)methyl]-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine]

Following the general procedure III as described above, the alkylation of l’-[(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (prepared according to
Example 69 above) with 1,4-dibenzyl-2-chloromethyl-piperazine (described in Journal of Medicinal Chemistry (1999), 42(9), 1587-1603) gave the title compound.

ES-MS m/e (%): 645.3(M+H+).

Example 234

5 l'-(6-Chloro-1-(pyrazin-2-ylmethyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure III as described above, the alkylation of l'-(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69 above) with methanesulfonic acid pyrazin-2-ylmethyl ester (preparation described in WO 2002064574) gave the title compound.

ES-MS m/e (%): 459.3(M+H+).

Example 235

r-(6-Chloro-1-(pyrimidin-5-ylmethyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine
Following the general procedure III as described above, the alkylation of l'-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with methanesulfonic acid pyrimidin-5-ylmethyl ester (preparation described herein) gave the title compound.

ES-MS m/e (%): 459.3(M+H+).

Example 236

l'-{(6-Chloro-l-[(5-methylisoxazol-3-yl)methyl]-1H-indol-3-yl)carbonyl}-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure III as described above, the alkylation of l'-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with methanesulfonic acid 5-methyl-isoxazol-3-ylmethyl ester (described in WO 2004092172) gave the title compound.

ES-MS m/e (%): 462.2(M+H+).
Example 237

\[ \text{r-} \{ \text{6-Chloro-1-[} \{ \text{5-cyclopropyl-2-methyl-1,3-oxazol-4-yl} \} \text{methyl]-1H-indol-3-y1} \} \text{carbonyl}-3H\text{-} \text{spiro[2-benzofuran-1,4'-piperidine} \]

Following the general procedure III as described above, the alkylation of \( \text{l'-} \{ \text{(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine} \) (prepared according to example 69 above) with methanesulfonic acid 5-cyclopropyl-2-methyl-oxazol-4-ylmethyl ester (prepared herein) gave the title compound.

ES-MS m/e (%): 502.2(M+H+).

Example 238

\[ \text{r-} \{ \text{6-Chloro-1-[} \{ \text{l-methyl-1H-imidazol-5-yl} \} \text{methyl]-1H-indol-3-yl} \} \text{carbonyl}-3H\text{-} \text{spiro[2-benzofuran-1,4'-piperidine} \]

Following the general procedure III as described above, the alkylation of \( \text{l'-} \{ \text{(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine} \) (prepared according to
example 69) with methanesulfonic acid 3-methyl-3H-imidazol-4-ylmethyl ester (prepared by mesylation of the commercially available (3-methyl-3H-imidazol-4-yl)-methanol) gave the title compound.

ES-MS m/e (%): 461.2(M+H+).

Example 239

l'-(6-Chloro-1-[(3-methylisoxazol-5-yl)methyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure III as described above, the alkylation of l'-(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with methanesulfonic acid 3-methyl-isoxazol-5-ylmethyl ester (described in Heterocycles, 23(3), 571-83; 1985) gave the title compound.

ES-MS m/e (%): 462.2(M+H+).

Example 240

r-(6-Chloro-1-[(1,5-dimethyl-1H-pyrazol-3-yl)methyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine
Following the general procedure III as described above, the alkylation of l'-(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran,4'-piperidine] (prepared according to example 69) with methanesulfonic acid 1,5-dimethyl-1H-pyrazol-3-ylmethyl ester (prepared by mesylation of commercially available methanesulfonic acid 1,5-dimethyl-1H-pyrazol-3-ylmethyl ester) gave the title compound.

ES-MS m/e (%): 475.2(M+H+).

Example 241

l'-(6-Chloro-1-[(3,5-dimethylisoxazol-4-yl)methyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran,4'-piperidine

Following the general procedure III as described above, the alkylation of l'-(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran,4'-piperidine] (prepared according to example 69) with methanesulfonic acid 3,5-dimethyl-isoxazol-4-ylmethyl ester (prepared by mesylation of the commercially available 1,5-dimethyl-1H-pyrazol-3-yl)-methanol) gave the title compound.
Example 242

\[ r-(\text{6-Chloro-l-[(2,5-dimethyl-l,3-oxazol-4-yl)methyl]-lH-indol-3-yl} \text{carbonyl})-3H-spiro[2-benzofuran-l,4'-piperidine} \]

Following the general procedure III as described above, the alkylation of \( l'-(\text{6-chloro-lH-indol-3-yl} \text{carbonyl})-3H-spiro[2-benzofuran-l,4'-piperidine} \) (prepared according to example 69) with methanesulfonic acid 2,5-dimethyl-oxazol-4-ylmethyl ester (prepared by mesylation of 2,5-dimethyl-oxazol-4-yl)-methanol, described in Organic Letters 1999, 1(1), 87-90 gave the title compound.

ES-MS m/e (%): \( 476.2(M+H^+) \).

Example 243

\[ l'-(\text{6-Chloro-l-[(3-fluorooxetan-3-yl)methyl]-lH-indol-3-yl} \text{carbonyl})-3H-spiro[2-benzofuran-l,4'-piperidine} \]
Following the general procedure III as described above, the alkylation of \( l'-(6\text{-chloro-1H-indol-3-yl})\text{carbonyl}-3\text{H-spiro[2-benzofuran-1,4'-piperidine]} \) (prepared according to example 69) with 3-bromomethyl-3-fluoro-oxetane (described in US2005215599) gave the title compound.

\[ \text{ES-MS m/e (\%): } 455.2(\text{M}+\text{H})^+. \]

Example 244

\( l'-(6\text{-Chloro-l-}\{l-(\text{methoxymethyl})\text{cyclopropyl})\text{methyl}\}-1\text{H-indol-3-yl})\text{carbonyl}-3\text{H-spiro[2-benzofuran-1,4'-piperidine]} \)

Following the general procedure III as described above, the alkylation of \( l'-(6\text{-chloro-1H-indol-3-yl})\text{carbonyl}-3\text{H-spiro[2-benzofuran-1,4'-piperidine]} \) (prepared according to example 69) with 1-bromomethyl-l-methoxymethyl-cyclopropane (described in WO 2001032633) gave the title compound.

\[ \text{ES-MS m/e (\%): } 465.5(\text{M}+\text{H})^+. \]

Example 245

\( (l-\{6\text{-Chloro-3-(rH,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl}\text{methyl})\text{cyclopropyl})\text{acetonitrile} \)
Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (prepared according to example 69) with 1-bromomethyl-cyclopropanecarbonitrile (described in EP 148004) gave the title compound.

ES-MS m/e (%): 460.5(M+H+).

Example 246

r-(6-Chloro-1-[2-(tetrahydro-2H-pyran-4-yl)ethyl]-1H-indol-3-yl)carbonyl)-3H-spiro[2-benzofuran-1,4’-piperidine]

Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (prepared according to example 69) with 4-(2-bromo-ethyl)-tetrahydro-pyran (described in US 2004220214) gave the title compound.
ES-MS m/e (%): 479.5(M+H+).

Example 247

tert-Butyl 2-[(6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-ylcarbonyl)-1H-indol-1-yl]methyl)morpholine-4-carboxylate

Following the general procedure III as described above, the alkylation of l'-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with 2-chloromethyl-morpholine-4-carboxylic acid tert-butyl ester (described in WO 2006020415) gave the title compound.

ES-MS m/e (%): 566.3(M+H+).

Example 248

l'-[(6-Chloro-1-(morpholin-2-ylmethyl)-1H-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] hydrochloride
tert-Butyl 2-\{[6-chloro-3-(1\,\textit{H},3\textit{H}-\textit{spiro}[2\textit{-}benzofuran-1,4'-\textit{piperidin}]-\textit{r-ylcarbonyl})-1\textit{H}-indol-1-yl\}methyl\}morpholine-4-carboxylate (prepared in example 247 above) was dissolved in a solution of HCl (5 eq) in dioxane and stirred at room temperature for 5h. A further portion of HCl (5 eq) in dioxane was added and the solution stirred for an additional 5h at room temperature. Evaporation gave the desired product in quantitative yield. ES-MS m/e (%): 466.2 (M+H\textsuperscript{+}).

Example 249

2-[6-Chloro-3-(1\textit{H},3\textit{H}-\textit{spiro}[2\textit{-}benzofuran-1,4'-\textit{piperidin}]-1'-ylcarbonyl)-1\textit{H}-indol-1-yl]-N-[2-(dimethylamino)ethyl]acetamide

A solution of [6-chloro-3-(1\,\textit{H},3\textit{H}-\textit{spiro}[2\textit{-}benzofuran-1,4'-\textit{piperidin}]-r-ylcarbonyl)-1\textit{H}-indol-1-yl] acetic acid (prepared by treatment of the sodium salt of l'-[(6-chloro-1\textit{H}-indol-3-yl)carbonyl]-3\textit{H}-spiro[2-benzofuran-1,4'-piperidine] with bromoacetic acid at room temperature in DMF), EDC (1 eq), HOBT (1 eq) and Et\textsubscript{3}N (1 eq) were stirred together at room temperature in DMF for 15 min. N,N-Dimethyl-ethane-1,2-diamine (1
eq) was added and the solution stirred at room temperature for 16h. Purification by prep. HPLC gave 37% of product.

ES-MS m/e (%): 495.6(M+H+).

Example 250

5 2-[6-Chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl]acetamide

A solution of [6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl] acetic acid (prepared as for 2-[6-chloro-3-(1'H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-ylcarbonyl)-1H-indol-1-yl]-N-[2-(dimethylamino)ethyl]acetamide), O-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate (1.05 eq) and Et3N (1.05 eq) were stirred together at room temperature in dry DMF for 15 min. A solution of ammonia (5 eq) in dioxane was added and the solution stirred at room temperature for 16 h. Purification by prep. HPLC gave 31% of product.

15 ES-MS m/e (%): 424.3(M+H+).

Example 251

2-[6-Chloro-3-(1'H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-ylcarbonyl)-1H-indol-1-yl]-N-[2-(methylamino)ethyl]acetamide
A solution of [6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-IH-indol-1-yl] acetic acid (prepared by treatment of the sodium salt of 1'-[6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] with bromoacetic acid at room temperature in DMF), O-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate (1.05 eq) and Et₃N (1.05 eq) were stirred together at room temperature in dry DMF for 15 min. Commercially available N-(2-aminoethyl)-N-methylcarbamic acid tert-butyl ester (1.5 eq) was added and the solution stirred at room temperature for 2 h, then a solution of HCl (15 eq) in dioxane was added and the solution stirred for 2 h. Evaporation and purification by prep. HPLC gave 37% of product.

ES-MS m/e (%): 481.3(M+H+).

Example 252

N-(2-Aminoethyl)-2-[6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-IH-indol-1-yl]acetamide
A solution of [6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl] acetic acid (prepared by treatment of the sodium salt of 1'-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] with bromoacetic acid at room temperature in DMF), O-(7-azabenzotriazol-l-yl)-N,N,N',N' -tetramethyluronium hexafluorophosphate (1.05 eq) and Et₃N (1.05 eq) were stirred together at room temperature in dry DMF for 15 min. Commercially available N-(tert-Butoxycarbonyl)-1,2-diaminoethane (1.5 eq) was added and the solution stirred at room temperature for 2 h, then a solution of HCl (15 eq) in dioxane was added and the solution stirred for 2 h. Evaporation and purification by prep. HPLC gave 39% of product.

ES-MS m/e (%): 467.4(M+H⁺).

Example 253

2-[6-Chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-1-yl]ethan amine
Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-
IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (prepared according to
example 69 above) with commercially available 2-chloro-ethylamine gave the title
compound in 11% yield.

ES-MS m/e (%): 410.2(M+H+).

Example 254

2-[6-Chloro-3-(rH,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-ylcarbonyl)-1H-indol-1-
yl]-N-methylethan amine

Following the general procedure III as described above, the alkylation of 1’-[(6-chloro-
IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (prepared according to
example 69) with commercially available (2-chloro-ethyl)-methyl-amine gave the title
compound in 42% yield.

ES-MS m/e (%): 424.2(M+H+).

Example 255

2-[6-Chloro-3-(rH,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-1H-indol-l-
yl]-N-methylacetamide
Following the general procedure III as described above, the alkylation of \( l'-(6\text{-chloro-}lH\text{-indol-3-yl})\text{carbonyl}\)-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with commercially available 2-chloro-\( N\)-methyl-acetamide gave the title compound in 19% yield.

ES-MS m/e (%): 438.2(M+H+).

Example 256

\( l'-(6\text{-Chloro-}l-(2\text{-morpholin-4-ylethyl})\text{-}lH\text{-indol-3-yl})\text{carbonyl}\)-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure III as described above, the alkylation of \( l'-(6\text{-chloro-lH-indol-3-yl})\text{carbonyl}\)-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to example 69) with commercially available 4-(2-chloro-ethyl)-morpholine gave the title compound in 50% yield.
Example 257

l’-[(6-Chloro-l-(3-morpholin-4-ylpropyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine]

Following the general procedure III as described above, the alkylation of l’-[((6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] (prepared according to example 69) with commercially available 4-(3-chloro-propyl)-morpholine gave the title compound in 52% yield.

Example 258

l’-[(6-Chloro-l-(oxiran-2-ylmethyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine]
Following the general procedure III as described above, the alkylation of l′-[(6-chloro-
IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4′-piperidine] (prepared according to
eexample 69) with commercially available 2-bromomethyl-oxirane gave the title compound in 47% yield.

ES-MS m/e (%): 423.4(M+H+).

Example 259

2-[6-Chloro-3-(rH,3H-spiro[2-benzofuran-1,4′-piperidin]-l′-ylcarbonyl)-lH-indol-l-
yl]ethanol

Following the general procedure III as described above, the alkylation of l′-[(6-chloro-
IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4′-piperidine] (prepared according to
eexample 69) with commercially available 2-chloro-ethanol gave the title compound in
47% yield.

ES-MS m/e (%): 411.4(M+H+).

Example 260

r-(l′-[(2-methylpyridin-4-yl)methyl]-lH-indol-3-yl)carbonyl]-3H-spiro[2-
benzofuran-1,4′-piperidine]
Following the general procedure III as described above, the alkylation of \( l'-(6\text{-chloro-}H\text{-indol-3-yl})\text{carbonyl}-3\text{H-spiro}[2\text{-benzofuran-1,4}'\text{-piperidine}] \) (prepared according to example 69) with 4-chloromethyl-2-methyl-pyridine (described in WO 2006023707) gave the title compound in 56% yield.

ES-MS m/e (%): 472.2(M+H+).

**Example 261**

\[ r-(6\text{-Chloro-}l-(3\text{S})\text{-piperidin-3-ylmethyl}-H\text{-indol-3-yl})\text{carbonyl}-3\text{H-spiro}[2\text{-benzofuran-1,4}'\text{-piperidine}] \]

A solution of the \( r-(6\text{-chloro-}H\text{-indol-3-yl})\text{carbonyl}-3\text{H-spiro}[2\text{-benzofuran-1,4}'\text{-piperidine}] \) (prepared according to example 69) in dry DMF was treated with NaH (1.1 eq) and stirred for 10 min at room temperature, then treated with (S)-3-methanesulfonyloxyethyl-piperidine-1-carboxylic acid tert-butyl ester (1.1 eq) (JP 2001278872) and stirred at room temperature for 16h then 70° for 4h. Concentration and
treatment with excess HCl in dioxane gave after purification by preparative HPLC the desired product in 39% yield.

ES-MS m/e (%): 464.2(M+H+).

Example 262

5 2-[6-Chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-1-ylcarbonyl)-1H-indol-1-yl]-N-hydroxyethan amine

A solution of the [6-chloro-3-(H,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-ylcarbonyl)-1H-indol-1-yl]acetaldehyde in MeOH was treated with KOAc (1.5 eq) and hydroxylamine hydrochloride (1.2 eq) and stirred for 1 h at room temperature, then treated with NaCNBH₃ (1.1 eq) and stirred at room temperature for 5 h. Concentration and purification by preparative HPLC the desired product in 19% yield.

ES-MS m/e (%): 426.1(M+H+).

Example 263

15 l'-(6-Chloro-l-(tetrahydro-2H-pyran-4-ylmethyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]
Following the general procedure III as described above, the alkylation of \( l'-(6\text{-chloro-lH-indol-3-yl})\text{carbonyl}-3\text{H-spiro[2-benzofuran-l,4'-piperidine]} \) (prepared according to example 69) with commercially available methanesulfonic acid 2-\((\text{tetrahydro-pyran-4-y})\)-ethyl ester (described in US 2004220214) gave the title compound in 55% yield.

\[
\text{ES-MS } m/e (\%) : 465.2(\text{M+H}^+) .
\]

Example 264

\( l'-(6\text{-Chloro-l-[(l-methylpyrrolidin-3-yl)methyl]-lH-indol-3-yl})\text{carbonyl}-3\text{H-spiro[2-benzofuran-l,4'-piperidine]} \)

A solution of \( l'-(6\text{-Chloro-l-[(pyrrolidin-3-ylmethyl]-lH-indol-3-yl})\text{carbonyl}-3\text{H-spiro[2-benzofuran-l,4'-piperidine]} \) in MeOH was treated with aq. \( \text{H}_2\text{CO} \) (1.5 eq), \( \text{AcOH} \) (1.1 eq) and stirred for 15 min at room temperature, then treated with \( \text{NaCNBH}_3 \) (1.1 eq) and stirred at room temperature for 1h. Concentration and purification by preparative HPLC gave the desired product.

\[
\text{ES-MS } m/e (\%) : 464.2(\text{M+H}^+) .
\]
Example 265

r-[(6-Chloro-l-{[(3S)-l-methylpiperidin-3-yl)methyl]-lH-indol-3-yl}carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine]

A solution of l'-{(6-chloro-l-{(3S)-piperidin-3-ylmethyl}-lH-indol-3-yl}carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidine] in MeOH was treated with aq. H₂CO (1.5 eq), AcOH (1.1 eq) and stirred for 15 min at room temperature, then treated with NaCNBH₃ (1.1 eq) and stirred at room temperature for 1h. Concentration and purification by preparative HPLC the desired product.

ES-MS m/e (%): 478.2(M+H+).

Example 266

l'-{(6-Chloro-l-{pyrrolidin-3-ylmethyl}-lH-indol-3-yl}carbonyl)-3H-spiro[2-benzofuran-1,4'-piperidine]

Following the general procedure III as described above, the alkylation of l'-{(6-chloro-lH-indol-3-yl}carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (prepared according to
example 69) with methanesulfonic acid pyrrolidin-3-ylmethyl ester (described in WO 9742189) gave the title compound.

ES-MS m/e (%): 450.2(M+H+).

Example 267

5 r-{(6-Chloro-l-{[(2S)-pyrrolidin-2-ylmethyl]-lH-indol-3-yl}carbonyl)-3H-spiro[2-benzofuran-l,4'-piperidine

Following the general procedure III as described above, the alkylation of l'-(6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidine] (prepared according to example 69) with methanesulfonic acid (S)-1-pyrrolidin-2-ylmethyl ester (described in Tetrahedron: Asymmetry (1997), 8(13), 2209-2213) gave the title compound.

ES-MS m/e (%): 450.2(M+H+).

Example 268

r-{(6-Chloro-l-{{(2S)-l-methylpyrrolidin-2-yl)methyl}-lH-indol-3-yl}carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidine
l'-(6-Chloro-l-[(2S)-pyrrolidin-2-ylmethyl]-lH-indol-3-yl)carbonyl)-3H-spiro[2-
benzofuran-l,4'-piperidine was treated with a 37% aq. formaldehyde (1.05 eq.), acetic
acid (1.05 eq.) and sodium cyanoborohydride (1.0 eq.) in MeOH at room temperature
for 2h to give after purification by preparative HPLC the title compound.

ES-MS m/e (%): 464.2(M+H+).

Example 269

r-[(6-Chloro-l-[(2R)-l-methylpyrrolidin-2-yl]methyl]-lH-indol-3-yl)carbonyl]-3H-
spiro[2-benzofuran-l,4'-piperidine

1'-(6-Chloro-l-[(2S)-pyrrolidin-2-ylmethyl]-lH-indol-3-yl)carbonyl)-3H-spiro[2-
benzofuran-l,4'-piperidine (prepared according to the procedure described above for the
preparation of l'-(6-chloro-l-[(2S)-pyrrolidin-2-ylmethyl]-lH-indol-3-yl)carbonyl)-
3H-spiro[2-benzofuran-l,4'-piperidine using methanesulfonic acid (R)-l-pyrrolidin-2-
ylmethyl ester instead of methanesulfonic acid (S)-l-pyrrolidin-2-ylmethyl ester) was
treated with a 37% aq. formaldehyde (1.05 eq.), acetic acid (1.05 eq.) and sodium
cyanoborohydride (1.0 eq.) in MeOH at room temperature for 2h to give after purification by prep.
HPLC the title compound.

ES-MS m/e (%): 464.2(M+H+).

Example 270

N-2-[6-Chloro-3-(rH,3H-spiro[2-benzofuran-l,4'-piperidin]-r-ylcarbonyl]-lH-indol-
1-yl]ethyl Jacetamide
2-[6-Chloro-3-(1 H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-yljethanamine was treated with acetylchloride (1.05 eq) and triethylamine (1.05 eq) in dichloromethane under argon at room temperature for 2 h to give after purification by prep. HPLC the title compound.

ES-MS m/e (%): 452.2(M+H+).

Example 271

N-{2-[6-Chloro-3-(1 H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-yljethyl}methanesulfonamide

2-[6-Chloro-3-(1 H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-indol-yljethanamine was treated with mesylchloride (1.05 eq) and triethylamine (1.05 eq) in dichloromethane under argon at room temperature to give after purification by prep. HPLC the title compound.

ES-MS m/e (%): 488.1(M+H+).

Example 272
N-{2-[6-Chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylicarbonyl)-1H-indol-1-yl]ethyl}acetamide

was treated with NaH (1.05 eq), MeI (1.05 eq) in dry DMF under argon at room temperature for 2h to give after purification by prep. HPLC the title compound.

ES-MS m/e (%): 466.2(M+H+).

Example 273

N-{2-[6-Chloro-3-(1H,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylicarbonyl)-1H-indol-1-yl]ethyl}methanesulfonamide

was treated with NaH (1.05 eq), MeI (1.05 eq) in dry
DMF under argon at room temperature for 2 h to give after purification by prep. HPLC the title compound.

ES-MS m/e (%): 502.1(M+H+).

Example 274

5 tert-Butyl 1O-(rH,3H-spiro[2-benzofuran-1,4’-piperidin]-r-ylcarbonyl)-3,4-
dihydropyrazino[1,2-a]indole-2(1H)-carboxylate

a) 10-(2,2,2-Trifluoro-acetyl)-3,4-dihydro-1H-pyrazinor 1,2-alindole-2-carboxylic acid tert-butyl ester

To a stirred solution of 0.21 ml (1.5 mmol) trifluoroacetic anhydride in 7 ml 1,2-
dichloroethane was added at 0 °C a solution of 0.37 g (1.4 mmol) 3,4-dihydro-1H-
pyrazino[1,2-a]indole-2-carboxylic acid tert-butyl ester and a solution of 0.23 ml (1.63
mmol) triethylamine in 3 ml 1,2-dichloroethane. After stirring for 30 min the reaction
mixture was quenched with saturated aqueous sodium hydrogen carbonate solution and
extracted with dichloromethane (2 x 100 ml). The combined organic layers were dried
over sodium sulfate, concentrated in vacuo and purified by flash chromatography (n-
heptane / ethyl acetate) to give the title compound (0.288 g, 58%) as a light yellow solid.

MS m/e (%): 369 (M+H+, 27).

b) 3,4-Dihydro-1H-pyrazinor,2-alindole-2J0-dicarboxylic acid 2-tert-butyl ester

To a stirred solution of 0.21 ml (1.5 mmol) trifluoroacetic anhydride in 7 ml 1,2-
dichloroethane was added at 0 °C a solution of 0.37 g (1.4 mmol) 3,4-dihydro-1H-
pyrazino[1,2-a]indole-2-carboxylic acid tert-butyl ester and a solution of 0.23 ml (1.63
mmol) triethylamine in 3 ml 1,2-dichloroethane. After stirring for 30 min the reaction
mixture was quenched with saturated aqueous sodium hydrogen carbonate solution and
extracted with dichloromethane (2 x 100 ml). The combined organic layers were dried
over sodium sulfate, concentrated in vacuo and purified by flash chromatography (n-
heptane / ethyl acetate) to give the title compound (0.288 g, 58%) as a light yellow solid.

MS m/e (%): 369 (M+H+, 27).
To a solution of 0.29 g (0.77 mmol) 10-(2,2,2-trifluoro-acetyl)-3,4-dihydro-1H-pyrazino[1,2-a]indole-2-carboxylic acid tert-butyl ester in 7 ml N,N-dimethylformamide were subsequently added 0.22 g (4.6 mmol) sodium hydride (50% in oil) and a solution of 0.070 ml (3.9 mmol) water in 1 ml N,N-dimethylformamide at room temperature. The reaction mixture was diluted with tert-butyl methyl ether after 2 h and extracted with 1 M sodium hydroxide solution (2 x 30 ml). The combined aqueous layers were acidified (pH 1-2) with 2 M hydrochloric acid at 0 °C and extracted with tert-butyl methyl ether (3 x 50 ml). The combined organic layers were dried over sodium sulfate and concentrated in vacuo to give the title compound (0.21 g, 86%) as a light brown solid.

MS m/e (%): 315 (M-H+ + , 100).

c) tert-Butyl 10-d H ,3H-spiror2-benzofuran-1,4'-piperidinl-3-ylcarbonyl)-3,4-dihydropyrainor,2-a1indole-2(1H)-carboxylate

To a solution of 0.10 g (0.32 mmol) 3,4-dihydro-1H-pyrazino[1,2-a]indole-2,10-dicarboxylic acid 2-tert-butyl ester, 0.066 g (0.35 mmol) spiroisobenzofuran-1(3H)-4'-piperidine and 0.051 g (0.38 mmol) 1-hydroxybenzotriazole in 3.5 ml N,N-dimethylformamide were added 0.073 g (0.38 mmol) N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride at room temperature. After stirring for 3 h the reaction mixture was diluted with saturated aqueous ammonium chloride solution and extracted with tert-butyl methyl ether (2 x 50 ml). The combined organic layers were washed with 1 M sodium hydroxide solution (1 x 30 ml) and water (1 x 30 ml), dried over sodium sulfate, concentrated in vacuo and purified by flash-chromatography (aminopropyl-modified silica gel, n-heptane / ethyl acetate) to give the title compound (0.097 g, 63%) as a light yellow solid.

MS m/e (%): 488 (M+H+, 81).
Example 275

l’-(1,2,3,4-Tetrahydropyrazino[1,2-a]indol-10-ylcarbonyl)-3H-spiro[2-benzofuran-1,4’-piperidine] hydrochloride

A mixture of 0.095 g (0.19 mmol) tert-butyl 10-(l’H,3H-spiro[2-benzofuran-1,4’-piperidin]-r-ylcarbonyl)-3,4-dihydropyrazino[1,2-a]indole-2(lH)-carboxylate and 1.56 ml of a 1.25 M solution of hydrochloric acid (1.95 mmol) in methanol was stirred for 15 min at 50 °C. The reaction mixture was concentrated in vacuo to give the title compound (0.084 g, 100%) as a light yellow solid.

MS m/e (%): 388 (M+H^+, 100).

Example 276

l’-[(2-Methyl-1,2,3,4-tetrahydropyrazino[1,2-a]indol-10-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine]

A solution of 0.050 g (0.12 mmol) l’-(1,2,3,4-tetrahydropyrazino[1,2-a]indol-10-ylcarbonyl)-3H-spiro[2-benzofuran-1,4’-piperidine] hydrochloride, 0.033 ml (0.24 mmol) triethylamine and 0.028 g (0.94 mmol) paraformaldehyde in 2 ml methanol was heated at reflux for 1 h. The reaction mixture was cooled to 0 °C on an ice-water bath and
treated with 0.011 g (0.18 mmol) sodium cyanoborohydride. After completed addition the mixture was allowed to warm to room temperature and stirred for 2 h. Quenching with water and dilution with 2 M aqueous sodium carbonate solution was followed by extraction with dichloromethane (2 x 50 ml). The combined organic layers were dried over sodium sulfate, concentrated in vacuo and purified by flash-chromatography (aminopropyl-modified silica gel, n-heptane / ethyl acetate) to give the title compound (0.036 g, 76%) as an off-white solid.

MS m/e (%): 402 (M+H⁺, 100).

Example 277

r-[(6-Chloro-2-methyl-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine]

a) (6-Chloro-1H-indol-2-yl)-methanol

To a solution of 2.00 g (8.94 mmol) 6-chlorindole-2-carboxylic acid ethyl ester in 50 ml diethyl ether were added 0.475 g (12.5 mmol) lithium aluminum hydride at 0 °C. The reaction mixture was heated at reflux for 45 min and quenched by consecutive addition of 10 ml water, 10 ml aqueous 2 M sodium hydroxide solution and 10 ml water at 0 °C. The aqueous layer was extracted with tert-butyl methyl ether (3 x 100 ml). The combined organic layers were dried over sodium sulfate and concentrated in vacuo to give the crude title compound (1.64 g; 100%) as a white solid.

MS m/e (%): 180 (M-H⁺, 100).

b) 6-Chloro-2-methyl-1H-indole

A solution of 1.60 g (8.81 mmol) (6-chloro-1H-indol-2-yl)-methanol in 5 ml 1,2-dichloroethane was added to a mixture of 80.0 ml trifluoroacetic acid and 32.0 ml
triethylsilane at 65 °C. After 5 min, the reaction mixture was cooled to room temperature and quenched with water. The pH was adjusted to 14 by the addition of aqueous sodium hydroxide solution (32%). The aqueous layer was extracted with tert-butyl methyl ether (3 x 200 ml). The combined organic layers were dried over sodium sulfate and concentrated in vacuo. The residue was purified by flash-chromatography (aminopropyl-modified silica gel, n-heptane / ethyl acetate) to give the title compound (0.39 g; 27%) as a white solid.

MS m/e (%): 164 (M-H+, 100).

c) 1-(6-Chloro-2-methyl-lH-indol-3-yl)-2,2,2-trifluoro-ethanone

\[
\text{CH}_2=\text{C}
\]

To a solution of 0.38 g (2.3 mmol) 6-chloro-2-methyl-lH-indole in 20 ml 1,2-dichloroethane at 0 °C were added 0.35 ml (2.5 mmol) trifluoroacetic anhydride. The reaction mixture was quenched with aqueous 2 M sodium carbonate solution after 30 min and extracted with dichloromethane (3 x 100 ml). The combined organic layers were dried over sodium sulfate and concentrated in vacuo to give the title compound (0.57 g; 95%) as an off-white solid.

MS m/e (%): 260 (M-H+, 100).

d) 6-Chloro-2-methyl-lH-indole-3-carboxylic acid

\[
\text{CH}_2=\text{C}
\]

A solution of 0.57 g (2.2 mmol) 1-(6-chloro-2-methyl-lH-indol-3-yl)-2,2,2-trifluoro-ethanone in 21.7 ml (86.8 mmol) aqueous 4 M sodium hydroxide solution was heated at reflux for 45 min. After cooling to room temperature the reaction mixture was diluted with water and extracted with tert-butyl methyl ether (2 x 50 ml). The aqueous layer was cooled to 0-5 °C, acidified (pH 1-2) with concentrated aqueous hydrochloric acid solution and extracted with ethyl acetate (3 x 100 ml). The combined ethyl acetate layers
were dried over sodium sulfate and concentrated in vacuo to give the title compound (0.14 g, 31%) as an off-white solid.

MS m/e (%): 208 (M-H+, 100).

e) \textit{r-}(6-Chloro-2-methyl-lH-indol-3-yl)carbonyl-3H-spiror2-benzofuran-l,4'-piperidine

To a solution of 0.040 g (0.19 mmol) 6-chloro-2-methyl-lH-indole-3-carboxylic acid, 0.069 ml (0.40 mmol) N,N-diisopropylethylamine and 0.061 g (0.19 mmol) 2-(1H-benzotriazole-1-yl)-1,1,3,3-tetramethyluronium tetrafluoroborate in 1 ml dry N,N-dimethylformamide were added 0.036 g (0.19 mmol) spiro[isobenzofuran-l(3H),4'-piperidine] at room temperature. After stirring for 1 h the reaction mixture was quenched with 0.5 M aqueous sodium hydroxide solution (20 ml) and extracted with ethyl acetate (2 x 30 ml). The combined organic layers were washed with water (2 x 30 ml) and brine (1 x 30 ml), dried over sodium sulfate and concentrated in vacuo. The residue was purified by flash chromatography (n-heptane / ethyl acetate) to give the title compound (0.036 g, 49%) as a white solid.

MS m/e (%): 379 (M-H+, 100).

Example 278

\textit{r-}(1-[3,5-Bis(trifluoromethyl)benzyl]-6-chloro-2-methyl-lH-indol-3-yl)carbonyl)-3H-spiro[2-benzofuran-l,4'-piperidine]
To a solution of 0.031 g (0.080 mmol) l'-[(6-chloro-2-methyl-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine] (5) in 1 ml dry N,N-dimethylformamide were added 0.004 g (0.08 mmol) sodium hydride (50% in oil). After stirring for 20 min 0.015 ml (0.08 mmol) 3,5-bis(trifluoromethyl)benzyl bromide were added. After stirring for 16 h the reaction mixture was quenched with water and extracted with ethyl acetate (2 x 50 ml). The combined organic layers were washed with water (2 x 30 ml) and brine (1 x 30), dried over sodium sulfate and concentrated in vacuo. The residue was purified by flash chromatography (n-heptane / ethyl acetate) to give the title compound (0.020 g; 41%) as a white solid.

MS m/e (%): 607 (M+H+ 100).

Examples of compounds of formula (I-f)

Example 279

5-Bromo-r-(1H-indol-3-ylcarbonyl)spiro[indole-3,4'-piperidin]-2(IH)-one

a) 5-Bromo-1,2-dihydro-2-oxosphior3H-indole-3,4'-piperidinel^-methyl

A solution of 1,2-benzo-8-methyl-3,8-diazaspiro [4,5] decane-4-one (described in Organic Preparations and Procedures International (1995), 27(6), 691-4) (6.3 g, 29.1
mmol) in CH₂CN (100 ml) and MeOH (5 ml) was cooled to -5 °C and NBS (7.8 g, 44 mmol) was slowly added with stirring. The reaction mixture was stirred for 3.5 h at 0°C. Solvent was removed by vacuo. The residue was purified by silica gel chromatography (2 - 20 % MeOH/dichloromethane) to give 6 g of a solid. The solid compound was dissolved in ethyl acetate (600 ml) and washed with saturated aqueous NaHCO₃ solution, dried (Na₂SO₄). Evaporation of the solvent in vacuo gave 4.2 g (47%) of the desired product.

1HNMR (CD₃OD, 400MHz) δ 7.51(d, J = 1.8 Hz, IH), 7.35(dd, J = 1.9 and 8.2 Hz, IH), 6.81(d, J = 8.2 Hz, IH), 2.93(m, 2H), 2.67(m, 2H), 2.41(s, 3H), 1.86(m, 4H).

b) 5-Bromo-1,2-dihydro-2-oxospiror3H-indole-3,4'-piperidinel-r-cvano

5-Bromo-l, 2-dihydro-2-oxospiro [3H-indole-3, 4'-piperidine]-l '/-methyl (4.6 g, 15.6 mmol) was dissolved in chloroform (700 ml) and treated with CNBr (22 g, 209.5 mmol) at room temperature. The mixture was heated to reflux for 24h. The reaction mixture was cooled, diluted with methylene chloride (300 ml) and washed with 10 % aqueous K₂CO₃ solution (2 x 100 ml). After the mixture was dried (Na₂SO₄) and concentrated, the residue was purified by silica gel chromatography (0 - 5% MeOH/dichloromethane) to give the desired product as a solid (3.9 g, 82%). 1HNMR (CDCl₃, 400MHz) δ 7.52 (d, J = 1.8Hz, IH), 7.37(dd, J = 1.8 and 8.2 Hz, IH), 6.82 (d, J = 8.2 Hz, IH), 3.83(m, 2H), 3.41(m, 2H), 2.00(m, 2H), 1.86(m, 2H).

c) 5-Bromospirorindole-3,4'-piperidinl-2(lH )-one
5-Bromo-1,2-dihydro-2-oxospiro[3H-indole-3,4′-piperidine]-1′-cyano (3.3 g, 10.8 mmol) was suspended in ethylene glycol (10 ml). The mixture was treated in NaOH (1.8 g, 45 mmol) and heated to 130°C for 15 min. It was diluted with methylene chloride (500 ml) and washed with 10% aqueous K₂CO₃ (2 x 100 ml). The organic layer was dried (Na₂SO₄) and concentrated and residue purified by silica gel chromatography (30% MeOH/dichloromethane) to give the desired product as a light ceramic white solid (1.8 g, 60%).

Mp 256 - 258°C. ¹HNMR (DMSO-d₆, 400MHz) δ 10.6(br s, IH, NH), 7.57(d, J = 1.84Hz, IH), 7.36(d, J = 8.2Hz, IH), 6.79(d, J = 8.2 Hz, IH), 4.05 (br s, IH, NH), 3.06(m, 2H), 2.84(m, 2H), 1.64(m, 2H), 1.55(m, 2H). ¹³C NMR (DMSO-d₆, 100MHz) δ 180.93, 140.64, 137.98, 130.42, 126.75, 113.20, 111.45, 46.24, 40.92, 32.94. Anal.Calcd for C₁₂H₁₃BrN₂O: C, 51.26; H, 4.66; N, 9.9. Found: C, 50.87; H, 4.91; N, 9.67.

Following the general procedure I as described above, the acylation of 5-bromospiro[indole-3,4′-piperidin]-2(1H)-one with 1H-indole-3-carboxylic acid (commercially available), gave the title compound.

ES-MS m/e (%): 424.3(M+H +).

Example 280

5-Bromo-r-(6-chloro-1H-indol-3-yl)carbonyl]spiro[indole-3,4′-piperidin]-2(1H)-one
Following the general procedure I as described above, the acylation of 5-bromospiro[indole-3,4'-piperidin]-2(1H)-one with 6-chloro-1H-indole-3-carboxylic acid (preparation described in example 5 above), gave the title compound.

ES-MS m/e (%): 458.3(M+H+)

Example of compounds of formula (I-g)

Example 281

(lR,3'R,5'S)-r-[(l-Benzyl-2-methyl-lH-indol-3-yl)carbonyl]-3',5'-dimethyl-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one

Following the general procedure I as described above, the acylation of (lR,3'R,5'S)-l'-(6-chloro-lH-indol-3-yl)carbonyl]-3',5'-dimethyl-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one (described in WO 9929696) with l-benzyl-2-methyl-lH-indole-3-carboxylic acid (preparation described in example 1), gave the title compound.

ES-MS m/e (%): 479.6(M+H+).
Examples of compounds of formula (I-h)

Example 282

r-[(6-Chloro-lH-indol-3-yl)carbonyl]spiro[isoindole-1,4'-piperidin]-3(2H)-one

a) Bis-(2-chloro-ethyl)-carbamic acid tert-butyl ester

\[
\text{Cl-} \bigg|\bigg|_{\text{N}} \text{Cl} \bigg|\bigg|_{\text{O}} \text{C}
\]

To a solution of 10.0 g (56.0 mmol) bis(2-chloroethyl) amine hydrochloride and 14.7 g (67.2 mmol) di-tert-butyl dicarbonate in 60 ml dichloromethane were added dropwise 9.37 ml (67.2 mmol) triethylamine at room temperature. After stirring for 4 h the solvent was evaporated. The residual oil was redissolved in 300 ml tert-butyl methyl ether and washed with saturated aqueous ammonium chloride solution (1 x 100 ml) and water (1 x 100 ml). The combined aqueous layers were extracted with tert-butyl methyl ether (1 x 200 ml). The combined organic layers were dried over sodium sulfate, concentrated in vacuo and purified by flash chromatography (n-heptane / ethyl acetate) to give the title compound (9.6 g, 71%) as a colourless oil.

MS m/e (%): 186 (M-C\textsubscript{4}H\textsubscript{8}H+, 100).

b) 4-(2-Bromo-phenyl)-4-cyano-piperidine-1-carboxylic acid tert-butyl ester

\[
\text{Br} \bigg|\bigg|_{\text{N}} \bigg|\bigg|_{\text{O}} \text{C}
\]

A mixture of 4.50 g (23.0 mmol) 3-bromophenylacetonitrile and 0.78 g (2.3 mmol) tetrabutylammonium hydrogen sulfate in 27 ml tetrahydrofuran and 45 ml of a 50 % aqueous sodium hydroxide solution was heated at reflux for 10 min. Thereafter 6.11 g (25.3 mmol) bis-(2-chloro-ethyl)-carbamic acid tert-butyl ester were added at room
temperature. The reaction mixture was heated at reflux for 4 h. Cooling to room temperature was followed by dilution with 60 ml water and extraction with tert-butyl methyl ether (3 x 100 ml). The combined organic layers were washed with brine (1 x 100 ml), dried over sodium sulfate and concentrated in vacuo. The residual crude product was purified by flash chromatography (n-heptane / ethyl acetate) to give the title compound (6.72 g, 80%) as a pale yellow oil.

MS m/e (%): 265, 267 (M+BOC+H+ , 82, 100).

c) 4-(2-Bromo-phenyl)-piperidine-1,4-dicarboxylic acid mono-tert-butyl ester

A mixture of 5.7 g (15.6 mmol) 4-(2-bromo-phenyl)-4-cyano-piperidine-1-carboxylic acid tert-butyl ester and 235 ml (936 mmol) of a 4 M aqueous hydrochloric acid solution was heated at reflux for 96 h. After cooling to room temperature the reaction mixture was basified with 93.90 ml (1014 mmol) of a 10.8 M aqueous sodium hydroxide solution and diluted with 200 ml 1,4-dioxane. A solution of 5.11 g (23.4 mmol) di-tert-butyl dicarbonate in 50 ml 1,4-dioxane was added quickly at 0 °C. After stirring for 2 h the reaction mixture was extracted with tert-butyl methyl ether (2 x 100 ml). The combined organic layers were washed with 1 M sodium hydroxide solution (1 x 100 ml). The combined aqueous layers were cooled by the addition of 100 g ice, acidified (pH 1-2) with ice-cold 2 M aqueous hydrochloric acid solution and extracted with ethyl acetate (3 x 150 ml). The combined ethyl acetate layers were dried over sodium sulfate and concentrated in vacuo to give the title compound (5.51 g, 92%) as a light yellow solid.

MS m/e (%): 382, 384 (M+H+ , 90, 100).

d) 4-Azidocarbonyl-4-(2-bromo-phenyl)-piperidine-1-carboxylic acid tert-butyl ester
To a solution of 2.00 g (5.20 mmol) 4-(2-bromo-phenyl)-piperidine-1,4-dicarboxylic acid mono-tert-butyl ester in 26 ml dichloromethane were added 0.76 ml (5.73 mmol) 1-chloro-N,N,2-trimethylpropenylamine at room temperature. After stirring for 45 min the reaction mixture was concentrated in vacuo. The residual oil was redissolved in 26 ml dry N,N-dimethylformamide and treated with 0.51 g (7.8 mmol) sodium azide. After stirring for 1 h the reaction mixture was diluted with 200 ml tert-butyl methyl ether and washed with 0.5 M aqueous sodium carbonate solution (2 x 50 ml). The combined aqueous layers were extracted with tert-butyl methyl ether (2 x 100 ml). The combined organic layers were washed with water (1 x 50 ml) and brine (1 x 50 ml), dried over sodium sulfate and concentrated in vacuo to give the crude title compound (1.76 g, 83%) as a light yellow solid.

e) tert-Butyl 3-oxo-2,3-dihydro-1'H-spirisoindole-1,4'-piperidinel-r-carboxylate

A solution of 1.60 g (3.91 mmol) 4-azidocarbonyl-4-(2-bromo-phenyl)-piperidine-1-carboxylic acid tert-butyl ester in 40 ml toluene was stirred at 90 °C for 1 h. The reaction mixture was concentrated in vacuo and the residue was redissolved in 40 ml dry tetrahydrofuran. Cooling of the solution to -100 °C was followed by slow dropwise addition of 4.6 ml (7.8 mmol) of a 1.7 M solution of tert-butyl lithium in pentane. The cooling bath was removed after 10 min and the reaction mixture was allowed to warm to 0 °C. After quenching with 5 ml saturated aqueous ammonium chloride solution the
mixture was extracted with tert-butyl methyl ether (3 x 50 ml). The combined organic layers were washed with brine (1 x 50 ml), dried over sodium sulfate and concentrated in vacuo to give the title compound (1.15 g, 97%) as a light yellow solid.

**MS:** m/e (%): 247 (M-C₆H₄+H⁺, 100).

5. **Spiroisoindole-1,4'-piperidinl-3(2H)-one**

To a solution of 0.10 g (0.33 mmol) tert-butyl 3-oxo-2,3-dihydro-l'H-spiro[isoindole-1,4'-piperidine]-l'-carboxylate in 3.3 ml dichloromethane were added 0.25 ml (3.3 mmol) trifluoroacetic acid at room temperature. After stirring for 4 h the reaction mixture was diluted with 20 ml aqueous 1 M sodium hydroxide solution (20 ml) and extracted with ethyl acetate (3 x 50 ml). The combined organic layers were dried over sodium sulfate and concentrated in vacuo to give the crude title compound (0.095 g) as a light yellow solid.

**MS:** m/e (%): 203 (M+H⁺, 100).

6. **1'-[(6-ChlTo-lH-indol-3-yl)carbonvHspirorisoindole-1,4'-piperidin1-3(2H)-one**

To a solution of 0.060 g (0.31 mmol) 6-chloro-lH-indole-S-carboxylic acid in 3 ml dichloromethane were added 0.045 ml (0.38 mmol) l-chloro-N,N,2-trimethyl-propenylamine at room temperature. After stirring for 1 h the reaction mixture was concentrated in vacuo. The residue was redissolved in 2 ml dry N,N-dimethylformamide. A solution of 0.062 g (0.31 mmol) spiro[isoindole-1,4'-piperidin]-3(2H)-one and 0.064
ml (0.46 mmol) triethylamine in 1 ml dry N,N-dimethylformamide was added at room temperature. After stirring for 2 h the reaction mixture was quenched with 1 M aqueous sodium hydroxide solution (30 ml) and extracted with ethyl acetate (3 x 50 ml). The combined organic layers were washed with water (2 x 30 ml), 0.5 M aqueous hydrochloric acid solution (1 x 30 ml) and brine (1 x 30 ml), dried over sodium sulfate and concentrated in vacuo. The crude product was triturated in warm diethyl ether, filtrated and dried in vacuo to give the title compound (0.031 g, 22%) as light brown solid with a purity of approx. 80 % by LC-MS.

**MS m/e (%):** 378 (M-H+, 100).

**Example 283**

![Chemical Structure](image)

2-{6-Chloro-3-[(3-oxo-2,3-dihydro-rH-spiro[isoindole-l,4'-piperidin]-l'-yl)carbonyl]-lH-indol-l-yl}-N-methylacetamide

To a solution of 0.35 g (0.13 mmol) 6-chloro-l-methylcarbamoylmethyl-lH-indole-S-carboxylic acid, 0.025 ml (0.14 mmol) N,N-diisopropylethylamine and 0.055 g (0.14 mmol) O-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate in 2 ml dry N,N-dimethylformamide were added 0.029 g (0.14 mmol) spiro[isoindole-l,4'-piperidin]-3(2H)-one at room temperature. After stirring for 2 h the reaction mixture was quenched with 0.5 M aqueous sodium hydroxide solution (20 ml) and extracted with ethyl acetate (2 x 30 ml). The combined organic layers were washed with water (2 x 30 ml) and brine (1 x 30 ml), dried over sodium sulfate and concentrated in vacuo. The residue was purified by flash chromatography (aminopropyl-modified silica gel, dichloromethane / methanol) to give the title compound (0.026 g, 44%) as a white solid.
Example 284

1' - [(6-Chloro-1H-indol-S-y carbonyl-1'^-dihydrospiroisoindole-1'^-piperidine]  

5  a) tert-Butyl 2,3-dihydro-1H-spiroisoindole-1,4'-piperidine-1'-carboxylate

To a solution of 0.20 g (0.66 mmol) tert-butyl 3-oxo-2,3-dihydro-1'H-spiro[isoindole-1,4'-piperidine]-1'-carboxylate in 6.6 ml toluene were added 0.33 ml (0.66 mmol) of a 2 M solution of borane dimethylsulfide complex in tetrahydrofuran. After stirring at reflux for 5 h the mixture was cooled to room temperature, treated with 1.5 ml of methanol and reheated to reflux for 15 min. The reaction mixture was then concentrated in vacuo to give the crude title compound (0.23 g, 84.%) as a light yellow solid with a purity of 70 % by LC-MS.

MS m/e (%): 289 (M+H +, 83).

15  b) 2,3-Dihydrospiroisoindole-1,4'-piperidine dihydrochloride

A solution of 0.16 g (0.56 mmol) tert-butyl 2,3-dihydro-1'H-spiro[isoindole-1,4'-piperidine]-1'-carboxylate in 4.5 ml (5.6 mmol) of a 1.25 M solution of hydrochloric acid in methanol was stirred at 50 °C for 30 min. The reaction mixture was concentrated in
vacuo. The residue was triturated in hot tetrahydrofuran, filtrated and dried in vacuo to give the title compound (0.15 g, 100%) as a light yellow solid.

MS m/e (%): 189 (M+H+, 100).

c) r-rC6-Chloro-lH-indol-B-vDcarbonyll-l^'-dihydrospiroisoindole-l^'-piperidinel

\[
\text{Example 285}
\]

To a solution of 0.10 g (0.51 mmol) 6-chloro-lH-indole-3-carboxylic acid in 5 ml dichloromethane were added 0.074 ml (0.56 mmol) l-chloro-N,N,2-trimethylpropenylamine at room temperature. The reaction mixture was concentrated in vacuo after 1 h. The residue was redissolved in 2 ml dry N,N-dimethylformamide. A suspension of 0.147 g (0.56 mmol) 2,3-dihydrospiro[isoindole-l,4'-piperidine] dihydrochloride and 0.285 ml (2.04 mmol) triethylamine in 2 ml dry N,N-dimethylformamide was added at room temperature. After stirring for 16 h the reaction mixture was quenched with 1 M aqueous sodium hydroxide solution (30 ml) and extracted with ethyl acetate (3 x 50 ml). The combined organic layers were washed with water (2 x 30 ml) and brine (1 x 30 ml), dried over sodium sulfate and concentrated in vacuo. The residue was purified by flash chromatography (dichloromethane / methanol) to give the title compound (0.045 g; 24%) as an off-white solid.

MS m/e (%): 366 (M+H+, 100).

\[
\text{Example 285}
\]

\[
\text{l'-[(6-Chloro-lH-indol-3-yl)carbonyl]-2-methyl-2,3-dihydrospiro[isoindole-l,4'-piperidine]}
\]
a) tert-Butyl 2-methyl-3-oxo-2,3-dihydro-lH-spirorisoindole-l,4'-piperidinel-carboxylate
To a solution of 0.310 g (1.03 mmol) spiro[isoindole-1,4'-piperidin]-3(2H)-one-1-carboxylic acid tert-butyl ester in 10 ml N,N-dimethylformamide were added 0.054 g (1.1 mmol) sodium hydride (50% in oil) at room temperature. After stirring for 30 min 0.067 ml (1.1 mmol) iodomethane were added. The reaction mixture was quenched with water after 1 h and extracted with tert-butyl methyl ether (2 x 100). The combined organic layers were washed with water (2 x 50 ml) and brine (1 x 50 ml), dried over sodium sulfate and concentrated in vacuo to give the title compound (0.32 g, 99%) as light yellow solid. MS m/e (%): 317 (M+H+, 21).

b) tert-Butyl 2-methyl-2,3-dihydro-1H-spiroisoindole-1,4'-piperidin-1'-carboxylate

To a solution of 0.10 g (0.32 mmol) tert-butyl 2-methyl-3-oxo-2,3-dihydro-1'H-spiro[isoindole-1,4'-piperidin]-1'-carboxylate in 3.2 ml toluene were added 0.16 ml (0.32 mmol) of a 2 M solution of borane dimethylsulfide complex in tetrahydrofuran. After heating at reflux for 4 h the reaction mixture was cooled to room temperature, quenched with 1.5 ml of methanol and reheated to reflux for 15 min. The reaction mixture was concentrated in vacuo and the residue was purified by flash chromatography (n-heptane / ethyl) to give the title compound (0.057 g, 60%) as light yellow solid.

MS m/e (%): 303 (M+H+, 100).

c) 2-Methyl-2,3-dihydrospiropiperidin-1,4'-piperidin1
A solution of 0.057 g (0.33 mmol) tert-butyl 2-methyl-2,3-dihydro-lH-spiro[isoindole-1,4'-piperidine]-l'-carboxylate in 1.32 ml (1.65 mmol) of a 1.25 M solution of hydrochloric acid in methanol was stirred at 50 °C for 15 min. The reaction mixture was concentrated in vacuo. The residue was dissolved in 2 M aqueous sodium hydroxide solution and extracted with dichloromethane (3 x 50 ml). The combined organic layers were dried over sodium sulfate and concentrated in vacuo to give the crude title compound (0.041 g) as a pale yellow amorphous solid.

MS m/e (%): 203 (M+H+, 100).

To a solution of 0.040 g (0.20 mmol) 6-chloro-lH-indole-S-carboxylic acid in 3 ml dichloromethane were added 0.03 ml (0.22 mmol) l-chloro-N,N,2-trimethylpropenylamine at room temperature. After stirring for 1 h the reaction mixture was concentrated in vacuo. The residue was redissolved in 2 ml dry N,N-dimethylformamide. A solution of 0.041 g (0.20 mmol) 2-methyl-2,3-dihydrospiro[isoindole-1,4'-piperidine] and 0.043 ml (0.31 mmol) triethylamine in 1 ml dry N,N-dimethylformamide was added at room temperature. After stirring for 2 h the reaction mixture was quenched with 1 M aqueous sodium hydroxide solution (30 ml) and extracted with ethyl acetate (3 x 50 ml). The combined organic layers were washed with water (2 x 30 ml) and brine (1 x 30 ml), dried over sodium sulfate and concentrated in vacuo. The residue was purified by flash
chromatography (n-heptane / ethyl acetate) to give the title compound (0.022 g, 23%) as an off-white solid with a purity of approx. 83% by LC-MS.

MS m/e (%): 380 (M+H+, 100).
1. Compounds of the general formula (I)

![Chemical structures]

wherein

A is selected from the following groups (a), (b), (c), (d), (e), (f), (g) and (h):
wherein in (a) the dotted line is either nil or a double bond;

R$^1$ is H,
or is C$_1$-$C_6$-alkyl optionally substituted by CN,
or is aryl, 5 or 6 membered heteroaryl or sulfonylaryl which are optionally substituted by one or more B,
or is -(CH$_2$)$_m$-$R^a$ wherein R$^a$ is:

CN, OR$^1$, NR$^i$$^R^j$,

C$_3$-$C_6$-cycloalkyl, 3 to 7 membered-heterocycloalkyl (3 from compound 261), aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
or is -(CH$_2$)$_n$-(CO)-R$^b$ or -(CH$_2$)$_n$-(SO$_2$)-R$^b$, wherein R$^b$ is:

c$_1$-$c_6$-alkyl,

C$_1$-$c_6$-alkoxy,

C$_3$-$c_6$-cycloalkyl,

-(CH$_2$)$_m$-NR$^m$$R^iv$,

NR$^i$$R^j$,

C$_3$-$c_6$-cycloalkyl, 4 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
or R$^1$ and R$^3$ together with the indole ring to which they are attached form a 5 or 6 membered heterocycloalkyl which can be substituted by =0, C(O)O-C$_1$-$c_6$-alkyl or C$_1$-$c_6$-alkyl;

R$^2$ is one or more of H, OH, halo, CN, nitro, C$_1$-$c_6$-alkyl optionally substituted by -

NR$^m$$R^iv$, C$_1$-$c_6$-alkoxy, -O-CH$_2$-C$_2$-$c_6$-alkenyl, benzyloxy,
or two R$^2$ may form an oxo or dioxo bridge together with the indole ring to which they are attached;

R$^3$ is H,
or is halo,
or is -(CO)-R$^c$ wherein R$^c$ is:

C$_1$-$c_6$-alkyl,

-(CH$_2$)$_m$-N$R^i$$R^j$,

-(CH$_2$)$_n$-NR$^m$$R^iv$,

5 or 6 membered heterocycloalkyl optionally substituted by C$_1$-$c_6$-alkyl,
or is C$_1$-$c_6$-alkyl or aryl, which are optionally substituted by halo,
-O(CO)-C₁₋₆-alkyl,
or by-NH(CO)R^d, wherein R^d is C^a-alkyl optionally substituted by halo
or nitro, or R^d is aryl or a 5 or 6 membered heteroaryl, which are
optionally substituted by halo, nitro, C^a-alkyl! or C₁₋₆-haloalkyl;

R^4 is one or more of H, halo, C₁₋₆-alkyl or C₁₋₆-alkoxy optionally substituted by OH,
or two R^4 may form an oxo or dioxo bridge together with the phenyl ring to
which they are attached;

R^5 is H, C₁₋₆-alkyl or aryl;

R^6 is H or C₁₋₆-alkyl;

R^7 is H or -SO₂-R^8 wherein R^8 is C₁₋₆-alkyl or aryl;

R^8 is H or C₁₋₆-alkyl;

X is CH₂ or C=O;

B is halo, CN, NR'R''R''', C₁₋₆-alkyl optionally substituted by CN, halo or C₁₋₆-alkoxy,
C₁₋₆-alkoxy, C₁₋₆-haloalkoxy, C₃₋₆-cycloalkyl, -C(O)O-C₁₋₆-alkyl, -C(O) NR'R''R''', -
C(O)-C₁₋₆-alkyl, -S(O)₂-C₁₋₆-alkyl, -S(O)₂-C₁₋₆-alkyl, -S(O)₂-C₁₋₆-alkyl, -S(O)₂-
(CRᵐR''')ₙ-phenyl, or

-alkyl;  
R^1 and R'^ are H, C₁₋₆-alkyl, C₁₋₆-alkyl-CNCR''R''', -(CO)O-C₁₋₆-alkyl, -C(O)-NR'R''R''', -
C(O)-C₁₋₆-alkyl, -S(O)₂-C₁₋₆-alkyl, -S(O)₂-C₁₋₆-alkyl, -S(O)₂-
NR'R''R''' or OH;

R'' and R''' are H or C₁₋₆-alkyl;

m is 1 to 6;

n is Oto 4;

as well as pharmaceutically acceptable salts thereof.

2. Compounds of the general formula (I) according to claim 1, wherein

A is selected from (a), (b), (c), (d), (e), (f), (g) or (h) and wherein

R^1 is H,
or is C₁₋₆-alkyl optionally substituted by CN,
or is aryl, 5 or 6 membered heteroaryl or sulfonylaryl which are optionally
substituted by one or more B,

or is -(CH₂)ₘ-R^a wherein R^a is:

OR^1,
CN,
NR\(R')_1,
C\(_3\)-\(_6\)-cycloalkyl, 3 to 7 membered heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,

or is -(CH\(_2\))\(_n\)-(CO)-R\(^b\) or -(CH\(_2\))\(_n\)-(SO\(_2\))-R\(^b\), wherein R\(^b\) is:

Ci-6-alkoxy,
NR\(R')_1,
4 to 7 membered heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,

or R\(^1\) and R\(^3\) together with the indole ring to which they are attached form a 5 or 6 membered heterocycloalkyl which can be substituted by =0, C(O)O-C\(_1\)-\(_6\)-alkyl or C\(_1\)-\(_6\)-alkyl;

R\(^2\) is one or more of H, OH, halo, C\(_1\)-\(_6\)-alkyl optionally substituted by -NR\(^m\)R\(^v\), C\(_1\)-\(_6\)-alkoxy;

R\(^3\) is H,
or is halo
or is -(CO)-R\(^c\), wherein R\(^c\) is:

Ci-6-alkyl
-(CH\(_2\))\(^n\)-VR\(^n\),
-(CH\(_2\))\(^n\)-NR\(^m\)R\(^v\),
5 or 6 membered heterocycloalkyl optionally substituted by C\(_1\)-\(_6\)-alkyl, or is C\(_1\)-\(_6\)-alkyl or aryl, which are optionally substituted by halo,

R\(^4\) is one or more of H, halo, or C\(_1\)-\(_6\)-alkoxy optionally substituted by OH, or two R\(^4\) may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;

R\(^5\) is H or aryl;
R\(^6\) is H;
R\(^7\) is H or -SO\(_2\)-R\(^e\) wherein R\(^e\) is C\(_1\)-\(_6\)-alkyl or aryl;
R\(^8\) is H or Ci-e-alkyl;

X is CH\(_2\) or C=O;

B is halo, CN, NH\(_2\), C\(_1\)-\(_6\)-alkyl optionally substituted by CN or C\(_1\)-\(_6\)-alkoxy, C\(_1\)-\(_6\)-alkoxy, Q-e-haloalkoxy, C\(_3\)-\(_6\)-cycloalkyl, -C(O)O-C\(_1\)-\(_6\)-alkyl, -(CR\(^m\)R\(^v\))\(^n\)-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:

halo, C\(_1\)-\(_6\)-alkyl optionally substituted by CN or halo, C\(_1\)-\(_6\)-alkoxy;
R¹ and R¹¹ are H, C₁₋₆-alkyl, Ci₋₆-alkyl-NRᵐRˡV, -(CO)O-C₁₋₆-alkyl, -(C(O)-NRᵐRˡV, -(C(O)-
Ci₋₆-alkyl, -S(O)₂Ci₋₆-alkyl or -S(O)₂⁻NRᵐRˡV or OH;
R⁶ and R⁶¹ are H or C₁₋₆-alkyl;
m is 1 to 6;
> n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

3. Compounds of the general formula (I) according to claim 1 or 2,
wherein A is selected from the following groups (a), (b), (c), (d) and (e); and

R¹ is H,
or is Ci₋₆-alkyl optionally substituted by CN,
or is aryl, 5 or 6 membered heteroaryl or sulfonylaryl which are optionally
substituted by one or more B,
or is -(CH₂)ₘ⁻Rᵃ where Rᵃ is:
CN,
OR¹,
NR²⁻Rⁿ,
C₃₋₆-cycloalkyl, 4 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered
heteroaryl which are optionally substituted by one or more B,
or is -(CH₂)ₙ -(CO)-Rᵇ or -(CH₂)ₙ -(SO₂)-Rᵇ, wherein Rᵇ is:
ci₋₆-alkyl,
Ci₋₆-alkoxy,
C₃₋₆-cycloalkyl,
-(CH₂)ₘ⁻NRᵐRˡV,
NR²⁻Rⁿ,
C₃₋₆-cycloalkyl, 4 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered
heteroaryl which are optionally substituted by one or more B,
or R¹ and R³ together with the indole ring to which they are attached form a 5 or
6 membered heterocycloalkyl which can be substituted by (CO);
R² is one or more of H, OH, halo, CN, nitro, C⁻-alkyl! optionally substituted by -
NRᵐRˡV, Ci₋₆-alkoxy, -O-CH₂⁻C₂₋₆-alkenyl, benzylxoy,
or two R² may form an oxo or dioxo bridge together with the indole ring to
which they are attached;
R³ is H,
or is halo,
or is -(CO)-R^c wherein R^c is:

-\(\text{Ci-}_6^-\text{alkyl}\),
-\((\text{CH}_2\text{VNR}^d\text{R}^d)\),
-\((\text{CH}_2)_n\text{-NR}^m\text{R}^l\),

5

or 6 membered heterocycloalkyl optionally substituted by \(\text{Ci-}_6^-\text{alkyl}\),

or is \(\text{Ci-}_6^-\text{alkyl}\) or aryl, which are optionally substituted by

halo,

-\((\text{O})\text{-Ci-}_6^-\text{alkyl}\),

or by \(\text{NH}^-\text{(CO)R}^-\text{d}\), wherein R^d is \(\text{C}^-\text{alky!}\) optionally substituted by halo

or nitro, or R^d is aryl or a 5 or 6 membered heteroaryl, which are

optionally substituted by halo, nitro, \(\text{C}^-\text{alky!}\) or \(\text{C}_1^-\text{-haloalkyl}\);

\(R^4\) is one or more of \(\text{H}, \text{halo}, \text{C}^-\text{alky!}\) or \(\text{C}_1^-\text{-alkoxy}\) or two \(R^4\) may form an oxo

or dioxo bridge together with the phenyl ring to which they are attached;

\(R^5\) is \(\text{H}, \text{Ci-}_6^-\text{alkyl}\) or aryl;

15 \(R^6\) is \(\text{H}\) or \(\text{C}_1^-\text{-alkyl}\);

\(R^7\) is \(\text{H}\) or \(-\text{SO}_{2}^-\text{R}^-\text{e}\) wherein R^e is \(\text{Ci-}_6^-\text{alkyl}\) or aryl;

B is

halo, \(\text{CN}, \text{NR}^-\text{R}^-\text{v}, \text{Ci-}_6^-\text{alkyl}\) optionally substituted by \(\text{CN}, \text{halo}\) or \(\text{Ci-}_6^-\text{alkoxy},\)

\(\text{Ci-}_6^-\text{alkoxy}\), \(\text{Ci-}\text{-e-haloalkoxy}, \text{C}_5^-\text{-cycloalkyl}, \text{-C}(\text{O})\text{-O-C}_1^-\text{-alkyl}, \text{-C}(\text{O})\text{-NR}^-\text{R}^-\text{v}, \text{-}

\(\text{C}(\text{O})\text{-Ci-}_6^-\text{alkyl}, \text{-S}(\text{O})_2^-\text{Ci-}_6^-\text{alkyl}, \text{-S}(\text{O})_2^-\text{-NR}^-\text{R}^-\text{v}, (\text{CR}^-\text{R}^-\text{v})_n\text{-phenyl}, \text{or}

20

\(\text{(CR}^-\text{R}^-\text{v})_n\text{-}5\) or 6 membered heteroaryl wherein the phenyl or 5 or 6 membered

heteroaryl moiety is optionally substituted by one or more substituent(s)

selected from the group consisting of:

-\(\text{halo}, \text{CN}, \text{NR}^-\text{R}^-\text{v}, \text{Ci-}_6^-\text{alkyl}\) optionally substituted by \(\text{CN}\) or \(\text{Ci-}_6^-\text{alkoxy},\)

\(\text{Ci-}_6^-\text{alkoxy}\), \(\text{Ci-}\text{-e-haloalkoxy}, \text{C}_5^-\text{-cycloalkyl}, \text{-C}(\text{O})\text{-O-C}_1^-\text{-alkyl}, \text{-C}(\text{O})\text{-}

\(\text{NR}^-\text{R}^-\text{v}, \text{-C}(\text{O})\text{-C}_1^-\text{-alkyl}, \text{-S}(\text{O})_2^-\text{C}_1^-\text{-alkyl}, \text{-S}(\text{O})_2^-\text{-NR}^-\text{R}^-\text{v};\)

\(R^1\) and \(R^v\) are \(\text{H}, \text{Ci-}_6^-\text{alkyl}, \text{Ci-}_6^-\text{alkyl-NR}^-\text{R}^-\text{v}, \text{-CO-O-Ci-}_6^-\text{alkyl}, \text{-C}(\text{O})\text{-NR}^-\text{R}^-\text{v}, \text{-C}(\text{O})\text{-}

\(\text{Ci-}_6^-\text{alkyl}, \text{-S}(\text{O})_2^-\text{Ci-}_6^-\text{alkyl} \text{or -S}(\text{O})_2^-\text{-NR}^-\text{R}^-\text{v};\)

\(R^w\) and \(R^u\) are \(\text{H}\) or \(\text{Ci-}_6^-\text{alkyl};\)

\(m\) is 1 to 6;

30 \(n\) is Otto 4;

as well as pharmaceutically acceptable salts thereof.

4. Compounds of the general formula (I) of claim 1 to 3,

wherein A is selected from the groups (a), (b), (c), (d) and (e); and

\(R^1\) is \(\text{H}\) or,

35 \(\text{Ci-}_6^-\text{alkyl}\) optionally substituted by \(\text{CN}\) or,
Ci\textsubscript{6}-alkoxy or, aryl or, 5 or 6 membered heteroaryl or, sulfonylaryl or, 

\[(CH_2)_m-R^a\] wherein \(R^a\) is C\textsubscript{3-6}-cycloalkyl, 5 or 6 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more substituents selected from the group consisting of: halo, CN, C\textsubscript{1-6}-alkyl, C\textsubscript{1-6}-alkoxy, C\textsubscript{1-6}-haloalkoxy, -C(O)O-C\textsubscript{1-6}-alkyl, and phenyl optionally substituted by halo, C\textsubscript{1-6}-alkyl, Ci\textsubscript{6}-haloalkyl or Ci\textsubscript{6}-alkoxy, 

\[(CH_2)_m-NR'R''\] or, 

\[(CH_2)_n-(CO)-R^b\] wherein \(R^b\) is aryl or 5 or 6 membered-heterocycloalkyl; 

\(R^2\) is one or more of H, halo, CN, nitro, C\textsubscript{1-0}-alkyl, C\textsubscript{1-6}-alkoxy, -O-CH\textsubscript{2}-C\textsubscript{2-6}-alkenyl, benzyloxy, or two \(R^2\) may form an oxo or dioxo bridge together with the indole ring to which they are attached; 

\(R^3\) is H or, halo or, -(CO)-R\textsuperscript{c}, wherein \(R^c\) is C\textsubscript{1-6}-alkyl, 5 or 6 membered heterocycloalkyl optionally substituted by C\textsubscript{1-6}-alkyl, or \(R^c\) is -(CH\textsubscript{2})\textsubscript{n}-NR'R'' or, 

C\textsubscript{1-6}-alkyl or aryl, which are optionally substituted by: -O(CO)-C\textsubscript{1-6}-alkyl, or by-NH(CO)R\textsuperscript{d}, wherein \(R^d\) is C\textsubscript{1-6}-alkyl optionally substituted by halo or nitro, or \(R^d\) is aryl or a 5 or 6 membered heteroaryl, which are optionally substituted by halo, nitro, C\textsubscript{1-6}-alkyl or C\textsubscript{1-6}-haloalkyl; 

\(R^4\) is one or more of H, halo, C\textsubscript{1-6}-alkyl or C\textsubscript{1-6}-alkoxy or two \(R^4\) may form an oxo or dioxo bridge together with the phenyl ring to which they are attached; 

\(R^5\) is H, C\textsubscript{1-6}-alkyl or aryl; 

\(R^6\) is H or Ci-e-alkyl; 

\(R^7\) is H or -SO\textsubscript{2}-R\textsuperscript{e} wherein \(R^e\) is Q-e-alkyl or aryl; 

\(R^r\) and \(R^r\) are independently selected from H, Ci-e-alkyl or -(CO)O-C\textsubscript{1-6}-alkyl; 

\(m\) is 1 to 6; 

\(n\) is 0 to 4; 
as well as pharmaceutically acceptable salts thereof.
5. The compounds of formula (I-a) according to any one of claims 1 to 4:

wherein the dotted line is either nil or a double bond and R\textsuperscript{1} to R\textsuperscript{6} are as defined in any one of claims 1 to 4.

6. The compounds of formula (I-a) according to claim 5, wherein:

the dotted line is either nil or a double bond;

R\textsuperscript{1} is H,

or is C\textsubscript{1-6}-alkyl optionally substituted by CN,

or is sulfonylaryl,

or is -(CH\textsubscript{2})\textsubscript{m}-R\textsuperscript{a} wherein R\textsuperscript{a} is:

OR\textsuperscript{1},

CN,

NR\textsubscript{1}R\textsubscript{2},

C\textsubscript{3-6}-cycloalkyl, 3 to 6 membered-heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,

or is -(CH\textsubscript{2})\textsubscript{n}-(CO)-R\textsuperscript{b}, wherein R\textsuperscript{b} is:

C\textsubscript{1-6}-alkoxy,

NR\textsubscript{1}R\textsubscript{2},

6 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B;

R\textsuperscript{2} is one or more of H, halo, C\textsubscript{1-6}-alkyl;

R\textsuperscript{3} is H,

or is C\textsubscript{1-6}-alkyl,

or is -(CO)-R\textsuperscript{c} wherein R\textsuperscript{c} is:

C\textsubscript{1-6}-alkyl
- (CH₂)n - NR R⁺,

R⁴ is one or more of H, halo, or C₁₋₆-alkoxy optionally substituted by OH, or two R⁴ may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;

5 R⁵ is H;

R⁶ is H;

B is halo, CN, Q-o-alkyl optionally substituted by CN or C₁₋₆-alkoxy, C₁₋₆-alkoxy, C₁₋₆-haloalkoxy, C₁₋₆-cycloalkyl, -C(O)O-C₁₋₆-alkyl, -(CRᵐRˡ)ₙ-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:

halo, C₁₋₆-alkyl optionally substituted by CN or halo, C₁₋₆-alkoxy;

R¹ and R⁺ are H, C₁₋₆-alkyl, Ci₋₆-alkyl-NRᵐRˡ, -C(O)-C₁₋₆-alkyl, -S(O)₂-Ci₋₆-alkyl or OH;

R⁺ and R⁺ are H or C₁₋₆-alkyl;

m is 1 to 6;

n is 0 to 4;

as well as pharmaceutically acceptable salts thereof.

7. The compounds of formula (I-a) according to claim 5 or 6, wherein:

the dotted line is either nil or a double bond;

R¹ is H,

or is -(CH₂)ₗ-Rᵃ wherein Rᵃ is aryl which is optionally substituted by one or more substituents selected from the group consisting of:

halo, CN, Ci-β-alkyl, Ci₋₆-alkoxy, Ci-e-haloalkoxy, -C(O)O-C₁₋₆-alkyl and phenyl optionally substituted by halo, C₁₋₆-alkyl, Ci₋₆-haloalkyl or Ci₋₆-alkoxy;

20 R² is H or halo;

R³ is H or Q₋₆-alkyl; and

R⁴, R⁵ and R⁶ are H;

m is 1 to 6;

as well as pharmaceutically acceptable salts thereof.

8. The compounds of formula (I-a) according to claims 5 to 7, wherein said compounds are selected from the group consisting of:

r-[(1-benzyl-2-methyl-lH-indol-3-yl)carbonyl]spiro[indene-1,4’-piperidine];

1’-[(1-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-2,3-dihydrospiro[indene-1,4’-piperidine];

35 1’’-[(1-benzyl-lH-indol-3-yl)carbonyl]spiro[indene-1,4’-piperidine];
9. The compounds of formula (I-b) according to any one of claims 1 to 4:

\[
\begin{array}{c}
\text{R}^1 \text{H} \\
\text{or is Ci-6-alkyl optionally substituted by CN,}
\end{array}
\begin{array}{c}
\text{or is sulfonylaryl,}
\text{or is -(CH}_2)_m-\text{Ra where in R}^a \text{is:}
\text{OR,}
\text{CN,}
\text{NRR},
\text{C}_3\text{-6-cycloalkyl, 3 to 6 membered-heterocycloalkyl, aryl or 5 or 6 membered}
\text{heteroaryl which are optionally substituted by one or more B,}
\text{or is -(CH}_2)_n-(\text{CO})-\text{R}^b \text{wherein R}^b \text{is:}
\text{Ci-6-alkoxy,}
\text{NRR},
\text{5 or 6 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which}
\text{are optionally substituted by one or more B;}
\end{array}
\]

10. The compounds of formula (I-b) according to claim 9, wherein

\[
\begin{array}{c}
\text{R}^2 \text{is one or more of H, halo, C}_1\text{-6-alkyl;}
\text{R}^3 \text{is H,}
\text{or is Ci-6-alkyl,}
\text{or is -(CO) R}^c \text{wherein R}^c \text{is:}
\end{array}
\]
Ci-alkyl
-(CHz)n-NR R,
R4 is is one or more of H, halo, or C1-alkoxy optionally substituted by OH, or two R4 may form an oxo or dioxo bridge together with the phenyl ring to which they are
attached;
R6 is H;
R7 is H or -SO2-R wherein R is Ci-alkyl or aryl;
B is halo, CN, Q-o-alkyl optionally substituted by CN, or C1-alkoxy, C1-alkoxy, C1-6-haloalkoxy, C1-6-cycloalkyl, -C(O)O-C1-6-alkyl, -(CRmRn)-phenyl, wherein
the phenyl is optionally substituted by one or more substituent(s) selected from the
halo, C1-alkyl optionally substituted by CN or halo, C1-alkoxy;
R1 and R are H, C1-alkyl, C1-6-alkyl-NR R, -C(O)-C1-6-alkyl, -S(O)2-C1-6-alkyl or OH;
R and R are H or C1-alkyl;
m is 1 to 6;
n is Oto 4;
as well as pharmaceutically acceptable salts thereof.

11. The compounds of formula (I-b) according to claim 9 or 10, wherein:
R1 is H,
or is -(CHz)m-R wherein R is aryl, or 5 or 6 membered heteroaryl which are
optionally substituted by one or more substituents selected from the group
consisting of:
halo, CN, C1-alkyl, C1-alkoxy, C1-haloalkoxy, -C(O)O-C1-6-alkyl
and phenyl optionally substituted by halo, Ci-6-haloalkyl or C1-6-alkoxy,
R2 is H or halo;
R3 is H or C1-alkyl;
R4 is H or halo;
R6 is H;
R7 is H or -SO2-R wherein R is Ci-alkyl or aryl;
B is halo, NH2, C1-alkyl optionally substituted by CN, or C1-alkoxy, C1-alkoxy,
Ci-alkoxy, C1-6-cycloalkyl, -C(O)O-C1-alkyl, -(CRmRn)-phenyl, wherein
the phenyl is optionally substituted by one or more substituent(s) selected from
the group consisting of:
halo, C\textsubscript{1-6}-alkyl optionally substituted by CN or halo, C\textsubscript{1-6}-alkoxy;
R\textsuperscript{1}, R\textsuperscript{1} are independently selected from H or C\textsubscript{1-6}-alkyl;
m is 1 to 6;
n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

12. The compounds of formula (I-b) according to any one of claims 9 to 11, wherein said compounds are selected from the group consisting of:
\begin{align*}
\mathrm{r-} \{ \mathrm{l}-(\mathrm{1-benzyl-2-methyl-lH-indol-3-yl})\mathrm{carbonyl}\}-\mathrm{l}\-(\mathrm{methylsulfonyl})-1,2-
dihydrospiro[indole-3,4'-piperidine];\\
\mathrm{r-} \{ \mathrm{[6-chloro-l-(3-fluorobenzoyl)-lH-indol-3-yl]carbonyl}\}-1,2-
dihydrospiro[indole-3,4'-piperidine];\\
\mathrm{l'}-\{ \mathrm{[6-chloro-l-(2-fluorobenzoyl)-lH-indol-3-yl]carbonyl}\}-1,2-
dihydrospiro[indole-3,4'-piperidine];\\
\mathrm{l'}-\{ \mathrm{[6-chloro-l-(3,5-difluorobenzoyl)-lH-indol-3-yl]carbonyl}\}-1,2-
dihydrospiro[indole-3,4'-piperidine];\\
\mathrm{r-} \{ \mathrm{[6-chloro-l-(2,3-difluorobenzoyl)-lH-indol-3-yl]carbonyl}\}-1,2-
dihydrospiro[indole-3,4'-piperidine];\\
\mathrm{l'}-\{ \mathrm{[6-chloro-l-(3,5-difluorophenyl)sulfonyl]-lH-indol-3-yl]carbonyl}\}-1,2-
dihydrospiro[indole-3,4'-piperidine];\\
\mathrm{2-}\{ \mathrm{[6-chloro-3-(1,2-dihydro-rH-spiro[indole-3,4'-piperidin]-l'-ylcarbonyl]-lH-indol-1-yl}\}-1-(3,5-difluorophenyl)ethanone;\\
\mathrm{2-}\{ \mathrm{[6-chloro-3-(1,2-dihydro-rH-spiro[indole-3,4'-piperidin]-l'-ylcarbonyl]-lH-indol-1-yl}\}-1-(3,4-difluorophenyl)ethanone;\\
\mathrm{2-}\{ \mathrm{[6-chloro-3-(1,2-dihydro-rH-spiro[indole-3,4'-piperidin]-l'-ylcarbonyl]-lH-indol-1-yl}\}-1-(2-fluorophenyl)ethanone;\\
\mathrm{2-}\{ \mathrm{[6-chloro-3-(1,2-dihydro-rH-spiro[indole-3,4'-piperidin]-l'-ylcarbonyl]-lH-indol-1-yl}\}-N,N-diethylethan amine; and\\
\mathrm{l'}-\{ \mathrm{[6-chloro-l-(pyridin-2-ylmethyl)-lH-indol-3-yl]carbonyl}\}-1,2-
dihydrospiro[indole-3,4'-piperidine].
\end{align*}

13. The compounds of formula (I-c) according to any one of claims 1 to 4:
wherein R\textsuperscript{1} to R\textsuperscript{4} are as defined in any one of claims 1 to 4.

14. The compounds of formula (I-c) according to claim 13, wherein

\begin{itemize}
  \item R\textsuperscript{1} is H,
  \item or is Ci-\textsubscript{6}-alkyl optionally substituted by CN,
  \item or is aryl, 5 or 6 membered heteroaryl or sulfonylaryl which are optionally substituted by one or more B,
  \item or is -(CH\textsubscript{2})\textsuperscript{m}-R\textsuperscript{a} wherein R\textsuperscript{a} is:
    \begin{itemize}
      \item OR\textsuperscript{1},
      \item CN,
      \item NR\textsuperscript{R}\textsuperscript{R}',
      \item C\textsubscript{3-6}-cycloalkyl,
      \item 4 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
    \end{itemize}
  \item or is -(CH\textsubscript{2})\textsuperscript{n}-(CO)-R\textsuperscript{b} wherein R\textsuperscript{b} is:
    \begin{itemize}
      \item Ci-\textsubscript{6}-alkoxy,
      \item NR\textsuperscript{R}\textsuperscript{R}',
      \item 4 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
    \end{itemize}
  \item or R\textsuperscript{1} and R\textsuperscript{3} together with the indole ring to which they are attached form a 5 or 6 membered heterocycloalkyl which can be substituted by =0;
  \item R\textsuperscript{2} is one or more of H, halo, Ci-\textsubscript{6}-alkyl optionally substituted by -NR\textsuperscript{m}R\textsuperscript{b}, C\textsubscript{1-6}-alkoxy;
  \item R\textsuperscript{3} is H,
  \item or is Ci-\textsubscript{6}-alkyl,
  \item or is halo,
  \item or is -(CO)-R\textsuperscript{c}, wherein R\textsuperscript{c} is:
\end{itemize}
Ci-6-alkyl

-(CHz)n-NR'R'

-(CH2)n-NRmR'l

5 or 6 membered heterocycloalkyl optionally substituted by C1-6-alkyl;

R4 is one or more of H, halo, or C1-6-alkoxy optionally substituted by OH, or two R4 may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;

B is halo, NH2, Q-o-alkyl optionally substituted by CN or C1-6-alkoxy, C1-6-alkoxy, Ci-e-haloalkoxy, C3-6-cycloalkyl, -C(O)O-C1-6-alkyl, -(CRmRl)n-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:

halo, Ci-6-alkyl optionally substituted by CN or halo, C1-6-alkoxy;

R1 and R'u are H, C1-6-alkyl, Ci-6-alkyl-NR'mR'l, -(CO)O-C1-6-alkyl, -(C(O)-NR'mR'l, -(C(O)-Ci-6-alkyl, -S(O)2-Ci-6-alkyl, -S(O)2-NR'mR'l;

R111 and R11V are H or C1-6-alkyl;

m is 1 to 6;

n is 0 to 4;

as well as pharmaceutically acceptable salts thereof.

15. The compounds of formula (I-c) according to claim 13 or 14, wherein said compounds are selected from the group consisting of:

r-[(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]-6-chloro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-[(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]-4-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-[(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]-6-methoxy-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-[(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]-5-methoxy-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-[(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]-7-chloro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-[(6-chloro-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-[(1-benzyl-2-methyl-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
6-chloro-3-{(5-fluoro-3-oxo-rH,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl} -N,N-dimethyl-lH-indole-2-carboxamide;

tert-butyl 2-{[(6-chloro-3-{(5-fluoro-3-oxo-lH,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl} -lH-indol-2-yl]carbonyl}amino}ethylJmethylcarbamate;

6-chloro-N,N-diethyl-3-{[(5-fluoro-3-oxo-lH,3H-spiro[2-benzofuran-1,4'-piperidin]-1'-yl)carbonyl} -lH-indole-2-carboxamide;

r-lf 6-chloro\^\^piperidin-l-ylcarbony\^\^lH-indol-S-ylJlcarbonyl\^\^S-fluoro-SH-spiro[2-benzofuran-1,4'-piperidin]-7-one;

r-[l benzyl-2-methyl-lH-indol-3-yl]carbonyl]-7H-spiro[furo[3,4-

[1,3]benzodioxole-5,4'-piperidin]-7-one;

3-{6-chloro-3-{[(3-oxo-lH,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl]-lH-indol-1-yl}propanenitrile;

{6-chloro-3-{[(3-oxo-lH ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-yl)carbonyl]-lH-indol-1-yl}acetonitrile;

l'-[(l-benzyl-2-methyl-lH-indol-3-yl]carbonyl]-6-(2-hydroxyethoxy)-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-[(l-benzyl-2-methyl-lH-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-[[6 chloro-l-(3,5-difluorophenyl]-lH-indol-3-yl]carbonyl] -3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-[[6 chloro-l-(3,5-difluorobenzoyl]-lH-indol-3-yl]carbonyl] -3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-[[6 chloro-l-(3,5-difluorobenzyl]-lH-indol-3-yl]carbonyl] -3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-[[6 chloro-l-(3-fluorobenzyl]-lH-indol-3-yl]carbonyl] -3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-[[6 chloro-l-(3-fluorobenzyl]-lH-indol-3-yl]carbonyl] -3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-(l benzyl-l-C\^\^S^\^S-difluorobenzy\^\^lH-indol-S-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-lfo-chloro-l-CS^\^S-difluorobenzy\^\^lH-indol-S-ylJlcarbonyl\^\^S-fluoro-SH-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-[[6 chloro-l-(3-fluorobenzyl]-lH-indol-3-yl]carbonyl] -3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-(l benzyl-l-[2-(3-fluorophenyl)-2-oxoethyl]-lH-indol-3-yl]carbonyl] -3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-[(l-benzyl-l-[2-(2,5-difluorophenyl)-2-oxoethyl]-lH-indol-3-yl]carbonyl] -3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-lfo-chloro-l-CS^\^S^\^S-difluorobenzy\^\^lH-indol-S-ylJlcarbonyl\^\^S-fluoro-SH-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-(l benzyl-l-[2-(3-fluorophenyl)-2-oxoethyl]-lH-indol-3-yl]carbonyl] -3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
5-bromo-1'-(6-chloro-1-(3,5-difluorobenzyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
5-bromo-1'-(6-chloro-1-(3-fluorobenzyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-(6-chloro-1-[2-(2-fluorophenyl)-2-oxoethyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-(6-chloro-1-[2-(3-fluorophenyl)-2-oxoethyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-(6-chloro-1-[2-(3,4-difluorophenyl)-2-oxoethyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-(6-chloro-1-[2-(3-fluorophenyl)-2-oxoethyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-(6-chloro-1-[2-(2-morpholin-4-yl-2-oxoethyl)-1H-indol-3-yl]carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-(6-chloro-1-[2-(2-methylpyridin-4-yl)methyl]-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-(6-chloro-1-[2-(6-chloropyridin-3-yl)methyl]-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-(6-chloro-1-[2-(3-chloro-6-methylpyridazin-4-yl)methyl]-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
1'-(6-chloro-1-(pyridin-4-ylmethyl)-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
l'-{(6-chloro-l-(2-pyridin-4-ylethyl)-lH-indol-3-yl)carbonyl}-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-(6-chloro-l-Cpyridin-4-ylmethyl^lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-{(6-chloro-l-(2-oxo-2-pyridin-2-ylethyl)-lH-indol-3-yl)carbonyl}-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-{(6-chloro-l-[2-(5-methyl-2-phenyl-1,3-oxazol-4-yl)-2-oxoethyl]-lH-indol-3-yl)carbonyl}-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-[(6-chloro-1-(pyridin-2-ylmethyl)-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

2-{6-chloro-3-[(5-fluoro-3-oxo-l'H,3H-spiro[2-benzofuran-l,4'-piperidin]-1'-yl)carbonyl]-lH-indol-3-yl}N,N-dimethylacetamide;

r-[(6-chloro-l-[2-(dimethylamino)ethyl]-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-[(6-chloro-l-(pyridin-3-ylmethyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-[(6-chloro-l-(pyrazin-2-ylmethyl)-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-[(6-chloro-l-(pyrimidin-5-ylmethyl)-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

3-{6-chloro-3-[(5-fluoro-3-oxo-l'H,3H-spiro[2-benzofuran-l,4'-piperidin]-1'-yl)carbonyl]-lH-indol-1-yl}propanenitrile;

carbonyl]-1H-indol-1-yl)acetate;

l'-{(6-chloro-l-(2-morpholin-4-yl-2-oxoethyl)-lH-indol-3-yl)carbonyl}-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-(l-[4-benzylmorpholin-2-yl)methyl]-6-chloro-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-(6-chloro-l-[5-methylisoxazol-3-yl)methyl]-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

l'-(6-chloro-l-[2-(4-methylpiperazin-1-yl)-2-oxoethyl]-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-[(6-chloro-l-(pyridin-2-ylmethyl)-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;

r-[(6-chloro-l-[5-cyclopropyl-2-methyl-1,3-oxazol-4-yl)methyl]-lH-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4'-piperidin]-3-one;
l’-({6-chloro-1-[(1-methyl-1H-imidazol-5-yl)methyl]-1H-indol-3-yl}carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one;  
1’-({6-chloro-1-[(3-methylisoxazol-5-yl)methyl]-1H-indol-3-yl}carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one;  
1’-({6-chloro-1-[(1,5-dimethyl-1H-pyrazol-3-yl)methyl]-1H-indol-3-yl}carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one;  
1’-({6-chloro-1-[(3,5-dimethylisoxazol-4-yl)methyl]-1H-indol-3-yl}carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one;  
1’-({6-chloro-1-[(2,5-dimethyl-1,3-oxazol-4-yl)methyl]-1H-indol-3-yl}carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one;  
1’-({6-chloro-1-[(3-fluoro-oxetan-3-yl)methyl]-1H-indol-3-yl}carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one;  
1’-({6-chloro-1-[(3-fluoro-oxetan-3-yl)methyl]-1H-indol-3-yl}carbonyl)-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one;  
l’-[(6-chloro-1-[(methoxymethyl)-cyclopropyl]-1H-indol-3-yl)carbonyl]-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one;  
l’-[(6-chloro-1-[(methoxymethyl)-cyclopropyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one;  
1’-[(6-chloro-1-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4’-piperidin]-yl)carbonyl]-1H-indol-1-yl]methyl)cyclopropyl]acetonitrile;  
1’-[(6-chloro-1-[(3-oxo-1H,3H-spiro[2-benzofuran-1,4’-piperidin]-yl)carbonyl]-1H-indol-1-yl]methyl)cyclopropyl]acetonitrile;  
l’-[(6-chloro-1-[(maorolin-2-ylmethyl)-1H-indol-3-yl]carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one dihydrochloride;  
l’-[(6-chloro-1-[(morhoxin-2-ylmethyl)-1H-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one hydrochloride;  
tl’-[(6-chloro-1-[(trahydro-2H-pyran-4-yl)ethyl]-1H-indol-3-yl]carbonyl)-5-fluoro-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one hydrochloride;  
tert-butyl 2-([6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4’-piperidin]-yl)carbonyl]-1H-indol-1-yl]methyl)morpholine-4-carboxylate;  
tert-butyl 2-([6-chloro-3-[(3-oxo-1H,3H-spiro[2-benzofuran-1,4’-piperidin]-yl)carbonyl]-1H-indol-1-yl]methyl)morpholine-4-carboxylate;  
l’-[(6-chloro-1-[(morhoxin-2-ylmethyl)-1H-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one dihydrochloride;  
l’-[(6-chloro-1-[(morphism-2-ylmethyl)-1H-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-1,4’-piperidin]-3-one hydrochloride;  
2’-([6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4’-piperidin]-yl)carbonyl]-1H-indol-1-yl]acetamide;
2-\{6-chloro-3-[(5-fluoro-3-oxo-1H,3H-spiro[2-benzofuran-1,4′-piperidin]-r-yl)carbonyl]-1H-indol-1-yl\}-N-methylacetamide;
 l'-\{[6-chloro-l-(2-oxo-2-piperazin-1-ylethyl)-1H-indol-3-yl]carbonyl\}-5-fluoro-3H-spiro[2-benzofuran-1,4′-piperidin]-3-one;
 l'-\{[l-(3,5-difluorobenzyl)-1H-indol-3-yl]carbonyl\}-5-fluoro-3H-spiro[2-benzofuran-1,4′-piperidin]-3-one;
 N,N-diethyl-2-\{3-(3-oxo-1H,3H-spiro[2-benzofuran-1,4′-piperidin]-r-yl)carbonyl\}-1H-indol-1-yl\}-N,N-dimethylacetamide.

16. The compounds of formula (I-d) according to any one of claims 1 to 4:

\[ R^1 \text{ to } R^5 \text{ are as defined in any one of claims 1 to 4.} \]

17. The compounds of formula (I-d) according to claim 16, wherein

R^1 \text{ is } H, \text{ or is C}_{i-6}-\text{alkyl optionally substituted by CN, or is aryl, 5 or 6 membered heteroaryl or sulfonylaryl which are optionally substituted by one or more B, or is -(CH}_2_\text{)}_m-R^a \text{ where } R^a \text{ is: OR}^1, \text{ CN, NR}R^1, \]
C₅₋₆-cycloalkyl, 3 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
or is -(CH₂)ₙ-(CO)-R² wherein R² is:
Ci₋₆-alkoxy,
NR岑⁻,
4 to 7 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
or R¹ and R³ together with the indole ring to which they are attached form a 5 or 6 membered heterocycloalkyl which can be substituted by C(O)O-C₁₋₆-alkyl
or C₁₋₆-alkyl;
R² is one or more of H, halo, C₁₋₆-alkyl, C₁₋₆-alkoxy;
R³ is H,
or is C₁₋₆-alkyl,
or is -(CO)-RC wherein RC is C₁₋₆-alkyl, or -(CH₂VNR R⁵; 
R⁴ is one or more of H, halo, or C₁₋₆-alkoxy optionally substituted by OH, or two R⁴ may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;
R⁵ is H or aryl;
B is halo, NH₂, Q-o-alkyl optionally substituted by CN or C₁₋₆-alkoxy, C₁₋₆-alkoxy, Ci-e-haloalkoxy, C₅₋₆-cycloalkyl, -C(O)O-C₁₋₆-alkyl, -(CRⁿRᵣ)ₙ-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:
halo, Ci₋₆-alkyl optionally substituted by CN or halo, C₁₋₆-alkoxy;
R¹ and Rᵣ are H, C₁₋₆-alkyl, Ci₋₆-alkyl-NRᵐRᵣ, -(CO)O-C₁₋₆-alkyl, -C(O)-NRᵐRᵣ, -C(O)-
C₁₋₆-alkyl, -S(O)₂-C₁₋₆-alkyl, -S(O)₂⁻NRᵐRᵣ or OH;
Rᵣ and Rᵣ are H or C₁₋₆-alkyl;
m is 1 to 6;
n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

18. The compounds of formula (I-d) according to claim 16 or 17, wherein said compounds are selected from the group consisting of:
-r-[(1-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-l,4'-piperidine];
r-[(1-benzyl-2-methyl-lH-indol-3-yl)carbonyl]-6-chloro-3H-spiro[2-benzofuran-l,4'-piperidine];
r-[(2-methyl-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
r-[(1-benzoyl-2-methyl-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-
piperidine];
r-[(2-methyl-1-(phenylsulfonyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-
benzofuran-1,4'-piperidine];
r-[(1-benzyl-1H-indol-3-yl)carbonyl]-3H-spiro[2-
benzofuran-1,4'-piperidine];
r-[(6-chloro-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];
N,N-dimethyl-2-[3-(1H ,3H-spiro[2-benzofuran-1,4'-piperidine]-r-
ylcarbonyl)-1H-indol-1-yl]ethanamine;
2-methyl-1-[3-(rH,3H-spiro[2-benzofuran-1,4'-piperidine]-l'-ylcarbonyl)-1H-
indol-2-yl]butan-1-one;
[6-chloro-3-(1H ,3H-spiro[2-benzofuran-1,4'-piperidine]-r-
ylcarbonyl)-1H-indol-
1-yl]acetonitrile;
l'-
[6-chloro-1-(3-fluorobenzoyl)-1H-indol-3-yl]carbonyl]-3H-spiro[2-
benzofuran-1,4'-piperidine];

l'-lf 6-chloro-1-(3-fluorobenzoyl)-1H-indol-S-yl]carbonylJ-SH-spiroP-
benzofuran-1,4'-piperidine];

5  l'-
[6-chloro-1-(3,5-difluorobenzoyl)-1H-indol-3-yl]carbonyl]-3H-spiro[2-
benzofuran-1,4'-piperidine];

r-lfo-chloro-1-Cl^-difluorobenzoy^-1H-indol-S-yl]carbonylJ-SH-spirofl-
benzofuran-1,4'-piperidine];

l'-
[6-chloro-1-(3,5-difluorobenzyl)-1H-indol-3-yl]carbonyl]-3H-spiro[2-
benzofuran-1,4'-piperidine];

r-lfo-chloro-1-CS-fluorobenzy^-1H-indol-S-yl]carbonylJ-SH-spirofl-benzofuran-
1,4'-piperidine];

l'-
[6-chloro-1-(2-oxo-2-piperidin-1-ylethyl)-1H-indol-3-yl]carbonyl]-3H-
spiro[2-benzofuran-1,4'-piperidine];

l'-
[6-chloro-1-(2-morpholin-4-yl-2-oxoethyl)-1H-indol-3-yl]carbonyl]-3H-
spiro[2-benzofuran-1,4'-piperidine];

2-[6-chloro-3-(1 H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-
indol- 1-yl]-N,N-dimethylacetamide;

2-[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4'-piperidin]-l'-ylcarbonyl)-lH-
indol- 1-yl]-N,N-diethylacetamide;

tert-butyl [6-chloro-3-(1 H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-
indol- 1-yl] acetate;

25  l'-
[6-chloro-1-(3,5-difluorophenyl)-1H-indol-3-yl]carbonyl]-3H-spiro[2-
benzofuran-1,4'-piperidine];

l'-
[6-chloro-1-(3-fluorophenyl)-1H-indol-3-yl]carbonyl]-3H-spiro[2-
benzofuran-1,4'-piperidine];

2-[6-chloro-3-(1 H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-
indol- 1-yl]-l-(2-fluorophenyl)ethanone;

r-l[(6-chloro-1-pyridin-2-yl-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-
piperidine];

r-[(6-chloro-1-(pyridin-4-ylmethyl)-1H-indol-3-yl]carbonyl]-3H-spiro[2-
benzofuran-1,4'-piperidine];

35  2-[6-chloro-3-(1 H ,3H-spiro[2-benzofuran-1,4'-piperidin]-r-ylcarbonyl)-lH-
indol- 1-yl]-l-pyridin-2-y lethanone;
r-[(6-chloro-1-(pyridin-3-ylmethyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

r-lf 6-chloro-l-Cpyridin-l-ylmethyl^-lH-indol-S-yllcarbonylj-SH-spirofl-benzofuran-1,4'-piperidine];

5 1'-(6-chloro-1-(pyrazin-2-ylmethyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

r-[(6-chloro-1-(pyrimidin-5-ylmethyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

2-[6-chloro-3-(l H ,3H-spiro[2-benzofuran-1,4'-piperidine]-r-ylcarbonyl)-IH-indol-1-yl]-N,N-dimethylethanamine;

l'-(6-chloro-1-(2-oxo-2-piperazin-l-ylethyl)-IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

l'-(1-[4-benzylmorpholin-2-yl)methyl]-6-chloro-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

15 r-d 6-chloro-l-FCS-methylisoxazol-S-y^methyy-lH-indol-SylJcarbonyl^-SH-spiro[2-benzofuran-1,4'-piperidine];

l'-(6-chloro-l-[2-(4-methylpiperazin-l-yl)-2-oxoethyl]-IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

4-(l'-(6-chloro-1-[(2-cyclopropyl-4-methylcyclopenta-1,4-dien-3-yl)viny]-2',3'-dihydrospiro[cyclohexane-1,r-indene] ];

l'-(6-chloro-1-[(1-methyl-1H-imidazol-5-yl)methyl]-IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

l'-(6-chloro-l-[(3-methylisoxazol-5-yl)methyl]-lH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

l'-(6-chloro-l-[(1,5-dimethyl-1H-pyrazol-3-yl)methyl]-IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

l'-(6-chloro-l-[(3,5-dimethylisoxazol-4-yl)methyl]-IH-indol-3-yl}carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

l'-(6-chloro-l-[(2,5-dimethyl-1,3-oxazol-4-yl)methyl]-IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

30 3H-spiro[2-benzofuran-1,4'-piperidine];

l'-(6-chloro-l-[(3-fluorooxetan-3-yl)methyl]-IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

1'-[(6-chloro-1-[1-(methoxymethyl)-cyclopropyl] methyl ]-IH-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4'-piperidine];

(l-[6-chloro-3-(1 H ,3H-spiro[2-benzofuran-1,4'-piperidine]-r-ylcarbonyl)-IH-indol-l-yl)methyljycyclopropy^acetonitrile;
l’-([6-chloro-1-[2-(tetrahydro-2H-pyran-4-yl)ethyl]-1H-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-1,4’-piperidine];
tert-butyl 2-([6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidine]-1’-ylcarbonyl]-1H-indol-1-yl)methyl)morpholine-4-carboxylate;
r-([6-chloro-1-(morpholin-2-ylmethyl)-1H-indol-3-yl]carbonyl)-3H-spiro[2-benzofuran-1,4’-piperidine] hydrochloride;
2-[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4’-piperidine]-1’-ylcarbonyl]-1H-indol-1-yl]-N-[2-(dimethylamino)ethyl] acetamide;
2-[6-chloro-5-methyl-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidine]-r-ylcarbonyl]-1H-indol-1-yl]-N,N-dimethylacetamide;
2-[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4’-piperidine]-1’-ylcarbonyl)]-1H-indol-1-yl)acetamide;
2-[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4’-piperidine]-1’-ylcarbonyl)]-1H-indol-1-yl)ethanamine;
2-[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4’-piperidine]-1’-ylcarbonyl)]-1H-indol-1-yl)N-methylacetamide;
N-(2-aminoethyl)-2-[6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidine]-r-ylcarbonyl)]-1H-indol-1-yl)acetamide;
2-[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4’-piperidine]-1’-ylcarbonyl)]-1H-indol-1-yl)ethanamine;
2-[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4’-piperidine]-1’-ylcarbonyl)]-1H-indol-1-yl)N-methylaceta
2-[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4’-piperidine]-1’-ylcarbonyl)]-1H-indol-1-yl)N-methylacetamide;
l’-([6-chloro-1-(2-morpholin-4-yl)ethyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];
l’-([6-chloro-1-(3-morpholin-4-yl)propyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];
l’-([6-chloro-1-(oxiran-2-ylmethyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];
2-[6-chloro-3-(rH,3H-spiro[2-benzofuran-1,4’-piperidine]-1’-ylcarbonyl)]-1H-indol-1-yl)ethanol;
l’-([6-chloro-1-((2-methylpyridin-4-yl)methyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];
r-([6-chloro-1-((3S)-piperidin-3-ylmethyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];
2-[6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidine]-r-ylcarbonyl)]-1H-indol-1-yl)N-hydroxyethanamine;
l’-[(6-chloro-1-(tetrahydro-2H-pyran-4-ylmethyl)-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];
	r-[(6-chloro-1-[1-methylpyrrolidin-3-yl)methyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];

5  r-[(6-chloro-1-[(3S)-1-methylpiperidin-3-yl)methyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];

l’-[(6-chloro-1-[(2S)-pyrrolidin-2-ylmethyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];

r-[(6-chloro-2-methyl-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];

l’-[(2,3,4-tetrahydropyrazino[1,2-a]indol-10-ylcarbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine] hydrochloride;

10  r-[(2-methyl-1,2,3,4-tetrahydropyrazino[1,2-a]indol-10-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];

l’-[(6-chloro-2-methyl-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];

N-[2-(6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidine]-r-yl)carbonyl)-1H-indol-1-yl]ethylacetamide;

15  N-[2-(6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidine]-r-yl)carbonyl)-1H-indol-1-yl]ethyl methanesulfonamide;

N-[2-(6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidine]-r-yl)carbonyl)-1H-indol-1-yl]ethyl-N-methylacetamide;

20  N-[2-(6-chloro-3-(1H,3H-spiro[2-benzofuran-1,4’-piperidine]-r-yl)carbonyl)-1H-indol-1-yl]ethyl-N-methylmethanesulfonamide;

r-[(6-chloro-1-[(2S)-1-methylpyrrolidin-2-yl)methyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine];

and

r-[(6-chloro-1-[(2R)-1-methylpyrrolidin-2-yl)methyl]-1H-indol-3-yl)carbonyl]-3H-spiro[2-benzofuran-1,4’-piperidine].

19. The compounds of formula (I-e) according to any one of claims 1 to 4:
wherein $R_1$, $R_2$, $R_3$, $R_4$ and $R_6$ are as defined in any one of claims 1 to 4.

20. The compounds of formula (I-e) according to claim 19, wherein

$R_1$ is $H$, or is $C_{1-6}$-alkyl optionally substituted by CN,

or is sulfonylaryl,

or is $-(CH_2)_n-R^a$ wherein $R^a$ is:

$OR^1$,

$CN$,

$NR^1R^2$,

$C_{3-6}$-cycloalkyl, 3 to 6 membered-heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more $B$,

or is $-(CH_2)_n-(CO)-R^b$, wherein $R^b$ is:

$C_{1-6}$-alkoxy,

$NR^1R^2$,

5 or 6 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more $B$;

$R_2$ is one or more of $H$, halo, $C_{1-6}$-alkyl;

$R_3$ is $H$, or is $C_{1-6}$-alkyl,

or is $-(CO)-R^c$, wherein $R^c$ is:

$C_{1-6}$-alkyl

$-(CH_2)_n-NR^1R^2$,

$R_4$ is one or more of $H$, halo, or $C_{1-6}$-alkoxy optionally substituted by OH, or two $R_4$ may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;

$R_6$ is $H$;
B is halo, CN, Ci-6-alkyl optionally substituted by CN or C1-6-alkoxy, C1-6-alkoxy, C1-6-haloalkoxy, C1-6-cycloalkyl, -(O)O-C1-6-alkyl, -(CRmRn)-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:
halo, Ci-6-alkyl optionally substituted by CN or halo, C1-6-alkoxy;
R1 and R1 are H, C1-6-alkyl, Q-e-alkyl-NR11, -C(O)-C1-6-alkyl, -S(O)2-C1-6-alkyl or OH;
R11 and R1V are H or C1-6-alkyl;
m is 1 to 6;
n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

21. The compounds of formula (I-e) according to claim 19 or 20, wherein
R1 is H or,
-(CH2)m-Ra wherein Ra is 5 or 6 membered-heterocycloalkyl, aryl, or 5 or 6
membered heteroaryl,
-(CH2)m-NR'R or,
-(CH2)m-(CO)-Rb wherein Rb is 5 or 6 membered-heterocycloalkyl;
R2 is one or more of H or halo;
R3 is H,
R4 is one or more of H or halo;
R6 is H;
R1 and R1 are C1-6-alkyl;
m is 1 to 6;
n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

22. The compounds of formula (I-e) according to any one of claims 19 to 21, wherein said compounds are selected from the group consisting of:
r-[(6-chloro-lH-indol-3-yl)carbonyl]spiro[1-benzofuran-3,4'-piperidine];
r-[(1-benzyl-2-methyl-lH-indol-3-yl)carbonyl]spiro[1-benzofuran-3,4'-
piperidine];
r-[(6-chloro-5-fluoro-lH-indol-3-yl)carbonyl]spiro[1-benzofuran-3,4'-
piperidine];
r-[(6-chloro-1H-indol-3-yl)carbonyl]spiro[1-benzofuran-3,4'-
piperidine].
2- [6-chloro-3- (1H-spiro[1-benzofuran-3,4'-piperidin]-ylcarbonyl)-1H-indol-1-yl] -N,N-dimethylethan amine;
   r-Ir 6-chloro-l-Cl-pyrrolidin-l-ylethy^H-indol-S-yllcarbonylJspirofl-
benzofuran-3,4'-piperidine] ;
5 3- [6-chloro-3- (1H-spiro[1-benzofuran-3,4'-piperidin]-ylcarbonyl)-1H-indol-1-yl] -N,N-dimethylpropan- l-amine;
   l'-{[6-chloro-l-(2-morpholin-4-ylethyl)-1H-indol-3-yl]carbonyl}spiro[1-
benzofuran-3,4'-piperidine] ;
   2- [6-chloro-3- (1H-spiro[1-benzofuran-3,4'-piperidin]-ylcarbonyl)-1H-indol-1-yl] -N,N-diethylethanamine;
   r-{(6-chloro-l-[2-(H-pyrrol-l-yl)ethyl]-lH-indol-3-yl]carbonyl}spiro[1-
benzofuran-3,4'-piperidine]; and
   r-{[6-chloro-l-(2-oxo-2-piperidin-l-ylethyl)-1H-indol-3-yl]carbonyl}spiro[1-
benzofuran-3,4'-piperidine] .

23. The compounds of formula (I-f) according to any one of claims 1 to 4:

   \[ \begin{align*}
   &\text{R}^1, \text{R}^2, \text{R}^3, \text{and} \text{R}^4 \text{ are as defined in any one of claims 1 to 4.}
   \end{align*} \]

24. The compounds of formula (I-f) according to claim 23, wherein

   \[ \begin{align*}
   &\text{R}^1 \text{ is } \text{H}, \text{ or is } \text{C}_{i-g} \text{-alkyl optionally substituted by CN,} \text{ or is sulfonylaryl,} \text{ or is } -(\text{CH}_2)_m \text{-R}^a \text{ wherein R}^a \text{ is:} \\
   &\text{OR,} \text{ CN,}
   \end{align*} \]
NR\(\text{R}^1\),
C\(\text{3-6}\)-cycloalkyl, 3 to 6 membered-heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
or is -(CH\(_2\))\(_n\)-(CO)-R\(^b\), wherein R\(^b\) is:
\(\text{C}^{-6}\)-alkoxy,
NR\(\text{R}^1\),
6 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B;

R\(^2\) is one or more of H, halo, C\(_{1-6}\)-alkyl;
R\(^3\) is H,
or is C\(_{-6}\)-alkyl,
or is -(CO)-R\(^c\), wherein R\(^c\) is:
C\(_{-6}\)-alkyl
-(CH\(_2\))\(_n\)-NR\(\text{R}^1\),
R\(^4\) is one or more of H, halo, or C\(_{1-6}\)-alkoxy optionally substituted by OH, or two R\(^4\) may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;

B is halo, CN, C\(_{-6}\)-alkyl optionally substituted by CN or C\(_{1-6}\)-alkoxy, C\(_{1-6}\)-alkoxy, C\(_{1-6}\)-haloalkoxy, C\(_{3-6}\)-cycloalkyl, -C(O)O-C\(_{1-6}\)-alkyl, -(CR\(^m\)R\(^b\))\(_n\)-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:
halo, C\(_{-6}\)-alkyl optionally substituted by CN or halo, C\(_{1-6}\)-alkoxy;
R\(^1\) and R\(^{\prime}w\) are H, C\(_{1-6}\)-alkyl, C\(_{-6}\)-alkyl-NR\(_m\)R\(^{\prime}w\), -C(O)-C\(_{1-6}\)-alkyl, -S(O)\(_2\)-C\(_{-6}\)-alkyl or OH;
R\(^{\prime}w\) and R\(^w\) are H or C\(_{1-6}\)-alkyl;
m is 1 to 6;
n is 0 to 4;
as well as pharmaceutically acceptable salts thereof.

25. The compounds of formula (I-f) according to claim 23 or 24, wherein
R\(^1\) is H;
R\(^2\) is one or more of H or halo;
R\(^3\) is H;
R\(^4\) is one or more of H or halo;
as well as pharmaceutically acceptable salts thereof.

26. The compounds of formula (I-f) according to any one of claims 23 to 25, wherein said compounds are selected from the group consisting of
5-bromo-\(r\)-(lH-indol-3-ylcarbonyl)spiro[indole-3,4'-piperidin]-2(lH)-one; and
5-bromo-\(r\)-[(6-chloro-lH-indol-3-yl)carbonyl]spiro[indole-3,4'-piperidin]-2(lH)-one.

27. The compounds of formula (I-g) according to any one of claims 1 to 4:

\[
\begin{align*}
\text{R}^1 & \quad \text{R}^2 \\
\text{R}^3 & \quad \text{R}^4 \\
\text{R}^1 & \quad \text{R}^3 \\
\text{R}^2 & \quad \text{R}^4
\end{align*}
\]

wherein \(\text{R}^1\), \(\text{R}^2\), \(\text{R}^3\), and \(\text{R}^4\) are as defined in any one of claims 1 to 4.

28. The compounds of formula (I-g) according to claim 27, wherein
\(\text{R}^1\) is
\(\text{H}\),
or is \(\text{Ci-6-alkyl}\) optionally substituted by CN,
or is sulfonaryl,
or is \(-(\text{CH}_2)_m\)-\(\text{R}^a\) wherein \(\text{R}^a\) is:
\(\text{OR}^1\),
\(\text{CN}\),
\(\text{NR}^1\text{R}^2\),
\(\text{C}_3\text{-6-cycloalkyl}\), 3 to 6 membered-heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,
or is \(-(\text{CH}_2)_n\)-(CO)-\(\text{R}^b\) wherein \(\text{R}^b\) is:
\(\text{Ci-6-alkoxy}\),
\(\text{NR}^1\text{R}^2\),
6 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B;
\(\text{R}^2\) is
one or more of H, halo, \(\text{Ci-6-alkyl}\);
\(\text{R}^3\) is
\(\text{H}\),
or is \(\text{Ci-6-alkyl}\),
or is -(CO)-R c, wherein R c is:

\[ \text{Ci}_6\text{-alkyl} \]

\[ -(\text{CH}_2)_n\text{-NR}_1\text{R}_1\text{1}, \]

R 4 is one or more of H, halo, or C 1-6-alkoxy optionally substituted by OH, or two R 4 may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;

R 3 and R 4 v are H, C 1-6-alkyl, C 1-6-alkyl-NR 4, -C(O)-C 1-6-alkyl, -S(O) 2-Ci 6-alkyl or OH;

m is 1 to 6;

n is 0 to 4;

as well as pharmaceutically acceptable salts thereof.

29. The compounds of formula (I-g) according to claim 27 or 28, wherein

R 1 is H or,

-(CH 2)_m-R a wherein R a is aryl;

R 2 is one or more of H or halo;

R 3 is H, or is C 1-6-alkyl;

R 4 is H;

as well as pharmaceutically acceptable salts thereof.

30. The compounds of formula (I-g) according to any one of claims 27 to 29, wherein said compounds are selected from the group consisting of

(SS,RR)-r-[1-benzyl-2-methyl-1H-indol-3-yl]carbonyl]-3',5'-dimethyl-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;

(RS,SR)-r-[1-benzyl-2-methyl-1H-indol-3-yl]carbonyl]-3',5'-dimethyl-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one;

(RS,SR)-r-[1-benzyl-2-methyl-1H-indol-3-yl]carbonyl]-3',5'-dimethyl-3H-spiro[2-benzofuran-l,4'-piperidin]-3-one; and
The compounds of formula (I-h) according to any one of claims 1 to 4:

wherein \( R^1, R^2, R^3, R^4, R^8 \), and X are as defined in any one of claims 1 to 4.

The compounds of formula (I-h) according to claim 31, wherein

\( R^1 \) is \( H \),

or is \( \text{Ci-}6\)-alkyl optionally substituted by CN,

or is sulfonylaryl,

or is \(-(\text{CH}_2)_m\)-R\(^a\) wherein R\(^a\) is:

\( \text{OR}^1 \),

\( \text{CN} \),

\( \text{NR}^1\text{R}^1 \),

\( \text{C}_3\text{e-cycloalkyl} \), 3 to 6 membered-heterocycloalkyl, aryl or 5 or 6 membered heteroaryl which are optionally substituted by one or more B,

or is \(-(\text{CH}_2)_n\)-(CO)-R\(^b\) wherein R\(^b\) is:

\( \text{Ci-}6\)-alkoxy,

\( \text{NR}^1\text{R}^1 \),

6 membered-heterocycloalkyl, aryl, or 5 or 6 membered heteroaryl which are optionally substituted by one or more B;

\( R^2 \) is one or more of H, halo, \( \text{Ci-}1\text{-}6\)-alkyl;

\( R^3 \) is \( H \),

or is \( \text{Ci-}6\)-alkyl,

or is \(-(\text{CO})\)-R\(^c\) wherein R\(^c\) is:

\( \text{Ci-}6\)-alkyl
-(CH\textsubscript{2})\textsubscript{n}-NR\textsubscript{u},

R\textsuperscript{4} is one or more of H, halo, or C\textsubscript{1-6}-alkoxy optionally substituted by OH, or two R\textsuperscript{4} may form an oxo or dioxo bridge together with the phenyl ring to which they are attached;

5 \hspace{0.5cm} R\textsuperscript{8} is H or C\textsubscript{1-6}-alkyl;

X is CH\textsubscript{2} or C=O;

B is halo, CN, Q-o-alkyl optionally substituted by CN or C\textsubscript{1-6}-alkoxy, C\textsubscript{1-6}-haloalkoxy, C\textsubscript{3-6}-cycloalkyl, -C(0)\textsubscript{0}-C\textsubscript{1-6}-alkyl, -(CR\textsubscript{m}R\textsubscript{1})\textsubscript{n}-phenyl, wherein the phenyl is optionally substituted by one or more substituent(s) selected from the group consisting of:

- halo, C\textsubscript{1-6}-alkyl optionally substituted by CN or halo, C\textsubscript{1-6}-alkoxy;

- R\textsuperscript{1} and R\textsuperscript{u} are H, C\textsubscript{1-6}-alkyl, C\textsubscript{1-6}-alkyl-NR\textsubscript{m}R\textsubscript{1}, -(C(O)\textsubscript{0})\textsubscript{0}-C\textsubscript{1-6}-alkyl, -(S(O))\textsubscript{2}-C\textsubscript{1-6}-alkyl or OH;

- R\textsuperscript{u} and R\textsuperscript{v} are H or C\textsubscript{1-6}-alkyl;

- m is 1 to 6;

- n is Oto 4;

as well as pharmaceutically acceptable salts thereof.

33. The compounds of formula (I-h) according to claim 31 or 32, wherein

- R\textsuperscript{1} is H,

20 or is -(CH\textsubscript{2})\textsubscript{n}-(CO)-R\textsuperscript{b}, wherein R\textsuperscript{b} is NR\textsubscript{R}\textsubscript{u};

- R\textsuperscript{2} is one or more of H or halo;

- R\textsuperscript{3} is H;

- R\textsuperscript{4} is H;

- R\textsuperscript{8} is H or C\textsubscript{1-6}-alkyl;

25 - X is CH\textsubscript{2} or C=O;

- R\textsuperscript{1} and R\textsuperscript{u} are H or C\textsubscript{1-6}-alkyl;

as well as pharmaceutically acceptable salts thereof.

34. The compounds of formula (I-h) according to any one of claims 31 to 33, wherein said compounds are selected from the group consisting of

- r-[(6-chloro-1H-indol-3-yl)carbonyl]spiro[isoindole-1,4'-piperidine]-3(2H)-one;

- 1'-[(6-chloro-1H-indol-3-yl)carbonyl]-2-methyl-2,3-dihydrospiro[isoindole-1,4'-piperidine] ;

- 1'-[(6-chloro-1H-indol-3-yl)carbonyl]-2,3-dihydrospiro[isoindole-1,4'-piperidine] ; and
2-\{6-chloro-3-[(3-oxo-2,3-dihydro-1\,H\text{-}spiro[indole-1,4\text{-}piperidin]-l\text{-}y1}carbonyl\} -IH-indol- 1-yl\}-N-methylacetamide.

35. A process for the preparation of the compounds of formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h) according to any one of claims 1 to 34 comprising the step of reacting a compound of formula (II):

\[
\begin{array}{c}
\text{HO} \\
\text{R}^2 \\
\text{R}^3 \\
\text{R}^1 \\
\text{II}
\end{array}
\]

with a compound of formula A-H in order to obtain the compound of formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h) wherein A, R^1, R^2 and R^3 are as defined in any one of claims 1 to 34.

36. A process for the preparation of the compounds of formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h) according to any one of claims 1 to 34 comprising the step of reacting a compound of formula (III):

\[
\begin{array}{c}
\text{A} \\
\text{R}^2 \\
\text{R}^3 \\
\text{H} \\
\text{III}
\end{array}
\]

with an electrophile compound of formula R^1-Y in order to obtain the compound of formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h), wherein R^1, R^2 and R^3 are as defined in any one of claims 1 to 34 and Y is halo.

37. A process for the preparation of the compounds of formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h) according to any one of claims 1 to 34 comprising the step of reacting a compound of formula (IV):
with an amine of formula HNR\textsubscript{1}R\textsubscript{1} in order to obtain the compound of formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h), wherein R\textsubscript{1}, R\textsubscript{2}, R\textsubscript{3}, R\textsubscript{4} and R\textsubscript{v} are as defined in any one of claims 1 to 34.

38. A compound formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h) obtainable by a process according to any one of claims 35 to 37.

39. A compound of formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h) according to any one of claims 1 to 34 for a use in the prevention or treatment of dysmenorrhea, hypertension, chronic heart failure, inappropriate secretion of vasopressin, liver cirrhosis, nephrotic syndrome, obsessive compulsive disorder, anxious and depressive disorders.

40. A pharmaceutical composition comprising one or more compounds of formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h) according to any one of claims 1 to 34 and a pharmaceutically acceptable carrier.

41. A pharmaceutical composition according to claim 40, wherein it is useful against dysmenorrhea, hypertension, chronic heart failure, inappropriate secretion of vasopressin, liver cirrhosis, nephrotic syndrome, obsessive compulsive disorder, anxious and depressive disorders.

42. Use of a compound of formula (I), (I-a), (I-b), (I-c), (I-d), (I-e), (I-f), (I-g) or (I-h) according to any one of claims 1 to 34 for the preparation of a medicament.

43. Use according to claim 42, wherein the medicament is useful against dysmenorrhea, hypertension, chronic heart failure, inappropriate secretion of vasopressin, liver cirrhosis, nephrotic syndrome, obsessive compulsive disorder, anxious and depressive disorders.

44. The invention as described hereinabove.

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INTERNATIONAL SEARCH REPORT

A. CLASSIFICATION OF SUBJECT MATTER

INV. C07D401/06 C07D471/10 C07D491/10 A61K31/454 A61P9/00

According to International Patent Classification (IPC) or to both national classification and IPC.

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
C07D A61K A61P

Documentation searched other than minimum documentation is the extent to which such documents are included in the fields searched.

Electronic data base consulted during the international search (name of data base and where practical, search terms used)
EPO-Internal, CHER ABS Data, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

<table>
<thead>
<tr>
<th>Category</th>
<th>Citation of document, with indication, where appropriate of the relevant passages</th>
<th>Relevant to claim No</th>
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<tr>
<td>A</td>
<td>US 5 686 624 A (DI MALTA ET AL) 11 November 1997 (1997-11-11) example 73</td>
<td>1,35-37, 40</td>
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</table>

Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents

'A' document defining the general state of the art which is not considered to be of particular relevance
'E' earlier document but published on or after the priority filing date
'L' document which may throw doubts on novelty claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
'O' document referring to an oral disclosure, use, exhibition or other means
'P' document published prior to the international filing date but later than the priority date claimed

T' later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
'X' document of particular relevance, the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
'Y' document of particular relevance, the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
'&' member of the same patent family

Date of the actual completion of the international search: 27 September 2006

Date of mailing of the international search report: 05/10/2006

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Authorized officer

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<tr>
<th>Patent document cited in search report</th>
<th>Publication date</th>
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<tr>
<td>US 5686624</td>
<td>11-11-1997</td>
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