

Piperidine and Piperazine Derivatives as Autotaxin Inhibitors

Description

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Technical field

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The present invention relates to piperidine and piperazine derivatives as autotaxin inhibitors and the use of such compounds for the treatment and/or prophylaxis of physiological and/or pathophysiological conditions, which are caused, mediated and/or propagated by increased lysophosphatic acid levels and/or the activation of autotaxin, in particular of different cancers.

Prior art

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Autotaxin (ATX) is the enzyme apparently responsible for increased lysophosphatidic acid (LPA) levels in ascites and plasma of ovarian cancer patients (Xu et al., *Clinical Cancer Research* 1995, 1: 1223; Xu et al., *Biochem. J.* 1995, 309: 933), because it transforms lysophatidylcholine (LPC) to LPA (Tokumura et al., *J. Biol. Chem.* 2002, 277: 39436; Umezue-Gozo et al., *J. Biol. Chem.* 2002, 158: 227).

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LPA is an intracellular lipid mediator, which influences a multiplicity of biologic and biochemical processes such as smooth muscle contraction, thrombocyte aggregation and apoptosis (Tigyi et al., *Prog. Lipid Res.* 2003, 42: 498; Mills et al., *Nat. Rev. Cancer* 2003, 3: 582; Lynch et al. *Prost. Lipid Med.* 2001, 64: 33).

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Furthermore, LPA is found in increased concentrations in plasma and ascites fluid of ovarian cancer patients of early and late phase.

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LPA has been shown to promote tumor cell proliferation, survival, migration and invasion into neighboring tissues, which can result in the formation of metastases (Xu et al., *Clinical Cancer Research* 1995, 1: 1223; Xu et al., *Biochem. J.* 1995, 309: 933). These biological and pathobiological processes are switched on through the activation of G-protein coupled receptors by LPA (Contos et al., *Mol. Pharm.* 2000, 58: 1188).

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Increased levels of LPA, altered receptor expression and altered responses to LPA may contribute to the initiation, progression or outcome of ovarian cancer.

Furthermore, LPA is potentially also involved in prostate, breast, melanoma, head and neck, bowel and thyroid cancers.

For all these reasons in the course of treating tumor patients it is desirable to lower the LPA level. This can be achieved through the inhibition of enzymes which are involved in LPA biosynthesis, such as ATX (Sano et al., J. Biol. Chem. 2002, 277: 21197; Aoki et al., J. Biol. Chem. 2003, 277: 48737).

ATX belongs to the family of nucleotide pyrophosphatases and phosphodiesterases (Goding et al., Immunol. Rev. 1998, 161: 11). It represents an important starting point for anti-tumor therapy (Mills et al. Nat. Rev. Cancer 2003, 3: 582; Goto et al. J. Cell. Biochem. 2004, 92: 1115), since it is increasingly expressed in tumors and effects tumor cell proliferation and invasion into neighboring tissues both of which can lead to the formation of metastases (Nam et al. 2000, Oncogene, Vol. 19 Seite 241). In addition, in the course of angiogenesis ATX together with other anti-angiogenetic factors brings about blood vessel formation (Nam et al. Cancer Res. 2001, 61: 6938). Angiogenesis is an important process during tumor growth as it secures supply of the tumor with nutrients. Therefore, the inhibition of angiogenesis is an important starting point of cancer and tumor therapy, by means of which the tumor is to be starved (Folkman, Nature Reviews Drug Discovery 2007, 6: 273-286).

Mutagenesis studies suggest an essential function of the PDE domain of ATX for LPA generation. Though this particular PDE domain shares little homology with other known PDEs, it is considered to be druggable by NCEs.

No severe adverse effects are expected for the inhibition of ATX as LPA involved in wound healing in this context is produced by another pathway.

Since ATX is a relatively novel target, the amount of preclinical data on protein production, *in vitro* and *in vivo* assays is rather limited. No target-dependent cell model has been described but LPA itself is an excellent biomarker to follow ATX inhibition *in vitro* and *in vivo*. Neither structural information nor reference compounds are available.

Compounds that are capable of inhibiting ATX are described in Peng et al. (Bioorganic & Medicinal Chemistry Letters 2007, 17: 1634-1640). The there described compounds represent lipid analogues, which structurally share no similarities with the compounds of the present invention.

Further prior art documents are as follows:

WO 2002/102380 describes monocyclic or bicyclic carbocycles and heterocycles as factor Xa inhibitors. The patent application does not mention the inhibition of autotaxin.

5 WO 2003/097615 is directed to treatment of fibroproliferative diseases, such as diabetic neuropathy, involving identifying a non-peptide small molecule, selectively binding to a transforming growth factor beta kinase receptor and administering the molecule to subjects. The patent application does not mention the inhibition of autotaxin.

10 US 2003/0139431 relates to the use of quinazoline- and quinolino-guanidine derivatives for treating urge incontinence, pain, memory disorders, endocrine disorders, psychotic behaviour, diabetes, hypertension and gastrointestinal disorders. The patent application does not mention the inhibition of autotaxin.

15 WO 2004/099192 describes heterocycle substituted carboxylic acids which can be used in the treatment of metabolic disorders. The patent application does not mention the inhibition of autotaxin.

20 WO 2005/003100 deals with the use of quinazoline derivatives for the treatment of tubulin inhibitor mediated diseases, such as cancer, autoimmune diseases, autoimmune lymphoproliferative syndrome, inflammation and viral infections. The patent application does not mention the inhibition of autotaxin.

25 WO 2006/062972 discloses heterocyclic compounds that function as selective inhibitors of serine protease enzymes of the coagulation cascade and can be used for the treatment of arterial cardiovascular thromboembolic disorders, thromboembolic disorders, unstable angina and acute coronary syndrome. The patent application does not mention the inhibition of autotaxin.

30 WO 2006/072828 is directed to heteroaromatic quinoline compounds that serve as PDE inhibitors, in particular PDE10 inhibitors. These compounds can be used for the treatment of central nervous system disorders, such as psychotic disorders, anxiety disorders, movement disorders, mood disorders and neurodegenerative disorders. The patent application does not mention the inhibition of autotaxin.

35 WO 2006/074147 describes 4-aryl-amino-quinazoline as caspase-3 cascade activators that can be used for the treatment of cancer, autoimmune diseases,

autoimmune lymphoproliferative syndrome, synovial cell hyperplasia, inflammation and viral infections. The patent application does not mention the inhibition of autotaxin.

5 WO 2006/108107 deals with diarylamine derivatives that are steroid hormone nuclear receptor modulators and can be used for the treatment of hypokalemia, hypertension, congestive heart failure, renal failure, arteriosclerosis and obesity. The patent application does not mention the inhibition of autotaxin.

10 WO 2007/030582 relates to alkyl amine compounds as potassium channel 1 function inhibitors that are useful for the treatment of arrhythmia, atrial fibrillation, atrial flutter, supraventricular arrhythmias, gastrointestinal disorders, esophagitis and asthma. The patent application does not mention the inhibition of autotaxin.

15 WO 2007/076034 describes fused bicyclic arene compounds that function as hepatitis C virus replication inhibitors and can be used for the treatment hepatitis C or other viral infections. The patent application does not mention the inhibition of autotaxin.

20 WO 2007/110868 discloses novel heterocyclic compounds that exhibit a dopamine receptor, preferably D4 receptor, antagonistic activity and/or a PDE5 inhibitory activity. These compounds can be used for the treatment of decreased libido, orgasm disorder and erectile dysfunction. The patent application does not mention the inhibition of autotaxin.

25 WO 2008/060621 is directed to aminopyrrolidines as chemokine receptor antagonists. The patent application does not mention the inhibition of autotaxin.

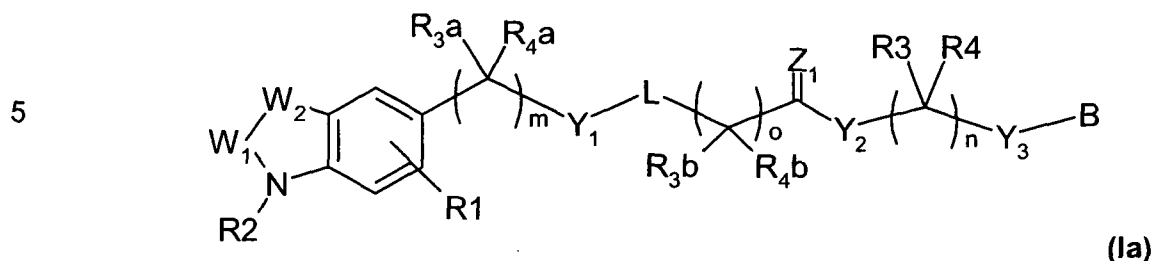
30 WO 2008/091580 relates to fungicidal amides and methods for controlling plant diseases caused by a fungal pathogen. The patent application does not mention the inhibition of autotaxin.

The citation of any reference in this application is not an admission that the reference is relevant prior art to this application.

Description of the invention

The present invention has the object to provide novel autotaxin inhibitors.

The object of the present invention has surprisingly been solved in one aspect by providing compounds according to formula (Ia)



wherein:

- 10 W_1, W_2 together independently form “-N=N-, -C(O)-O-, -C(O)-S-, -C(O)-N(R5)-, -C(S)-N(R5a)-, -C(O)-C(R6)(R7)-, -N=C[N(R8)(R9)]-”;
- Y_1 is independently selected from the group consisting of “-C(O)-, -C(S)-, -S(O)₂-, -N(R10)-C(O)-, -C(O)-N(R11)-, -OC(O)-, single bond”;
- 15 Y_2 is independently selected from the group consisting of “-C(R12)(R13)-, -O-, -N(R14)-, -C(O)-, -C(O)-NH-, single bond”;
- Y_3 is independently selected from the group consisting of “-O-, -C(O)-, single bond”;
- 20 Z_1 is independently selected from the group consisting of “O, S, N(R15)”;
- L is independently selected from the group consisting of the group consisting of:
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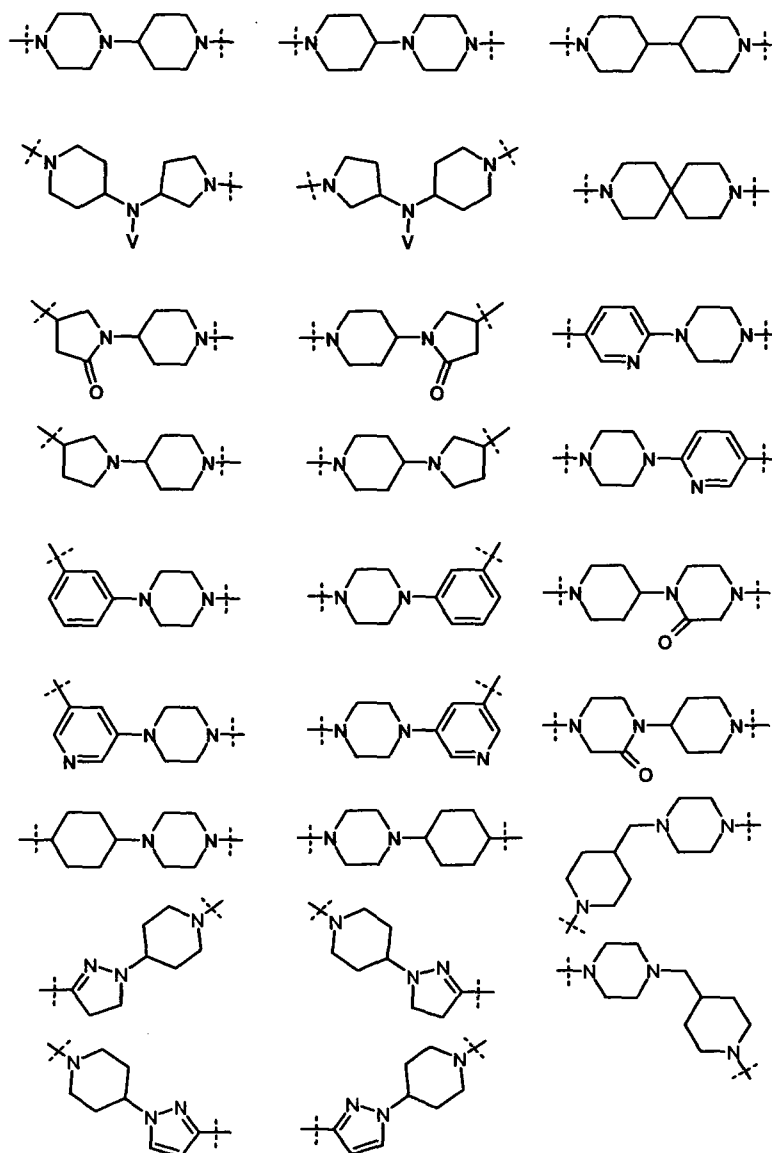
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is independently selected from the group consisting of "cycloalkyl, heterocyclyl, aryl, heteroaryl", wherein "cycloalkyl, heterocyclyl, aryl, heteroaryl" can be independently substituted with one or more identical or different substituents selected from the group consisting of: "(i) „hydrogen, alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl, halogen, -F, -Cl, -Br, -I, -CN, -CF₃, -SF₃, -N₃, -NH₂, -NHX₁, -NX₂X₃, -NO₂, -OH, =O, -OCF₃, -SCF₃, -OCHF₂, -SCHF₂, -SH, -O-SO₃H, -OP(O)(OH)₂, -CHO, -COOH, -C(O)NH₂, -SO₃H, -P(O)(OH)₂, -C(O)-X₄, -C(O)O-X₅, -C(O)NH-X₆, -C(O)NX₇X₈, -O-X₉, -O(-X₁₀-O)_a-H (a = 1, 2, 3, 4, 5), -O(-X₁₁-

$O)_b-X12$ ($b = 1, 2, 3, 4, 5$), $-OC(O)-X13$, $-OC(O)-O-X14$, $-OC(O)-NHX15$, $-O-C(O)-NX16X17$, $-OP(O)(OX18)(OX19)$, $-OSi(X20)(X21)(X22)$, $-OS(O_2)-X23$, $-NHC(O)-NH_2$, $-NHC(O)-X24$, $-NX25C(O)-X26$, $-NH-C(O)-O-X27$, $-NH-C(O)-NH-X28$, $-NH-C(O)-NX29X30$, $-NX31-C(O)-O-X32$, $-NX33-C(O)-NH-X34$, $-NX35-C(O)-NX36X37$, $-NHS(O_2)-X38$, $-NX39S(O_2)-X40$, $-S-X41$, $-S(O)-X42$, $-S(O_2)-X43$, $-S(O_2)NH-X44$, $-S(O_2)NX45X46$, $-S(O_2)O-X47$, $-P(O)(OX48)(OX49)$, $-Si(X50)(X51)(X52)$, $-C(NH)-NH_2$, $-C(NX53)-NH_2$, $-C(NH)-NHX54$, $-C(NH)-NX55X56$, $-C(NX57)-NHX58$, $-C(NX59)-NX60X61$, $-NH-C(O)-NH-O-X62$, $-NH-C(O)-NX63-O-X64$, $-NX65-C(O)-NX66-O-X67$, $-N(-C(O)-NH-O-X68)_2$, $-N(-C(O)-NX69-O-X70)_2$, $-N(-C(O)-NH-O-X71)(-C(O)-NX72-O-X73)$, $-C(S)-X74$, $-C(S)-O-X75$, $-C(S)-NH-X76$, $-C(S)-NX77X78$, $-C(O)-NH-O-X79$, $-C(O)-NX80-O-X81$, $-C(S)-NH-O-X82$, $-C(S)-NX83-O-X84$, $-C(O)-NH-NH-X85$, $-C(O)-NH-NX86X87$, $-C(O)-NX88-NX89X90$, $-C(S)-NH-NH-X91$, $-C(S)-NH-NX92X93$, $-C(S)-NX94-NX95X96$, $-C(O)-C(O)-O-X97$, $-C(O)-C(O)-NH_2$, $-C(O)-C(O)-NHX98$, $-C(O)-C(O)-NX99X100$, $-C(S)-C(O)-O-X101$, $-C(O)-C(S)-O-X102$, $-C(S)-C(S)-O-X103$, $-C(S)-C(O)-NH_2$, $-C(S)-C(O)-NHX104$, $-C(S)-C(O)-NX105X106$, $-C(S)-C(S)-NH_2$, $-C(S)-C(S)-NHX107$, $-C(S)-C(S)-NX108X109$, $-C(O)-C(S)-NH_2$, $-C(O)-C(S)-NHX110$, $-C(O)-C(S)-NX111X112$ ";

wherein X1, X2, X3, X4, X5, X6, X7, X8, X9, X10, X11, X12, X13, X14, X15, X16, X17, X18, X19, X20, X21, X22, X23, X24, X25, X26, X27, X28, X29, X30, X31, X32, X33, X34, X35, X36, X37, X38, X39, X40, X41, X42, X43, X44, X45, X46, X47, X48, X49, X50, X51, X52, X53, X54, X55, X56, X57, X58, X59, X60, X61, X62, X63, X64, X65, X66, X67, X68, X69, X70, X71, X72, X73, X74, X75, X76, X77, X78, X79, X80, X81, X82, X83, X84, X85, X86, X87, X88, X89, X90, X91, X92, X93, X94, X95, X96, X97, X98, X99, X100, X101, X102, X103, X104, X105, X106, X107, X108, X109, X110, X111, X112 are independently from each other selected from the group consisting of: „alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl“ and wherein alternatively X7, X8 and/or X16, X17 and/or X29, X30 and/or

X36, X37 and/or X45, X46 and/or X55, X56 and/or X60, X61 and/or X77, X78 and/or X86, X87 and/or X89, X90 and/or X92, X93 and/or X95, X96 and/or X99, X100 and/or X105, X106 and/or X108, X109 and/or X111, X112 respectively together can also form

5 „heterocycl“;

wherein optionally above substituents of substituents group (i) can in turn independently from each other be substituted with one or more identical or different substituents V;

10 R1, R2, R3, R3a, R3b, R4, R4a, R4b, R5, R5a, R6, R7, R8, R9, R10, R11, R12, R13, R14, R15 are independently from each other selected from the group consisting of: “V”; alternatively, R3a and R4a, R3b and R4b as well as R3 and R4 together can form “cycloalkyl” or “heterocycl“;

15 V is independently selected from the group consisting of: “(i)
 „hydrogen, alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocycl, heterocyclalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl, halogen, -F, -Cl, -Br, -I, -CN, -CF₃, -SF₃, -N₃, -NH₂, -NHA1, -NA2A3, -NO₂, -OH, =O, -OCF₃, -SCF₃, -OCHF₂, -SCHF₂, -SH, -O-SO₃H, -OP(O)(OH)₂, -CHO, -COOH, -C(O)NH₂, -SO₃H, -P(O)(OH)₂, -C(O)-A4, -C(O)O-A5, -C(O)NH-A6, -C(O)NA7A8, -O-A9, -O(-A10-O)_a-H (a = 1, 2, 3, 4, 5), -O(-A11-O)_b-A12 (b = 1, 2, 3, 4, 5), -OC(O)-A13, -OC(O)-O-A14, -OC(O)-NHA15, -O-C(O)-NA16A17, -OP(O)(OA18)(OA19), -OSi(A20)(A21)(A22), -OS(O₂)-A23, -NHC(O)-NH₂, -NHC(O)-A24, -NA25C(O)-A26, -NH-C(O)-O-A27, -NH-C(O)-NH-A28, -NH-C(O)-NA29A30, -NA31-C(O)-O-A32, -NA33-C(O)-NH-A34, -NA35-C(O)-NA36A37, -NHS(O₂)-A38, -NA39S(O₂)-A40, -S-A41, -S(O)-A42, -S(O₂)-A43, -S(O₂)NH-A44, -S(O₂)NA45A46, -S(O₂)O-A47, -P(O)(OA48)(OA49), -Si(A50)(A51)(A52), -C(NH)-NH₂, -C(NA53)-NH₂, -C(NH)-NHA54, -C(NH)-NA55A56, -C(NA57)-NHA58, -C(NA59)-NA60A61, -NH-C(O)-NH-O-A62, -NH-C(O)-NA63-O-A64, -NA65-C(O)-NA66-O-A67, -N(-C(O)-NH-O-A68)₂, -N(-C(O)-NA69-O-A70)₂, -N(-C(O)-NH-O-A71)(-C(O)-NA72-O-A73), -C(S)-A74, -C(S)-O-A75, -C(S)-NH-A76, -

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C(S)-NA77A78, -C(O)-NH-O-A79, -C(O)-NA80-O-A81, -C(S)-NH-O-A82, -C(S)-NA83-O-A84, -C(O)-NH-NH-A85, -C(O)-NH-NA86A87, -C(O)-NA88-NA89A90, -C(S)-NH-NH-A91, -C(S)-NH-NA92A93, -C(S)-NA94-NA95A96, -C(O)-C(O)-O-A97, -C(O)-C(O)-NH₂, -C(O)-C(O)-NHA98, -C(O)-C(O)-NA99A100, -C(S)-C(O)-O-A101, -C(O)-C(S)-O-A102, -C(S)-C(S)-O-A103, -C(S)-C(O)-NH₂, -C(S)-C(O)-NHA104, -C(S)-C(O)-NA105A106, -C(S)-C(S)-NH₂, -C(S)-C(S)-NHA107, -C(S)-C(S)-NA108A109, -C(O)-C(S)-NH₂, -C(O)-C(S)-NHA110, -C(O)-C(S)-NA111A112";

wherein A1, A2, A3, A4, A5, A6, A7, A8, A9, A10, A11, A12, A13, A14, A15, A16, A17, A18, A19, A20, A21, A22, A23, A24, A25, A26, A27, A28, A29, A30, A31, A32, A33, A34, A35, A36, A37, A38, A39, A40, A41, A42, A43, A44, A45, A46, A47, A48, A49, A50, A51, A52, A53, A54, A55, A56, A57, A58, A59, A60, A61, A62, A63, A64, A65, A66, A67, A68, A69, A70, A71, A72, A73, A74, A75, A76, A77, A78, A79, A80, A81, A82, A83, A84, A85, A86, A87, A88, A89, A90, A91, A92, A93, A94, A95, A96, A97, A98, A99, A100, A101, A102, A103, A104, A105, A106, A107, A108, A109, A110, A111, A112 are independently from each other selected from the group consisting of: „alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl“ and wherein alternatively A7, A8 and/or A16, A17 and/or A29, A30 and/or A36, A37 and/or A45, A46 and/or A55, A56 and/or A60, A61 and/or A77, A78 and/or A86, A87 and/or A89, A90 and/or A92, A93 and/or A95, A96 and/or A99, A100 and/or A105, A106 and/or A108, A109 and/or A111, A112 respectively together can also form „heterocyclyl“;

wherein optionally above substituents of substituents group (i) can in turn independently from each other be substituted with one or more identical or different substituents V;

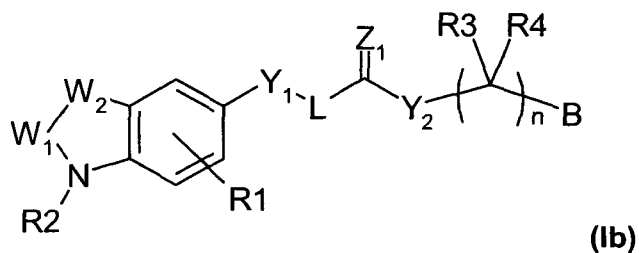
m is independently 0, 1, 2, or 3;

n is independently 0, 1, 2, or 3;

o is independently 0, 1, 2, or 3;

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

The object of the present invention has surprisingly been solved in one aspect by providing compounds according to formula (Ib)



wherein:

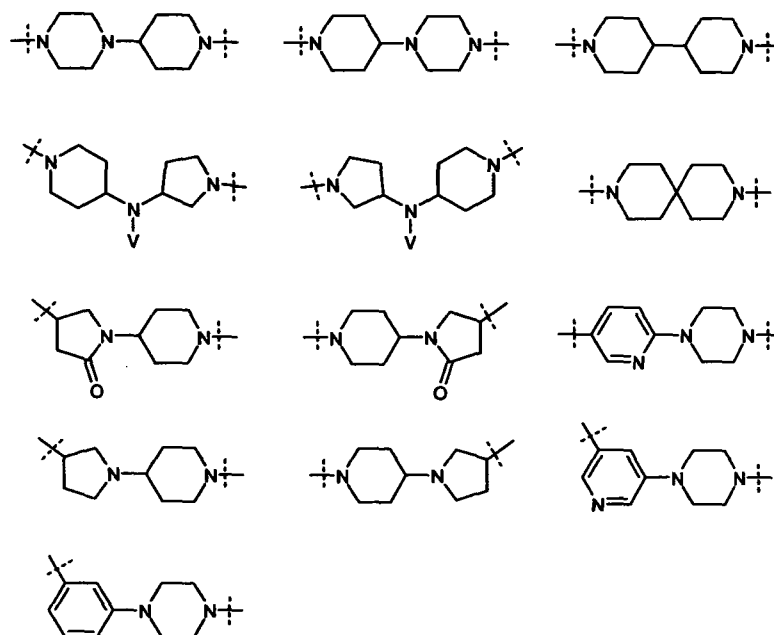
W_1, W_2 together independently form “ $-N=N-$, $-C(O)-O-$, $-C(O)-S-$, $-C(O)-N(R_5)-$, $-C(O)-C(R_6)(R_7)-$, $-N=C[N(R_8)(R_9)]-$ ”;

Y_1 is independently selected from the group consisting of “ $-C(O)-$, $-C(S)-$, $-N(R_{10})-C(O)-$, $-C(O)-N(R_{11})-$, $-OC(O)-$, single bond”;

Y_2 is independently selected from the group consisting of “ $-C(R_{12})(R_{13})-$, $-O-$, $-N(R_{14})-$, $-C(O)-NH-$, single bond”;

Z_1 is independently selected from the group consisting of “O, S, N(R₁₅)”;

L is independently selected from the group consisting of the group consisting of:



B is independently selected from the group consisting of "cycloalkyl, heterocyclyl, aryl, heteroaryl", wherein "cycloalkyl, heterocyclyl, aryl, heteroaryl" can be independently substituted with one or more identical or different substituents selected from the group consisting of: "(i) „hydrogen, alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl, halogen, -F, -Cl, -Br, -I, -CN, -CF₃, -N₃, -NH₂, -NHX₁, -NX₂X₃, -NO₂, -OH, -OCF₃, -SCF₃, -SH, -O-SO₃H, -OP(O)(OH)₂, -CHO, -COOH, -C(O)NH₂, -SO₃H, -P(O)(OH)₂, -C(O)-X₄, -C(O)O-X₅, -C(O)NH-X₆, -C(O)NX₇X₈, -O-X₉, -O(-X₁₀-O)_a-H (a = 1, 2, 3, 4, 5), -O(-X₁₁-O)_b-X₁₂ (b = 1, 2, 3, 4, 5), -OC(O)-X₁₃, -OC(O)-O-X₁₄, -OC(O)-NHX₁₅, -O-C(O)-NX₁₆X₁₇, -OP(O)(OX₁₈)(OX₁₉), -OSi(X₂₀)(X₂₁)(X₂₂), -OS(O₂)-X₂₃, -NHC(O)-NH₂, -NHC(O)-X₂₄, -NX₂₅C(O)-X₂₆, -NH-C(O)-O-X₂₇, -NH-C(O)-NH-X₂₈, -NH-C(O)-NX₂₉X₃₀, -NX₃₁-C(O)-O-X₃₂, -NX₃₃-C(O)-NH-X₃₄, -NX₃₅-C(O)-NX₃₆X₃₇, -NHS(O₂)-X₃₈, -NX₃₉S(O₂)-X₄₀, -S-X₄₁, -S(O)-X₄₂, -S(O₂)-X₄₃, -S(O₂)NH-X₄₄, -S(O₂)NX₄₅X₄₆, -S(O₂)O-X₄₇, -P(O)(OX₄₈)(OX₄₉), -Si(X₅₀)(X₅₁)(X₅₂), -C(NH)-NH₂, -C(NX₅₃)-NH₂, -C(NH)-NHX₅₄, -C(NH)-NX₅₅X₅₆, -C(NX₅₇)-NHX₅₈, -C(NX₅₉)-NX₆₀X₆₁, -NH-

C(O)-NH-O-X62, -NH-C(O)-NX63-O-X64, -NX65-C(O)-NX66-
 O-X67, -N(-C(O)-NH-O-X68)₂, -N(-C(O)-NX69-O-X70)₂, -N(-
 C(O)-NH-O-X71)(-C(O)-NX72-O-X73), -C(S)-X74, -C(S)-O-
 X75, -C(S)-NH-X76, -C(S)-NX77X78, -C(O)-NH-O-X79, -C(O)-
 5 NX80-O-X81, -C(S)-NH-O-X82, -C(S)-NX83-O-X84, -C(O)-
 NH-NH-X85, -C(O)-NH-NX86X87, -C(O)-NX88-NX89X90, -
 C(S)-NH-NH-X91, -C(S)-NH-NX92X93, -C(S)-NX94-NX95X96, -
 C(O)-C(O)-O-X97, -C(O)-C(O)-NH₂, -C(O)-C(O)-NHX98, -C(O)-
 C(O)-NX99X100, -C(S)-C(O)-O-X101, -C(O)-C(S)-O-X102, -
 10 C(S)-C(S)-O-X103, -C(S)-C(O)-NH₂, -C(S)-C(O)-NHX104, -
 C(S)-C(O)-NX105X106, -C(S)-C(S)-NH₂, -C(S)-C(S)-NHX107, -
 C(S)-C(S)-NX108X109, -C(O)-C(S)-NH₂, -C(O)-C(S)-NHX110, -
 C(O)-C(S)-NX111X112";

wherein X1, X2, X3, X4, X5, X6, X7, X8, X9, X10, X11, X12, X13,
 15 X14, X15, X16, X17, X18, X19, X20, X21, X22, X23, X24, X25, X26,
 X27, X28, X29, X30, X31, X32, X33, X34, X35, X36, X37, X38, X39,
 X40, X41, X42, X43, X44, X45, X46, X47, X48, X49, X50, X51, X52,
 X53, X54, X55, X56, X57, X58, X59, X60, X61, X62, X63, X64, X65,
 X66, X67, X68, X69, X70, X71, X72, X73, X74, X75, X76, X77, X78,
 20 X79, X80, X81, X82, X83, X84, X85, X86, X87, X88, X89, X90, X91,
 X92, X93, X94, X95, X96, X97, X98, X99, X100, X101, X102, X103,
 X104, X105, X106, X107, X108, X109, X110, X111, X112 are
 independently from each other selected from the group consisting of:
 „alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl,
 25 heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl“ and
 wherein alternatively X7, X8 and/or X16, X17 and/or X29, X30 and/or
 X36, X37 and/or X45, X46 and/or X55, X56 and/or X60, X61 and/or
 X77, X78 and/or X86, X87 and/or X89, X90 and/or X92, X93 and/or
 X95, X96 and/or X99, X100 and/or X105, X106 and/or X108, X109
 30 and/or X111, X112 respectively together can also form
 „heterocyclyl“;

wherein optionally above substituents of substituents group (i) can in
 turn independently from each other be substituted with one or more
 35 identical or different substituents V;

R1, R2, R3, R4, R5, R6, R7, R8, R9, R10, R11, R12, R13, R14, R15 are independently from each other selected from the group consisting of: "V";

V is independently selected from the group consisting of: "(i)
 „hydrogen, alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl, halogen, -F, -Cl, -Br, -I, -CN, -CF₃, -N₃, -NH₂, -NHA1, -NA2A3, -NO₂, -OH, -OCF₃, -SCF₃, -SH, -O-SO₃H, -OP(O)(OH)₂, -CHO, -COOH, -C(O)NH₂, -SO₃H, -P(O)(OH)₂, -C(O)-A4, -C(O)O-A5, -C(O)NH-A6, -C(O)NA7A8, -O-A9, -O(-A10-O)_a-H (a = 1, 2, 3, 4, 5), -O(-A11-O)_b-A12 (b = 1, 2, 3, 4, 5), -OC(O)-A13, -OC(O)-O-A14, -OC(O)-NHA15, -O-C(O)-NA16A17, -OP(O)(OA18)(OA19), -OSi(A20)(A21)(A22), -OS(O₂)-A23, -NHC(O)-NH₂, -NHC(O)-A24, -NA25C(O)-A26, -NH-C(O)-O-A27, -NH-C(O)-NH-A28, -NH-C(O)-NA29A30, -NA31-C(O)-O-A32, -NA33-C(O)-NH-A34, -NA35-C(O)-NA36A37, -NHS(O₂)-A38, -NA39S(O₂)-A40, -S-A41, -S(O)-A42, -S(O₂)-A43, -S(O₂)NH-A44, -S(O₂)NA45A46, -S(O₂)O-A47, -P(O)(OA48)(OA49), -Si(A50)(A51)(A52), -C(NH)-NH₂, -C(NA53)-NH₂, -C(NH)-NHA54, -C(NH)-NA55A56, -C(NA57)-NHA58, -C(NA59)-NA60A61, -NH-C(O)-NH-O-A62, -NH-C(O)-NA63-O-A64, -NA65-C(O)-NA66-O-A67, -N(-C(O)-NH-O-A68)₂, -N(-C(O)-NA69-O-A70)₂, -N(-C(O)-NH-O-A71)(-C(O)-NA72-O-A73), -C(S)-A74, -C(S)-O-A75, -C(S)-NH-A76, -C(S)-NA77A78, -C(O)-NH-O-A79, -C(O)-NA80-O-A81, -C(S)-NH-O-A82, -C(S)-NA83-O-A84, -C(O)-NH-NH-A85, -C(O)-NH-NA86A87, -C(O)-NA88-NA89A90, -C(S)-NH-NH-A91, -C(S)-NH-NA92A93, -C(S)-NA94-NA95A96, -C(O)-C(O)-O-A97, -C(O)-C(O)-NH₂, -C(O)-C(O)-NHA98, -C(O)-C(O)-NA99A100, -C(S)-C(O)-O-A101, -C(O)-C(S)-O-A102, -C(S)-C(S)-O-A103, -C(S)-C(O)-NH₂, -C(S)-C(O)-NHA104, -C(S)-C(O)-NA105A106, -C(S)-C(S)-NH₂, -C(S)-C(S)-NHA107, -C(S)-C(S)-NA108A109, -C(O)-C(S)-NH₂, -C(O)-C(S)-NHA110, -C(O)-C(S)-NA111A112";

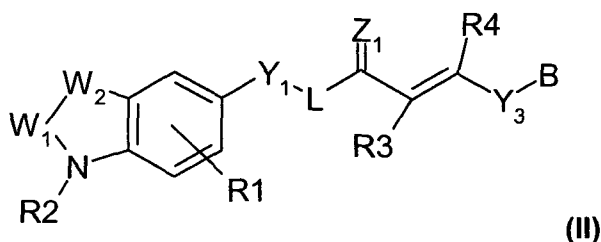
wherein A1, A2, A3, A4, A5, A6, A7, A8, A9, A10, A11, A12, A13, A14, A15, A16, A17, A18, A19, A20, A21, A22, A23, A24, A25, A26, A27, A28, A29, A30, A31, A32, A33, A34, A35, A36, A37, A38, A39, A40, A41, A42, A43, A44, A45, A46, A47, A48, A49, A50, A51, A52, A53, A54, A55, A56, A57, A58, A59, A60, A61, A62, A63, A64, A65, A66, A67, A68, A69, A70, A71, A72, A73, A74, A75, A76, A77, A78, A79, A80, A81, A82, A83, A84, A85, A86, A87, A88, A89, A90, A91, A92, A93, A94, A95, A96, A97, A98, A99, A100, A101, A102, A103, A104, A105, A106, A107, A108, A109, A110, A111, A112 are independently from each other selected from the group consisting of: „alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl“ and wherein alternatively A7, A8 and/or A16, A17 and/or A29, A30 and/or A36, A37 and/or A45, A46 and/or A55, A56 and/or A60, A61 and/or A77, A78 and/or A86, A87 and/or A89, A90 and/or A92, A93 and/or A95, A96 and/or A99, A100 and/or A105, A106 and/or A108, A109 and/or A111, A112 respectively together can also form „heterocyclyl“;

wherein optionally above substituents of substituents group (i) can in turn independently from each other be substituted with one or more identical or different substituents V;

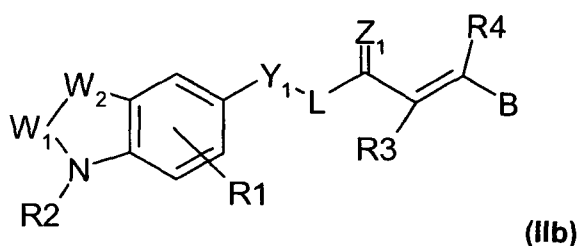
n is independently 0, 1, 2, or 3;

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

In a preferred embodiment, compounds according to formula (II) or (IIb) are provided,



15



wherein according to formulae (Ia) or (Ib) as defined supra:

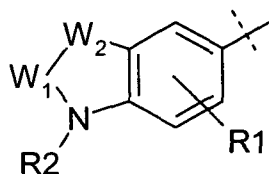
Y_2 independently is a single bond;

n independently is 2;

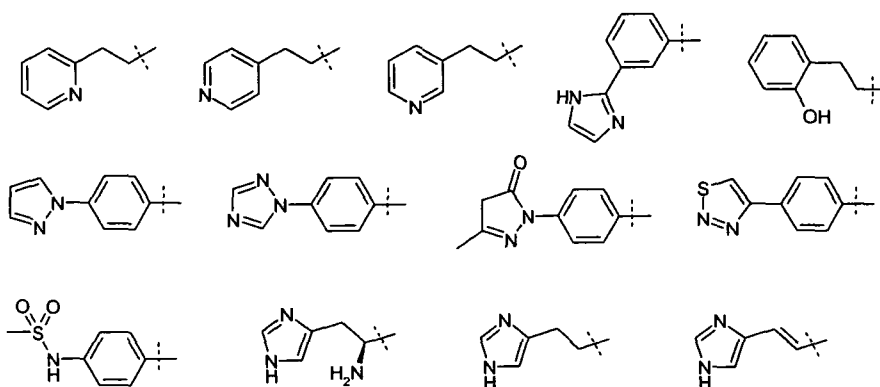
W_1 , W_2 , Y_1 , Y_3 , L , Z_1 , B , R_1 , R_2 , R_3 , R_4 have the meanings according to formulae (Ia) or (Ib) as defined supra;

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

In a preferred embodiment, compounds according to formulae (Ia), (Ib) and (II) or (IIb) as defined supra and above preferred embodiments are provided, wherein:



is independently substituted by a chemical group selected from the group consisting of:



and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

In a further preferred embodiment, compounds according to formulae (Ia), (Ib) and (II) or (IIb) as defined supra and above preferred embodiments are provided, wherein:

W₁, W₂ together independently form “-N=N-, -C(O)-O-, -C(O)-S-, -C(S)-N(R5a)-”;

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

In a further preferred embodiment, compounds according to formulae (Ia), (Ib) and (II) or (IIb) as defined supra and above preferred embodiments are provided, wherein:

Y₁ is independently selected from the group consisting of “-C(O)-, -N(R10)-C(O)-, -C(O)-N(R11)-, -OC(O)-, -S(O)₂-, single bond”;

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

In a further preferred embodiment, compounds according to formulae (Ia), (Ib) and (II) or (IIb) as defined supra and above preferred embodiments are provided, wherein:

Z₁ is independently “O”;

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

In a further preferred embodiment, compounds according to formulae (Ia), (Ib) and (II) or (IIb) as defined supra and above preferred embodiments are provided, wherein:

B is independently selected from the group consisting of “(4-chloro-phenyl)-1H-[1,2,3]triazol-4-yl, [3,3']bithiophenyl-5-yl, 1H-benzotriazol-5-yl, 1H-imidazol-4-yl, 2-(4-chloro-phenyl)-4-methyl-thiazol-5-yl, 2-(4-chloro-phenyl)-cyclopropane-yl, 2-(4-trifluoromethyl-phenyl)-thiazol-

5-yl, 2,3,5,6-tetrafluoro-4-trifluoromethyl-phenyl, 2,3-difluoro-4-trifluoromethyl-phenyl, 2,3-dihydro-benzo[1,4]dioxin-6-yl, 2,4-dichloro-phenyl, 2-chloro-phenyl, 2-fluoro-4-trifluoromethyl-phenyl, 2-fluoro-5-trifluoromethyl-phenyl, 2-methyl-2H-indazol-3-yl, 2-methyl-5-phenyl-furan-3-yl, 2-pyridin-2-yl, 3-(4-chloro-phenyl)-2-oxo-oxazolidin-5-yl, 3,4,5-trifluoro-phenyl, 3,4,5-trimethoxy-phenyl, 3,4-dichloro-phenyl, 3,4-difluoro-5-trifluoromethyl-phenyl, 3,4-dimethyl-phenyl, 3,5-bis-trifluoromethyl-phenyl, 3,5-dibromo-4-methyl-phenyl, 3,5-dibromo-phenyl, 3,5-dichloro-4-fluoro-phenyl, 3,5-dichloro-phenyl, 3,5-dimethoxy-phenyl, 3,5-dimethyl-phenyl, 3'-trifluoromethyl-biphenyl-2-yl, 3-bromo-4-trifluoromethoxy-phenyl, 3-bromo-5-chloro-phenyl, 3-bromo-5-fluoro-phenyl, 3-chloro-4,5-difluoro-phenyl, 3-chloro-4-fluoro-phenyl, 3-chloro-4-trifluoromethoxy-phenyl, 3-chloro-4-trifluoromethyl-phenyl, 3-chloro-5-fluoro-phenyl, 3-chloro-5-trifluoromethyl-phenyl, 3-chloro-phenyl, 3-fluoro-4-trifluoromethoxy-phenyl, 3-fluoro-4-trifluoromethyl-phenyl, 3-fluoro-5-trifluoromethyl-phenyl, 3-trifluoromethoxy-phenyl, 3-trifluoromethyl-phenyl, 4-(1,2,3-thiadiazol-4-yl)-phenyl, 4-(1,2,4-triazol-1-yl)-phenyl, 4-(3-methyl-5-oxo-4,5-dihydropyrazol-1-yl)-phenyl, 4'-methyl-biphenyl-2-yl, 4'-methyl-biphenyl-3-yl, 4-bromo-2,6-difluoro-phenyl, 4-bromo-2-fluoro-phenyl, 4-bromo-phenyl, 4-chloro-2-fluoro-phenyl, 4-chloro-3-fluoro-phenyl, 4-chloro-3-trifluoromethoxy-phenyl, 4-chloro-3-trifluoromethyl-phenyl, 4-chloro-phenyl, 4-cyano-phenyl, 4-difluoromethoxy-phenyl, 4-difluoromethylsulfanyl-phenyl, 4-ethyl-phenyl, 4-fluoro-3,5-dimethyl-phenyl, 4-fluoro-3-trifluoromethyl-phenyl, 4-isopropylphenyl, 4-methanesulfonyl-phenyl, 4-methoxy-3,5-dimethyl-phenyl, 4-methyl-2-(4-trifluoromethyl-phenyl)-thiazol-5-yl, 4-methyl-3-trifluoromethyl-phenyl, 4-methyl-phenyl, 4-methylsulfanyl-phenyl, 4-nitro-phenyl, 4-trifluoromethoxy-phenyl, 4'-trifluoromethyl-biphenyl-2-yl, 4-trifluoromethyl-phenyl, 4-trifluoromethylsulfanyl-phenyl, 5-bromo-benzofuran-2-yl, 5-chloro-2-fluoro-3-trifluoromethyl-phenyl, 5-chloro-2-methoxy-phenyl, 5-trifluoromethyl-1H-benzimidazole-2-yl, benzo[1,3]dioxol-5-yl, phenyl, tetrahydrofuran-2-yl";

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

5 In a further preferred embodiment, compounds according to formulae (Ia), (Ib) and (II) or (IIb) as defined supra and above preferred embodiments are provided, wherein:

R1, R2, R3, R3a, R3b, R4, R4a, R4b, R5, R5a, R6, R7, R8, R9, R10, R11, R12, R13, R14, R15 are independently from each other selected
10 from the group consisting of: "hydrogen, alkyl, methyl, ethyl, isopropyl, halogen, -F, -Br, -Cl, -CN, -CF₃, -SF₃, -OCF₃, -SCF₃, -OCHF₂, -SCHF₂, -O-alkyl, -O-methyl, -S-alkyl, -S-methyl, -NO₂, -S(O)₂-methyl"

15 and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

In a further preferred embodiment, compounds according to formulae (Ia), (Ib) and (II) or (IIb) as defined supra and above preferred embodiments are provided,
20 wherein:

V is independently selected from the group consisting of "hydrogen, alkyl, methyl, ethyl, isopropyl, halogen, -F, -Br, -Cl, -CN, -CF₃, -SF₃, -OCF₃, -SCF₃, -OCHF₂, -SCHF₂, =O, -O-alkyl, -O-methyl, -S-alkyl, -S-methyl, -NO₂, -S(O)₂-methyl";

25 and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

In a further preferred embodiment, compounds according to formulae (Ia), (Ib) and (II) or (IIb) as defined supra and above preferred embodiments are provided,
30 wherein:

m is independently 0, 1 or 2;

n is independently 0, 1 or 2;

35 o is independently 0, 1 or 2;

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

In a further preferred embodiment, compounds according to formulae (Ia), (Ib) and (II) or (IIb) as defined supra and above preferred embodiments are provided, wherein:

W_1, W_2 together independently form “-N=N-, -C(O)-O-, -C(O)-S-, -C(S)-N(R5a)-”;

Y_1 is independently selected from the group consisting of “-C(O)-, -N(R10)-C(O)-, -C(O)-N(R11)-, -OC(O)-, -S(O)₂-, single bond”;

Z_1 is independently “O”;

B is independently selected from the group consisting of “(4-chloro-phenyl)-1H-[1,2,3]triazol-4-yl, [3,3']bithiophenyl-5-yl, 1H-benzotriazol-5-yl, 1H-imidazol-4-yl, 2-(4-chloro-phenyl)-4-methyl-thiazol-5-yl, 2-(4-chloro-phenyl)-cyclopropane-yl, 2-(4-trifluoromethyl-phenyl)-thiazol-5-yl, 2,3,5,6-tetrafluoro-4-trifluoromethyl-phenyl, 2,3-difluoro-4-trifluoromethyl-phenyl, 2,3-dihydro-benzo[1,4]dioxin-6-yl, 2,4-dichloro-phenyl, 2-chloro-phenyl, 2-fluoro-4-trifluoromethyl-phenyl, 2-fluoro-5-trifluoromethyl-phenyl, 2-methyl-2H-indazol-3-yl, 2-methyl-5-phenyl-furan-3-yl, 2-pyridin-2-yl, 3-(4-chloro-phenyl)-2-oxo-oxazolidin-5-yl, 3,4,5-trifluoro-phenyl, 3,4,5-trimethoxy-phenyl, 3,4-dichloro-phenyl, 3,4-difluoro-5-trifluoromethyl-phenyl, 3,4-dimethyl-phenyl, 3,5-bis-trifluoromethyl-phenyl, 3,5-dibromo-4-methyl-phenyl, 3,5-dibromo-phenyl, 3,5-dichloro-4-fluoro-phenyl, 3,5-dichloro-phenyl, 3,5-dimethoxy-phenyl, 3,5-dimethyl-phenyl, 3'-trifluoromethyl-biphenyl-2-yl, 3-bromo-4-trifluoromethoxy-phenyl, 3-bromo-5-chloro-phenyl, 3-bromo-5-fluoro-phenyl, 3-chloro-4,5-difluoro-phenyl, 3-chloro-4-fluoro-phenyl, 3-chloro-4-trifluoromethoxy-phenyl, 3-chloro-4-trifluoromethyl-phenyl, 3-chloro-5-fluoro-phenyl, 3-chloro-5-trifluoromethyl-phenyl, 3-chloro-phenyl, 3-fluoro-4-trifluoromethoxy-phenyl, 3-fluoro-4-trifluoromethyl-phenyl, 3-fluoro-5-trifluoromethyl-phenyl, 3-trifluoromethoxy-phenyl, 3-trifluoromethyl-phenyl, 4-(1,2,3-thiadiazol-4-yl)-phenyl, 4-(1,2,4-triazol-1-yl)-phenyl,

4-(3-methyl-5-oxo-4,5-dihydropyrazol-1-yl)-phenyl, 4'-methyl-biphenyl-2-yl, 4'-methyl-biphenyl-3-yl, 4-bromo-2,6-difluoro-phenyl, 4-bromo-2-fluoro-phenyl, 4-bromo-phenyl, 4-chloro-2-fluoro-phenyl, 4-chloro-3-fluoro-phenyl, 4-chloro-3-trifluoromethoxy-phenyl, 4-chloro-3-trifluoromethyl-phenyl, 4-chloro-phenyl, 4-cyano-phenyl, 4-difluoromethoxy-phenyl, 4-difluoromethylsulfanyl-phenyl, 4-ethyl-phenyl, 4-fluoro-3,5-dimethyl-phenyl, 4-fluoro-3-trifluoromethyl-phenyl, 4-isopropylphenyl, 4-methanesulfonyl-phenyl, 4-methoxy-3,5-dimethyl-phenyl, 4-methyl-2-(4-trifluoromethyl-phenyl)-thiazol-5-yl, 4-methyl-3-trifluoromethyl-phenyl, 4-methyl-phenyl, 4-methylsulfanyl-phenyl, 4-nitro-phenyl, 4-trifluoromethoxy-phenyl, 4'-trifluoromethyl-biphenyl-2-yl, 4-trifluoromethyl-phenyl, 4-trifluoromethylsulfanyl-phenyl, 5-bromo-benzofuran-2-yl, 5-chloro-2-fluoro-3-trifluoromethyl-phenyl, 5-chloro-2-methoxy-phenyl, 5-trifluoromethyl-1H-benzoimidazole-2-yl, benzo[1,3]dioxol-5-yl, phenyl, tetrahydrofuran-2-yl";

R1, R2, R3, R3a, R3b, R4, R4a, R4b, R5, R5a, R6, R7, R8, R9, R10, R11, R12, R13, R14, R15 are independently from each other selected from the group consisting of: "hydrogen, alkyl, methyl, ethyl, isopropyl, halogen, -F, -Br, -Cl, -CN, -CF₃, -SF₃, -OCF₃, -SCF₃, -OCHF₂, -SCHF₂, -O-alkyl, -O-methyl, -S-alkyl, -S-methyl, -NO₂, -S(O)₂-methyl"

V is independently selected from the group consisting of "hydrogen, alkyl, methyl, ethyl, isopropyl, halogen, -F, -Br, -Cl, -CN, -CF₃, -SF₃, -OCF₃, -SCF₃, -OCHF₂, -SCHF₂, =O, -O-alkyl, -O-methyl, -S-alkyl, -S-methyl, -NO₂, -S(O)₂-methyl";

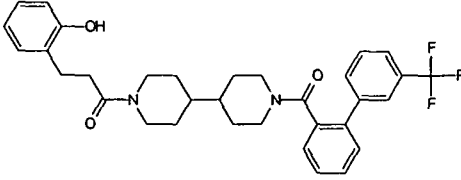
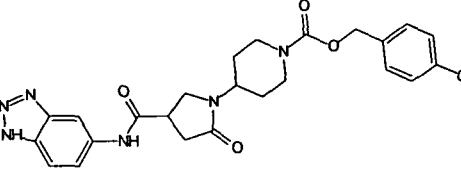
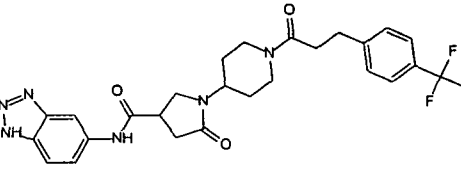
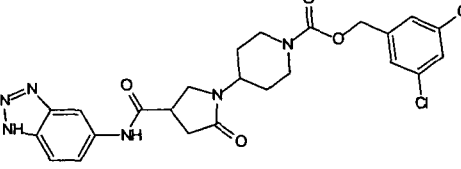
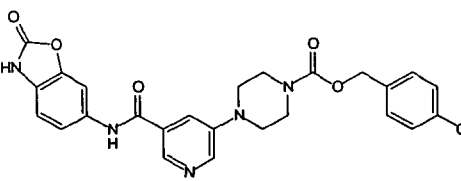
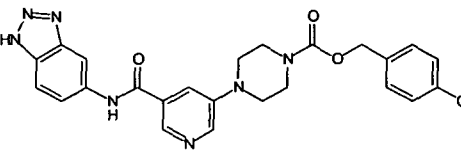
m is independently 0, 1 or 2;

n is independently 0, 1 or 2;

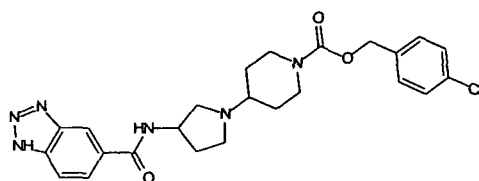
o is independently 0, 1 or 2;

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

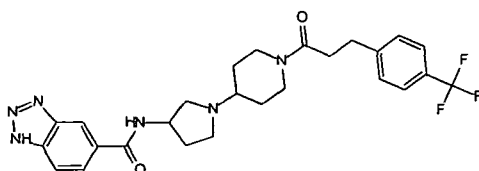
In another aspect, the object of the invention has surprisingly been solved by providing a compound selected from the group consisting of:

- 5
- Compound 1**
- 
- 3-(2-Hydroxy-phenyl)-1-[1'-(3'-trifluoromethyl-biphenyl-2-carbonyl)-[4,4']bipiperidiny-1-yl]-propan-1-one
- 10
- Compound 2**
- 
- 4-[4-(1H-Benzotriazol-5-ylcarbamoyl)-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 4-chloro-benzyl ester
- 15
- Compound 3**
- 
- 5-Oxo-1-{1-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-pyrrolidine-3-carboxylic acid (1H-benzotriazol-5-yl)-amide
- 20
- Compound 4**
- 
- 4-[4-(1H-Benzotriazol-5-ylcarbamoyl)-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester
- 25
- Compound 5**
- 
- 4-[5-(2-Oxo-2,3-dihydro-benzooxazol-6-ylcarbamoyl)-pyridin-3-yl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester
- 30
- Compound 6**
- 
- 4-[5-(1H-Benzotriazol-5-ylcarbamoyl)-pyridin-3-yl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester

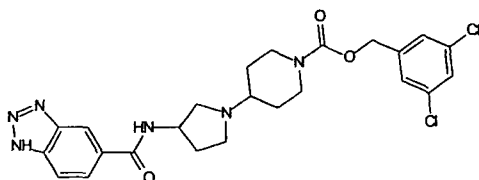
22

**Compound
7**

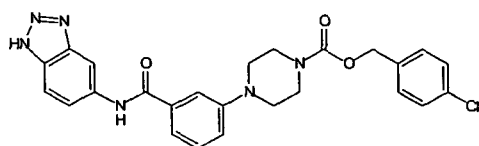
4-{3-[(1H-Benzotriazole-5-carbonyl)-amino]-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 4-chloro-benzyl ester

**Compound
8**

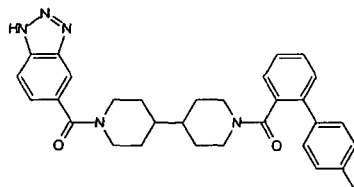
1H-Benzotriazole-5-carboxylic acid (1-{1-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-pyrrolidin-3-yl)-amide

**Compound
9**

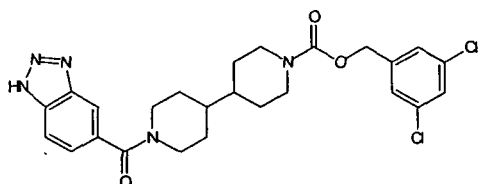
4-{3-[(1H-Benzotriazole-5-carbonyl)-amino]-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

**Compound
10**

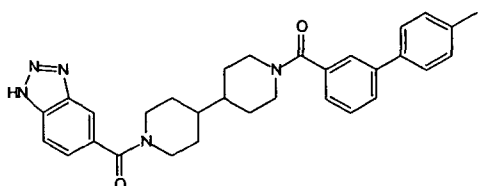
4-[3-(1H-Benzotriazol-5-ylcarbonyl)-phenyl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester

**Compound
11**

[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidiny-1-yl]-(4'-methylbiphenyl-2-yl)-methanone

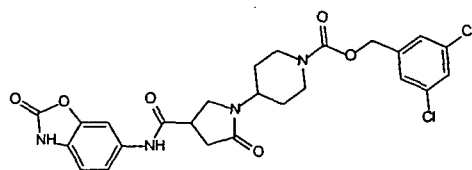
**Compound
12**

1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidiny-1-carboxylic acid 3,5-dichloro-benzyl ester

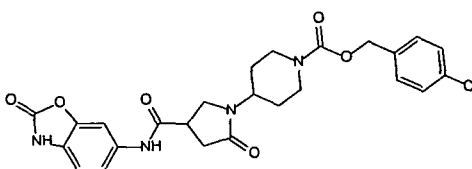
**Compound
13**

[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidiny-1-yl]-(4'-methylbiphenyl-3-yl)-methanone

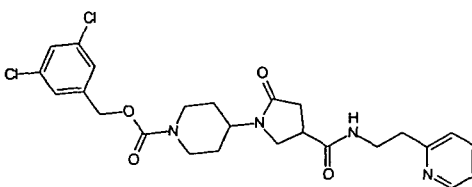
23

Compound 15

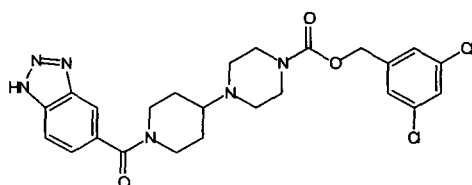
4-[2-Oxo-4-(2-oxo-2,3-dihydro-benzooxazol-6-ylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

Compound 16

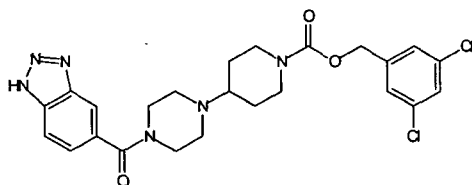
4-[2-Oxo-4-(2-oxo-2,3-dihydro-benzooxazol-6-ylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 4-chloro-benzyl ester

Compound 17

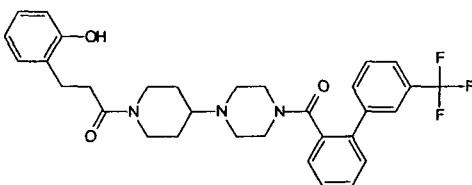
4-[2-Oxo-4-(2-pyridin-2-yl-ethylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

Compound 19

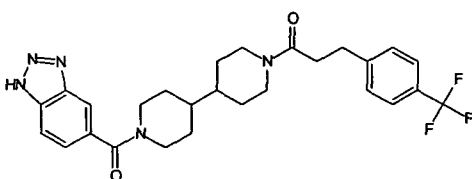
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dichloro-benzyl ester

Compound 20

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

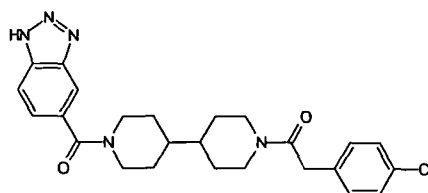
Compound 21

3-(2-Hydroxy-phenyl)-1-{4-[4-(3'-trifluoromethyl-biphenyl-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-propan-1-one

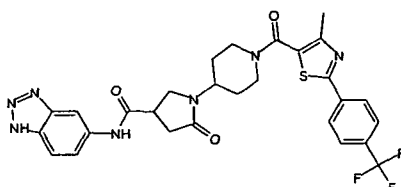
Compound 22

1-[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidinyl-1-yl]-3-(4-trifluoromethyl-phenyl)-propan-1-one

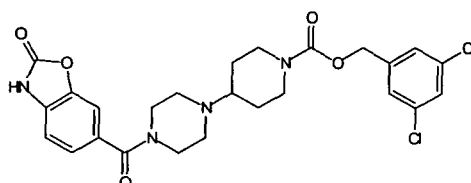
24

**Compound
23**

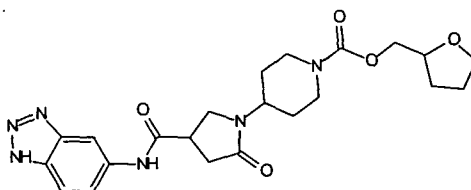
1-[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidiny-1-yl]-2-(4-chloro-phenyl)-ethanone

**Compound
24**

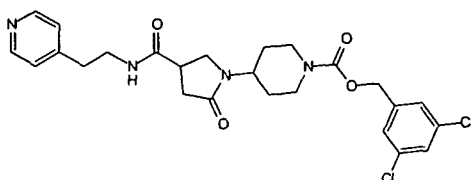
1-{1-[4-Methyl-2-(4-trifluoromethyl-phenyl)-thiazole-5-carbonyl]-piperidin-4-yl}-5-oxo-pyrrolidine-3-carboxylic acid (1H-benzotriazol-5-yl)-amide

**Compound
25**

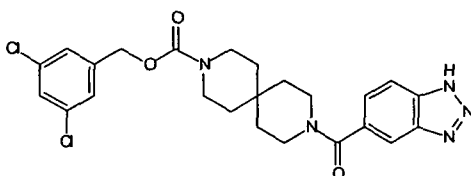
4-[4-(2-Oxo-2,3-dihydro-benzooxazole-6-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

**Compound
27**

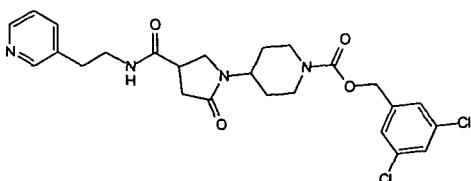
4-[4-(1H-Benzotriazol-5-ylcarbonyl)-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid tetrahydro-furan-2-ylmethyl ester

**Compound
28**

4-[2-Oxo-4-(2-pyridin-4-ylethylcarbonyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

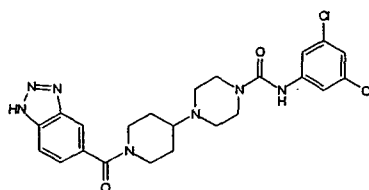
**Compound
29**

9-(1H-Benzotriazole-5-carbonyl)-3,9-diaza-spiro[5.5]undecane-3-carboxylic acid 3,5-dichloro-benzyl ester

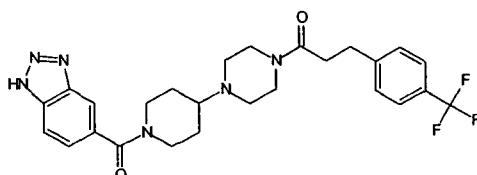
**Compound
30**

4-[2-Oxo-4-(2-pyridin-3-ylethylcarbonyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

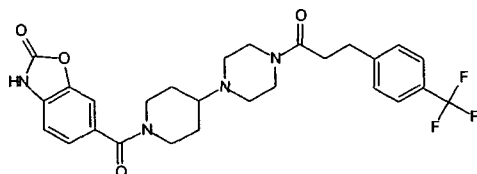
25

Compound 31

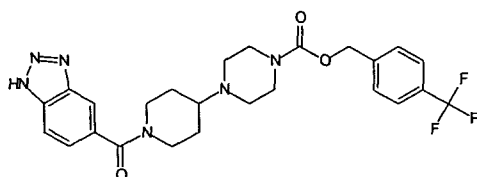
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (3,5-dichloro-phenyl)-amide

Compound 32

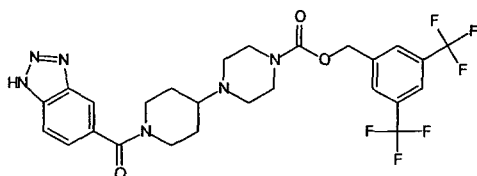
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-trifluoromethyl-phenyl)-propan-1-one

Compound 33

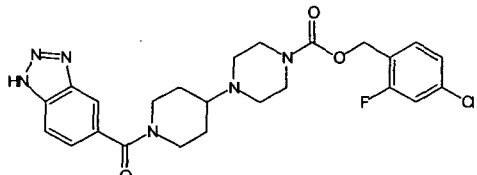
6-(4-{4-[3-(4-Trifluoromethyl-phenyl)-propionyl]-piperazin-1-yl}-piperidine-1-carbonyl)-3H-benzoxazol-2-one

Compound 34

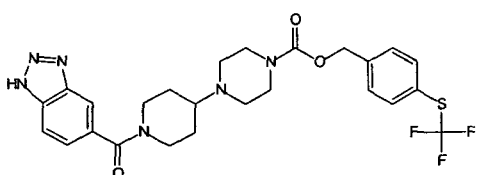
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

Compound 35

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

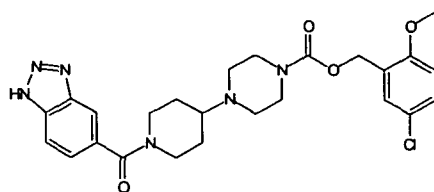
Compound 36

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester

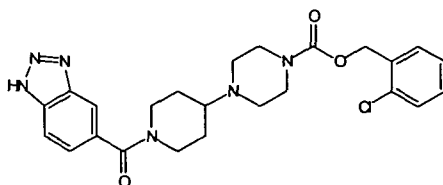
Compound 37

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzyl ester

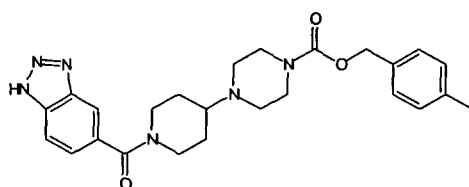
26

**Compound
38**

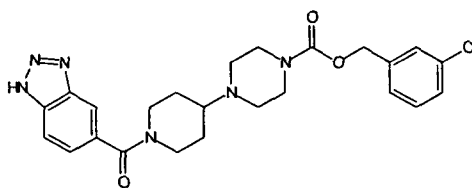
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 5-
chloro-2-methoxy-benzyl ester

**Compound
39**

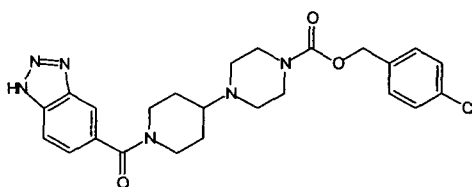
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 2-
chloro-benzyl ester

**Compound
40**

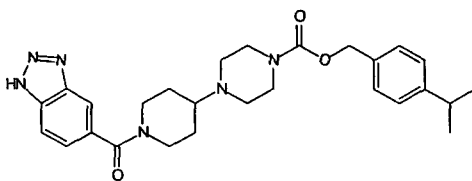
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 4-
methyl-benzyl ester

**Compound
41**

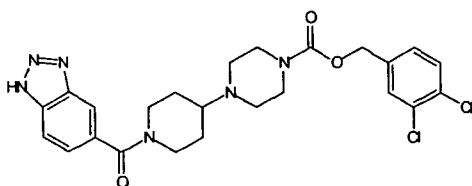
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 3-
chloro-benzyl ester

**Compound
42**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 4-
chloro-benzyl ester

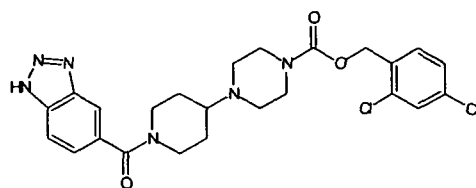
**Compound
43**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 4-
isopropyl-benzyl ester

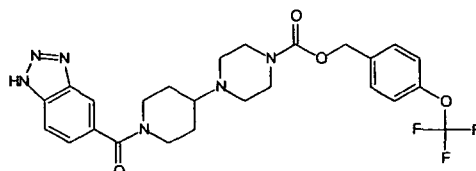
**Compound
45**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 3,4-
dichloro-benzyl ester

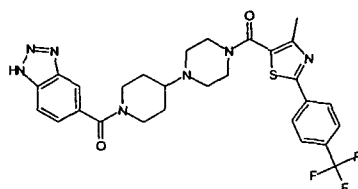
27

**Compound
46**

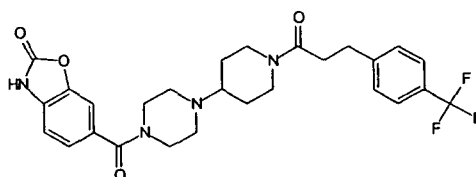
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,4-dichloro-benzyl ester

**Compound
47**

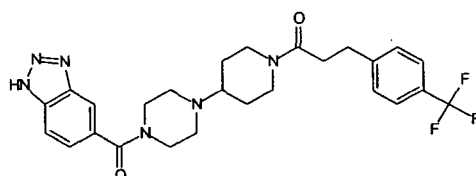
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethoxy-benzyl ester

**Compound
48**

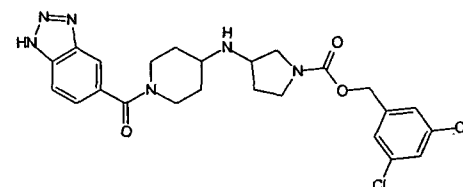
(1H-Benzotriazol-5-yl)-(4-[4-(4-methyl-2-(4-trifluoromethyl-phenyl)-thiazole-5-carbonyl]-piperazin-1-yl)-piperidin-1-yl)-methanone

**Compound
49**

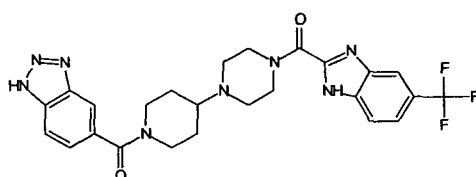
6-(4-[1-[3-(4-Trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl]-piperazine-1-carbonyl)-3H-benzooxazol-2-one

**Compound
50**

1-[4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl]-3-(4-trifluoromethyl-phenyl)-propan-1-one

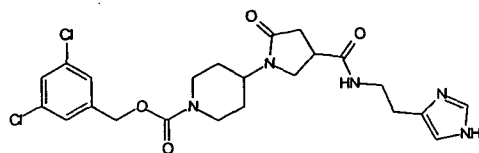
**Compound
51**

3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidine-1-carboxylic acid 3,5-dichloro-benzyl ester

**Compound
52**

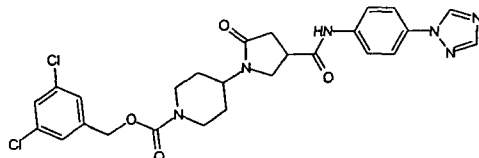
(1H-Benzotriazol-5-yl)-(4-[4-(5-trifluoromethyl-1H-benzimidazole-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl)-methanone

28

Compound 53

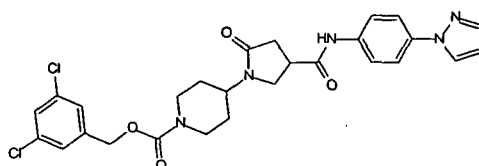
4-{4-[2-(1H-imidazol-4-yl)-ethylcarbamoyl]-2-oxo-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

5

Compound 54

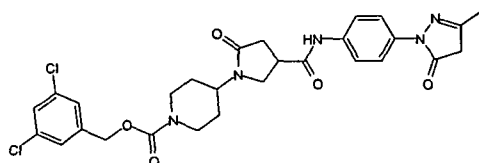
4-[2-Oxo-4-(4-[1,2,4]triazol-1-yl)-phenylcarbamoyl]-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

10

Compound 55

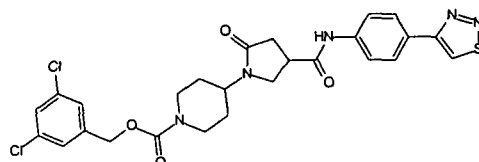
4-[2-Oxo-4-(4-pyrazol-1-yl)-phenylcarbamoyl]-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

15

Compound 56

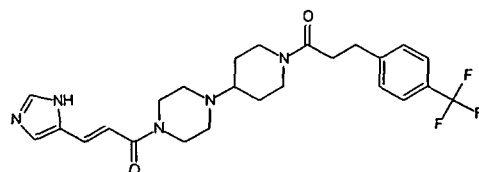
4-{4-[4-(3-Methyl-5-oxo-4,5-dihydro-pyrazol-1-yl)-phenylcarbamoyl]-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

20

Compound 57

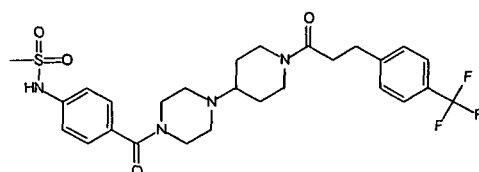
4-[2-Oxo-4-(4-[1,2,3]thiadiazol-4-yl)-phenylcarbamoyl]-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

25

Compound 58

(E)-3-(3H-imidazol-4-yl)-1-(4-{1-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-piperazin-1-yl)-propenone

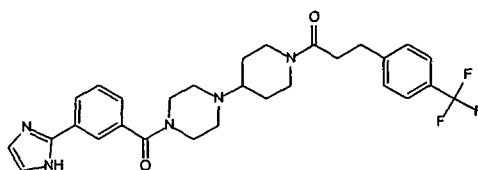
30

Compound 59

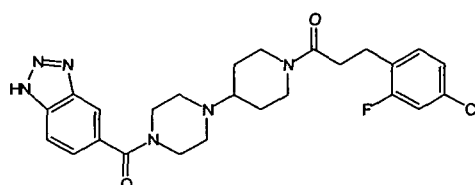
N-[4-(4-{1-[3-(4-Trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-piperazine-1-carbonyl)-phenyl]-methanesulfonamide

35

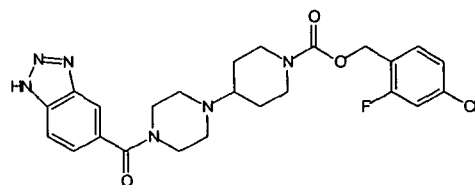
29

Compound 60

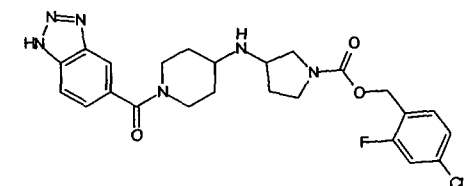
1-(4-{4-[3-(1H-Imidazol-2-yl)-benzoyl]-piperazin-1-yl}-piperidin-1-yl)-3-(4-trifluoromethyl-phenyl)-propan-1-one

Compound 61

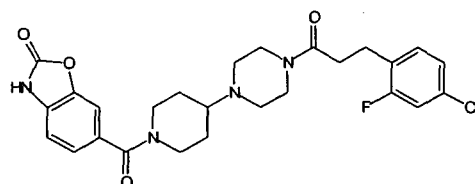
1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-chloro-2-fluoro-phenyl)-propan-1-one

Compound 62

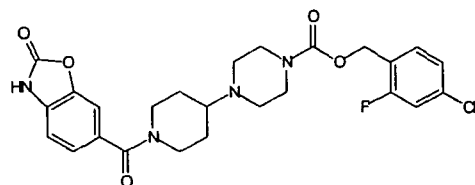
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester

Compound 63

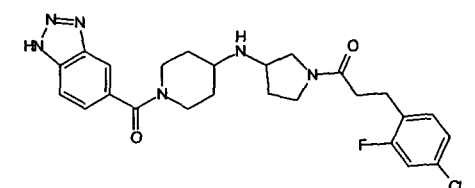
3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester

Compound 64

6-(4-{4-[3-(4-Chloro-2-fluoro-phenyl)-propionyl]-piperazin-1-yl}-piperidine-1-carbonyl)-3H-benzooxazol-2-one

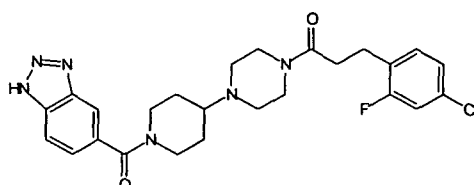
Compound 65

4-[1-(2-Oxo-2,3-dihydro-benzooxazole-6-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester

Compound 66

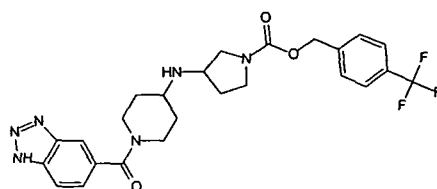
1-{3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidin-1-yl}-3-(4-chloro-2-fluoro-phenyl)-propan-1-one

5

**Compound
67**

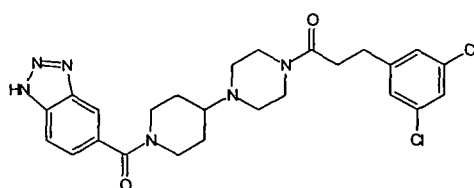
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-chloro-2-fluoro-phenyl)-propan-1-one

10

**Compound
68**

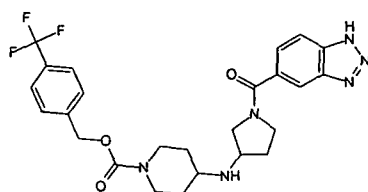
3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

15

**Compound
69**

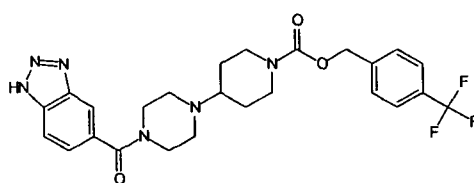
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,5-dichloro-phenyl)-propan-1-one

20

**Compound
70**

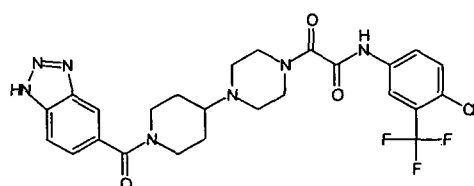
4-[1-(1H-Benzotriazole-5-carbonyl)-pyrrolidin-3-ylamino]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

25

**Compound
71**

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

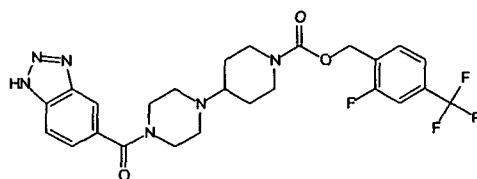
30

**Compound
72**

2-[4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl]-N-(4-chloro-3-trifluoromethyl-phenyl)-2-oxo-acetamide

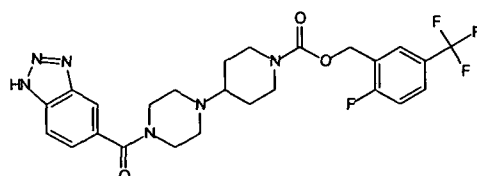
35

31

**Compound
73**

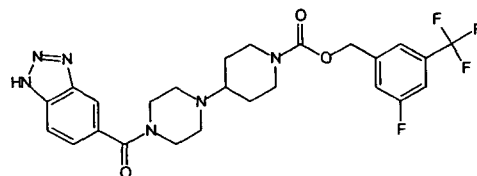
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 2-fluoro-4-trifluoromethyl-benzyl ester

5

**Compound
74**

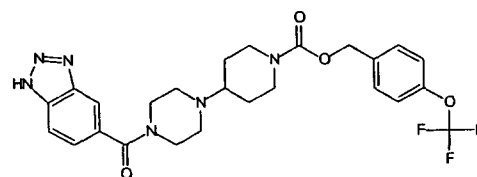
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 2-fluoro-5-trifluoromethyl-benzyl ester

10

**Compound
75**

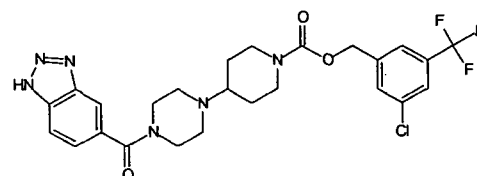
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-fluoro-5-trifluoromethyl-benzyl ester

15

**Compound
76**

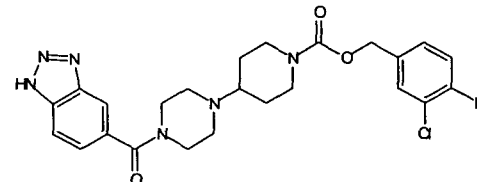
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethoxy-benzyl ester

20

**Compound
77**

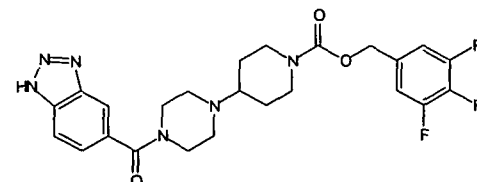
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester

25

**Compound
78**

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-4-fluoro-benzyl ester

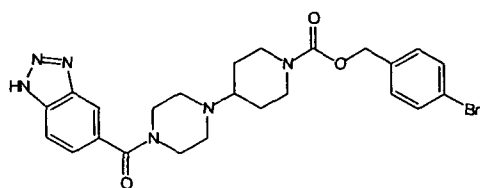
30

**Compound
79**

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,4,5-trifluoro-benzyl ester

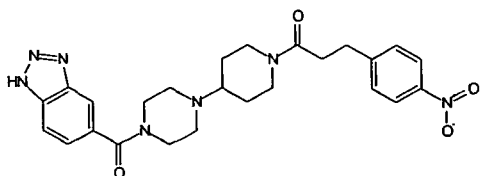
35

32

Compound 80

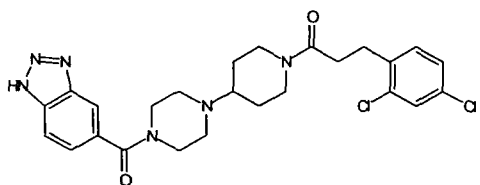
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-bromo-benzyl ester

5

Compound 81

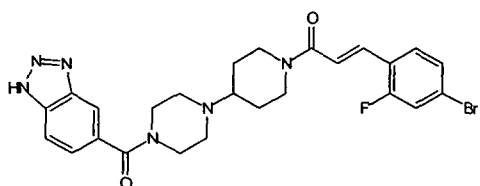
1-[4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl]-3-(4-nitro-phenyl)-propan-1-one

10

Compound 82

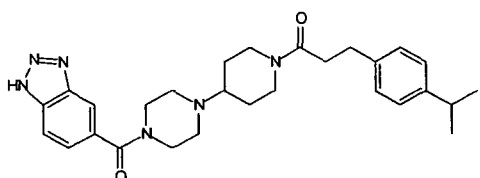
1-[4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl]-3-(2,4-dichloro-phenyl)-propan-1-one

15

Compound 83

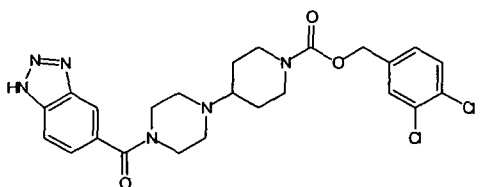
(E)-1-[4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl]-3-(4-bromo-2-fluoro-phenyl)-propenone

20

Compound 84

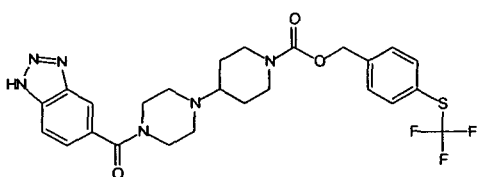
1-[4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl]-3-(4-isopropyl-phenyl)-propan-1-one

25

Compound 85

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,4-dichloro-benzyl ester

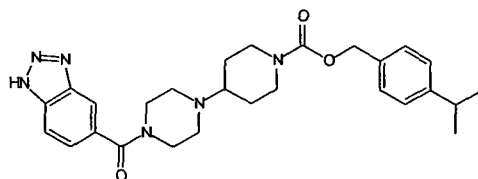
30

Compound 86

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzyl ester

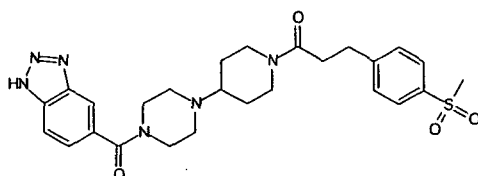
35

33

**Compound
87**

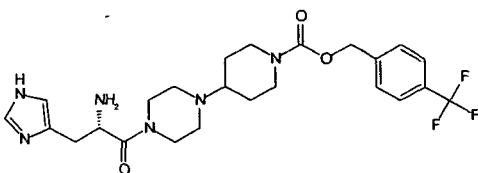
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-isopropyl-benzyl ester

5

**Compound
88**

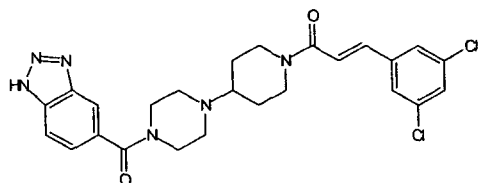
1-[4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl]-3-(4-methanesulfonyl-phenyl)-propan-1-one

10

**Compound
89**

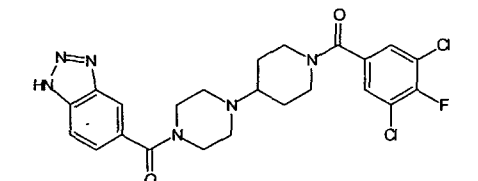
4-{4-[(S)-2-Amino-3-(1H-imidazol-4-yl)-propionyl]-piperazin-1-yl}-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

15

**Compound
90**

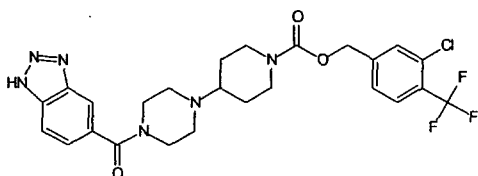
(E)-1-[4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl]-3-(3,5-dichloro-phenyl)-propenone

20

**Compound
91**

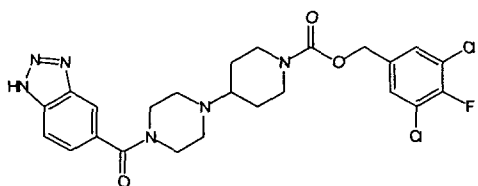
(1H-Benzotriazol-5-yl)-{4-[1-(3,5-dichloro-4-fluoro-benzoyl)-piperidin-4-yl]-piperazin-1-yl}-methanone

25

**Compound
92**

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-4-trifluoromethyl-benzyl ester

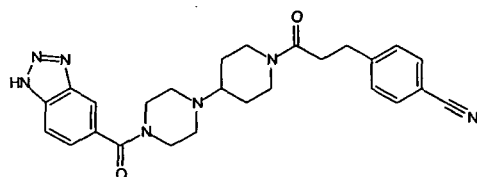
30

**Compound
93**

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-4-fluoro-benzyl ester

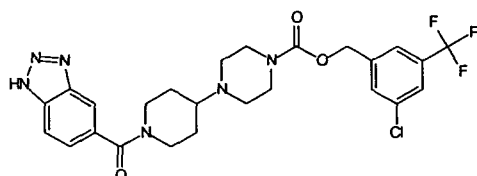
35

34

**Compound
94**

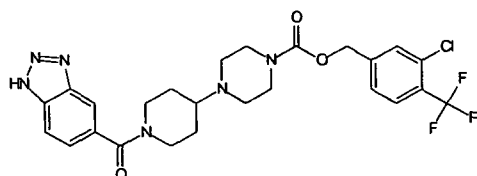
4-(3-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-oxo-propyl)-benzonitrile

5

**Compound
95**

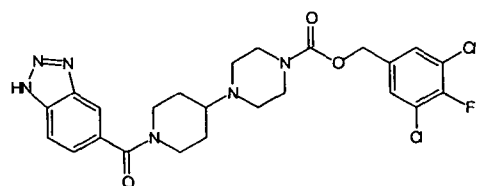
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester

10

**Compound
96**

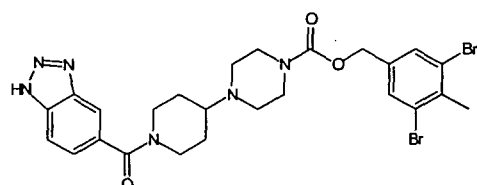
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-4-trifluoromethyl-benzyl ester

15

**Compound
97**

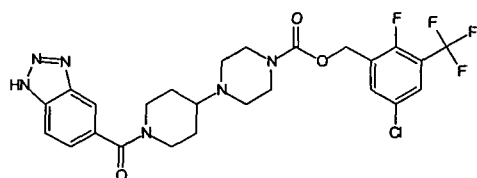
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dichloro-4-fluoro-benzyl ester

20

**Compound
98**

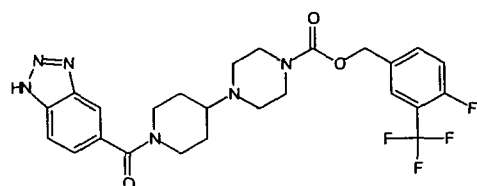
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dibromo-4-methyl-benzyl ester

25

**Compound
99**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 5-chloro-2-fluoro-3-trifluoromethyl-benzyl ester

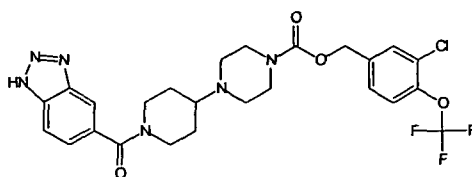
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**Compound
100**

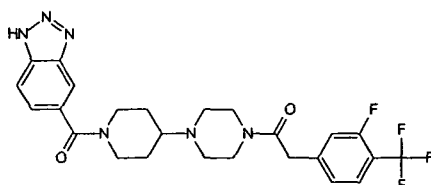
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-fluoro-3-trifluoromethyl-benzyl ester

35

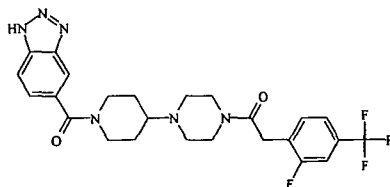
35

**Compound
101**

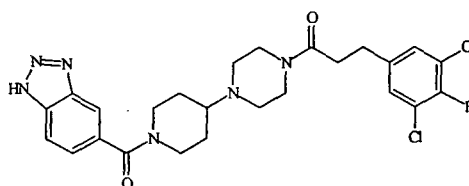
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-4-trifluoromethoxy-benzyl ester

**Compound
102**

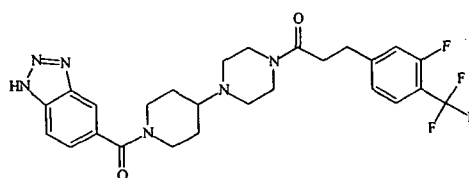
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(3-fluoro-4-trifluoromethyl-phenyl)-ethanone

**Compound
103**

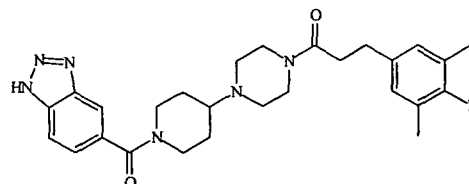
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(2-fluoro-4-trifluoromethyl-phenyl)-ethanone

**Compound
104**

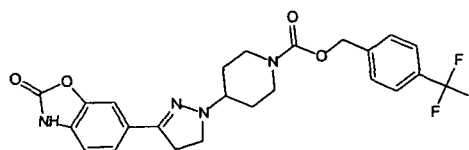
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,5-dichloro-4-fluoro-phenyl)-propan-1-one

**Compound
105**

1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3-fluoro-4-trifluoromethyl-phenyl)-propan-1-one

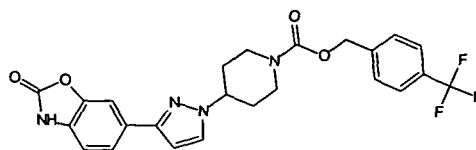
**Compound
106**

1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-fluoro-3,5-dimethyl-phenyl)-propan-1-one

**Compound
107**

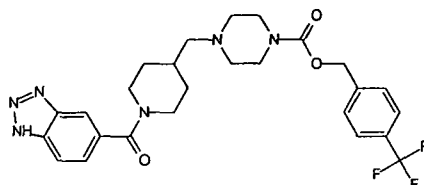
4-[3-(2-Oxo-2,3-dihydro-benzooxazol-6-yl)-4,5-dihydro-pyrazol-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

36

**Compound
108**

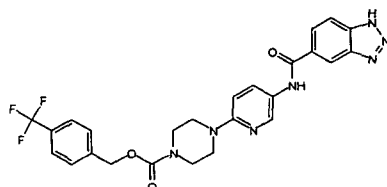
4-[3-(2-Oxo-2,3-dihydro-benzooxazol-6-yl)-pyrazol-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

5

**Compound
109**

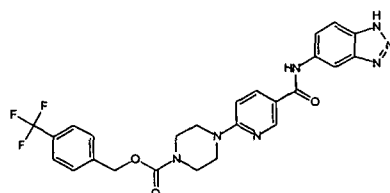
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylmethyl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

10

**Compound
110**

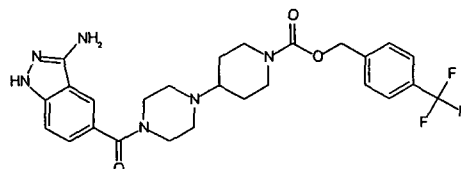
4-{5-[(1H-Benzotriazole-5-carbonyl)-amino]-pyridin-2-yl}-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

15

**Compound
111**

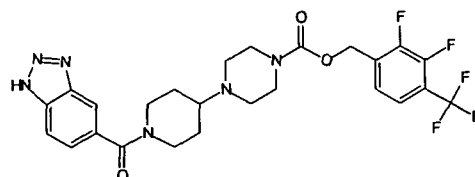
4-[5-(1H-Benzotriazol-5-ylcarbonyl)-pyridin-2-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

20

**Compound
112**

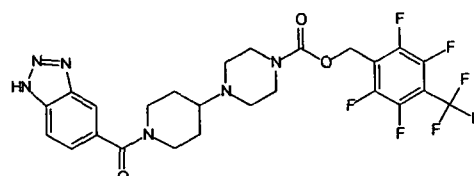
4-[4-(3-Amino-1H-indazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

25

**Compound
113**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,3-difluoro-4-trifluoromethyl-benzyl ester

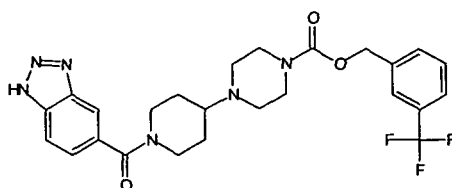
30

**Compound
114**

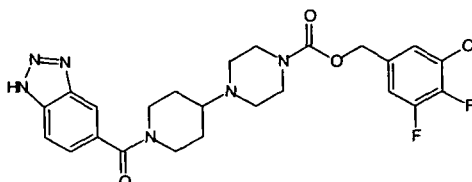
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,3,5,6-tetrafluoro-4-trifluoromethyl-benzyl ester

35

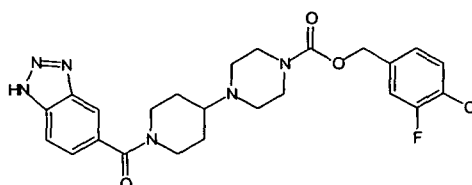
37

**Compound
115**

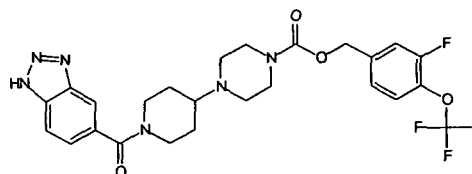
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 3-
trifluoromethyl-benzyl ester

**Compound
116**

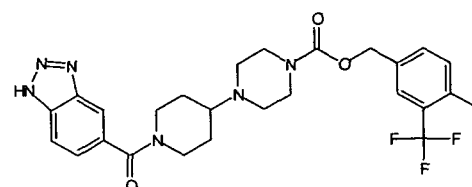
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 3-
chloro-4,5-difluoro-benzyl ester

**Compound
117**

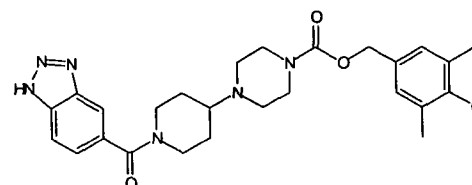
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 4-
chloro-3-fluoro-benzyl ester

**Compound
118**

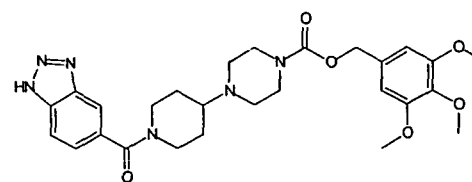
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 3-
fluoro-4-trifluoromethoxy-benzyl
ester

**Compound
119**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 4-
methyl-3-trifluoromethyl-benzyl
ester

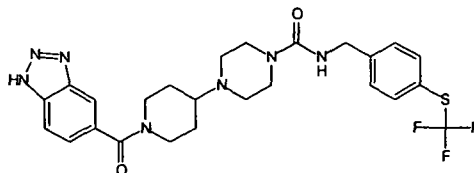
**Compound
120**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 4-
methoxy-3,5-dimethyl-benzyl
ester

**Compound
121**

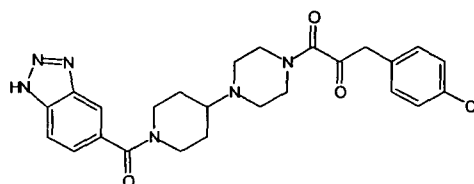
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid
3,4,5-trimethoxy-benzyl ester

5

**Compound
122**

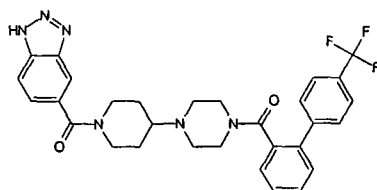
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzylamide

10

**Compound
123**

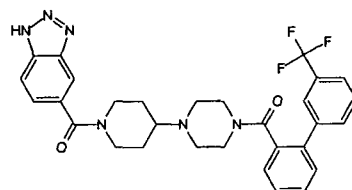
1-[4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl]-3-(4-chloro-phenyl)-propane-1,2-dione

15

**Compound
124**

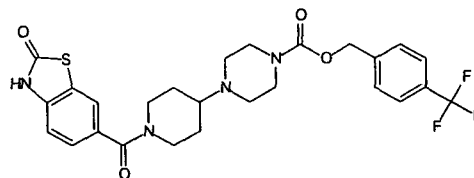
(1H-Benzotriazol-5-yl)-{4-[4-(4'-trifluoromethyl-biphenyl-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-methanone

20

**Compound
125**

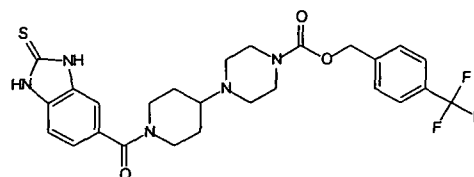
(1H-Benzotriazol-5-yl)-{4-[4-(3'-trifluoromethyl-biphenyl-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-methanone

25

**Compound
126**

4-[1-(2-Oxo-2,3-dihydro-benzothiazole-6-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

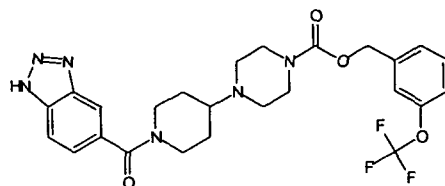
30

**Compound
127**

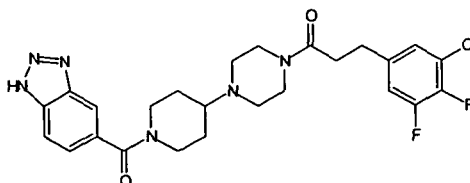
4-[1-(2-Thioxo-2,3-dihydro-1H-benzoimidazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

35

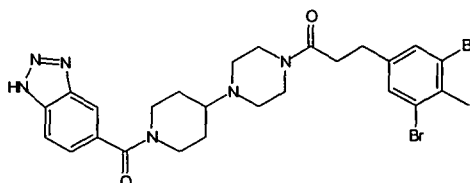
39

**Compound
128**

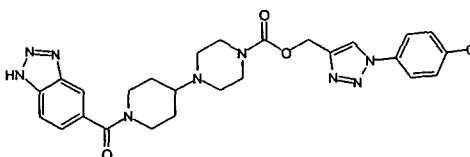
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-trifluoromethoxy-benzyl ester

**Compound
129**

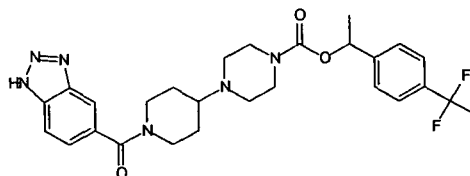
1-[4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl]-3-(3-chloro-4,5-difluorophenyl)propan-1-one

**Compound
130**

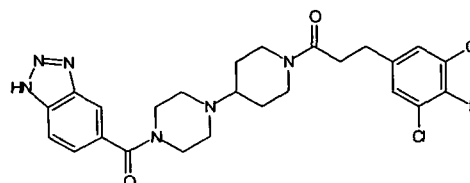
1-[4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl]-3-(3,5-dibromo-4-methylphenyl)propan-1-one

**Compound
131**

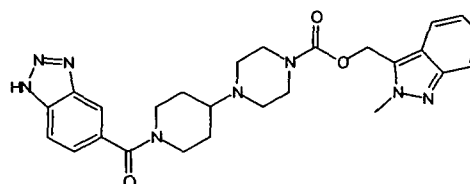
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(4-chloro-phenyl)-1H-[1,2,3]triazol-4-ylmethyl ester

**Compound
132**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(4-trifluoromethyl-phenyl)-ethyl ester

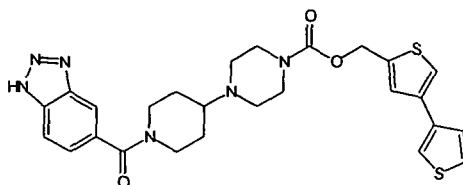
**Compound
133**

1-[4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl]-3-(3,5-dichloro-4-fluorophenyl)propan-1-one

**Compound
134**

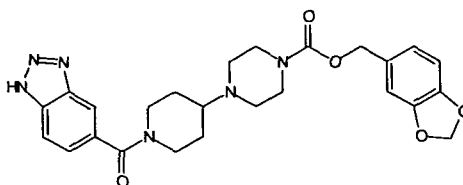
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-methyl-2H-indazol-3-ylmethyl ester

5

**Compound
135**

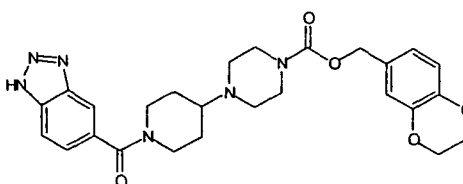
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid [3,3']bithiophenyl-5-ylmethyl ester

10

**Compound
136**

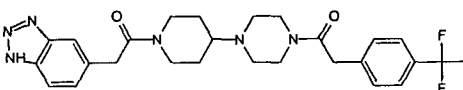
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid benzo[1,3]dioxol-5-ylmethyl ester

15

**Compound
137**

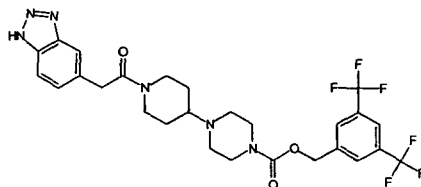
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,3-dihydro-benzo[1,4]dioxin-6-ylmethyl ester

20

**Compound
138**

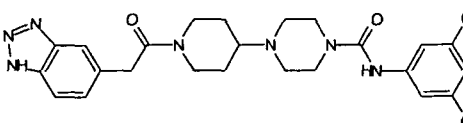
2-(1H-Benzotriazol-5-yl)-1-(4-{4-[2-(4-trifluoromethyl-phenyl)-acetyl]-piperazin-1-yl}-piperidin-1-yl)-ethanone

25

**Compound
139**

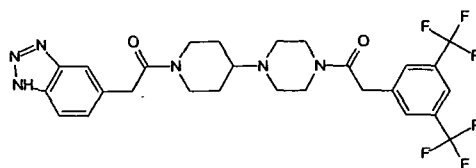
4-[1-(2-1H-Benzotriazol-5-yl)-acetyl]-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

30

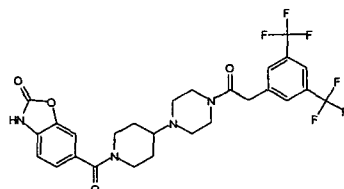
**Compound
140**

4-[1-(2-1H-Benzotriazol-5-yl)-acetyl]-piperidin-4-yl]-piperazine-1-carboxylic acid (3,5-dichloro-phenyl)-amide

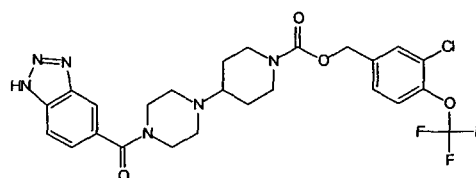
35

**Compound
141**

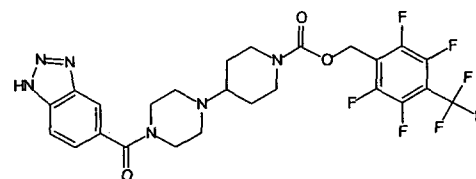
2-(1H-Benzotriazol-5-yl)-1-(4-{4-[2-(3,5-bis-trifluoromethyl-phenyl)-acetyl]-piperazin-1-yl}-piperidin-1-yl)-ethanone

**Compound
142**

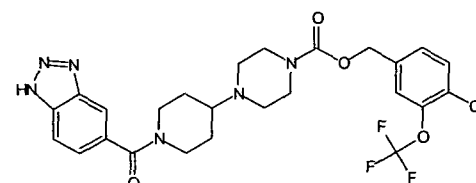
6-(4-{4-[2-(3,5-Bis-trifluoromethyl-phenyl)-acetyl]-piperazin-1-yl}-piperidine-1-carbonyl)-3H-benzooxazol-2-one

**Compound
143**

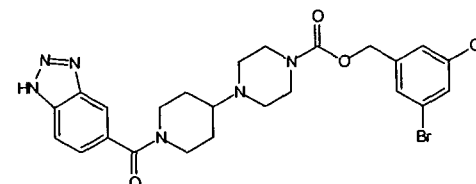
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-4-trifluoromethoxy-benzyl ester

**Compound
144**

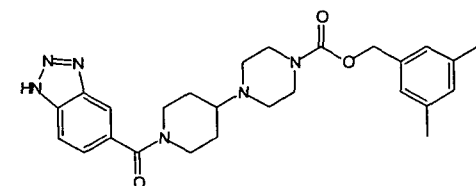
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 2,3,5,6-tetrafluoro-4-trifluoromethyl-benzyl ester

**Compound
145**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-3-trifluoromethoxy-benzyl ester

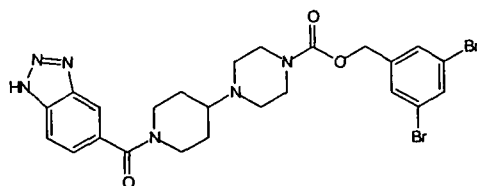
**Compound
146**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-bromo-5-chloro-benzyl ester

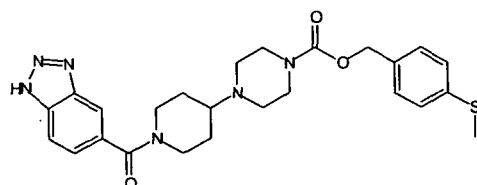
**Compound
147**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dimethyl-benzyl ester

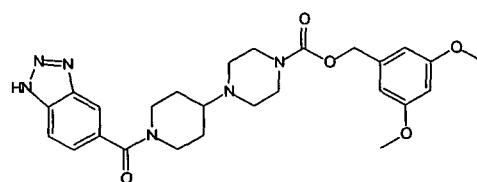
42

**Compound
148**

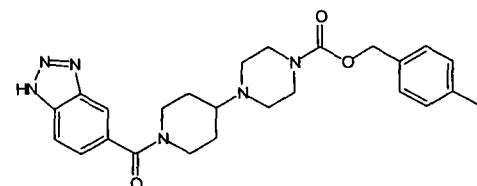
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dibromo-benzyl ester

**Compound
149**

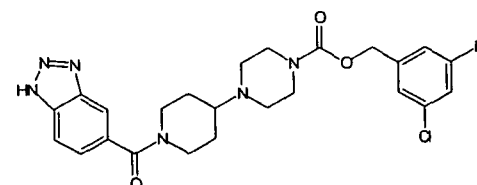
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-methylsulfanyl-benzyl ester

**Compound
150**

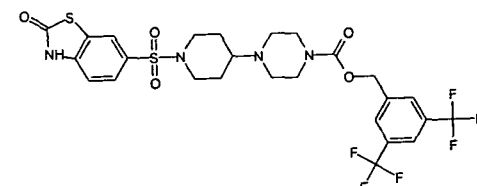
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dimethoxy-benzyl ester

**Compound
151**

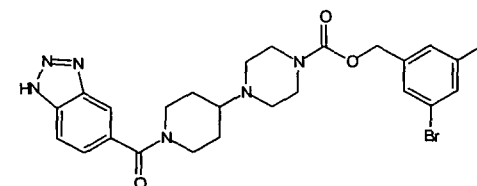
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-ethyl-benzyl ester

**Compound
152**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-fluoro-benzyl ester

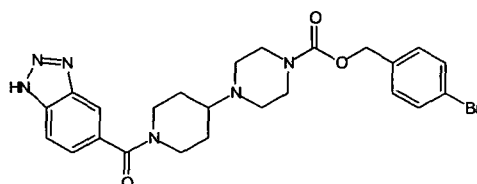
**Compound
153**

4-[1-(2-Oxo-2,3-dihydro-benzothiazole-6-sulfonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

**Compound
154**

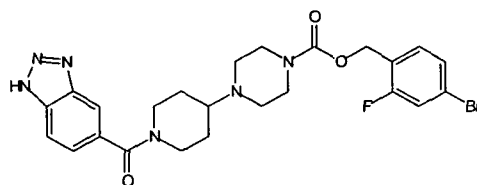
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-bromo-5-fluoro-benzyl ester

5

**Compound
155**

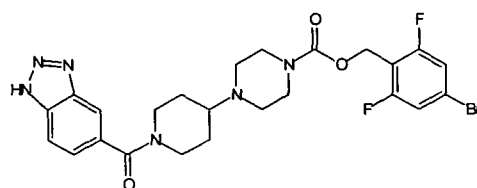
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-bromo-benzyl ester

10

**Compound
156**

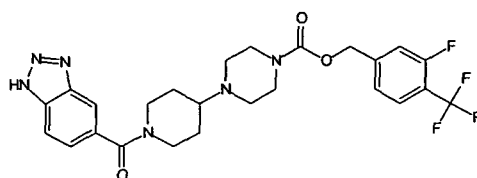
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-bromo-2-fluoro-benzyl ester

15

**Compound
157**

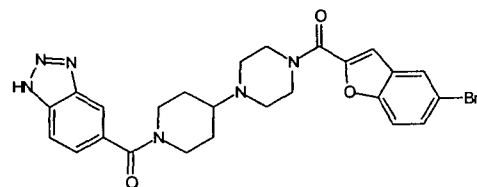
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-bromo-2,6-difluoro-benzyl ester

20

**Compound
158**

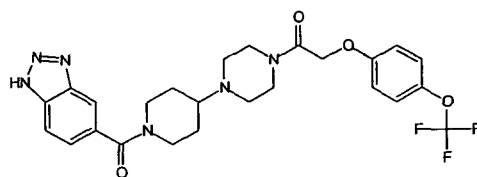
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-fluoro-4-trifluoromethyl-benzyl ester

25

**Compound
159**

(1H-Benzotriazol-5-yl)-{4-[4-(5-bromo-benzofuran-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-methanone

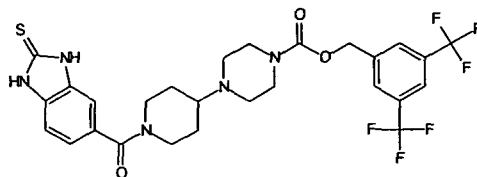
30

**Compound
160**

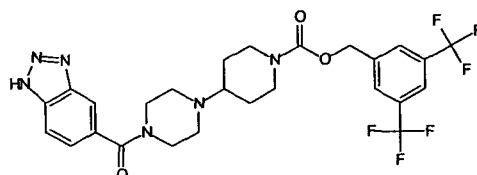
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(4-trifluoromethoxyphenoxy)-ethanone

35

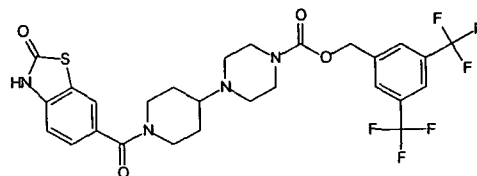
44

Compound 161

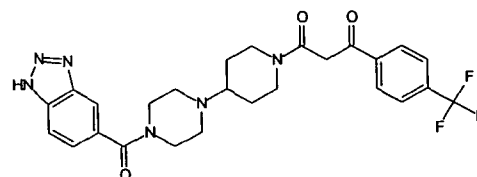
4-[1-(2-Thioxo-2,3-dihydro-1H-benzimidazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

Compound 162

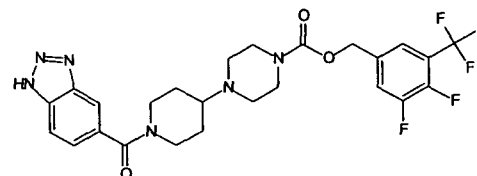
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

Compound 163

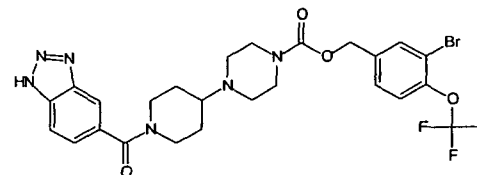
4-[1-(2-Oxo-2,3-dihydro-benzothiazole-6-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

Compound 164

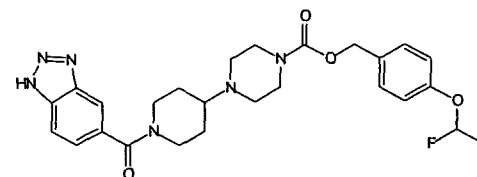
1-[4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl]-3-(4-trifluoromethyl-phenyl)-propane-1,3-dione

Compound 165

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,4-difluoro-5-trifluoromethyl-benzyl ester

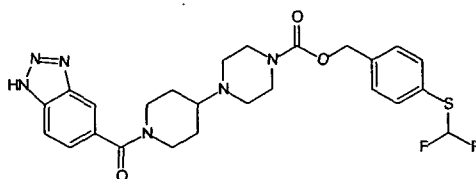
Compound 166

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-bromo-4-trifluoromethoxy-benzyl ester

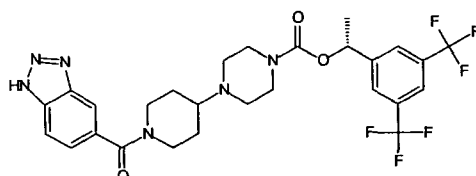
Compound 167

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-difluoromethoxy-benzyl ester

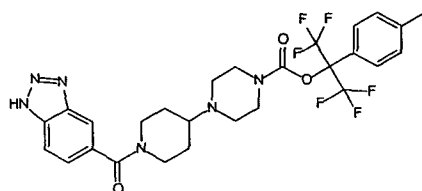
45

**Compound
168**

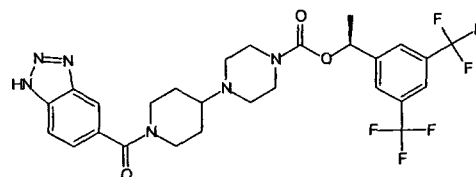
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-difluoromethylsulfanyl-benzyl ester

**Compound
169**

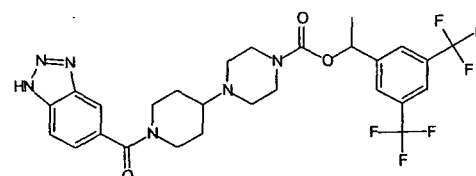
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (R)-1-(3,5-bis-trifluoromethyl-phenyl)-ethyl ester

**Compound
170**

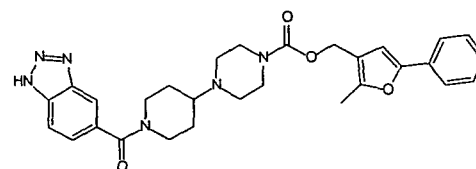
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,2,2-trifluoro-1-p-tolyl-1-trifluoromethyl-ethyl ester

**Compound
171**

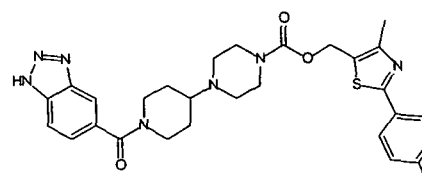
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (S)-1-(3,5-bis-trifluoromethyl-phenyl)-ethyl ester

**Compound
172**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(3,5-bis-trifluoromethyl-phenyl)-ethyl ester

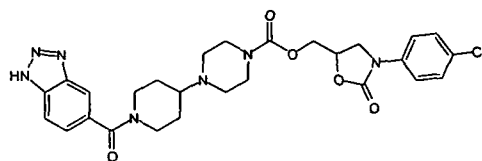
**Compound
173**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-methyl-5-phenyl-furan-3-ylmethyl ester

**Compound
174**

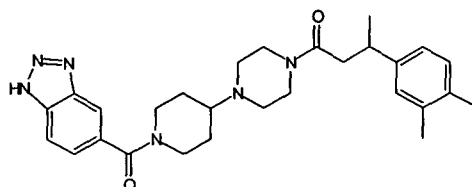
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-(4-chloro-phenyl)-4-methyl-thiazol-5-ylmethyl ester

46

**Compound
175**

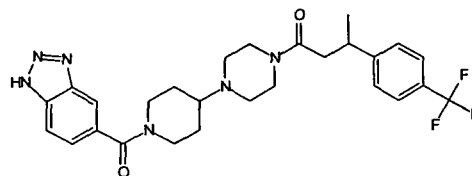
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-(4-chloro-phenyl)-2-oxo-oxazolidin-5-ylmethyl ester

5

**Compound
176**

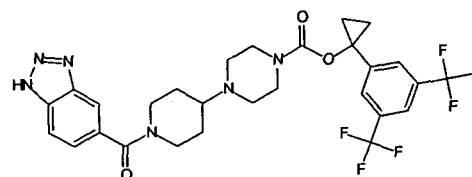
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,4-dimethyl-phenyl)-butan-1-one

10

**Compound
177**

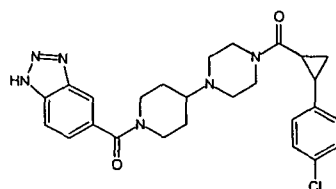
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-trifluoromethyl-phenyl)-butan-1-one

15

**Compound
178**

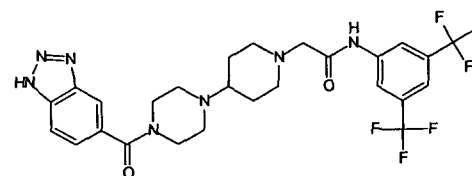
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(3,5-bis-trifluoromethyl-phenyl)-cyclopropyl ester

20

**Compound
179**

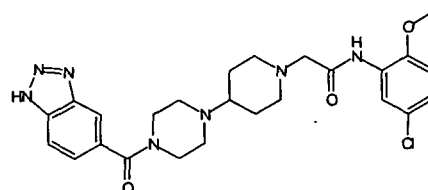
(1H-Benzotriazol-5-yl)-(4-{4-[2-(4-chloro-phenyl)-cyclopropanecarbonyl]-piperazin-1-yl}-piperidin-1-yl)-methanone

25

**Compound
180**

2-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-N-(3,5-bis-trifluoromethyl-phenyl)-acetamide

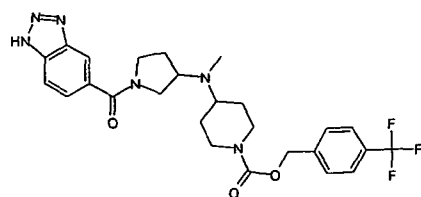
30

**Compound
181**

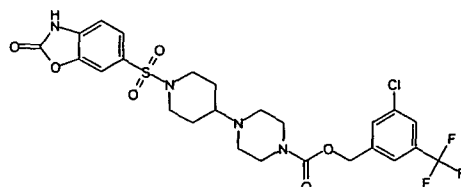
2-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-N-(5-chloro-2-methoxy-phenyl)-acetamide

35

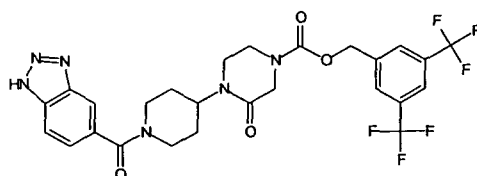
47

**Compound
182**

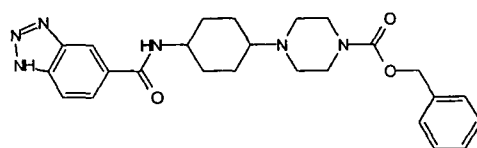
4-[[1-(1H-Benzotriazole-5-carbonyl)-pyrrolidin-3-yl]-methyl-amino]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

**Compound
183**

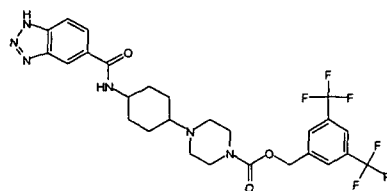
4-[1-(2-Oxo-2,3-dihydro-benzooxazole-6-sulfonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester

**Compound
184**

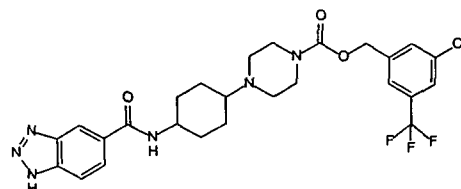
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-3-oxo-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

**Compound
185**

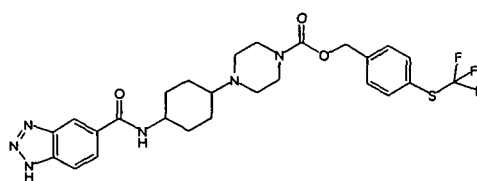
4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid benzyl ester

**Compound
186**

4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

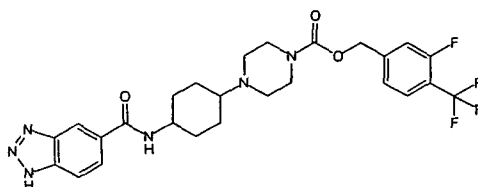
**Compound
187**

4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester

**Compound
188**

4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzyl ester

**Compound
189**



4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3-fluoro-4-trifluoromethyl-benzyl ester

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

For the avoidance of doubt, if chemical name and chemical structure of the above illustrated compounds do not correspond by mistake, the chemical structure is regarded to unambiguously define the compound.

All the above generically or explicitly disclosed compounds, including preferred subsets/embodiments of the herein disclosed formulae (Ia), (Ib) and (II) or (IIb) and Compounds 1 to 189, are hereinafter referred to as compounds of the (present) invention.

The nomenclature as used herein for defining compounds, especially the compounds according to the invention, is in general based on the rules of the IUPAC organisation for chemical compounds and especially organic compounds.

The terms indicated for explanation of the above compounds of the invention always, unless indicated otherwise in the description or in the claims, have the following meanings:

The term "unsubstituted" means that the corresponding radical, group or moiety has no substituents.

The term "substituted" means that the corresponding radical, group or moiety

has one or more substituents. Where a radical has a plurality of substituents, and a selection of various substituents is specified, the substituents are selected independently of one another and do not need to be identical.

The terms "alkyl" or "A" as well as other groups having the prefix "alk" for the purposes of this invention refer to acyclic saturated or unsaturated hydrocarbon radicals which may be branched or straight-chain and preferably have 1 to 8 carbon atoms, i.e. C₁-C₈-alkanyls, C₂-C₈-alkenyls and C₂-C₈-alkynyls. Alkenyls have at least one C-C double bond and alkynyls at least one C-C triple bond. Alkynyls may additionally also have at least one C-C double bond. Examples of suitable alkyl radicals are methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, n-pentyl, iso-pentyl, neo-pentyl, tert-pentyl, 2- or 3-methyl-pentyl, n-hexyl, 2-hexyl, isohexyl, n-heptyl, n-octyl, n-nonyl, n-decyl, n-undecyl, n-dodecyl, n-tetradecyl, n-hexadecyl, n-octadecyl, n-icosanyl, n-docosanyl, ethylenyl (vinyl), propenyl (-CH₂CH=CH₂; -CH=CH-CH₃, -C(=CH₂)-CH₃), butenyl, pentenyl, hexenyl, heptenyl, octenyl, octadienyl, octadecenyl, octadec-9-enyl, icosenyl, icos-11-enyl, (Z)-icos-11-enyl, docosanyl, docos-13-enyl, (Z)-docos-13-enyl, ethynyl, propynyl (-CH₂-C≡CH, -C≡C-CH₃), butynyl, pentynyl, hexynyl, heptynyl, octynyl. Especially preferred is C₁₋₄-alkyl. A C₁₋₄-alkyl radical is for example a methyl, ethyl, propyl, isopropyl, butyl, isobutyl, tert-butyl.

The term "(C₉-C₃₀)alkyl" for the purposes of this invention refers to acyclic saturated or unsaturated hydrocarbon radicals which may be branched or straight-chain and have 9 to 30 carbon atoms, i.e. C₉₋₃₀-alkanyls, C₉₋₃₀-alkenyls and C₉₋₃₀-alkynyls. C₉₋₃₀-Alkenyls have at least one C-C double bond and C₉₋₃₀-alkynyls at least one C-C triple bond. C₉₋₃₀-Alkynyls may additionally also have at least one C-C double bond. Examples of suitable (C₉-C₃₀)alkyl radicals are tetradecyl, hexadecyl, octadecyl, eicosanyl, cis-13-docosenyl (erucyl), trans-13-docosenyl (brassidyl), cis-15-tetracosenyl (nervonyl) and trans-15-tetracosenyl.

The term "cycloalkyl" for the purposes of this invention refers to saturated and partially unsaturated non-aromatic cyclic hydrocarbon groups/radicals, having 1 to 3 rings, that contain 3 to 20, preferably 3 to 12, most preferably 3 to 8 carbon atoms. The cycloalkyl radical may also be part of a bi- or polycyclic system, where, for example, the cycloalkyl radical is fused to an aryl, heteroaryl or heterocyclyl radical as defined herein by any possible and desired ring member(s). The bonding to the compounds of the general formula can be effected via any possible ring member of

the cycloalkyl radical. Examples of suitable cycloalkyl radicals are cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, cyclodecyl, cyclohexenyl, cyclopentenyl and cyclooctadienyl. Especially preferred are C₃-C₉-cycloalkyl and C₄-C₈-cycloalkyl. A C₄-C₈-cycloalkyl radical is for example a cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl.

The term "heterocyclyl" for the purposes of this invention refers to a mono- or polycyclic system of 3 to 20, preferably 5 or 6 to 14 ring atoms comprising carbon atoms and 1, 2, 3, 4, or 5 heteroatoms, in particular nitrogen, oxygen and/or sulfur which are identical or different. The cyclic system may be saturated, mono- or polyunsaturated but may not be aromatic. In the case of a cyclic system consisting of at least two rings the rings may be fused or spiro- or otherwise connected. Such "heterocyclyl" radicals can be linked via any ring member. The term "heterocyclyl" also includes systems in which the heterocycle is part of a bi- or polycyclic saturated, partially unsaturated and/or aromatic system, such as where the heterocycle is fused to an "aryl", "cycloalkyl", "heteroaryl" or "heterocyclyl" group as defined herein via any desired and possible ring member of the heterocyclyl radical. The bonding to the compounds of the general formula can be effected via any possible ring member of the heterocyclyl radical. Examples of suitable "heterocyclyl" radicals are pyrrolidinyl, thiapyrrolidinyl, piperidinyl, piperazinyl, oxapiperazinyl, oxapiperidinyl, oxadiazolyl, tetrahydrofuryl, imidazolidinyl, thiazolidinyl, tetrahydropyranyl, morpholinyl, tetrahydrothiophenyl, dihydropyranyl, indolinyl, indolinylmethyl, imidazolidinyl, 2-aza-bicyclo[2.2.2]octanyl.

The term "aryl" for the purposes of this invention refers to a mono- or polycyclic aromatic hydrocarbon systems having 3 to 14, preferably 5 to 14, more preferably 6 to 10 carbon atoms. The term "aryl" also includes systems in which the aromatic cycle is part of a bi- or polycyclic saturated, partially unsaturated and/or aromatic system, such as where the aromatic cycle is fused to an "aryl", "cycloalkyl", "heteroaryl" or "heterocyclyl" group as defined herein via any desired and possible ring member of the aryl radical. The bonding to the compounds of the general formula can be effected via any possible ring member of the aryl radical. Examples of suitable "aryl" radicals are phenyl, biphenyl, naphthyl, 1-naphthyl, 2-naphthyl and anthracenyl, but likewise indanyl, indenyl, or 1,2,3,4-tetrahydronaphthyl. The most preferred aryl is phenyl.

The term "heteroaryl" for the purposes of this invention refers to a 3 to 15,

preferably 5 to 14, more preferably 5-, 6- or 7-membered mono- or polycyclic aromatic hydrocarbon radical which comprises at least 1, where appropriate also 2, 3, 4 or 5 heteroatoms, preferably nitrogen, oxygen and/or sulfur, where the heteroatoms are identical or different. The number of nitrogen atoms is preferably 0, 1, 2, or 3, and that of the oxygen and sulfur atoms is independently 0 or 1. The term "heteroaryl" also includes systems in which the aromatic cycle is part of a bi- or polycyclic saturated, partially unsaturated and/or aromatic system, such as where the aromatic cycle is fused to an "aryl", "cycloalkyl", "heteroaryl" or "heterocyclyl" group as defined herein via any desired and possible ring member of the heteroaryl radical. The bonding to the compounds of the general formula can be effected via any possible ring member of the heteroaryl radical. Examples of suitable "heteroaryl" are acridinyl, benzodioxinyl, benzimidazolyl, benzisoxazolyl, benzodioxolyl, benzofuranyl, benzothiadiazolyl, benzothiazolyl, benzothieryl, benzoxazolyl, carbazolyl, cinnolinyl, dibenzofuranyl, dihydrobenzothieryl, furanyl, furazanyl, furyl, imidazolyl, indazolyl, indolinyl, indoliziny, indolyl, isobenzylfuranyl, isoindolyl, isoquinolinyl, isoquinolyl, isothiazolyl, isoxazolyl, naphthyridinyl, oxadiazolyl, oxazolyl, phenazinyl, phenothiazinyl, phenoxazinyl, phthalazinyl, pteridinyl, purinyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridinyl, pyridyl, pyrimidinyl, pyrimidyl, pyrrolyl, quinazolinyl, quinolinyl, quinolyl, quinoxalinyl, tetrazolyl, thiadiazolyl, thiazolyl, thienyl, thiophenyl, triazinyl, triazolyl.

For the purposes of the present invention, the terms "alkyl-cycloalkyl", "cycloalkylalkyl", "alkyl-heterocyclyl", "heterocyclylalkyl", "alkyl-aryl", "arylalkyl", "alkyl-heteroaryl" and "heteroarylalkyl" mean that alkyl, cycloalkyl, heterocycl, aryl and heteroaryl are each as defined above, and the cycloalkyl, heterocyclyl, aryl and heteroaryl radical is bonded to the compounds of the general formula via an alkyl radical, preferably C₁-C₈-alkyl radical, more preferably C₁-C₄-alkyl radical.

The term "alkyloxy" or "alkoxy" for the purposes of this invention refers to an alkyl radical according to above definition that is attached to an oxygen atom. The attachment to the compounds of the general formula is via the oxygen atom. Examples are methoxy, ethoxy and n-propyloxy, propoxy, isopropoxy. Preferred is "C₁-C₄-alkyloxy" having the indicated number of carbon atoms.

The term "cycloalkyloxy" or "cycloalkoxy" for the purposes of this invention refers to a cycloalkyl radical according to above definition that is attached to an oxygen atom. The attachment to the compounds of the general formula is via the

oxygen atom. Examples are cyclopropyloxy, cyclobutyloxy, cyclopentyloxy, cyclohexyloxy, cycloheptyloxy, cyclooctyloxy. Preferred is "C₃-C₉cycloalkyloxy" having the indicated number of carbon atoms.

5 The term "heterocyclyloxy" for the purposes of this invention refers to a heterocyclyl radical according to above definition that is attached to an oxygen atom. The attachment to the compounds of the general formulae is via the oxygen atom. Examples are pyrrolidinyloxy, thiapyrrolidinyloxy, piperidinyloxy, piperazinyloxy.

10 The term "aryloxy" for the purposes of this invention refers to an aryl radical according to above definition that is attached to an oxygen atom. The attachment to the compounds of the general formula is via the oxygen atom. Examples are phenyloxy, 2-naphthyloxy, 1-naphthyloxy, biphenyloxy, indanyloxy. Preferred is phenyloxy.

15 The term "heteroaryloxy" for the purposes of this invention refers to a heteroaryl radical according to above definition that is attached to an oxygen atom. The attachment to the compounds of the general formula is via the oxygen atom. Examples are pyrrolyloxy, thienyloxy, furyloxy, imidazolyloxy, thiazolyloxy.

20 The term "carbonyl" or "carbonyl moiety" for the purposes of this invention refers to a -C(O)- group.

The term "alkylcarbonyl" for the purposes of this invention refers to a "alkyl-C(O)-" group, wherein alkyl is as defined herein.

The term "alkoxycarbonyl" or "alkyloxycarbonyl" for the purposes of this invention refers to a "alkyl-O-C(O)-" group, wherein alkyl is as defined herein.

25 The term "alkoxyalkyl" for the purposes of this invention refers to a "alkyl-O-alkyl-" group, wherein alkyl is as defined herein.

30 The term "haloalkyl" for the purposes of this invention refers to an alkyl group as defined herein comprising at least one carbon atom substituent with at least one halogen as defined herein.

35 The term "halogen", "halogen atom", "halogen substituent" or "Hal" for the purposes of this invention refers to one or, where appropriate, a plurality of fluorine (F, fluoro), bromine (Br, bromo), chlorine (Cl, chloro), or iodine (I, iodo) atoms. The designations "dihalogen", "trihalogen" and "perhalogen" refer respectively to two, three and four substituents, where each substituent can be selected independently

from the group consisting of fluorine, chlorine, bromine and iodine. "Halogen" preferably means a fluorine, chlorine or bromine atom. Fluorine is most preferred, when the halogens are substituted on an alkyl (haloalkyl) or alkoxy group (e.g. CF_3 and CF_3O).

5 The term "hydroxyl" or "hydroxy" means an OH group.

10 The term "composition", as in pharmaceutical composition, for the purposes of this invention is intended to encompass a product comprising the active ingredient(s), and the inert ingredient(s) that make up the carrier, as well as any product which results, directly or indirectly, from combination, complexation or aggregation of any two or more of the ingredients, or from dissociation of one or more of the ingredients, or from other types of reactions or interactions of one or more of the ingredients. Accordingly, the pharmaceutical compositions of the present invention encompass any composition made by admixing a compound of the present invention and a pharmaceutically acceptable carrier.

15 The terms "administration of" and "administering a" compound should be understood to mean providing a compound of the invention or a prodrug of a compound of the invention to the individualist need.

20 As used herein, the term "effective amount" refers to any amount of a drug or pharmaceutical agent that will elicit the biological or medical response of a tissue, system, animal or human that is being sought, for instance, by a researcher or clinician. Furthermore, the term "therapeutically effective amount" means any amount which, as compared to a corresponding subject who has not received such amount, results in improved treatment, healing, prevention, or amelioration of a disease, disorder, or side effect, or a decrease in the rate of advancement of a disease or disorder. The term also includes within its scope amounts effective to enhance normal physiological function.

30 All stereoisomers of the compounds of the invention are contemplated, either in a mixture or in pure or substantially pure form. The compounds of the invention can have asymmetric centers at any of the carbon atoms. Consequently, they can exist in the form of their racemates, in the form of the pure enantiomers and/or diastereomers or in the form of mixtures of these enantiomers and/or diastereomers.

35 The mixtures may have any desired mixing ratio of the stereoisomers.

Thus, for example, the compounds of the invention which have one or more centers of chirality and which occur as racemates or as diastereomer mixtures can be fractionated by methods known per se into their optical pure isomers, i.e. enantiomers or diastereomers. The separation of the compounds of the invention can take place by column separation on chiral or nonchiral phases or by recrystallization from an optionally optically active solvent or with use of an optically active acid or base or by derivatization with an optically active reagent such as, for example, an optically active alcohol, and subsequent elimination of the radical.

The compounds of the invention may be present in the form of their double bond isomers as "pure" E or Z isomers, or in the form of mixtures of these double bond isomers.

Where possible, the compounds of the invention may be in the form of the tautomers, such as keto-enol tautomers.

It is likewise possible for the compounds of the invention to be in the form of any desired prodrugs such as, for example, esters, carbonates, carbamates, ureas, amides or phosphates, in which cases the actually biologically active form is released only through metabolism. Any compound that can be converted in vivo to provide the bioactive agent (i.e. compounds of the invention) is a prodrug within the scope and spirit of the invention.

Various forms of prodrugs are well known in the art and are described for instance in:

(i) Wermuth CG et al., Chapter 31: 671-696, The Practice of Medicinal Chemistry, Academic Press 1996;

(ii) Bundgaard H, Design of Prodrugs, Elsevier 1985; and

(iii) Bundgaard H, Chapter 5: 131-191, A Textbook of Drug Design and Development, Harwood Academic Publishers 1991.

Said references are incorporated herein by reference.

It is further known that chemical substances are converted in the body into metabolites which may where appropriate likewise elicit the desired biological effect - in some circumstances even in more pronounced form.

Any biologically active compound that was converted in vivo by metabolism from any of the compounds of the invention is a metabolite within the scope and spirit of

the invention.

5 The compounds of the invention can, if they have a sufficiently basic group such as, for example, a secondary or tertiary amine, be converted with inorganic and organic acids into salts. The pharmaceutically acceptable salts of the compounds of the invention are preferably formed with hydrochloric acid, hydrobromic acid, iodic acid, sulfuric acid, phosphoric acid, methanesulfonic acid, p-toluenesulfonic acid, carbonic acid, formic acid, acetic acid, sulfoacetic acid, trifluoroacetic acid, oxalic acid, malonic acid, maleic acid, succinic acid, tartaric acid, racemic acid, malic acid, 10 embonic acid, mandelic acid, fumaric acid, lactic acid, citric acid, taurocholic acid, glutaric acid, stearic acid, glutamic acid or aspartic acid. The salts which are formed are, inter alia, hydrochlorides, chlorided, hydrobromides, bromides, iodides, sulfates, phosphates, methanesulfonates, tosylates, carbonates, bicarbonates, formates, acetates, sulfoacetates, triflates, oxalates, malonates, maleates, succinates, 15 tartrates, malates, embonates, mandelates, fumarates, lactates, citrates, glutarates, stearates, aspartates and glutamates. The stoichiometry of the salts formed from the compounds of the invention may moreover be an integral or non-integral multiple of one.

20 The compounds of the invention can, if they contain a sufficiently acidic group such as, for example, the carboxy, sulfonic acid, phosphoric acid or a phenolic group, be converted with inorganic and organic bases into their physiologically tolerated salts. Examples of suitable inorganic bases are ammonium, sodium hydroxide, potassium hydroxide, calcium hydroxide, and of organic bases are 25 ethanolamine, diethanolamine, triethanolamine, ethylenediamine, t-butylamine, t-octylamine, dehydroabietylamine, cyclohexylamine, dibenzylethylene-diamine and lysine. The stoichiometry of the salts formed from the compounds of the invention can moreover be an integral or non-integral multiple of one.

30 It is likewise possible for the compounds of the invention to be in the form of their solvates and, in particular, hydrates which can be obtained for example by crystallization from a solvent or from aqueous solution. It is moreover possible for one, two, three or any number of solvate or water molecules to combine with the compounds of the invention to give solvates and hydrates.

35 By the term "solvate" is meant a hydrate, an alcoholate, or other solvate of

crystallization.

It is known that chemical substances form solids which exist in different order states which are referred to as polymorphic forms or modifications. The various modifications of a polymorphic substance may differ greatly in their physical properties. The compounds of the invention can exist in various polymorphic forms and certain modifications may moreover be metastable. All these polymorphic forms of the compounds are to be regarded as belonging to the invention.

The compounds of the invention are surprisingly characterized by a strong and/or selective inhibition of autotaxin.

Due to their surprisingly strong and/or selective enzyme inhibition, the compounds of the invention can be advantageously administered at lower doses compared to other less potent or selective inhibitors of the prior art while still achieving equivalent or even superior desired biological effects. In addition, such a dose reduction may advantageously lead to less or even no medicinal adverse effects. Further, the high inhibition selectivity of the compounds of the invention may translate into a decrease of undesired side effects on its own regardless of the dose applied.

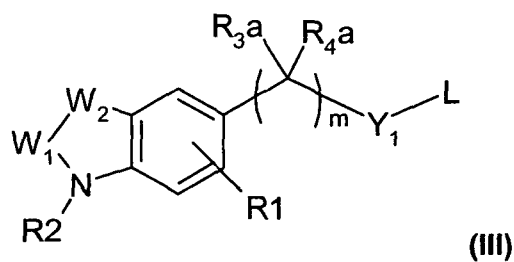
The compounds of the invention being autotaxin inhibitors generally have an inhibition constant IC_{50} of less than about 30 μM , and preferably less than about 5 μM .

The object of the present invention has surprisingly been solved in another aspect by providing the use of a compound of the invention as autotaxin inhibitor.

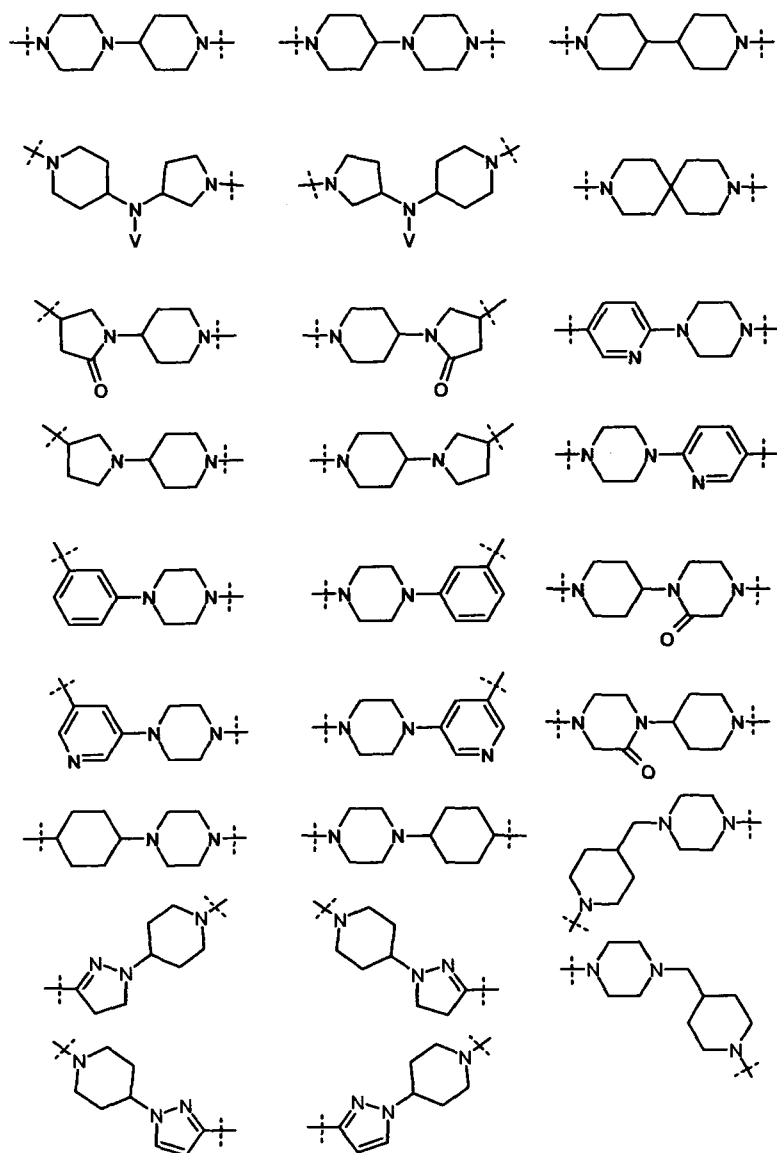
The terms "inhibiting, inhibition and/or retardation" are intended to refer for the purposes of the present invention to as follows: "partial or complete inhibiting, inhibition and/or retardation". In this case, it is within the specialist knowledge of the average person skilled in the art to measure and determine such inhibiting, inhibition, and/or retardation by means of the usual methods of measurement and determination. Thus, a partial inhibiting, inhibition and/or retardation, for example, can be measured and determined in relation to a complete inhibiting, inhibition and/or retardation.

The object of the present invention has surprisingly been solved in another aspect by providing a process for the preparation of a compound of the invention, comprising the steps:

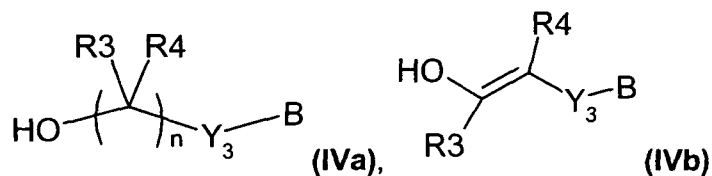
(a) reacting a compound of the formula (III),



wherein L is selected from the group consisting of:



and W_1 , W_2 , Y_1 , R_1 , R_2 , R_{3a} , R_{4a} , V , m have the meanings as indicated supra, with a compound of the formula (IVa) or (IVb),

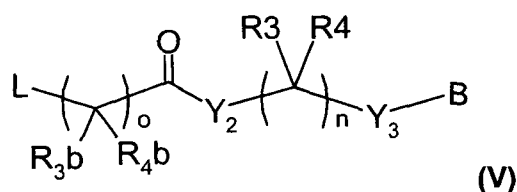


wherein B , Y_3 , R_3 , R_4 , n have the meanings as indicated supra, and a compound selected from the group consisting of: 1,1'-carbonyldiimidazole, phosgene, diphosgene, triphosgene to yield a compound according to formulae (Ia), (Ib) or (II) as indicated supra, in which Z_1 denotes "O" and Y_2 denotes "O";

10

or

(b) reacting a compound of the formula (V)



wherein L is selected from the group consisting of:

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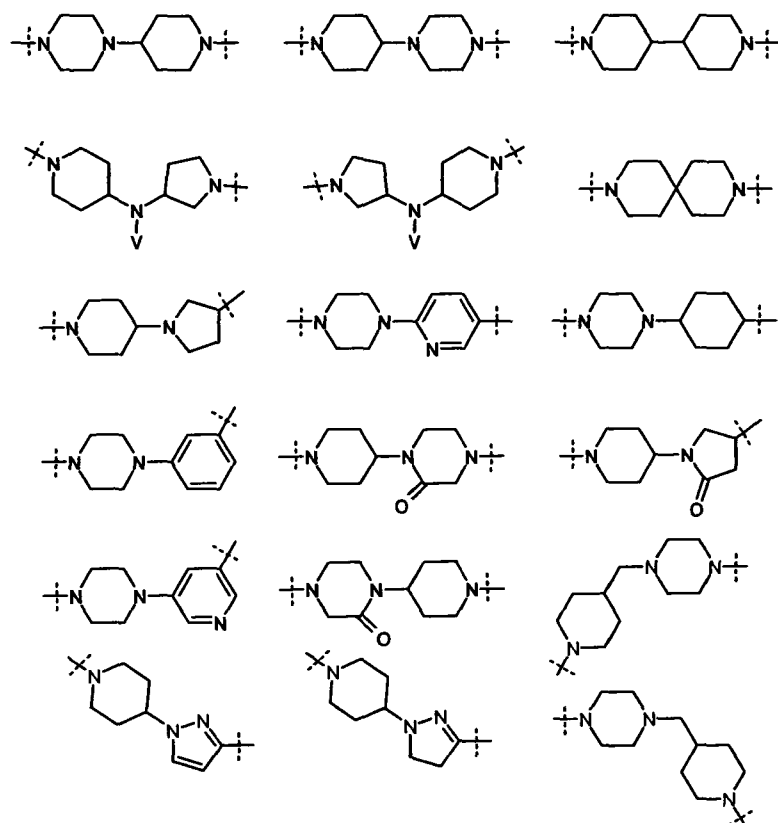
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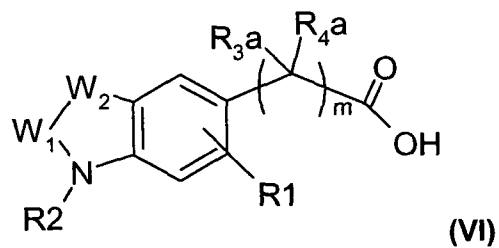
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15



and B, Y₂, Y₃, R₃, R₄, R_{3b}, R_{4b}, V, n, o have the meanings as indicated supra, with a compound of the formula (VI)

20



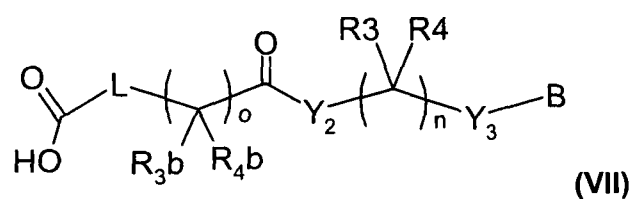
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wherein W₁, W₂, R₁, R₂, R_{3a}, R_{4a}, m have the meanings as indicated supra, to yield a compound according to formulae (Ia), (Ib) or (II) as indicated supra, in which Z₁ denotes "O", and Y₁ denotes "-C(O)-";

or

30

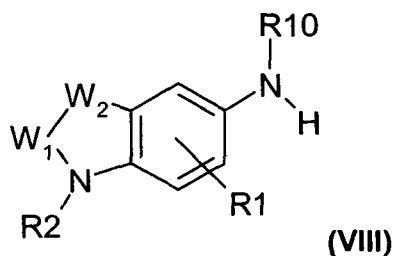
(c) reacting a compound of the formula (VII)



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wherein L, Y₂, Y₃, B, R₃, R₄, R_{3b}, R_{4b}, n, o have the meanings as indicated supra, with a compound of formula (VIII)



wherein W₁, W₂, R₁, R₂, R₁₀ have the meanings as indicated supra, to yield a compound according to formulae (Ia), (Ib) or (II) as indicated supra, in which Z₁ denotes "O" and Y₁ denotes "-N(R₁₀)-C(O)-";

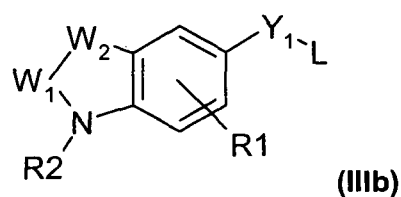
or

(d) liberating them from one of their functional derivatives (e.g. with protection groups) by treating them with an acidic, basic, solvolyzing or hydrogenolyzing agent.

and/or converting a base or an acid of a compound according to formulae (Ia), (Ib) or (II) into one of its salts.

The object of the present invention has surprisingly been solved in another aspect by providing a process for the preparation of a compound of the invention, comprising the steps:

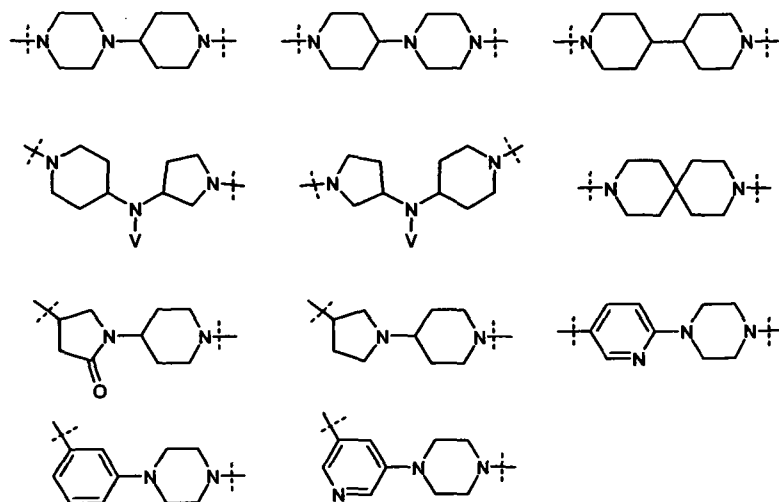
(a) reacting a compound of the formula (IIIb),



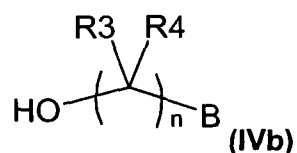
wherein L is selected from the group consisting of:

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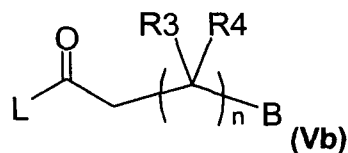
and W_1 , W_2 , Y_1 , R_1 , R_2 have the meanings as indicated supra, with a compound of the formula (IVb),



wherein B , R_3 , R_4 , n have the meanings as indicated supra, and a compound selected from the group consisting of: 1,1'-carbonyldiimidazole, phosgene, diphosgene, triphosgene to yield a compound according to formula (Ib) as indicated supra, in which Z_1 denotes "O" and Y_2 denotes "O";

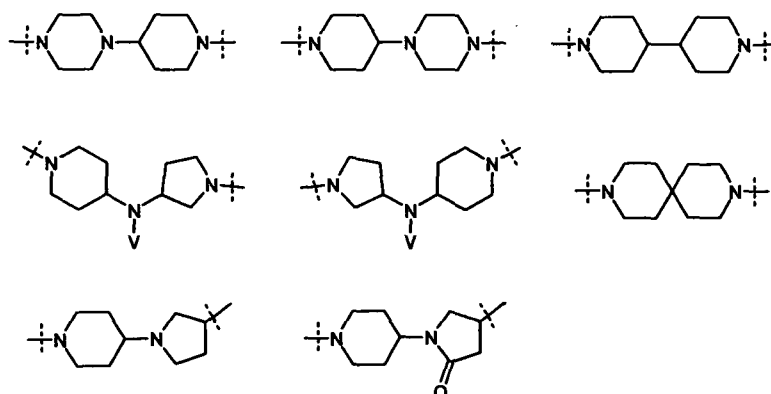
or

(b) reacting a compound of the formula (Vb)

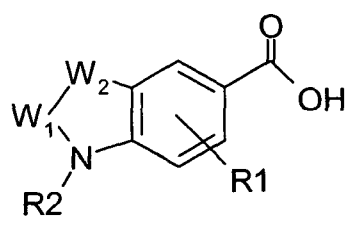


wherein L is selected from the group consisting of:

62



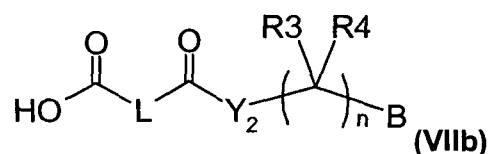
and B, R₃, R₄, V, n have the meanings as indicated supra, with a compound of the formula (VIb)



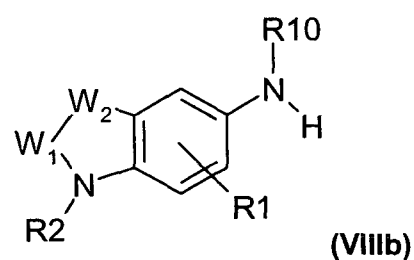
wherein W₁, W₂, R₁, R₂ have the meanings as indicated supra, to yield a compound according to formula (Ib) as indicated supra, in which Z₁ denotes "O", Y₁ denotes "-C(O)-", and Y₂ denotes "single bond" or "-CH₂-";

or

(c) reacting a compound of the formula (VIIb)



wherein L, Y₂, B, R₃, R₄, n have the meanings as indicated supra, with a compound of formula (VIIIb)



wherein W_1 , W_2 , R_1 , R_2 , R_{10} have the meanings as indicated supra, to yield a compound according to formula (Ib) as indicated supra, in which Z_1 denotes "O" and Y_1 denotes " $-N(R_{10})-C(O)-$ ".

5

All crude products were subjected to standard chromatography using solvent mixtures containing methanol, ethanol, isopropanol, n-hexane, cyclohexane or petrol ether, respectively.

10

For a further detailed description of the manufacturing processes, please refer also to the examples and the following general description of the preferred conditions.

15

A physiologically acceptable salt of a compound of the invention can also be obtained by isolating and/or treating the compound of the invention obtained by the described reaction with an acid or a base.

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The compounds of the invention and also the starting materials for their preparation are, are prepared by methods as described in the examples or by methods known per se, as described in the literature (for example in standard works, such as Houben-Weyl, Methoden der Organischen Chemie [Methods of Organic Chemistry], Georg Thieme Verlag, Stuttgart; Organic Reactions, John Wiley & Sons, Inc., New York), to be precise under reaction conditions which are known and suitable for the said reactions. Use can also be made here of variants which are known per se, but are not mentioned here in greater detail.

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The starting materials for the claimed process may, if desired, also be formed in situ by not isolating them from the reaction mixture, but instead immediately converting them further into the compounds of the invention. On the other hand, it is possible to carry out the reaction stepwise.

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Preferably, the reaction of the compounds is carried out in the presence of a suitable solvent, which is preferably inert under the respective reaction conditions. Examples of suitable solvents are hydrocarbons, such as hexane, petroleum ether, benzene, toluene or xylene; chlorinated hydrocarbons, such as trichlorethylene, 1,2-dichloroethane, tetrachloromethane, chloroform or dichloromethane; alcohols, such

as methanol, ethanol, isopropanol, n-propanol, n-butanol or tert-butanol; ethers, such as diethyl ether, diisopropyl ether, tetrahydrofuran (THF) or dioxane; glycol ethers, such as ethylene glycol monomethyl or monoethyl ether or ethylene glycol dimethyl ether (diglyme); ketones, such as acetone or butanone; amides, such as acetamide, dimethylacetamide, dimethylformamide (DMF) or N-methyl pyrrolidinone (NMP); nitriles, such as acetonitrile; sulfoxides, such as dimethyl sulfoxide (DMSO); nitro compounds, such as nitromethane or nitrobenzene; esters, such as ethyl acetate, or mixtures of the said solvents or mixtures with water. Polar solvents are in general preferred. Examples for suitable polar solvents are chlorinated hydrocarbons, alcohols, glycol ethers, nitriles, amides and sulfoxides or mixtures thereof. More preferred are amides, especially dimethylformamide (DMF).

As stated above, the reaction temperature is between about -100° C and 300° C, depending on the reaction step and the conditions used.

Reaction times are generally in the range between some minutes and several days, depending on the reactivity of the respective compounds and the respective reaction conditions. Suitable reaction times are readily determinable by methods known in the art, for example reaction monitoring. Based on the reaction temperatures given above, suitable reaction times generally lie in the range between 10 min and 48 hrs.

A base of a compound of the invention can be converted into the associated acid-addition salt using an acid, for example by reaction of equivalent amounts of the base and the acid in a preferably inert solvent, such as ethanol, followed by evaporation. Suitable acids for this reaction are, in particular, those which give physiologically acceptable salts. Thus, it is possible to use inorganic acids, for example sulfuric acid, sulfurous acid, dithionic acid, nitric acid, hydrohalic acids, such as hydrochloric acid or hydrobromic acid, phosphoric acids, such as, for example, orthophosphoric acid, sulfamic acid, furthermore organic acids, in particular aliphatic, alicyclic, araliphatic, aromatic or heterocyclic monobasic or polybasic carboxylic, sulfonic or sulfuric acids, for example formic acid, acetic acid, propionic acid, hexanoic acid, octanoic acid, decanoic acid, hexadecanoic acid, octadecanoic acid, pivalic acid, diethylacetic acid, malonic acid, succinic acid, pimelic acid, fumaric acid, maleic acid, lactic acid, tartaric acid, malic acid, citric acid, gluconic acid, ascorbic acid, nicotinic acid, isonicotinic acid, methane- or

ethanesulfonic acid, ethanedisulfonic acid, 2-hydroxyethanesulfonic acid, benzenesulfonic acid, trimethoxybenzoic acid, adamantanecarboxylic acid, p-toluenesulfonic acid, glycolic acid, embonic acid, chlorophenoxyacetic acid, aspartic acid, glutamic acid, proline, glyoxylic acid, palmitic acid,
5 parachlorophenoxyisobutyric acid, cyclohexanecarboxylic acid, glucose 1-phosphate, naphthalenemono- and -disulfonic acids or laurylsulfuric acid.

Salts with physiologically unacceptable acids, for example picrates, can be used to isolate and/or purify the compounds of the invention.

10 On the other hand, compounds of the invention can be converted into the corresponding metal salts, in particular alkali metal salts or alkaline earth metal salts, or into the corresponding ammonium salts, using bases (for example sodium hydroxide, potassium hydroxide, sodium carbonate or potassium carbonate). Suitable salts are furthermore substituted ammonium salts, for example the
15 dimethyl-, diethyl- and diisopropylammonium salts, monoethanol-, diethanol- and diisopropanolammonium salts, cyclohexyl- and dicyclohexylammonium salts, dibenzylethylenediammonium salts, furthermore, for example, salts with arginine or lysine.

If desired, the free bases of the compounds of the invention can be liberated
20 from their salts by treatment with strong bases, such as sodium hydroxide, potassium hydroxide, sodium carbonate or potassium carbonate, so long as no further acidic groups are present in the molecule. In the cases where the compounds of the invention have free acid groups, salt formation can likewise be achieved by treatment with bases. Suitable bases are alkali metal hydroxides,
25 alkaline earth metal hydroxides or organic bases in the form of primary, secondary or tertiary amines.

Every reaction step described herein can optionally be followed by one or more working up procedures and/or isolating procedures. Suitable such procedures are
30 known in the art, for example from standard works, such as Houben-Weyl, Methoden der organischen Chemie [Methods of Organic Chemistry], Georg-Thieme-Verlag, Stuttgart). Examples for such procedures include, but are not limited to evaporating a solvent, distilling, crystallization, fractionised crystallization, extraction procedures, washing procedures, digesting procedures, filtration procedures,
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chromatography, chromatography by HPLC and drying procedures, especially drying procedures in vacuo and/or elevated temperature.

5 The object of the present invention has surprisingly been solved in another aspect by providing a medicament comprising at least one compound of the invention.

10 The object of the present invention has surprisingly been solved in another aspect by providing a medicament comprising at least one compound of the invention for use in the treatment and/or prophylaxis of physiological and/or pathophysiological conditions, which are caused, mediated and/or propagated by increased lysophosphatic acid levels and/or the activation of autotaxin. A corresponding use for the preparation of a medicament for the treatment and/or
15 prophylaxis of the aforementioned conditions is intended to be comprised.

20 The object of the present invention has surprisingly been solved in another aspect by providing a medicament comprising at least one compound of the invention for use in the treatment and/or prophylaxis of physiological and/or pathophysiological conditions selected from the group consisting of: "cancer, tumour, malignant tumours, benign tumours, solid tumours, sarcomas, carcinomas, hyperproliferative disorders, carcinoids, Ewing sarcomas, Kaposi sarcomas, brain tumours, tumours originating from the brain and/or the nervous system and/or the
25 meninges, gliomas, glioblastomas, neuroblastomas, stomach cancer, kidney cancer, kidney cell carcinomas, prostate cancer, prostate carcinomas, connective tissue tumours, soft tissue sarcomas, pancreas tumours, liver tumours, head tumours, neck tumours, laryngeal cancer, oesophageal cancer, thyroid cancer, osteosarcomas, retinoblastomas, thymoma, testicular cancer, lung cancer, lung
30 adenocarcinoma, small cell lung carcinoma, bronchial carcinomas, breast cancer, mamma carcinomas, intestinal cancer, colorectal tumours, colon carcinomas, rectum carcinomas, gynaecological tumours, ovary tumours/ovarian tumours, uterine cancer, cervical cancer, cervix carcinomas, cancer of body of uterus, corpus carcinomas, endometrial carcinomas, urinary bladder cancer, urogenital tract
35 cancer, bladder cancer, skin cancer, epithelial tumours, squamous epithelial

carcinoma, basaliomas, spinaliomas, melanomas, intraocular melanomas, leukaemias, monocyte leukaemia, chronic leukaemias, chronic myelotic leukaemia, chronic lymphatic leukemia, acute leukaemias, acute myelotic leukaemia, acute lymphatic leukemia, lymphomas, angiogenesis, arteriosclerosis, ophthalmic diseases, choroidal neovascularization, diabetic retinopathy, inflammatory diseases, arthritis, neurodegeneration, restenosis, wound healing and/or transplant rejection". A corresponding use for the preparation of a medicament for the treatment and/or prophylaxis of the aforementioned conditions is intended to be comprised.

Compounds of the invention may be used in combination with one or more other active substances (ingredients, drugs) in the treatment, prevention, suppression or amelioration of diseases or conditions for which compounds of the invention or the other substances have utility. Typically the combination of the drugs is safer or more effective than either drug alone, or the combination is safer or more effective than would it be expected based on the additive properties of the individual drugs. Such other drug(s) may be administered, by a route and in an amount commonly used contemporaneously or sequentially with a compound of the invention. When a compound of the invention is used contemporaneously with one or more other drugs, a combination product containing such other drug(s) and the compound of the invention is preferred. However, combination therapy also includes therapies in which the compound of the invention and one or more other drugs are administered on different overlapping schedules. It is contemplated that when used in combination with other active ingredients, the compound of the present invention or the other active ingredient or both may be used effectively in lower doses than when each is used alone. Accordingly, the pharmaceutical compositions of the present invention include those that contain one or more other active ingredients, in addition to a compound of the invention.

Examples of other active substances (ingredients, drugs) that may be administered in combination with a compound of the invention, and either administered separately or in the same pharmaceutical composition, include, but are not limited to the compounds classes and specific compounds listed in Table 1:

Table 1

Alkylating agents	Cyclophosphamide Busulfane Ifosfamide Melphalane Hexamethylmelamine Thiotepa Chlorambucil Dacarbazine Carmustine	Lomustine Procarbazine Altretamine Estramustinephosphate Mechlorethamine Streptozocine Temozolomide Semustine
Platinum agents	Cisplatin Oxaliplatin Spiroplatin Carboxyphthalatoplatinum Tetraplatin Orniplatin Iproplatin	Carboplatin ZD-0473 (AnorMED) Lobaplatin (AeternaZentaris) Satraplatin (Johnson Matthey) BBR-3464 (Hoffmann-La Roche) SM-11355 (Sumitomo) AP-5280 (Access)
Antimetabolites	Azacytidine Gemcitabine Capecitabine 5-Fluoruracil Floxuridine 2-Chlordesoxyadenosine 6-Mercaptopurine 6-Thioguanine Cytarabine 2-Fluordesoxycytidine Methotrexate Idatrexate	Tomudex Trimetrexate Deoxycoformycine Fludarabine Pentostatine Raltitrexede Hydroxyurea Decitabine (SuperGen) Clofarabine (Bioenvision) Irofulven (MGI Pharma) DMDC (Hoffmann-La Roche) Ethynylcytidine (Taiho)
Topoisomerase inhibitors	Amsacrine Epirubicine Etoposide Teniposide or Mitoxantrone Irinotecane (CPT-11) 7-Ethyl-10-hydroxycamptothecine Topotecane Dexrazoxanet (TopoTarget) Pixantrone (Novuspharmna) Rebeccamycin-Analogue (Exelixis) BBR-3576 (Novuspharmna)	Rubitecane (SuperGen) Exatecanmesylate (Daiichi) Quinamed (ChemGenex) Gimatecane (Sigma- Tau) Diflomotecane (Beaufour-Ipsen) TAS-103 (Taiho) Elsamitrucine (Spectrum) J-107088 (Merck & Co) BNP-1350 (BioNumerik) CKD-602 (Chong Kun Dang) KW-2170 (Kyowa Hakko)
Antitumor antibiotics	Dactinomycin (Actinomycin D) Doxorubicin (Adriamycin) Deoxyrubicin	Amonafide Azonafide Anthrapyrazole Oxantrazole

5		Valrubicin Daunorubicin (Daunomycin) Epirubicin Therarubicin Idarubicin Rubidazone Plicamycinp Porfiromycin Cyanomorpholinodoxorubicin Mitoxantron (Novantron)	Losoxantrone Bleomycinsulfate (Blenoxan) Bleomycinacid Bleomycin A Bleomycin B Mitomycin C MEN-10755 (Menarini) GPX-100 (Gem Pharmaceuticals)
10	Antimitotic agents	Paclitaxel Docetaxel Colchicin Vinblastine Vincristine Vinorelbine Vindesine Dolastatine 10 (NCI) Rhizoxine (Fujisawa) Mivobuline (Warner-Lambert) Cemadotine (BASF) RPR 109881A (Aventis) TXD 258 (Aventis) Epothilon B (Novartis) T 900607 (Tularik) T 138067 (Tularik) Cryptophycin 52 (Eli Lilly) Vinflunine (Fabre) Auristatine PE (Teikoku Hormone) BMS 247550 (BMS) BMS 184476 (BMS) BMS 188797 (BMS) Taxoprexine (Protarga)	SB 408075 (GlaxoSmithKline) E7010 (Abbott) PG-TXL (Cell Therapeutics) IDN 5109 (Bayer) A 105972 (Abbott) A 204197 (Abbott) LU 223651 (BASF) D 24851 (ASTA Medica) ER-86526 (Eisai) Combretastatine A4 (BMS) Isohomohalichondrin-B (PharmaMar) ZD 6126 (AstraZeneca) PEG-Paclitaxel (Enzon) AZ10992 (Asahi) IDN-5109 (Indena) AVL B (Prescient NeuroPharma) Azaepothilon B (BMS) BNP- 7787 (BioNumerik) CA-4-Prodrug (OXiGENE) Dolastatin-10 (NrH) CA-4 (OXiGENE)
25	Aromatase inhibitors	Aminoglutethimide Letrozole Anastrozole Formestane	Exemestane Atamestane (BioMedicines) YM-511 (Yamanouchi)
30	Thymidylatesynthas e inhibitors	Pemetrexed (Eli Lilly) ZD-9331 (BTG)	Nolatrexed (Eximias) CoFactor™ (BioKeys)
35	DNA antagonists	Trabectedine (PharmaMar) Glufosfamide (Baxter International) Albumin + 32P (Isotope Solutions) Thymectacine (NewBiotics) Edotreotide (Novartis)	Mafosfamide (Baxter International) Apaziquone (Spectrum Pharmaceuticals) O6-Benzylguanine (Paligent)

	Farnesyltransferase inhibitors	Arglabin (NuOncology Labs) lonafarnibe (Schering-Plough) BAY-43-9006 (Bayer)	Tipifarnibe (Johnson & Johnson) Perillylalcohol (DOR BioPharma)
5	Pump inhibitors	CBT-1 (CBA Pharma) Tariquidar (Xenova) MS-209 (Schering AG)	Zosuquidar-Trihydrochloride (Eli Lilly) Biricodar-Dicitrate (Vertex)
	Histoneacetyltransferase inhibitors	Tacedinaline (Pfizer) SAHA (Aton Pharma) MS-275 (Schering AG)	Pivaloyloxymethylbutyrate (Titan) Depsipeptide (Fujisawa)
10	Metalloproteinase inhibitors / Ribonucleosidereductase inhibitors	Neovastat (Aeterna Laboratories) Marimastat (British Biotech) Galliummaltolate (Titan) Triapine (Vion)	CMT -3 (CollaGenex) BMS-275291 (Celltech) Tezacitabine (Aventis) Didox (Molecules for Health)
15	TNF-alpha agonists/antagonists	Virulizine (Lorus Therapeutics) CDC-394 (Celgene)	Revimide (Celgene)
	Endotheline-A receptor antagonists	Atrasentane (Abbot) ZD-4054 (AstraZeneca)	YM-598 (Yamanouchi)
20	Retinoic acid receptor agonists	Fenretinide (Johnson & Johnson) LGD-1550 (Ligand)	Alitretinoin (Ligand)
25	Immunomodulators	Interferon Oncophage (Antigenics) GMK (Progenics) Adenocarcinoma vaccine (Biomira) CTP-37 (AVI BioPharma) JRX-2 (Immuno-Rx) PEP-005 (Peplin Biotech) Synchrovax vaccine (CTL Immuno) Melanoma vaccine (CTL Immuno) p21-RAS vaccine (GemVax)	Dexosome therapy (Anosys) Pentrix (Australian Cancer Technology) JSF-154 (Tragen) Cancer vaccine (Intercell) Noreline (Biostar) BLP-25 (Biomira) MGV (Progenics) 13-Alethine (Dovetail) CLL-Thera (Vasogen)
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35	Hormonal and anti-hormonal agents	Estrogens Conjugated Estrogens Ethinylestradiol Chlorotrianisen Idenestrole Hydroxyprogesteroncaproate	Prednisone Methylprednisolone Prednisolone Aminoglutethimide Leuprolide Goserelin

5		Medroxyprogesterone Testosterone Testosteronpropionate Fluoxymesterone Methyltestosterone Diethylstilbestrole Megestrole Tamoxifen Toremofine Dexamethasone	Leuporelin Cetrorelix Bicalutamide Flutamide Octreotide Nilutamide Mitotane P-04 (Novogen) 2-Methoxyestradiol (EntreMed) Arzoxifen (Eli Lilly)
10	Photodynamic agents	Talaporfine (Light Sciences) Theralux (Theratechnologies) Motexafin Gadolinium (Pharmacyclics)	Pd-Bacteriopheophorbide (Yeda) Lutetium-Texaphyrine (Pharmacyclics) Hypericine
15	Tyrosinkinase inhibitors	Imatinib (Novartis) Leflunomid (Sugen/Pharmacia) ZD1839 (AstraZeneca) Erlotinib (Oncogene Science) Canertinib (Pfizer) Squalamin (Genaera) SU5416 (Pharmacia) SU6668 (Pharmacia) ZD4190 (AstraZeneca) ZD6474 (AstraZeneca) Vatalanib (Novartis) PKI166 (Novartis) GW2016 (GlaxoSmithKline) EKB-509 (Wyeth) EKB-569 (Wyeth)	Kahalid F (PharmaMar) CEP- 701 (Cephalon) CEP-751 (Cephalon) MLN518 (Millenium) PKC412 (Novartis) Phenoxodiol O Trastuzumab (Genentech) C225 (ImClone) rhu-Mab (Genentech) MDX-H210 (Medarex) 2C4 (Genentech) MDX-447 (Medarex) ABX-EGF (Abgenix) IMC-1C11 (ImClone)
25	Different agents	SR-27897 (CCK-A inhibitor, Sanofi-Synthelabo) Tocladesine (cyclic-AMP agonist, Ribapharm) Alvocidib (CDK inhibitor, Aventis) CV-247 (COX-2-Inhibitor, Ivy Medical) P54 (COX-2 inhibitor, Phytopharm) CapCell™ (CYP450 stimulans, Bavarian Nordic) GCS-100 (gal3 antagonist, GlycoGenesys) G17DT immunogen (Gastrin inhibitor, Aphton) Efaproxiral (Oxygenator,	BCX-1777 (PNP inhibitor, BioCryst) Ranpirnase (Ribonuclease stimulans, AlfaCell) Galarubicin (RNA synthesis inhibitor, Dong-A) Tirapazamin (reducing agent, SRI International) N-Acetylcystein (reducing agent, Zambon) R-Flurbiprofen (NF-kappaB inhibitor, Encore) 3CPA (NF-kappaB inhibitor, Active Biotech) Seocalcitol (Vitamin-D receptor agonist, Leo) 131-I-TM-601 (DNA
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	Allos Therapeutics)	antagonist, TransMolecular)
	PI-88 (Heparanase inhibitor, Progen)	Eflornithin (ODC inhibitor, ILEX Oncology)
	Tesmilifen (Histamine antagonist, YM BioSciences)	Minodronic acid (Osteoclasts inhibitor, Yamanouchi)
	Histamine (Histamine-H2 receptor agonist, Maxim)	Indisulam (p53 stimulans, Eisai)
	Tiazofurin (IMPDH inhibitor, Ribapharm)	Aplidin (PPT inhibitor, PharmaMar)
	Cilengitide (Integrine antagonist, Merck KGaA)	Rituximab (CD20 antibody, Genentech)
	SR-31747 (IL-1 antagonist, Sanofi-Synthelabo)	Gemtuzumab (CD33 antibody, Wyeth Ayerst)
	CCI-779 (mTOR kinase inhibitor, Wyeth)	PG2 (Hematopoiesis enhancer, Pharmagenesis)
	Exisulind (PDE-V inhibitor, Cell Pathways)	Immunol™ (Triclosan oral irrigation, Endo)
	CP-461 (PDE-V inhibitor, Cell Pathways)	Triacetyluridine (Uridine prodrug, Wellstat)
	AG-2037 (GART inhibitor, Pfizer)	SN-4071 (sarcoma agent, Signature BioScience)
	WX-UK1 (Plasminogen activator inhibitor, Willex)	TransMID-107™ (Immunotoxine, KS Biomedix)
	PBI-1402 (PMN stimulans, ProMetic LifeSciences)	PCK-3145 (Apoptosis enhancer, Procyon)
	Bortezomib (Proteasome inhibitor, Millennium)	Doranidazole (Apoptosis enhancer, Pola)
	SRL-172 (T-cell stimulans, SR Pharma)	CHS-828 (cytotoxic agent, Leo)
	TLK-286 (Glutathione-S-transferase inhibitor, Telik)	trans-Retinoic acid (Differentiator, NIH)
	PT-100 (Growth factor agonist, Point Therapeutics)	MX6 (Apoptosis enhancer, MAXIA)
	Midostaurin (PKC inhibitor, Novartis)	Apomin (Apoptosis enhancer, ILEX Oncology)
	Bryostatatin-1 (PKC stimulans, GPC Biotech)	Urocidine (Apoptosis enhancer, Bioniche)
	CDA-II (Apoptosis enhancer, Everlife)	Ro-31-7453 (Apoptosis enhancer, La Roche)
	SDX-101 (Apoptosis enhancer, Salmedix)	Brostallicin (Apoptosis enhancer, Pharmacia)
	Ceflatonin (Apoptosis enhancer, ChemGenex)	

In a preferred embodiment, a compound of the invention is administered in combination with one or more known anti-tumor agents, such as the following: estrogen receptor modulators, androgen receptor modulators, retinoid receptor modulators, cytotoxics, antiproliferative agents, prenyl proteintransferase inhibitors,

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HMG-CoA-reductase inhibitors, HIV protease inhibitors, reverse transcriptase inhibitors, angiogenesis inhibitors.

The compounds of the invention are in particular well suited for administration in combination with radiotherapy. The synergistic effects of VEGF inhibition in combination with radiotherapy are known to the skilled artisan (WO 00/61186).

The term "estrogen receptor modulators" in the course of the present invention refers to compounds that interfere with or inhibit the binding of estrogen to estrogen receptor – independently from the mode of action. Non-limiting examples of estrogen receptor modulators are tamoxifen, raloxifen, idoxifen, LY353381, LY 117081, toremifen, fulvestrant, 4-[7-(2,2-Dimethyl-1-oxopropoxy-4-methyl-2-[4-[2-(1-piperidinyl)ethoxy]phenyl]-2H-1-benzopyran-3-yl]phenyl-2,2-dimethyl-propanoate, 4,4'-Dihydroxybenzophenon-2,4-dinitrophenylhydrazone and SH646.

The term "androgen receptor modulators" in the course of the present invention refers to compounds that interfere with or inhibit the binding of androgens to androgen receptor – independently from the mode of action. Non-limiting examples of androgen receptor modulators are finasteride and other 5alpha-reductase inhibitors, nilutamide, flutamide, bicalutamide, liarozole and abirateron acetate.

The term "retinoid receptor modulators" in the course of the present invention refers to compounds that interfere with or inhibit the binding of retinoids to retinoid receptor – independently from the mode of action. Non-limiting examples of retinoid receptor modulators are bexaroten, tretinoin, 13-cis-retinoic acid, 9-cis-retinoic acid, alpha-difluoromethylornithine, ILX23-7553, trans-N-(4'-Hydroxyphenyl)retinamide and N-4-carboxyphenylretinamide.

The term "cytotoxics" in the course of the present invention refers to compounds that primarily trigger cell death through direct action on cell function(s) or which interfere with or inhibit cell myosis, such as alkylating agents, tumor necrosis factors, intercalating agents, microtubule inhibitors and topoisomerase inhibitors. Non-limiting examples of cytotoxics are tirapazimin, sertenef, cachectine, ifosfamide, tasonermine, lonidamine, carboplatin, altretamine, prednimustine, dibromodulcit, ranimustine, fotemustine, nedaplatin, oxaliplatin, temozolomide, heptaplatin, estramustin, improsulfan-tosylate, trofosfamide, nimustine, dibrospidium-chloride, pumitepa, lobaplatin, satraplatin, profiromycin, cisplatin, irofulven, dexifosfamide, cis-amindichloro(2-methylpyridine)platin, benzylguanine, glufosfamide, GPX100,

(trans,trans,trans)-bis-mu-(hexane-1,6-diamine)-mu-[diamine-platin(II)]bis-[diamine(chloro)platin(II)]-tetrachloride, diarizidinylspermine, arsenium trioxide, 1-(11-Dodecylamino-10-hydroxyundecyl)-3,7-dimethylxanthine, zorubicin, idarubicin, daunorubicin, bisantren, mitoxantron, pirarubicin, pinafide, valrubicine, amrubicine, antineoplaston, 3'-desamino-3'-morpholino-13-desoxo-10-hydroxycarminomycin, annamycin, galarubicin, elinafide, MEN10755 and 4-desmethoxy-3-desamino-3-aziridiny-4-methylsulfonyl-daunorubicin (WO 00/50032).

Non-limiting examples of microtubule inhibitors are paclitaxel, vindesine-sulfate, 3',4'-dideshydro-4'-desoxy-8'-norvincal leukoblastine, docetaxol, rhizoxine, dolastatine, mivobuline-isethionate, auristatine, cemadotone, RPR109881, BMS184476, vinflunine, cryptophycine, 2,3,4,5,6-pentafluoro-N-(3-fluoro-4-methoxyphenyl)-benzenesulfonamide, anhydrovinblastine, N,N-dimethyl-L-valyl-L-valyl-N-methyl-L-valyl-L-prolyl-L-proline-t-butylamide, TDX258 and BMS188797.

Non-limiting examples of topoisomerase inhibitors are topotecane, hycaptamine, irinotecane, rubitecane, 6-ethoxypropionyl-3',4'-O-exo-benzylidene-chartreusine, 9-methoxy-N,N-dimethyl-5-nitropyrazolo[3,4,5-kl]acridine-2-(6H)propanamine, 1-amino-9-ethyl-5-fluoro-2,3-dihydro-9-hydroxy-4-methyl-1H,12H-benzo-[de]-pyrano-[3',4':b,7]indolizino[1,2b]quinoline-10,13(9H,15H)-dione, lurtotecane, 7-[2-(N-isopropylamino)ethyl]-(20S)camptothecin, BNP1350, BNPI1100, BN80915, BN80942, etoposide-phosphate, teniposide, sobuzoxane, 2'-dimethylamino-2'-desoxy-etoposide, GL331, N-[2-(dimethylamino)ethyl]-9-hydroxy-5,6-dimethyl-6H-pyrido[4,3-b]carbazole-1-carboxamide, asulacrine, (5a,5aB,8aa,9b)-9-[2-[N-[2-(dimethylamino)ethyl]-N-methylamino]ethyl]-5-[4-hydroxy-3,5-dimethoxyphenyl]-5,5a,6,8,8a,9-hexahydrofuro(3',4':6,7)naphtho(2,3-d)-1,3-dioxol-6-one, 2,3-(methylenedioxy)-5-methyl-7-hydroxy-8-methoxybenzo[c]phenanthridinium, 6,9-bis[(2-aminoethyl)amino]-benzo[g]isoquinoline-5,10-dione, 5-(3-aminopropylamino)-7,10-dihydroxy-2-(2-hydroxyethylaminomethyl)-6H-pyrazolo[4,5,1-de]acridine-6-one, N-[1-[2(diethylamino)ethylamino]-7-methoxy-9-oxo-9H-thioxane-then-4-ylmethyl]formamide, N-(2-(dimethyl-amino)-ethyl)acridine-4-carboxamide, 6-[[2-(dimethylamino)-ethyl]amino]-3-hydroxy-7H-indeno[2,1-c]quinolin-7-one and dimesna.

Non-limiting examples of antiproliferative agents are antisense RNA- and antisense-DNA oligonucleotides, such as G3139, ODN698, RVASKRAS, GEM231 and INX3001, as well as antimetabolites such as enocitabine, carmofur, tegafur,

pentostatine, doxifluridine, trimetrexate, fludarabine, capecitabine, galocitabine, cytarabin-ocfosfate, fosteabine sodiumhydrate, raltitrexed, paltitrexide, emitetur, tiazofurine, decitabine, nolatrexed, pemetrexed, nelzarabine, 2'-desoxy-2'-methylidencytidine, 2'-fluoromethylen-2'-desoxycytidine, N-[5-(2,3-dihydrobenzofuryl)sulfonyl]-N'-(3,4-dichlorophenyl)urea, N6-[4-desoxy-4-[N2-[2(E),4(E)-tetradecadienoyl]glycylamino]-L-glycero-B-L-manno-heptopyranosyl]adenine, aplidine, ecteinascidine, troxacitabine, 4-[2-amino-4-oxo-4,6,7,8-tetrahydro-3H-pyrimidino[5,4-b][1,4]thiazine-6-yl-(S)-ethyl]-2,5-thienoyl-L-glutaminic acid, aminopterin, 5-fluorouracil, alanosine, 11-acetyl-8-(carbamoyloxymethyl)-4-formyl-6-methoxy-14-oxa-1,11-diaza-tetracyclo-(7.4.1.0.0)-tetradeca-2,4,6-trien-9-ylacetic acid ester, swainsonine, lometrexole, dexrazoxane, methioninase, 2'-cyan-2'-desoxy-N4-palmitoyl-1-B-D-arabinofuranosylcytosine and 3-aminopyridine-2-carboxaldehyde-thiosemicarbazone.

"Antiproliferative agents" also comprises monoclonal antibodies against growth factors that have not been listed under "angiogenesis inhibitors", such as trastuzumab, as well as tumor suppressor genes, such as p53.

In another aspect of the invention, a medicament according to above aspects and embodiments is provided, wherein in such medicament comprises at least one additional pharmacologically active substance (drug, ingredient).

In a preferred embodiment the at least one pharmacologically active substance is a substance as described herein.

In another aspect of the invention, a medicament according to above aspects and embodiments is provided, wherein the medicament is applied before and/or during and/or after treatment with at least one additional pharmacologically active substance.

In a preferred embodiment the at least one pharmacologically active substance is a substance as described herein.

In another aspect of the invention, a pharmaceutical composition comprising a therapeutically effective amount of at least one compound of the invention is provided.

5 In a preferred embodiment, the pharmaceutical composition contains at least one additional compound selected from the group consisting of physiologically acceptable excipients, auxiliaries, adjuvants, diluents, carriers and/or additional pharmaceutically active substance other than the compounds of the invention.

10 In another aspect of the invention, a pharmaceutical composition is disclosed which comprises at least one compound of the invention, at least one pharmacologically active substance other than the compounds of the invention as described herein; and a pharmaceutically acceptable carrier.

15 A further embodiment of the present invention is a process for the manufacture of said pharmaceutical compositions, characterized in that one or more compounds according to the invention and one or more compounds selected from the group consisting of solid, liquid or semiliquid excipients, auxiliaries, adjuvants, diluents, carriers and pharmaceutically active agents other than the compounds according to the invention, are converted in a suitable dosage form.

20 In another aspect of the invention, a kit is provided comprising a therapeutically effective amount of at least one compound of the invention and/or at least one pharmaceutical composition as described herein and a therapeutically effective amount of at least one further pharmacologically active substance other than the compounds of the invention.

30 The pharmaceutical compositions of the present invention may be administered by any means that achieve their intended purpose. For example, administration may be by oral, parenteral, topical, enteral, intravenous, intramuscular, inhalant, nasal, intraarticular, intraspinal, transtracheal, transocular, subcutaneous, intraperitoneal, transdermal, or buccal routes. Alternatively, or concurrently, administration may be by the oral route. The dosage administered will be dependent upon the age, health, and weight of the recipient, kind of concurrent treatment, if any, frequency of

35

treatment, and the nature of the effect desired. Parenteral administration is preferred. Oral administration is especially preferred.

Suitable dosage forms include, but are not limited to capsules, tablets, pellets, dragees, semi-solids, powders, granules, suppositories, ointments, creams, lotions, inhalants, injections, cataplasms, gels, tapes, eye drops, solution, syrups, aerosols, suspension, emulsion, which can be produced according to methods known in the art, for example as described below:

tablets: mixing of active ingredient/s and auxiliaries, compression of said mixture into tablets (direct compression), optionally granulation of part of mixture before compression.

capsules: mixing of active ingredient/s and auxiliaries to obtain a flowable powder, optionally granulating powder, filling powders/granulate into opened capsules, capping of capsules.

semi-solids (ointments, gels, creams): dissolving/dispersing active ingredient/s in an aqueous or fatty carrier; subsequent mixing of aqueous/fatty phase with complementary fatty/ aqueous phase, homogenization (creams only).

suppositories (rectal and vaginal): dissolving/dispersing active ingredient/s in carrier material liquified by heat (rectal: carrier material normally a wax; vaginal: carrier normally a heated solution of a gelling agent), casting said mixture into suppository forms, annealing and withdrawal suppositories from the forms.

aerosols: dispersing/dissolving active agent/s in a propellant, bottling said mixture into an atomizer.

In general, non-chemical routes for the production of pharmaceutical compositions and/or pharmaceutical preparations comprise processing steps on suitable mechanical means known in the art that transfer one or more compounds of the invention into a dosage form suitable for administration to a patient in need of such a treatment. Usually, the transfer of one or more compounds of the invention into such a dosage form comprises the addition of one or more compounds, selected from the group consisting of carriers, excipients, auxiliaries and pharmaceutical active ingredients other than the compounds of the invention.

Suitable processing steps include, but are not limited to combining, milling, mixing,

granulating, dissolving, dispersing, homogenizing, casting and/or compressing the respective active and non-active ingredients. Mechanical means for performing said processing steps are known in the art, for example from Ullmann's Encyclopedia of Industrial Chemistry, 5th Edition. In this respect, active ingredients are preferably at least one compound of the invention and one or more additional compounds other than the compounds of the invention, which show valuable pharmaceutical properties, preferably those pharmaceutical active agents other than the compounds of the invention, which are disclosed herein.

Particularly suitable for oral use are tablets, pills, coated tablets, capsules, powders, granules, syrups, juices or drops, suitable for rectal use are suppositories, suitable for parenteral use are solutions, preferably oil-based or aqueous solutions, furthermore suspensions, emulsions or implants, and suitable for topical use are ointments, creams or powders. The compounds of the invention may also be lyophilised and the resultant lyophilisates used, for example, for the preparation of injection preparations. The preparations indicated may be sterilised and/or comprise assistants, such as lubricants, preservatives, stabilisers and/or wetting agents, emulsifiers, salts for modifying the osmotic pressure, buffer substances, dyes, flavours and/or a plurality of further active ingredients, for example one or more vitamins.

Suitable excipients are organic or inorganic substances, which are suitable for enteral (for example oral), parenteral or topical administration and do not react with the compounds of the invention, for example water, vegetable oils, benzyl alcohols, alkylene glycols, polyethylene glycols, glycerol triacetate, gelatine, carbohydrates, such as lactose, sucrose, mannitol, sorbitol or starch (maize starch, wheat starch, rice starch, potato starch), cellulose preparations and/or calcium phosphates, for example tricalcium phosphate or calcium hydrogen phosphate, magnesium stearate, talc, gelatine, tragacanth, methyl cellulose, hydroxypropylmethylcellulose, sodium carboxymethylcellulose, polyvinyl pyrrolidone and/or vaseline.

If desired, disintegrating agents may be added such as the above-mentioned starches and also carboxymethyl-starch, cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof, such as sodium alginate. Auxiliaries include, without limitation, flow-regulating agents and lubricants, for example, silica, talc, stearic acid or salts thereof, such as magnesium stearate or calcium stearate, and/or polyethylene glycol. Dragee cores are provided with suitable coatings, which, if

desired, are resistant to gastric juices. For this purpose, concentrated saccharide solutions may be used, which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, polyethylene glycol and/or titanium dioxide, lacquer solutions and suitable organic solvents or solvent mixtures. In order to produce coatings resistant to gastric juices or to provide a dosage form affording the advantage of prolonged action, the tablet, dragee or pill can comprise an inner dosage and an outer dosage component the latter being in the form of an envelope over the former. The two components can be separated by an enteric layer, which serves to resist disintegration in the stomach and permits the inner component to pass intact into the duodenum or to be delayed in release. A variety of materials can be used for such enteric layers or coatings, such materials including a number of polymeric acids and mixtures of polymeric acids with such materials as shellac, acetyl alcohol, solutions of suitable cellulose preparations such as acetyl-cellulose phthalate, cellulose acetate or hydroxypropylmethyl-cellulose phthalate, are used. Dye stuffs or pigments may be added to the tablets or dragee coatings, for example, for identification or in order to characterize combinations of active compound doses.

Suitable carrier substances are organic or inorganic substances which are suitable for enteral (e.g. oral) or parenteral administration or topical application and do not react with the novel compounds, for example water, vegetable oils, benzyl alcohols, polyethylene glycols, gelatin, carbohydrates such as lactose or starch, magnesium stearate, talc and petroleum jelly. In particular, tablets, coated tablets, capsules, syrups, suspensions, drops or suppositories are used for enteral administration, solutions, preferably oily or aqueous solutions, furthermore suspensions, emulsions or implants, are used for parenteral administration, and ointments, creams or powders are used for topical application. The compounds of the invention can also be lyophilized and the lyophilizates obtained can be used, for example, for the production of injection preparations.

The preparations indicated can be sterilized and/or can contain excipients such as lubricants, preservatives, stabilizers and/or wetting agents, emulsifiers, salts for affecting the osmotic pressure, buffer substances, colorants, flavourings and/or aromatizers. They can, if desired, also contain one or more further active compounds, e.g. one or more vitamins.

Other pharmaceutical preparations, which can be used orally include push-fit capsules made of gelatine, as well as soft, sealed capsules made of gelatine and a

plasticizer such as glycerol or sorbitol. The push-fit capsules can contain the active compounds in the form of granules, which may be mixed with fillers such as lactose, binders such as starches, and/or lubricants such as talc or magnesium stearate and, optionally, stabilizers. In soft capsules, the active compounds are preferably dissolved or suspended in suitable liquids, such as fatty oils, or liquid paraffin. In addition, stabilizers may be added.

The liquid forms in which the novel compositions of the present invention may be incorporated for administration orally include aqueous solutions, suitably flavoured syrups, aqueous or oil suspensions, and flavoured emulsions with edible oils such as cottonseed oil, sesame oil, coconut oil or peanut oil, as well as elixirs and similar pharmaceutical vehicles. Suitable dispersing or suspending agents for aqueous suspensions include synthetic and natural gums such as tragacanth, acacia, alginate, dextran, sodium carboxymethylcellulose, methylcellulose, polyvinylpyrrolidone or gelatine.

Suitable formulations for parenteral administration include aqueous solutions of the active compounds in water-soluble form, for example, water-soluble salts and alkaline solutions. In addition, suspensions of the active compounds as appropriate oily injection suspensions may be administered. Suitable lipophilic solvents or vehicles include fatty oils, for example, sesame oil, or synthetic fatty acid esters, for example, ethyl oleate or triglycerides or polyethylene glycol-400 (the compounds are soluble in PEG-400).

Aqueous injection suspensions may contain substances, which increase the viscosity of the suspension, including, for example, sodium carboxymethyl cellulose, sorbitol, and/or dextran, optionally, the suspension may also contain stabilizers.

For administration as an inhalation spray, it is possible to use sprays in which the active ingredient is either dissolved or suspended in a propellant gas or propellant gas mixture (for example CO₂ or chlorofluorocarbons). The active ingredient is advantageously used here in micronized form, in which case one or more additional physiologically acceptable solvents may be present, for example ethanol. Inhalation solutions can be administered with the aid of conventional inhalers.

Possible pharmaceutical preparations, which can be used rectally include, for example, suppositories, which consist of a combination of one or more of the active

compounds with a suppository base. Suitable suppository bases are, for example, natural or synthetic triglycerides, or paraffin hydrocarbons. In addition, it is also possible to use gelatine rectal capsules, which consist of a combination of the active compounds with a base. Possible base materials include, for example, liquid
5 triglycerides, polyethylene glycols, or paraffin hydrocarbons.

For use in medicine, the compounds of the present invention will be in the form of pharmaceutically acceptable salts. Other salts may, however, be useful in the preparation of the compounds of the invention or of their pharmaceutically acceptable salts. Suitable pharmaceutically acceptable salts of the compounds of
10 this invention include acid addition salts which may, for example be formed by mixing a solution of the compound according to the invention with a solution of a pharmaceutically acceptable acid such as hydrochloric acid, sulphuric acid, methanesulphonic acid, fumaric acid, maleic acid, succinic acid, acetic acid, benzoic acid, oxalic acid, citric acid, tartaric acid, carbonic acid or phosphoric acid.
15 Furthermore, where the compounds of the invention carry an acidic moiety, suitable pharmaceutically acceptable salts thereof may include alkali metal salts, e.g. sodium or potassium salts; alkaline earth metal salts, e.g. calcium or magnesium salts; and salts formed with suitable organic bases, e.g. quaternary ammonium salts.

20 The pharmaceutical preparations can be employed as medicaments in human and veterinary medicine. As used herein, the term "effective amount" means that amount of a drug or pharmaceutical agent that will elicit the biological or medical response of a tissue, system, animal or human that is being sought, for instance, by a researcher or clinician. Furthermore, the term "therapeutically effective amount"
25 means any amount which, as compared to a corresponding subject who has not received such amount, results in improved treatment, healing, prevention, or amelioration of a disease, disorder, or side effect, or a decrease in the rate of advancement of a disease or disorder. The term also includes within its scope
30 amounts effective to enhance normal physiological function. Said therapeutic effective amount of one or more of the compounds of the invention is known to the skilled artisan or can be easily determined by standard methods known in the art.

The compounds of the invention and the additional active substances are generally administered analogously to commercial preparations. Usually, suitable
35 doses that are therapeutically effective lie in the range between 0.0005 mg and

1000 mg, preferably between 0.005 mg and 500 mg and especially between 0.5 mg and 100 mg per dose unit. The daily dose is preferably between about 0.001 mg/kg and 10 mg/kg of body weight.

5 Those of skill will readily appreciate that dose levels can vary as a function of the specific compound, the severity of the symptoms and the susceptibility of the subject to side effects. Some of the specific compounds are more potent than others. Preferred dosages for a given compound are readily determinable by those of skill in the art by a variety of means. A preferred means is to measure the physiological potency of a given compound.

10 For the purpose of the present invention, all mammalian species are regarded as being comprised. In a preferred embodiment, such mammals are selected from the group consisting of "primate, human, rodent, equine, bovine, canine, feline, domestic animals, cattle, livestock, pets, cow, sheep, pig, goat, horse, pony, donkey, 15 hinny, mule, hare, rabbit, cat, dog, guinea pig, hamster, rat, mouse". More preferably, such mammals are humans. Animal models are of interest for experimental investigations, providing a model for treatment of human diseases.

20 The specific dose for the individual patient depends, however, on the multitude of factors, for example on the efficacy of the specific compounds employed, on the age, body weight, general state of health, the sex, the kind of diet, on the time and route of administration, on the excretion rate, the kind of administration and the dosage form to be administered, the pharmaceutical combination and severity of the 25 particular disorder to which the therapy relates. The specific therapeutic effective dose for the individual patient can readily be determined by routine experimentation, for example by the doctor or physician, which advises or attends the therapeutic treatment.

30 In the case of many disorders, the susceptibility of a particular cell to treatment with the subject compounds may be determined by in vitro testing. Typically a culture of the cell is combined with a subject compound at varying concentrations for a period of time sufficient to allow the active agents to show a relevant reaction, usually between about one hour and one week. For in vitro testing, cultured cells 35 from a biopsy sample may be used.

Even without further details, it is assumed that a person skilled in the art will be able to utilise the above description in the broadest scope. The preferred embodiments should therefore merely be regarded as descriptive disclosure, which is absolutely not limiting in any way.

Above and below, all temperatures are indicated in °C. In the following examples, "conventional work-up" means that, if necessary, the solvent is removed, water is added if necessary, the pH is adjusted, if necessary, to between 2 and 10, depending on the constitution of the end product, the mixture is extracted with ethyl acetate or dichloromethane, the phases are separated, the organic phase is washed with saturated NaHCO₃ solution, if desired with water and saturated NaCl solution, is dried over sodium sulfate, filtered and evaporated, and the product is purified by chromatography on silica gel, by preparative HPLC and/or by crystallisation. The purified compounds are, if desired, freeze-dried.

Mass spectrometry (MS): ESI (electrospray ionisation) (M+H)⁺

List of Abbreviations and Acronyms:

AcOH acetic acid, anh anhydrous, atm atmosphere(s), BOC tert-butoxycarbonyl
CDI 1,1'-carbonyl diimidazole, conc concentrated, d day(s), dec decomposition,
DMAC NN-dimethylacetamide, DMPU 1,3-dimethyl-3,4,5,6-tetrahydro-2(IH)-
pyrimidinone, DMF NN-dimethylformamide, DMSO dimethylsulfoxide, DPPA
diphenylphosphoryl azide, EDCI 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide,
EtOAc ethyl acetate, EtOH ethanol (100%), Et₂O diethyl ether, Et₃N triethylamine, h
hour(s), MeOH methanol, pet. ether petroleum ether (boiling range 30-60°C), temp.
temperature, THF tetrahydrofuran, TFA trifluoroAcOH, Tf trifluoromethanesulfonyl.

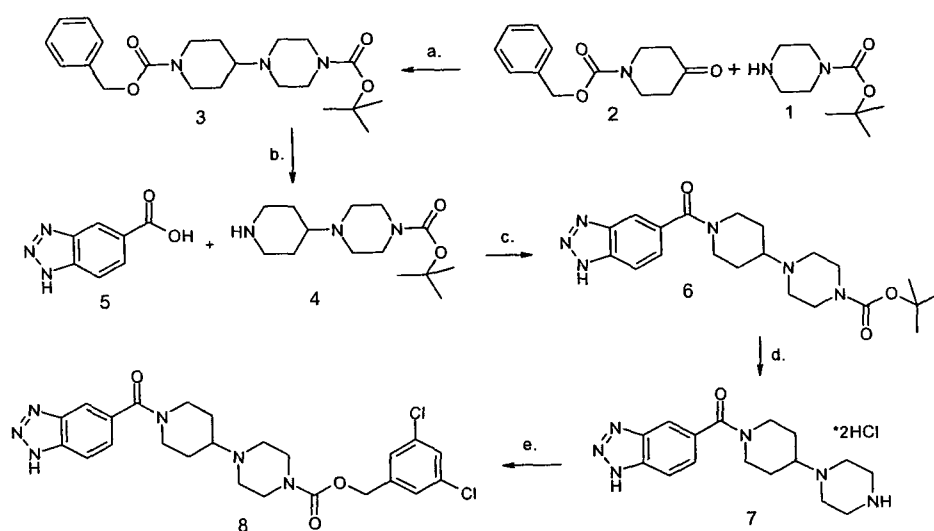
The contents of all cited references are hereby incorporated by reference in their entirety. The invention is explained in more detail by means of the following examples without, however, being restricted thereto.

Examples**I. Synthesis of selected compounds of the invention**

The following compounds were synthesized and characterized. However, it lies in the knowledge of a person skilled in the art to prepare and characterize these compounds differently.

Example 1

Synthesis of 4-[1-(1H-benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-carbonic-acid-3,5-dichloro-benzylester **8**



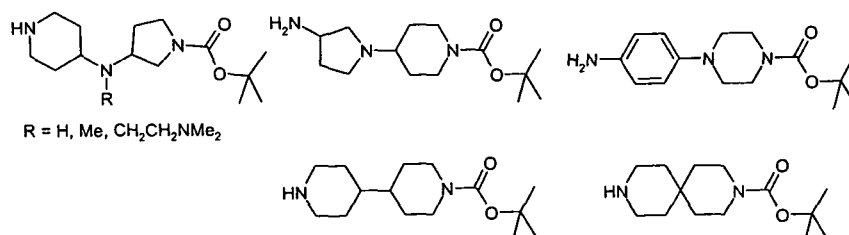
a. N-Boc-piperazine **1** (50.0 g, 268 mmol) and precursor **2** (70.0 g, 300 mmol) were given to MeOH (700 mL), sodium-acetoxaboronhydride (70.0 g, 330 mmol) was added at RT. It was stirred for 15 h at RT. The main part of MeOH was removed in vacuo via a rotating evaporator. The residue was taken up in ethylacetate (400 mL) and washed with water (300 mL). The aqueous phase was extracted with ethylacetate (200 mL), and the organic phases were pooled and dried with sodium sulphate. After filtration, the mixture was concentrated in vacuo until dryness and could be directly used without further processing (colourless oil **3**, 101 g, 255 mmol, 95%).

b. Intermediate **3** (101 g, 255 mmol) was given to THF (1 L), Pd/C-5% (22.0 g, 52,3 % water) was added. It was stirred under hydrogen atmosphere and

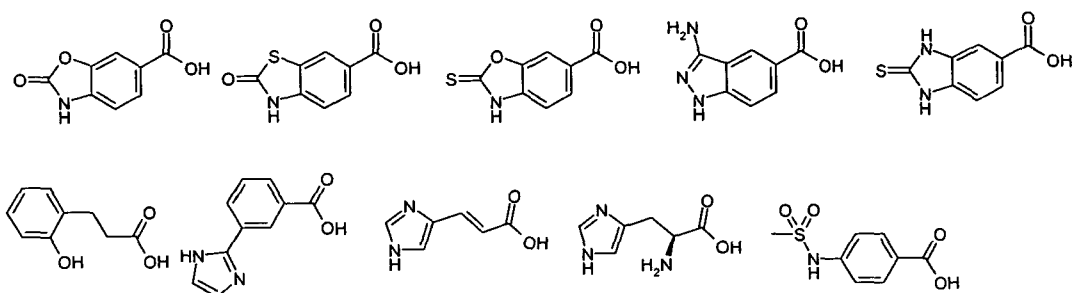
standard pressure for 16 h. Ca. 6 L hydrogen were used up. The reaction mixture was filtrated and concentrated in vacuo until dryness. The yellowish-oily product **4** crystallized slowly und was used without further processing (58.9 g, 219 mmol, 86%).

- c. 1H-benzotriazole-5-carbonic-acid **5** (5.00 g, 18.6 mmol) and intermediate **4** (3.00 g, 18.4 mmol) were added to DMF (30 mL), N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride (3.60 g, 18.8 mmol) and 4-methylmorpholine (2.60 mL, 23.6 mmol) were added at RT. It was stirred for 15 h at RT. The reaction mixture was concentrated in vacuo until dryness and directly purified via column chromatography (gradient with ethylacetate/methanol). A colourless solid was yielded (**6**, 3.40 g, 8.20 mmol, 44%).
- d. Intermediate **6** (3.40 g, 8.20 mmol) was taken up in 2-propanol (10 mL), 6N HCl in 2-Propanol (25 mL) were added and stirring was continued for 5 h at RT. The reaction mixture was concentrated in vacuo until dryness and the residue was comminuted with diethyl ether. The precipitate was filtrated, washed with diethyl ether and dried. Colourless solid **7** as dihydrochloride was yielded (3.15 g, 8.13 mmol, 99%).
- e. 3,5-dichlorobenzylalcohol (75.4 mg, 0.42 mmol) was dissolved in DMF (3 mL), 1,1'-carbonyldiimidazole (68,1 mg, 0.42 mmol) was added, and stirring was continued for 2 h at RT. To this mixture intermediate **7** (148 mg, 0.42 mmol) was added at RT. Stirring was continued for 18 h at RT. The reaction mixture was poured onto water (20 mL) and extracted twice with ethyl acetate. The pooled organic phases were dried with sodium sulphate, filtrated and concentrated in vacuo until dryness. The residue was purified via column chromatography (gradient with ethyl acetate/ methanol). The resulting colourless solid is product **8** (142 mg, 0.27 mmol, 65%).

Analogous to above description, the following amines were used instead of intermediate **4**:

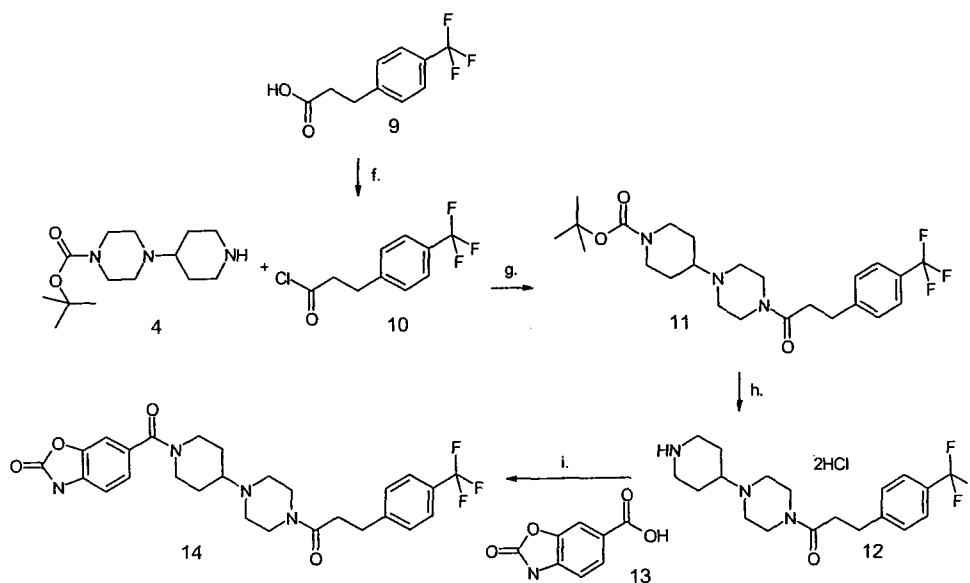


Analogously, intermediate **5** was replaced by the following acids:



Example 2

Synthesis of 6-(4-{4-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperazin-1-yl}-piperidine-1-carbonyl)-3H-benzooxazol-2-one **14**



f. 3-(4-trifluoromethyl-phenyl)-propionic acid **9** (4.70 g, 21.5 mmol) was taken up in thionylchloride (16 mL, 220 mmol) and boiled for 2h under reflux. The reaction mixture was concentrated in vacuo until dryness and was directly used without further purification. A reddish-brown oil was yielded (**10**, 4.87 g, 20.6 mmol, 96%).

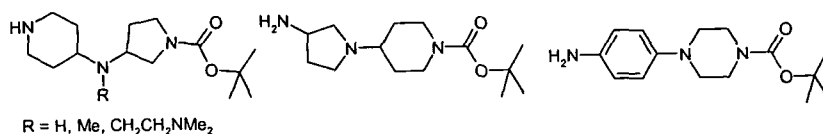
g. Intermediate **4** (515 mg, 1.91 mmol) was given to DMF (5 mL), triethylamine (0.80 mL, 5.74 mmol) was added at RT. Subsequently, at the same temperature the acid chloride **10** (905 mg, 3.83 mmol), dissolved in DMF (1 mL) was added

dropwise. It was stirred for 15 h at RT. The reaction mixture was poured onto water and extracted twice with ethyl acetate. The pooled organic phases were dried over sodium sulphate, filtrated and concentrated in vacuo until dryness. A colourless solid was yielded (**11**, 792 mg, 1.69 mmol, 88%), which was used without further purification.

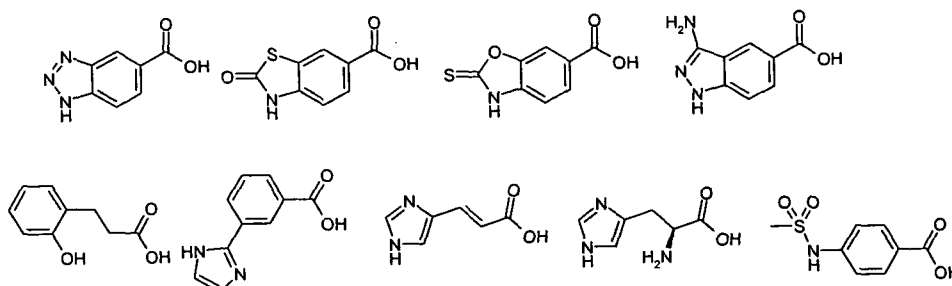
h. Intermediate **11** (0.75 g, 1.60 mmol) was taken up in 6N HCl in 2-propanol (10 mL) and stirring was continued for 1 h at RT. The reaction mixture was concentrated in vacuo until dryness and the residue was comminuted with diethyl ether. The precipitate was filtrated, washed with diethyl ether and dried. A colourless solid **12** was yielded as dihydrochloride (545 mg, 1.26 mmol, 79%).

i. 2-oxo-2,3-dihydro-benzooxazole-6-carbonic-acid **13** (38.6 mg, 0.22 mmol), intermediate **12** (95.5 mg, 0.22 mmol) and 4-methylmorpholine (0.72 mL, 0.66 mmol) were given to DMF (3 mL). N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride (41.4 mg, 0.22 mmol) and 1-hydroxybenzotriazolehydrate (29,2 mg, 0.2 mmol) were added at RT. It was stirred for 15 h at RT. The reaction mixture was poured onto water and extracted twice with ethyl acetate. The pooled organic phases were dried over sodium sulphate, filtrated and concentrated in vacuo until dryness. The oily residue was comminuted with diethyl ether, and the emerging solid was filtered off. A colourless solid (**14**, 50.4 mg, 0.09 mmol, 44%) in high purity was yielded.

Analogous to above description, the following amines were used instead of intermediate **4**:

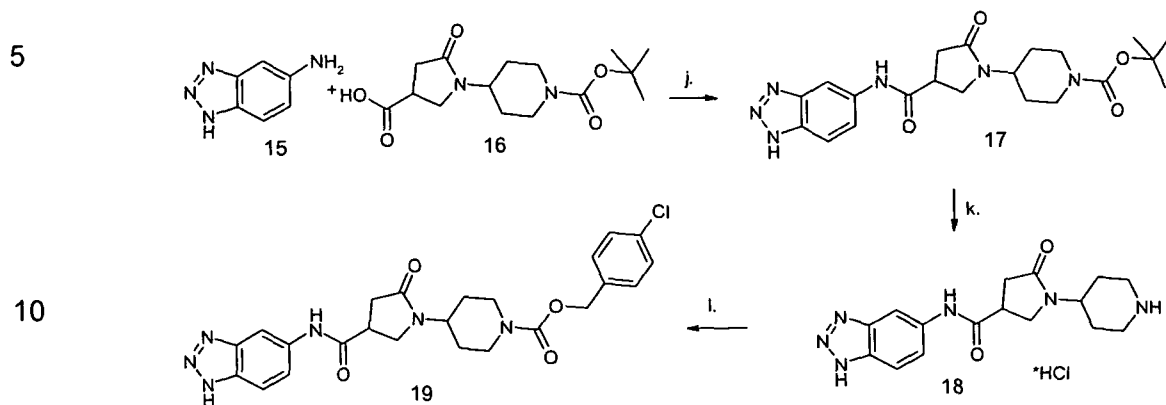


Analogously, intermediate **13** was replaced by the following acids:



Example 3:

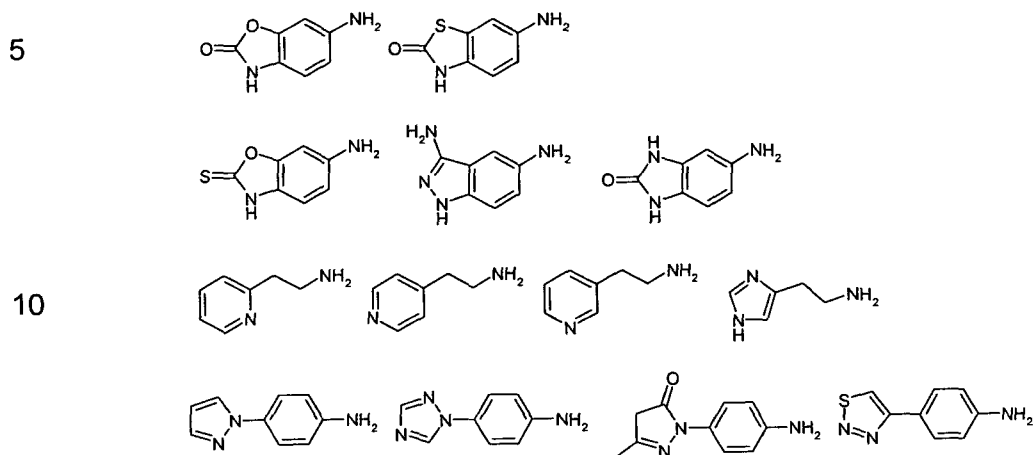
Synthesis of 4-[4-(1H-benzotriazol-5-ylcarbamoyl)-2-oxopyrrolidin-1-yl]-piperidine-1-carbonic-acid-4-chloro-benzylester **19**



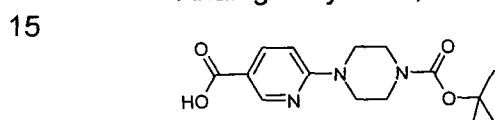
- j. 5-amino-1H-benzotriazole **15** (2.15 g, 16.0 mmol) and commercially available 1-[1-(tert-butoxycarbonyl)-4-piperidinyl]-5-oxo-3-pyrrolidincarbonic-acid **16** (5.00 g, 16.0 mmol) were given to DMF (30 mL). N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride (3.07 g, 16.0 mmol) and 1-hydroxybenzotriazolehydrate (2.16 g, 16.0 mmol) were added at RT. It was stirred for 15 h at RT. The reaction mixture was poured onto water and extracted twice with ethyl acetate. The pooled organic phases were dried over sodium sulphate, filtrated and concentrated in vacuo until dryness. The residue was recrystallized in a bit methanol/diethyl ether and dried. A colourless solid was yielded (**17**, 5.20 g, 12.1 mmol, 76%).
- k. Analogously to h. above, intermediate **17** (5.00 g, 11.7 mmol) in 6N HCl was reacted in 2-propanol (50 mL) (1 h at RT). The reaction mixture was concentrated in vacuo until dryness and the residue was recrystallized from methanol/diethyl ether and dried. A colourless solid **18** was yielded as hydrochloride (2.79 g, 7.65 mmol, 65%).
- l. Analogously to e. above, 4-chlorobenzylalcohol (86.8 mg, 0.61 mmol), 1,1'-carbonyldiimidazole (98,8 mg, 0.61 mmol) and intermediate **18** (222 mg, 0.61 mmol) were reacted at RT in DMF (3 mL). Stirring was continued for 18 h at RT. The reaction mixture was poured onto water and extracted twice with ethyl acetate. The pooled organic phases were dried over sodium sulphate, filtrated and concentrated in vacuo until dryness. The residue was purified via column

chromatography (gradient with ethyl acetate/methanol). The resulting colourless solid is product **19** (157 mg, 0.32 mmol, 52%).

Analogously, the following educts were used instead of educt **15**:

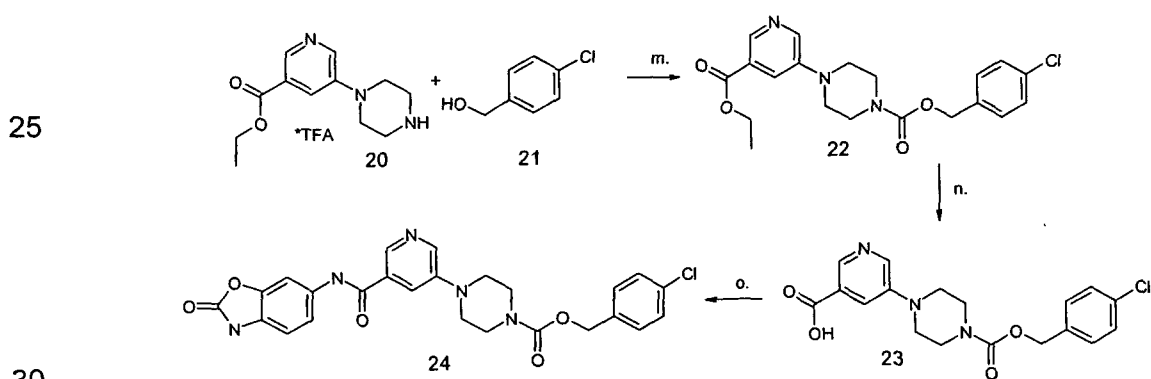


Analogously to **16**, the following commercially available compounds were used:



Example 4:

Synthesis of 4-[5-(2-Oxo-2,3-dihydro-benzooxazol-6-ylcarbamoyl)-pyridin-3-yl]-piperazin-1-carbonic-acid-4-chloro-benzylester **24**:



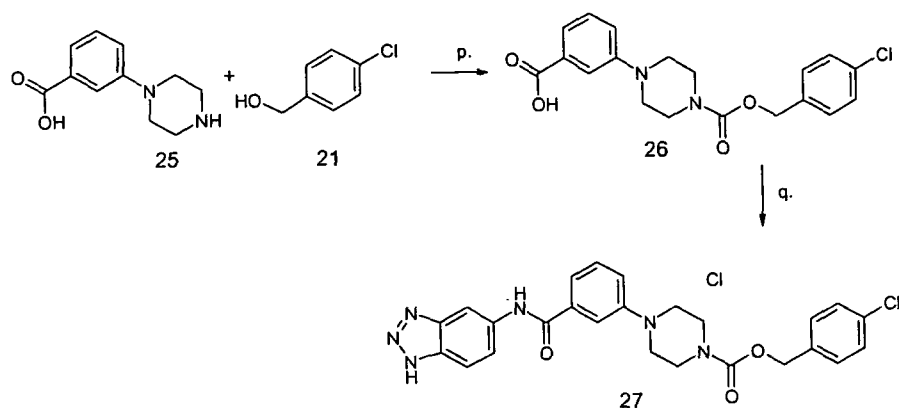
m. Analogously to e. 4-chlorobenzylalcohol (508 mg, 2.86 mmol), 1,1'-carbonyldiimidazole (464 mg, 2.86mmol) and intermediate **20** (1.00 g, 2.86 mmol) were reacted at RT in DMF (5 mL). Stirring was continued for 18 h bei RT

The reaction mixture was poured onto water and the emerging precipitate was filtrated and washed with water. The resulting colourless solid in high purity is intermediate **22** (986 mg, 2.44 mmol, 85%).

- n. Intermediate **22** was taken up in 20 ml ethanol and a solution of LiOH (117 mg, 4.88 mmol) in 2 mL water was added dropwise at RT. Stirring was continued over night at RT. The reaction mixture was concentrated in vacuo until dryness. The residue was taken up in water, adjusted to pH 5 with 2N HCl, the precipitating product filtrated and dried. Intermediate **23** was yielded as colourless solid (812 mg, 2.16 mmol, 88%).
- o. 6-amino-3H-benzooxazol-2-one (59.9 mg, 0.40 mmol) and intermediate **23** (150 mg, 0.40 mmol) were given to DMF (5 mL). N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride (76.5 g, 0.40 mmol) and 1-hydroxybenzotriazole (53.9 mg, 0.40 mmol) were added at RT. It was stirred for 15 h at RT. The reaction mixture was poured onto water and extracted twice with ethyl acetate. The pooled organic phases were dried over sodium sulphate, filtrated and concentrated in vacuo until dryness. The residue was recrystallized from ethyl acetate/diethyl ether and dried. A colourless solid was yielded as product (**24**, 189 mg, 0.37 mmol, 93%).

Example 5:

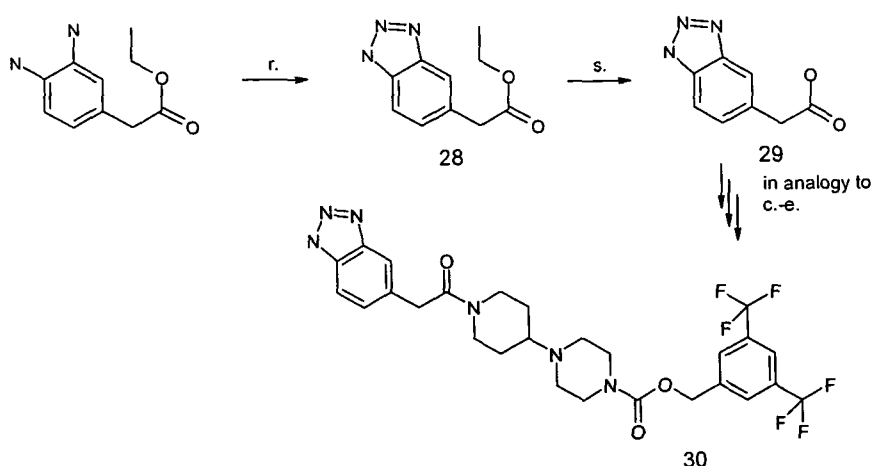
Synthesis of 4-[3-(1H-benzotriazol-5-ylcarbamoyl)-phenyl]-piperazin-1-carbosäure-4-chloro-benzylester **27**:



- p. Analogously to e. 4-chlorobenzylalcohol (200 mg, 1.40 mmol), 1,1'-carbonyldiimidazole (250 mg, 1.54 mmol) and intermediate **25** (289 mg, 1.40 mmol) were reacted at RT in dichloromethan (5 mL). Stirring was continued for 3 d at 50°C in a pressure piston. The reaction mixture was washed threetimes with water. The organic phase was dried over sodium sulphate, filtrated and concentrated in vacuo until dryness. The resulting yellowish oil is intermediate **26** (520 mg, 0.93 mmol, 66%), which was used without further purification.
- q. 5-amino-1H-benzotriazole **15** (40.0 mg, 0.30 mmol), intermediate **26** (230 mg, 0.41 mmol) and N-methylmorpholine (0.10 mL, 0.91 mmol) were given to DMF (3 mL). N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochlorid (80.0 g, 0.42 mmol) and 1-hydroxybenzotriazole (60.0 mg, 0.44 mmol) were added at RT. It was stirred for 15 h at RT. The reaction mixture was concentrated in vacuo and directly purified via column chromatography (gradient with ethyl acetate/methanol). The residue was taken up in a 0.1 M HCl solution in isopropanol and concentrated in vacuo until dryness. A colourless solid was yielded as hydrochloride (**27**, 108 mg, 0.20 mmol, 68%).

Example 6:

Synthesis of 4-[1-(2-1H-Benzotriazol-5-yl-acetyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester



- r. (3,4-Diamino-phenyl)-acetic acid ethyl ester (5.00 g, 25.7 mmol) was dissolved in a mixture of acetic acid (20 mL) and water (20 mL) and cooled to below 5°C in an ice bath. Sodium nitrite (2.66 g, 36.6 mmol) in 20 mL of water was added dropwise to the mixture, kept below 10°C. The reaction mixture was stirred for

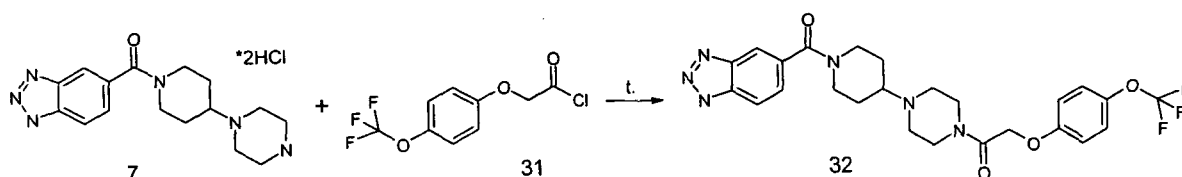
3h at RT. Ethyl acetate was added and the organic layer was washed with water and brine, dried over sodium sulfate and the solvent was removed to give 5,27 g (25.7 mmol, 100%) of a brownish oil, which was used without further purification.

- 5 s. Compound **28** 5,27 g (25.7 mmol) was dissolved in water (25 mL) and EtOH (10 mL) and NaOH (5.00 g, 125 mmol) dissolved in water (75 mL) was added at RT. The solution was stirred at reflux for 3 h and the solvent was removed under vacuum. The residue was dissolved in water and treated with HCl 5-6N in 2-propanol to pH 4. The mixture was stirred and brownish crystals were collected and identified as compound **29** (4.20 g, 19.7 mmol, 77%). The compound was used without further purification.

In analogy to the general procedures c-e compound **30** was synthesized.

15 Example 7:

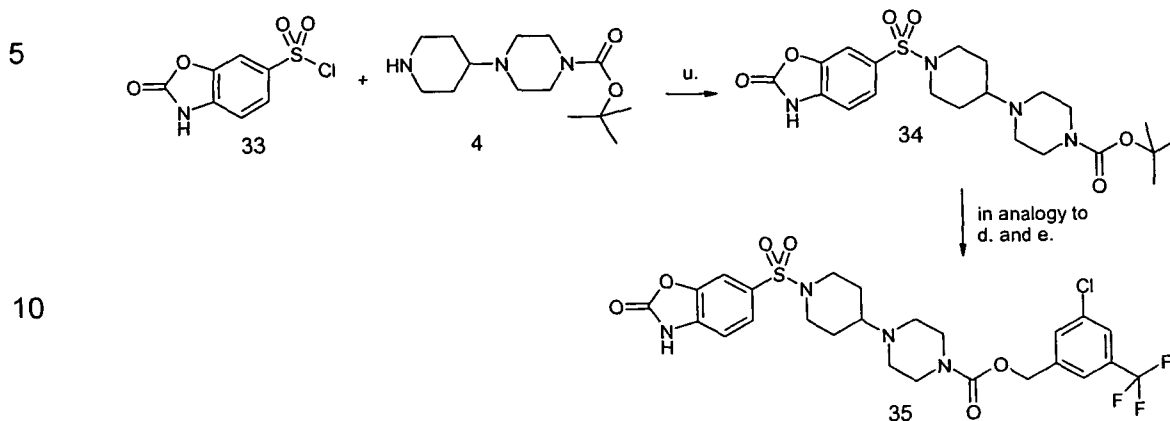
Synthesis of 1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(4-trifluoromethoxy-phenoxy)-ethanone



- 25 t. **7** (100 mg, 0.26 mmol) was dissolved in dichloromethan (2 mL), then triethyl amine (0.11 mL, 0.79 mmol) was added at 0°C followed by compound **31** (130 mg, 0.50 mmol). It was stirred at RT for 18 h. The mixture was diluted with ethyl acetate and washed with water 3 times and with brine once. The organic layer was dried over sodium sulfate, filtered and the solvent evaporated under reduced pressure. The residue was purified by chromatography (methanol/dichloromethane, gradient) to result in a colorless solid identified as **32** (70.9 mg, 0.12 mmol, 48%).

Example 8:

Synthesis of 4-[1-(2-Oxo-2,3-dihydro-benzooxazole-6-sulfonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester

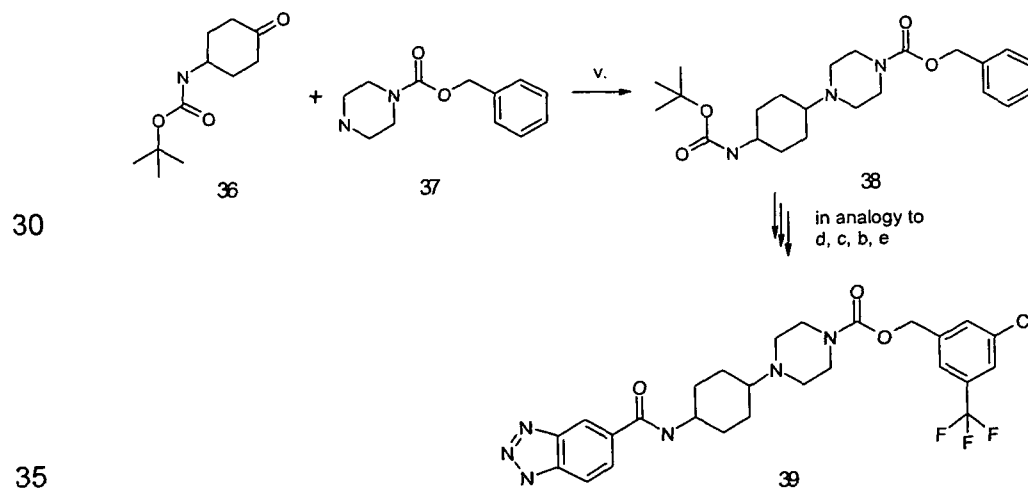


u. **4** (0.54 g, 2.00 mmol) and triethyl amine (0.28 mL, 2.00 mmol) was dissolved in dichloromethane (30 mL) and **33** (0.47 g, 2.00 mmol) was added at RT slowly. It was stirred for additional 16 h at RT. The mixture was washed with water, dried over sodium sulfate and the solvent was evaporated. The residue was purified by chromatography (ethyl acetate) to result in a colorless solid identified as **34** (0.29 g, 0.62 mmol, 31%).

In analogy to the general procedures d-e compound **35** was synthesized.

Example 9:

Synthesis of 4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester

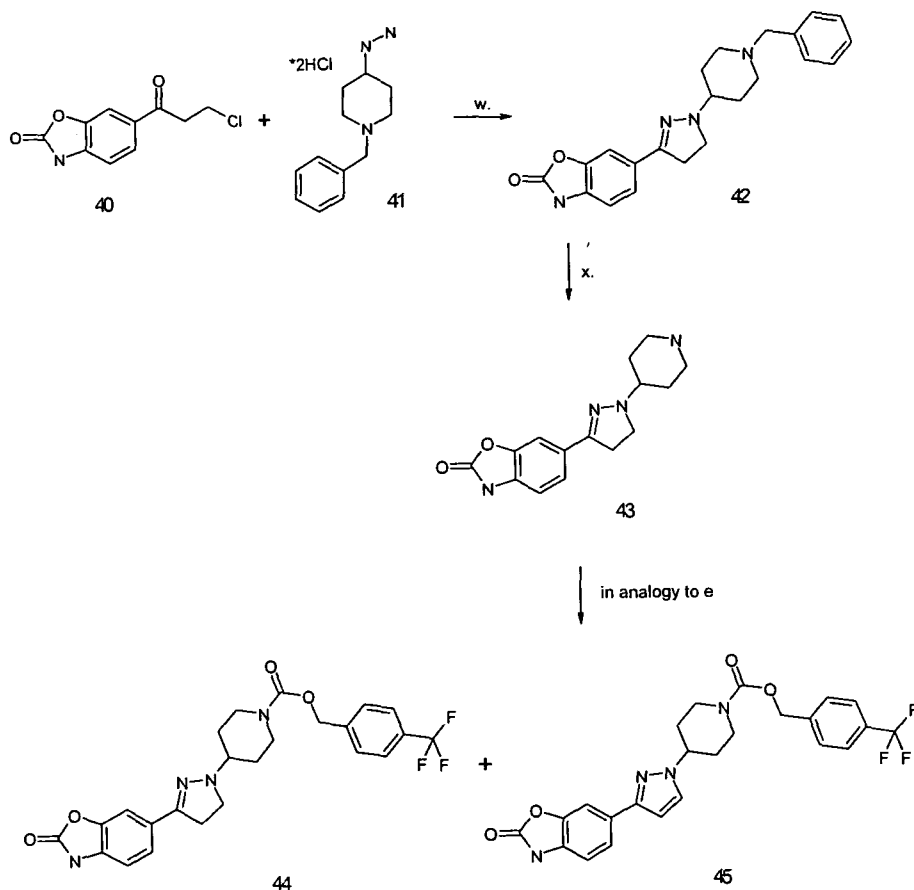


v. Compound **36** (1.20 g, 5.63 mmol) and compound **37** (1.50 g, 6.81 mmol) were dissolved in methanol (100 mL) and sodium triacetoxyborohydride (1.50 g, 7.07 mmol) was added at RT slowly. The mixture was stirred for 18 h at RT. The solvent was evaporated and the residue redissolved in dichloromethane. The organic phase was washed with saturated sodium hydrogencarbonate solution. This aqueous solution was extracted with dichloromethane. The combined organic layers were dried over sodium sulphate, filtered and evaporated to dryness. The crude solid compound **38** (1.90 g, 4.55 mmol, 81%) was used without further purification.

In analogy to the general procedures d, c, b. and e compound **39** was synthesized.

Example 10:

Synthesis of 4-[3-(2-Oxo-2,3-dihydro-benzooxazol-6-yl)-4,5-dihydro-pyrazol-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester and 4-[3-(2-Oxo-2,3-dihydro-benzooxazol-6-yl)-pyrazol-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester



w. **40** (480 mg, 2.13 mmol) and **41** (490 mg, 1.76 mmol) were suspended in ethanol (3 mL). To this suspension triethyl amine (0.73 mL, 5.28 mmol) were added dropwise at RT. It was stirred additional 6 h at RT. The precipitate was filtered off and washed with diethyl ether and dried in vacuo. Compound **42** was obtained as a yellow solid (346 mg, 0.92 mmol), which was used without further purification.

x. **42** (340 mg, 0.90 mmol) was dissolved in methanol (10 mL) Pd/C-5% (52.3 % water, 0.40 g) was added and the mixture was stirred under H₂ atmosphere for 18 h. at RT. The catalyst was filtered off and the solvent was evaporated. 220 mg (0.77 mmol, 85%) of a colorless solid identified as **43** were obtained, which was used without further purification.

In analogy to procedure e the carbamate of compound **43** (70.0 mg, 0.244 mmol) was formed. Besides the expected compound **44** (26 mg, 0.05 mmol, 22%) also compound **45** (26.0 mg, 0.05, 22 %) was isolated by chromatography (dichloromethane/methanol, 1:9).

An overview about further analogously synthesized compounds of the invention including physico-chemical parameters for all compounds of the invention is given in Table 2.

Table 2

Compound	Chemical Name	ESI [M+1] ⁺	HPLC Rt [min] ¹	HPLC/MS Rt [min] ²
1	3-(2-Hydroxy-phenyl)-1-[1'-(3'-trifluoromethyl-biphenyl-2-carbonyl)-[4,4']bipiperidinyl-1-yl]-propan-1-one	566	4,61	2,39
2	4-[4-(1H-Benzotriazol-5-ylcarbamoyl)-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 4-chloro-benzyl ester	498	3,79	1,88
3	5-Oxo-1-{1-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-pyrrolidine-3-carboxylic acid (1H-benzotriazol-5-yl)-amide	530	3,47	1,83
4	4-[4-(1H-Benzotriazol-5-ylcarbamoyl)-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	531 533	3,87	2,02
5	4-[5-(2-Oxo-2,3-dihydro-benzooxazol-6-ylcarbamoyl)-pyridin-3-yl]-piperazine-1-carboxylic acid 4-	509	3,52	1,87

	chloro-benzyl ester			
6	4-[5-(1H-Benzotriazol-5-ylcarbamoyl)-pyridin-3-yl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester	493	3,44	1,83
7	4-{3-[(1H-Benzotriazole-5-carbonyl)-amino]-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 4-chloro-benzyl ester	484	3,17	1,77
8	1H-Benzotriazole-5-carboxylic acid (1-{1-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-pyrrolidin-3-yl)-amide	516	3,23	1,80
9	4-{3-[(1H-Benzotriazole-5-carbonyl)-amino]-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	517 519	3,47	1,91
10	4-[3-(1H-Benzotriazol-5-ylcarbamoyl)-phenyl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester	492	4,32	2,14
11	[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidinyl-1-yl]-(4'-methyl-biphenyl-2-yl)-methanone	509	4,00	2,04
12	1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidinyl-1-carboxylic acid 3,5-dichloro-benzyl ester	516 518	4,69	2,32
13	[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidinyl-1-yl]-(4'-methyl-biphenyl-3-yl)-methanone	509	4,21	2,16
15	4-[2-Oxo-4-(2-oxo-2,3-dihydro-benzooxazol-6-ylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	547 549	3,81	2,05
16	4-[2-Oxo-4-(2-oxo-2,3-dihydro-benzooxazol-6-ylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 4-chloro-benzyl ester	514	3,76	1,89
17	4-[2-Oxo-4-(2-pyridin-2-yl-ethylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	520	3,33	1,90
19	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dichloro-benzyl ester	517 519	3,23	1,76
20	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	517 519	3,41	1,93
21	3-(2-Hydroxy-phenyl)-1-[4-[4-(3'-trifluoromethyl-biphenyl-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl]-propan-1-one	567	3,01	1,87

22	1-[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidinyl-1-yl]-3-(4-trifluoromethyl-phenyl)-propan-1-one	515	3,97	2,21
23	1-[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidinyl-1-yl]-2-(4-chloro-phenyl)-ethanone	467	3,68	2,10
24	1-{1-[4-Methyl-2-(4-trifluoromethyl-phenyl)-thiazole-5-carbonyl]-piperidin-4-yl}-5-oxo-pyrrolidine-3-carboxylic acid (1H-benzotriazol-5-yl)-amide	599	3,71	1,98
25	4-[4-(2-Oxo-2,3-dihydro-benzooxazole-6-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	533 535	3,55	1,83
27	4-[4-(1H-Benzotriazol-5-ylcarbamoyl)-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid tetrahydro-furan-2-ylmethyl ester	458	2,11	1,53
28	4-[2-Oxo-4-(2-pyridin-4-yl-ethylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	519 521	3,33	1,75
29	9-(1H-Benzotriazole-5-carbonyl)-3,9-diaza-spiro[5.5]undecane-3-carboxylic acid 3,5-dichloro-benzyl ester	502 504	4,51	2,25
30	4-[2-Oxo-4-(2-pyridin-3-yl-ethylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	519 521	2,93	1,75
31	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (3,5-dichloro-phenyl)-amide	502 504	3,07	1,71
32	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-trifluoromethyl-phenyl)-propan-1-one	516		1,69
33	6-(4-{4-[3-(4-Trifluoromethyl-phenyl)-propionyl]-piperazin-1-yl}-piperidine-1-carbonyl)-3H-benzooxazol-2-one	532	3,49	1,72
34	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	518	3,23	1,61
35	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	586	3,49	1,75
36	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester	502	3,04	1,43

37	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzyl ester	550	3,55	1,76
38	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 5-chloro-2-methoxy-benzyl ester	514	3,07	1,61
39	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-chloro-benzyl ester	484	2,88	1,62
40	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-methyl-benzyl ester	464	2,96	1,62
41	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-benzyl ester	484	2,99	1,63
42	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester	484	3,04	1,66
43	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-isopropyl-benzyl ester	492	3,41	1,81
45	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,4-dichloro-benzyl ester	517 519	3,25	1,75
46	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,4-dichloro-benzyl ester	517 519	3,25	1,74
47	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethoxy-benzyl ester	534	3,36	1,76
48	(1H-Benzotriazol-5-yl)-(4-[4-methyl-2-(4-trifluoromethyl-phenyl)-thiazole-5-carbonyl]-piperazin-1-yl)-piperidin-1-yl)-methanone	585	3,76	1,79
49	6-(4-{1-[3-(4-Trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-piperazine-1-carbonyl)-3H-benzooxazol-2-one	532	3,15	1,71
50	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-trifluoromethyl-phenyl)-propan-1-one	516	3,52	1,70
51	3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidine-1-carboxylic acid 3,5-dichloro-benzyl ester	517 519	3,36	1,79
52	(1H-Benzotriazol-5-yl)-(4-[4-(5-trifluoromethyl-1H-benzimidazole-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl)-methanone	528	2,91	1,63

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53	4-{4-[2-(1H-Imidazol-4-yl)-ethylcarbamoyl]-2-oxo-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	508 510	3,33	1,81
54	4-[2-Oxo-4-(4-[1,2,4]triazol-1-yl-phenylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	557 559	4,85	2,10
55	4-[2-Oxo-4-(4-pyrazol-1-yl-phenylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	556 558	5,33	2,28
56	4-{4-[4-(3-Methyl-5-oxo-4,5-dihydro-pyrazol-1-yl)-phenylcarbamoyl]-2-oxo-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	586 588	4,59	2,01
57	4-[2-Oxo-4-(4-[1,2,3]thiadiazol-4-yl-phenylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	574 576	5,49	2,29
58	(E)-3-(3H-Imidazol-4-yl)-1-(4-{1-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-piperazin-1-yl)-propanone	491	2,83	1,58
59	N-[4-(4-{1-[3-(4-Trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-piperazine-1-carbonyl)-phenyl]-methanesulfonamide	568	3,31	1,76
60	1-(4-{4-[3-(1H-Imidazol-2-yl)-benzoyl]-piperazin-1-yl}-piperidin-1-yl)-3-(4-trifluoromethyl-phenyl)-propan-1-one	541	2,91	1,63
61	1-(4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl)-3-(4-chloro-2-fluoro-phenyl)-propan-1-one	500		1,64
62	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester	502	3,20	1,72
63	3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester	502	3,12	1,66
64	6-(4-{4-[3-(4-Chloro-2-fluoro-phenyl)-propionyl]-piperazin-1-yl}-piperidine-1-carbonyl)-3H-benzooxazol-2-one	516	3,01	1,64
65	4-[1-(2-Oxo-2,3-dihydro-benzooxazole-6-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester	518	3,17	1,71
66	1-{3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidin-1-yl}-3-(4-chloro-2-fluoro-phenyl)-propan-1-one	500	2,96	1,64

67	1-[4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl]-3-(4-chloro-2-fluoro-phenyl)-propan-1-one	500	2,91	1,62
68	3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	518	3,29	1,57
69	1-[4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl]-3-(3,5-dichloro-phenyl)-propan-1-one	515 517	3,15	1,57
70	4-[1-(1H-Benzotriazole-5-carbonyl)-pyrrolidin-3-ylamino]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	518	3,23	1,58
71	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	518	3,23	1,63
72	2-[4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl]-N-(4-chloro-3-trifluoromethyl-phenyl)-2-oxo-acetamide	565	3,28	1,65
73	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 2-fluoro-4-trifluoromethyl-benzyl ester	536	3,41	1,80
74	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 2-fluoro-5-trifluoromethyl-benzyl ester	536	3,31	1,74
75	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-fluoro-5-trifluoromethyl-benzyl ester	536	3,41	1,76
76	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethoxy-benzyl ester	534	3,41	1,76
77	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester	552	3,57	1,90
78	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-4-fluoro-benzyl ester	502	3,23	1,61
79	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,4,5-trifluoro-benzyl ester	503	3,2	1,63
80	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-bromo-benzyl ester	528	3,28	1,69

81	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-nitro-phenyl)-propan-1-one	493	2,61	1,44
82	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(2,4-dichloro-phenyl)-propan-1-one	515 517	3,23	1,69
83	(E)-1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-bromo-2-fluoro-phenyl)-propenone	542	3,17	1,64
84	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-isopropyl-phenyl)-propan-1-one	490	3,36	1,72
85	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,4-dichloro-benzyl ester	517 519	3,33	1,78
86	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzyl ester	550	3,6	1,83
87	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-isopropyl-benzyl ester	492	3,47	1,81
88	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-methanesulfonyl-phenyl)-propan-1-one	526	1,36	1,20
89	4-{4-[(S)-2-Amino-3-(1H-imidazol-4-yl)-propionyl]-piperazin-1-yl}-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	510	2,83	1,44
90	(E)-1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(3,5-dichloro-phenyl)-propenone	513 515	3,28	1,74
91	(1H-Benzotriazol-5-yl)-{4-[1-(3,5-dichloro-4-fluoro-benzoyl)-piperidin-4-yl]-piperazin-1-yl}-methanone	505 507	3,01	1,56
92	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-4-trifluoromethyl-benzyl ester	552	3,55	1,76
93	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-4-fluoro-benzyl ester	535 537	3,44	1,76
94	4-(3-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-oxo-propyl)-benzonitrile	473	2,11	1,36
95	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-	552	3,41	1,78

	trifluoromethyl-benzyl ester			
96	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-4-trifluoromethyl-benzyl ester	552	3,39	1,77
97	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dichloro-4-fluoro-benzyl ester	535 537	3,31	1,74
98	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dibromo-4-methyl-benzyl ester	621	3,60	1,78
99	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 5-chloro-2-fluoro-3-trifluoromethyl-benzyl ester	570	3,36	1,72
100	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-fluoro-3-trifluoromethyl-benzyl ester	536	3,15	1,63
101	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-4-trifluoromethoxy-benzyl ester	568	3,44	1,77
102	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(3-fluoro-4-trifluoromethyl-phenyl)-ethanone	520	3,04	1,53
103	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(2-fluoro-4-trifluoromethyl-phenyl)-ethanone	520	3,01	1,53
104	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,5-dichloro-4-fluoro-phenyl)-propan-1-one	534	3,28	1,60
105	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3-fluoro-4-trifluoromethyl-phenyl)-propan-1-one	534	3,25	1,61
106	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-fluoro-3,5-dimethyl-phenyl)-propan-1-one	494	3,09	1,56
107	4-[3-(2-Oxo-2,3-dihydro-benzooxazol-6-yl)-4,5-dihydro-pyrazol-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	489	4,08	2,43
108	4-[3-(2-Oxo-2,3-dihydro-benzooxazol-6-yl)-pyrazol-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	487	4,69	

109	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylmethyl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	532	3,23	1,62
110	4-{5-[(1H-Benzotriazole-5-carbonyl)-amino]-pyridin-2-yl}-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	526		2,05
111	4-[5-(1H-Benzotriazol-5-ylcarbonyl)-pyridin-2-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	526		2,27
112	4-[4-(3-Amino-1H-indazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	532		1,653
113	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,3-difluoro-4-trifluoromethyl-benzyl ester	554	3,49	1,712
114	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,3,5,6-tetrafluoro-4-trifluoromethyl-benzyl ester	589	3,57	1,759
115	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-trifluoromethyl-benzyl ester	518	3,31	1,651
116	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-4,5-difluoro-benzyl ester	520	3,31	1,636
117	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-3-fluoro-benzyl ester	502	3,23	1,639
118	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-fluoro-4-trifluoromethoxy-benzyl ester	552	3,52	1,741
119	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-methyl-3-trifluoromethyl-benzyl ester	532	3,44	1,735
120	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-methoxy-3,5-dimethyl-benzyl ester	508	3,09	1,598
121	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,4,5-trimethoxy-benzyl ester	540	2,45	1,409
122	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzylamide	549	3,17	1,673

123	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-chloro-phenyl)-propane-1,2-dione	496	2,93	1,539
124	(1H-Benzotriazol-5-yl)-{4-[4-(4'-trifluoromethyl-biphenyl-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-methanone	564	3,15	1,625
125	(1H-Benzotriazol-5-yl)-{4-[4-(3'-trifluoromethyl-biphenyl-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-methanone	564	3,2	1,653
126	4-[1-(2-Oxo-2,3-dihydro-benzothiazole-6-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	550	3,33	1,732
127	4-[1-(2-Thioxo-2,3-dihydro-1H-benzoimidazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	549	3,17	1,627
128	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-trifluoromethoxy-benzyl ester	534	3,36	1,696
129	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3-chloro-4,5-difluoro-phenyl)-propan-1-one	518	3,2	1,601
130	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,5-dibromo-4-methyl-phenyl)-propan-1-one	617, 619, 621	3,6	1,84
131	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(4-chloro-phenyl)-1H-[1,2,3]triazol-4-ylmethyl ester	551	2,99	1,553
132	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(4-trifluoromethyl-phenyl)-ethyl ester	532	3,41	1,733
133	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(3,5-dichloro-4-fluoro-phenyl)-propan-1-one	534	3,28	1,694
134	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-methyl-2H-indazol-3-ylmethyl ester	504	1,71	1,352
135	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid [3,3']bithiophenyl-5-ylmethyl ester	538	3,31	1,66
136	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid benzo[1,3]dioxol-5-	494	2,48	1,378

	ylmethyl ester			
137	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,3-dihydro-benzo[1,4]dioxin-6-ylmethyl ester	508	2,51	1,445
138	2-(1H-Benzotriazol-5-yl)-1-(4-{4-[2-(4-trifluoromethyl-phenyl)-acetyl]-piperazin-1-yl}-piperidin-1-yl)-ethanone	516	3,09	1,529
139	4-[1-(2-1H-Benzotriazol-5-yl-acetyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	600	3,68	1,853
140	4-[1-(2-1H-Benzotriazol-5-yl-acetyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (3,5-dichloro-phenyl)-amide	517	3,23	1,659
141	2-(1H-Benzotriazol-5-yl)-1-(4-{4-[2-(3,5-bis-trifluoromethyl-phenyl)-acetyl]-piperazin-1-yl}-piperidin-1-yl)-ethanone	584	3,52	1,769
142	6-(4-{4-[2-(3,5-Bis-trifluoromethyl-phenyl)-acetyl]-piperazin-1-yl}-piperidine-1-carbonyl)-3H-benzooxazol-2-one	586	3,6	1,812
143	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-4-trifluoromethoxy-benzyl ester	568	3,65	1,751
144	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 2,3,5,6-tetrafluoro-4-trifluoromethyl-benzyl ester	589	3,6	1,753
145	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-3-trifluoromethoxy-benzyl ester	568	3,6	1,794
146	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-bromo-5-chloro-benzyl ester	561, 563	3,41	1,642
147	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dimethyl-benzyl ester	478	3,17	1,63
148	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dibromo-benzyl ester	605, 607, 609	3,41	1,732
149	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-methylsulfanyl-benzyl ester	496	3,04	1,592

150	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dimethoxy-benzyl ester	510	2,77	1,496
151	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-ethyl-benzyl ester	478	3,17	1,66
152	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-fluoro-benzyl ester	502	3,09	1,603
153	4-[1-(2-Oxo-2,3-dihydro-benzothiazole-6-sulfonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	654	4,03	1,978
154	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-bromo-5-fluoro-benzyl ester	545, 547	3,15	1,672
155	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-bromo-benzyl ester	527, 529	3,07	1,627
156	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-bromo-2-fluoro-benzyl ester	545, 547	3,12	1,667
157	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-bromo-2,6-difluoro-benzyl ester	563, 565	3,12	1,664
158	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-fluoro-4-trifluoromethyl-benzyl ester	536	3,31	1,687
159	(1H-Benzotriazol-5-yl)-(4-[4-(5-bromo-benzofuran-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl)-methanone	537, 539	3,09	1,564
160	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(4-trifluoromethoxy-phenoxy)-ethanone	534	3,09	1,645
161	4-[1-(2-Thioxo-2,3-dihydro-1H-benzimidazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	617	3,65	1,821
162	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	586	3,73	1,888
163	4-[1-(2-Oxo-2,3-dihydro-benzothiazole-6-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	618	3,79	1,851

164	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-trifluoromethyl-phenyl)-propane-1,3-dione	530	3,04	1,533
165	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,4-difluoro-5-trifluoromethyl-benzyl ester	554	3,41	1,695
166	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-bromo-4-trifluoromethoxy-benzyl ester	611, 613	3,63	1,806
167	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-difluoromethoxy-benzyl ester	516	3,09	1,549
168	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-difluoromethylsulfanyl-benzyl ester	532	3,28	1,649
169	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (R)-1-(3,5-bis-trifluoromethyl-phenyl)-ethyl ester	600	3,73	1,884
170	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,2,2-trifluoro-1-p-tolyl-1-trifluoromethyl-ethyl ester	600	3,73	1,914
171	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (S)-1-(3,5-bis-trifluoromethyl-phenyl)-ethyl ester	600	3,68	1,878
172	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(3,5-bis-trifluoromethyl-phenyl)-ethyl ester	600	3,63	1,856
173	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-methyl-5-phenyl-furan-3-ylmethyl ester	530	3,49	1,764
174	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-(4-chloro-phenyl)-4-methyl-thiazol-5-ylmethyl ester	581	3,39	1,757
175	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-(4-chloro-phenyl)-2-oxo-oxazolidin-5-ylmethyl ester	569	2,8	1,512
176	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,4-dimethyl-phenyl)-butan-1-one	490	2,8	1,604
177	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-trifluoromethyl-phenyl)-butan-1-one	530	3,07	1,672

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178	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(3,5-bis-trifluoromethyl-phenyl)-cyclopropyl ester	612	3,65	1,844
179	(1H-Benzotriazol-5-yl)-(4-{4-[2-(4-chloro-phenyl)-cyclopropanecarbonyl]-piperazin-1-yl}-piperidin-1-yl)-methanone	494	3,04	1,618
180	2-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-N-(3,5-bis-trifluoromethyl-phenyl)-acetamide	585	3,12	1,628
181	2-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-N-(5-chloro-2-methoxy-phenyl)-acetamide	513	3,97	1,391
182	4-[[1-(1H-Benzotriazole-5-carbonyl)-pyrrolidin-3-yl]-methyl-amino]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	532		
183	4-[1-(2-Oxo-2,3-dihydro-benzooxazole-6-sulfonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester	604	3,33	1,882
184	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-3-oxo-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	600	3,13	2,177
185	4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid benzyl ester	464	2,29	1,434
186	4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	600	2,61	1,855
187	4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester	566	3,65	1,754
188	4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzyl ester	564		1,725
189	4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3-fluoro-4-trifluoromethyl-benzyl ester	550		1,725

In the following ¹H-NMR data for selected compounds of the invention are displayed:

Compound 4, $C_{24}H_{24}Cl_2N_6O_4$ ($C_{24}H_{22}(D_2)Cl_2N_6O_4$)

1H NMR (500 MHz, DMSO- d_6 , d-TFA exchanged) δ [ppm] = 8.35 (d, J = 1.3, 1H), 7.84 (d, J = 8.9, 1H), 7.40 (dd, J = 9.0, 1.7, 2H), 7.34 (s, 2H), 5.04 (s, 2H), 4.11 (d, J = 13.0, 2H), 3.98 (dt, J = 10.6, 5.9, 1H), 3.59 (t, J = 9.2, 1H), 3.48-3.41 (m, 1H), 3.40-3.26 (m, 1H), 2.86 (s, 2H), 2.57 (d, J = 8.2, 2H), 1.57 (d, J = 17.6, 4H).

Compound 5, $C_{25}H_{22}ClN_5O_5$

1H NMR (500 MHz, DMSO- d_6) δ [ppm] = 11.57 (s, 1H), 10.35 (s, 1H), 8.51 (dd, J = 21.2, 2.2, 2H), 7.87 – 7.68 (m, 2H), 7.53 – 7.33 (m, 5H), 7.08 (d, J = 8.4, 1H), 5.11 (s, 2H), 3.58 (s, 4H), 3.34-2.99 (m, 4H, overlapped with water signal).

Compound 7, $C_{24}H_{27}ClN_6O_3$

1H NMR (500 MHz, DMSO- d_6) δ [ppm] = 8.60 (d, J = 6.9, 1H), 8.47 (s, 1H), 7.99-7.84 (m, 2H), 7.44 (d, J = 9.0, 2H), 7.35 (d, J = 9.0, 2H), 5.05 (s, 2H), 4.45-4.36 (m, 1H), 4.07 (s, 1H), 3.88 (d, J = 11.9, 2H), 3.01-2.86 (m, 3H), 2.79-2.67 (m, 1H), 2.60-2.52 (m, 2H), 2.24 (t, J = 9.6, 1H), 2.20-2.08 (m, 1H), 1.88-1.74 (m, 3H), 1.35-1.25 (m, 2H).

Compound 9, $C_{24}H_{26}Cl_2N_6O_3$

1H NMR (500 MHz, DMSO- d_6) δ [ppm] = 16.53-15.23 (m, 1H), 8.61 (d, J = 6.9, 1H), 8.48 (s, 1H), 7.93 (q, J = 8.6, 2H), 7.55 (t, J = 1.8, 1H), 7.41 (d, J = 1.8, 2H), 5.06 (s, 2H), 4.42 (d, J = 7.6, 1H), 3.89 (d, J = 10.0, 2H), 3.05-2.85 (m, 4H), 2.73 (s, 1H), 2.61-2.52 (d, 1H), 2.27 (s, 1H), 2.20-2.05 (m, 2H), 1.91-1.70 (m, 4H), 1.43-1.25 (m, 2H).

Compound 12, $C_{25}H_{27}Cl_2N_5O_3$

1H NMR (500 MHz, DMSO- d_6) δ [ppm] = 8.00-7.88 (m, 2H), 7.56 (t, J = 1.9, 1H), 7.45-7.35 (m, 3H), 5.07 (s, 2H), 4.51 (s, 1H), 4.04 (d, J = 12.9, 2H), 2.90-2.65 (5H), 1.88-1.50 (m, 5H), 1.46-1.28 (m, 2H), 1.28-1.12 (d, J = 26.3, 2H), 1.12-1.03 (m, 2H).

Compound 13, $C_{31}H_{33}N_5O_2$

1H NMR (500 MHz, DMSO- d_6) δ [ppm] = 15.83 (s, 1H), 7.93 (s, 2H), 7.70 (d, J = 7.7, 1H), 7.58 (d, J = 7.6, 3H), 7.50 (t, J = 7.6, 1H), 7.43 (s, 1H), 7.30 (dd, J = 15.3, 7.7, 3H), 4.54 (s, 2H), 3.64 (s, 2H), 3.15-2.84 (m, 2H), 2.63-2.82 (m, 2H), 2.35 (s, 3H), 1.92-1.50 (m, 4H), 1.41 (s, 2H), 1.08-1.28 (m, 4H).

Compound 16, $C_{25}H_{25}ClN_4O_6$

1H NMR (500 MHz, DMSO- d_6) δ [ppm] = 11.52 (s, 1H), 10.12 (s, 1H), 7.68 (d, J = 1.9, 1H), 7.44 (d, J = 8.5, 2H), 7.39 (d, J = 8.5, 2H), 7.24 (dd, J = 8.4, 1.9, 1H), 7.03 (d, J = 8.4, 1H), 5.07 (s, 2H), 4.08 (d, J = 12.5, 2H), 4.02-3.87 (m, 1H), 3.56 (t, J = 9.1, 1H), 3.41 (dd, J = 9.6, 6.2, 1H), 3.30-3.17 (m, 1H), 2.89 (s, 2H), 2.62-2.52 (m, 2H), 1.69-1.47 (m, 4H).

Compound 19, $C_{24}H_{26}Cl_2N_6O_3$

1H NMR (500 MHz, DMSO- d_6) δ [ppm] = 15.83 (s, 1H), 7.95 (s, 2H), 7.57 (t, J = 1.8, 1H), 7.45 (d, J = 8.4, 1H), 7.42 (d, J = 1.8, 2H), 5.08 (s, 2H), 4.69-4.32 (m, 1H), 3.70-3.35 (m, 5H), 3.14-2.70 (m, 3H), 2.62-2.52 (m, 4H), 1.91-1.60 (m, 2H), 1.50-1.37 (m, 2H).

Compound 25, $C_{25}H_{26}Cl_2N_4O_5$

1H NMR (500 MHz, DMSO- d_6) δ [ppm] = 11.75 (s, 1H), 7.55 (t, J = 1.9, 1H), 7.41 (d, J = 1.9, 2H), 7.32 (d, J = 1.4, 1H), 7.18 (dd, J = 8.0, 1.4, 1H), 7.11 (d, J = 8.0, 1H), 5.06 (s, 2H), 4.01 (d, J = 13.2, 2H), 3.46 (s, 3H), 2.95-2.71 (m, 3H), 2.48-2.38 (m, 4H), 1.74 (d, J = 10.9, 2H), 1.38-1.25 (m, 3H).

Compound 34, $C_{25}H_{27}F_3N_6O_3$

1H NMR (500 MHz, DMSO- d_6) δ [ppm] = 15.83 (s, 1H), 7.94 (s, 2H), 7.74 (d, J = 8.1, 2H), 7.57 (d, J = 8.0, 2H), 7.44 (d, J = 8.3, 1H), 5.17 (s, 2H), 4.41 (s, 1H), 3.60 (s, 1H), 3.47-3.34 (m, 5H), 3.14-2.68 (m, 2H), 2.60-2.53 (m, 4H), 1.94-1.58 (m, 2H), 1.50-1.35 (m, 2H).

Compound 51, $C_{24}H_{26}Cl_2N_6O_3$ ($C_{24}H_{24}(D_2)Cl_2N_6O_3$)

1H NMR (500 MHz, DMSO- d_6 , d-TFA exchanged) δ [ppm] = 8.01-7.96 (m, 2H),
7.53-7.51 (m, 1H), 7.47 (dd, J = 8.67, 1.00, 1H), 7.44 (s, 2H), 5.13 (d, J = 3.1, 2H),
4.60 (s, 1H), 4.07-3.97 (m, 1H), 3.82-3.70 (m, 1H), 3.68-3.28 (m, 4H), 3.28-2.81 (m,
3H), 2.37-2.23 (m, 1H), 2.20-2.07 (m, 3H), 1.55 (s, 2H).

Compound 63, $C_{24}H_{26}ClFN_6O_3$

1H NMR (500 MHz, DMSO- d_6) δ [ppm] = 7.95 (d, J = 8.5, 1H), 7.90 (s, 1H), 7.51-
7.39 (m, 3H), 7.31 (d, J = 8.1, 1H), 5.07 (s, 2H), 4.31 (s, 1H), 3.63-3.34 (m, 4H),
3.12-2.94 (m, 4H), 2.71 (s, 1H), 2.55-2.50 (m, 2H), 2.04-1.52 (m, 4H), 1.22 (s, 2H).

Compound 75, $C_{25}H_{26}F_4N_6O_3$

1H NMR (500 MHz, DMSO- d_6) δ [ppm] = 15.88 (s, 1H), 7.94 (s, 2H), 7.63 (d, J =
9.0, 1H), 7.58 (s, 1H), 7.54 (d, J = 9.0, 1H), 7.44 (d, J = 8.1, 1H), 5.16 (s, 2H), 4.05
(d, J = 9.2, 2H), 3.62 (s, 2H), 3.44-3.32 (m, 2H), 3.17 (d, J = 5.1, 1H), 2.95-2.72 (m,
2H), 2.60-2.50 (m, 4H), 1.75 (d, J = 11.8, 2H), 1.38-1.25 (m, 2H).

Compound 77, $C_{25}H_{26}ClF_3N_6O_3$, ($C_{25}H_{25}(D)ClF_3N_6O_3$)

1H NMR (500 MHz, DMSO- d_6) δ [ppm] = 15.88 (s, 1H), 7.95 (s, 2H), 7.82 (s, 1H),
7.75 (s, 1H), 7.69 (s, 1H), 7.45 (s, 1H), 5.15 (s, 2H), 4.05 (d, J = 8.8, 2H), 3.81-3.39
(m, 2H), 2.95-2.71 (m, 3H), 2.60-2.45 (m, 4H), 1.76 (d, J = 9.4, 2H), 1.48-1.17 (m,
3H).

1H NMR (500 MHz, DMSO- d_6 , d-TFA exchanged) δ [ppm] = 8.14 (s, 1H), 7.99 (d, J
= 8.5, 1H), 7.77 (s, 2H), 7.71 (s, 1H), 7.57 (dd, J = 8.5, 1.3, 1H), 5.20 (s, 2H), 4.21
(d, J = 13.5, 2H), 3.66-3.21 (m, 8H), 3.09-2.75 (m, 3H), 2.11 (d, J = 11.5, 2H), 1.67-
1.57 (m, 2H).

Compound 101, C₂₅H₂₆ClF₃N₆O₄

¹H NMR (500 MHz, DMSO-d₆) δ [ppm] = 15.83 (s, 1H), 7.93 (s, 2H), 7.67 (d, J = 1.9, 1H), 7.57 (dd, J = 8.4, 1.1, 1H), 7.48-7.41 (m, 2H), 5.09 (s, 2H), 4.51 (s, 1H), 3.61 (s, 1H), 3.39 (m, 5H), 3.01 (s, 1H), 2.85 (s, 1H), 2.60-2.49 (m, 4H), 1.90-1.58 (m, 2H), 1.52-1.31 (m, 2H).

Compound 109

¹H NMR (500 MHz, DMSO) δ = 15.86 (s, 1H), 7.96-7.89 (m, 2H), 7.74 (d, J=8.1, 2H), 7.57 (d, J=8.0, 2H), 7.42 (d, J=8.5, 1H), 5.17 (s, 2H), 4.48 (s, 1H), 3.65-3.35 (m, 4H), 3.12-2.75 (m, 2H), 2.56-2.51 (m, 1H), 2.35-2.31 (m, 4H), 2.18 (d, J=6.9, 2H), 1.87-1.58 (m, 3H), 1.17-1.15 (m, 2H).

Compound 110

¹H NMR (500 MHz, DMSO) δ = 15.77 (s, 1H), 10.30 (s, 1H), 8.60 (s, 1H), 8.51 (d, J=2.6, 1H), 8.04-7.93 (m, 2H), 7.74 (d, J=8.1, 2H), 7.61 (d, J=8.1, 2H), 6.90 (d, J=9.1, 1H), 5.22 (s, 2H), 3.66-3.28(m, 9H).

Compound 112

¹H NMR (500 MHz, DMSO) δ = 11.55 (s, 1H), 7.82 (s, 1H), 7.74 (d, J=8.1, 2H), 7.57 (d, J=8.1, 2H), 7.27-7.21 (m, 2H), 5.46 (s, 2H), 5.16 (s, 2H), 4.03 (d, J=12.8, 2H), 3.50 (s, 4H), 3.30-3.20 (m, 3H), 2.89 (s, 3H), 2.47-2.42 (m, 1H), 1.76 (d, J=11.9, 2H), 1.39-1.28 (m, 2H).

Compound 115

¹H NMR (500 MHz, DMSO) δ = 15.79 (s, 1H), 7.94 (s, 2H), 7.72-7.65 (m, 3H), 7.64-7.60 (m, 1H), 7.44 (d, J=8.5, 1H), 5.17 (s, 2H), 4.51 (s, 1H), 3.59 (s, 1H), 3.39 (s, 4H), 3.29 (s, 4H), 3.17-2.72 (m, 2H), 2.58-2.50 (m, 1H), 1.90-1.60 (m, 2H), 1.48-1.35 (m, 2H).

Compound 116

¹H NMR (500 MHz, DMSO) δ = 15.80 (m, 1H), 7.98-7.90 (m, 2H), 7.51-7.42 (m, 3H), 5.04 (s, 2H), 4.51 (s, 1H), 3.60 (s, 2H), 3.38 (s, 7H), 3.10-2.76 (m, 2H), 2.58-2.51 (m, 1H), 1.90-1.60 (m, 2H), 1.49-1.35 (m, 2H).

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Compound 118

¹H NMR (400 MHz, DMSO) δ = 15.85 (m, 1H), 7.98-7.92 (m, 2H), 7.58 (td, J=8.3, 1.1, 1H), 7.50 (dd, J=11.3, 1.9, 1H), 7.44 (dd, J=8.6, 0.9, 1H), 7.31 (d, J=9.0, 1H), 5.10 (s, 2H), 4.51 (s, 1H), 3.60 (s, 3H), 3.38 (s, 5H), 3.12-2.75 (m, 4H), 1.90-1.60 (m, 2H), 1.50-1.35 (m, 2H).

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Compound 123

¹H NMR (500 MHz, DMSO) δ = 15.83 (s, 1H), 7.94 (s, 2H), 7.46-7.39 (m, 3H), 7.29 (d, J=8.4, 2H), 4.51 (s, 1H), 4.08 (s, 2H), 3.72-3.38 (m, 3H), 3.18 (t, J=4.9, 2H), 3.10-2.72 (m, 2H), 2.58-2.51 (m, 1H), 2.41 (s, 2H), 2.20 (s, 2H), 1.83-1.53 (m, 2H), 1.45-1.30 (m, 2H).

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Compound 127

¹H NMR (500 MHz, DMSO) δ = 12.56 (s, 2H), 7.74 (d, J=8.1, 2H), 7.57 (d, J=8.1, 2H), 7.13 (s, 2H), 7.10 (s, 1H), 5.17 (s, 2H), 3.45-3.45 (m, 12H), 2.89 (s, 1H), 1.75 (s, 2H), 1.41-1.31 (m, 2H).

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Compound 139

¹H NMR (500 MHz, DMSO) δ = 15.57 (s, 1H), 8.08-8.03 (m, 3H), 7.84 (s, 1H), 7.70 (s, 1H), 7.29 (d, J=8.6, 1H), 5.25 (s, 2H), 4.39 (d, J=13.0, 1H), 4.02 (d, J=13.0, 1H), 3.88 (s, 2H), 3.45-3.25 (m, 5H), 2.99 (t, J=12.1, 1H), 2.55 (t, J=12.1, 1H), 2.47-2.38 (m, 4H), 1.77-1.62 (m, 2H), 1.26-1.12 (m, 2H).

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Compound 140

1H NMR (500 MHz, DMSO) δ = 15.50 (s, 1H), 8.81 (s, 1H), 7.85 (d, J=8.5, 1H), 7.72 (s, 1H), 7.59 (d, J=1.9, 2H), 7.30 (d, J=8.5, 1H), 7.11 (t, J=1.9, 1H), 4.41 (d, J=13.0, 1H), 4.04 (d, J=12.0, 1H), 3.90 (s, 2H), 3.45-3.35 (m, 5H), 3.01 (t, J=11.9, 1H), 2.59 (t, J=11.9, 1H), 2.48-2.41 (m, 4H), 1.80-1.69 (m, 2H), 1.29-1.14 (m, 2H).

Compound 144

1H NMR (400 MHz, DMSO) δ = 15.85 (s, 1H), 7.97-7.92 (m, 2H), 7.44 (dd, J=8.6, 1.1, 1H), 5.24 (s, 2H), 3.94 (s, 2H), 3.62 (s, 3H), 3.32 (s, 4H), 2.80 (s, 3H), 2.50-2.40 (m, 1H), 1.73 (d, J=12.3, 2H), 1.35-1.22 (m, 2H).

Compound 147

1H NMR (500 MHz, DMSO) δ = 15.70 (m, 1H), 7.97-7.92 (m, 2H), 7.43 (d, J=9.1, 1H), 6.94 (s, 3H), 4.98 (s, 2H), 4.50 (s, 1H), 3.60 (s, 2H), 3.35 (s, 5H), 3.10-2.65 (m, 2H), 2.55-2.45 (m, 3H), 2.26 (s, 6H), 1.92-1.60 (m, 2H), 1.48-1.35 (m, 2H).

Compound 152

1H NMR (500 MHz, DMSO) δ = 15.78 (s, 1H), 7.98-7.93 (m, 2H), 7.45 (d, J=8.8, 1H), 7.40 (dt, J=8.8, 2.1, 1H), 7.30 (s, 1H), 7.22 (d, J=9.4, 1H), 5.09 (s, 2H), 4.52 (s, 1H), 3.62 (s, 2H), 3.50-3.40 (s, 7H), 3.12-2.72 (m, 2H), 2.60-2.55 (m, 1H), 1.92-1.62 (m, 2H), 1.50-1.35 (m, 2H).

Compound 153

1H NMR (500 MHz, DMSO) δ = 12.33 (s, 1H), 8.07-8.03 (m, 4H), 7.62 (dd, J=8.4, 1.9, 1H), 7.28 (d, J=8.4, 1H), 5.24 (s, 2H), 3.63 (d, J=11.7, 2H), 3.35 (s, 4H), 2.43-2.37 (m, 4H), 2.24 (t, J=11.0, 3H), 1.75 (d, J=10.9, 2H), 1.48-1.38 (m, 2H).

Compound 158

1H NMR (500 MHz, DMSO) δ = 15.75 (m, 1H), 7.98-7.93 (m, 2H), 7.81 (t, J=7.9, 1H), 7.50 (d, J=11.8, 1H), 7.45 (d, J=9.3, 1H), 7.40 (d, J=8.1, 1H), 5.18 (s, 2H), 4.52

(s, 1H), 3.61 (s, 2H), 3.45 (s, 6H), 3.12-2.75 (m, 2H), 2.60-2.52 (m, 2H), 1.92-1.60 (m, 2H), 1.50-1.37 (m, 2H).

Compound 160

¹H NMR (500 MHz, DMSO) δ = 15.85 (s, 1H), 7.94 (s, 2H), 7.44 (d, J=8.5, 1H), 7.27 (d, J=8.7, 2H), 7.02-6.97 (m, 2H), 4.85 (s, 2H), 4.51 (s, 1H), 3.61 (s, 1H), 3.43 (s, 5H), 3.12-2.75 (m, 2H), 2.62-2.40 (m, H), 1.92-1.60 (s, 2H), 1.50-1.35 (m, 2H).

Compound 161

¹H NMR (500 MHz, DMSO) δ = 12.64 (s, 2H), 8.09-8.04 (m, 3H), 7.15 (s, 2H), 7.12 (s, 1H), 5.26 (s, 2H), 4.35 (s, 1H), 3.70-3.41 (m, 6H), 2.89 (s, 2H), 2.49-2.44 (m, 4H), 1.74 (s, 2H), 1.42-1.30 (m, 2H).

Compound 162

¹H NMR (500 MHz, DMSO) δ = 15.86 (s, 1H), 8.05 (s, 3H), 7.94 (d, J=7.6, 2H), 7.43 (d, J=8.4, 1H), 5.24 (s, 2H), 4.03 (d, J=12.8, 2H), 3.62 (s, 2H), 3.28 (s, 5H), 2.95-2.75 (s, 2H), 2.61-2.45 (m, 2H), 1.76 (d, J=11.0, 2H), 1.36-1.26 (m, 2H).

Compound 163

¹H NMR (500 MHz, DMSO) δ = 12.01 (s, 1H), 8.05 (s, 3H), 7.64 (d, J=1.5, 1H), 7.29 (dd, J=8.2, 1.6, 1H), 7.13 (d, J=8.2, 1H), 5.25 (s, 2H), 4.40 (s, 1H), 3.98-3.56 (m, 2H), 3.36 (s, 7H), 2.89 (s, 2H), 2.58-2.50 (m, 1H), 1.74 (s, 2H), 1.39 (qd, J=12.2, 4.2, 2H).

Compound 169

¹H NMR (500 MHz, DMSO) δ = 15.84 (s, 1H), 8.03 (s, 3H), 7.97-7.90 (m, 2H), 7.43 (d, J=8.7, 1H), 5.88 (q, J=6.6, 1H), 4.50 (s, 1H), 3.70-3.30 (m, 6H), 3.10-2.75 (m, 2H), 2.62 – 2.31 (m, 4H), 1.89-1.60 (m, 2H), 1.52 (d, J=6.6, 3H), 1.48-1.38 (m, 2H).

Compound 178

1H NMR (500 MHz, DMSO) δ = 15.85 (s, 1H), 7.90-7.91 (m, 3H), 7.77 (s, 2H), 7.44 (d, J=8.4, 1H), 4.52 (s, 1H), 3.72-3.30 (m, 7H), 3.12-2.75 (m, 2H), 2.60-2.53 (m, 1H), 2.48-2.38 (m, 2H), 1.92-1.60 (m, 2H), 1.52-1.35 (m, 6H).

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Compound 179

1H NMR (500 MHz, DMSO) δ = 15.83 (s, 1H), 7.97-7.92 (m, 2H), 7.44 (d, J=9.2, 1H), 7.32 (d, J=8.5, 2H), 7.22 (d, J=8.5, 2H), 4.52 (s, 1H), 3.69-3.5 (m, 3H), 3.48 (s, 3H), 3.12-2.77 (m, 2H), 2.60-2.41 (m, 4H), 2.31 (t, J=7.2, 2H), 1.96-1.60 (m, 2H), 1.51-1.36 (m, 3H), 1.22-1.15 (m, 1H).

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Compound 182

1H NMR (500 MHz, DMSO, d-TFA-exchanged) δ = 8.08 (s, 1H), 7.88 (t, J=8.2, 1H), 7.69-7.58 (m, 2H), 7.58-7.40 (m, 3H), 5.13 (s, 2H), 4.25-3.90 (m, 4H), 3.84-3.54 (m, 3H), 3.52 (s, 3H), 3.10-2.65 (m, 4H), 2.20-1.40 (m, 5H).

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Compound 183

1H NMR (400 MHz, DMSO) δ = 12.11 (s, 1H), 7.81 (s, 1H), 7.74 (s, 1H), 7.68 (s, 1H), 7.63 (d, J=1.5, 1H), 7.53 (dd, J=8.2, 1.7, 1H), 7.29 (d, J=8.2, 1H), 5.14 (s, 2H), 3.65 (d, J=11.6, 2H), 3.58-3.30 (m, 4H), 2.42 (s, 4H), 2.22 (t, J=11.0, 3H), 1.76 (d, J=11.5, 2H), 1.52-1.38 (m, 2H).

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Compound 184

1H NMR (500 MHz, DMSO) δ = 8.10 (s, 2H), 8.06 (s, 1H), 7.98 (d, J=12.5, 1H), 7.93 (d, J=8.4, 1H), 7.44 (d, J=8.5, 1H), 5.28 (s, 2H), 4.57-4.50 (m, 1H), 4.11-3.93 (m, 2H), 3.67-3.50 (m, 3H), 3.40-3.33 (m, 2H), 3.30-2.75 (m, 4H), 1.78-1.41 (m, 4H).

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Compound 187

1H NMR (500 MHz, DMSO) δ = 15.88 (s, 1H), 8.43 (s, 1H), 8.36 (d, J=7.8, 1H), 7.95-7.86 (m, 2H), 7.81 (s, 1H), 7.76 (s, 1H), 7.70 (s, 1H), 5.17 (s, 2H), 3.80-3.70

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(m, 1H), 3.50-3.24 (m, 8H), 2.35-2.27 (m, 1H), 1.93 (d, J=11.7, 2H), 1.83 (d, J=11.3, 2H), 1.48 – 1.27 (m, 4H).

Compound 188

¹H NMR (500 MHz, DMSO) δ = 8.43 (s, 1H), 8.36 (d, J=7.8, 1H), 7.90 (s, 2H), 7.73 (d, J=8.1, 2H), 7.51 (d, J=8.2, 2H), 5.15 (s, 2H), 3.79-3.70 (m, 1H), 3.50-3.15 (m, 9H), 2.35-2.27 (m, 1H), 1.93 (d, J=11.5, 2H), 1.83 (d, J=11.3, 2H), 1.45-1.29 (m, 4H).

The following analytical methods were use for determining above illustrated physico-chemical parameters:

ESI: Electrospray Ionization Mass Spectrometry (M+H)⁺

¹HPLC-Methode (non-polar)

Solvent A: water + 0,1% TFA

Solvent B: acetonitrile+ 0,08% TFA

Flow: 1,5 ml/min

Gradient:

0,0 min 20% B

5,0 min 100% B

5,5 min 100% B

6,0 min 20% B

6,5 min 20% B

Column: Chromolith Performance RP18e 100-3

²HPLC/MS-Methode (polar)

Solvent A: water + 0,05 % formic acid

Solvent B: acetonitriël + 0,04 % formic acid

Flow: 2,4 ml/min,

wavelength : 220nm

5 Gradient:

0,0 min 4 % B

2,8 min 100 % B

3,3 min 100% B

10 3,4 min 4% B

Column: Chromolith Speed ROD RP-18e 50-4.6 mm

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II. Autotaxin Assay

Assay Description

Autotaxin activity is measured indirectly by means of Amplex Red Reagent. In this course, Amplex Red is measured as fluorogenic indicator for generated H_2O_2 . Autotaxin converts substrate lysophosphatidylcholine (LPC) to phosphocholine and lysophosphatidyl acid (LPA). After the conversion phosphocholine is reacted with alkaline phosphatase to obtain inorganic phosphate and choline. During the next step choline is oxidized with choline oxidase to yield betaine, whereby H_2O_2 is generated. H_2O_2 reacts with Amplex Red Reagent in the presence of peroxidase (Horseradish peroxidase) in an 1:1 stoichiometry and yields highly fluorescent resorufin. The generated fluorescence is measured in a reaction-dependent kinetic mode in order to enable subtraction of fluorescence signals of possible other fluorescent compounds that are not part of the reaction from total measured fluorescence.

Performing the assay

1,5 μ l of a standard solution or of the compounds of the invention are dissolved in 20mM Hepes pH 7.2 with maximally 7.7% DMSO in individual concentrations. The resulting solution is pre-incubated together with 10 μ l (16 ng) of highly purified recombinant autotaxin in a black 384-hole microtiter plate for 30 min at 22°C.

Thereafter, the reaction is started through the addition of 5 μ l L- α -lysophosphatidylcholine (LPC), whereby the final concentration of LPC is 75 μ M. The mixture is incubated for 90 min. at 37°C. After the incubation Amplex Red Reagent, peroxidase (Horseradish peroxidase) and choline oxidase are added. The fluorescence is immediately measured at a wavelength of 612 nm with an excitation wavelength of 485 nm in a „Tecan Ultra multimode“ fluorescence reader. The activity of autotaxin is indirectly calculated via the amount of detected generated H_2O_2 .

For IC_{50} analysis, ten serial 1:3 dilutions starting at 30 μ M for each compound were run in duplicates.

IC₅₀ values were calculated on normalized data. For normalization, control wells were added to each assay plate and the signal of uninhibited control wells was set to 100%, whereas the signal inhibited by 500µM C14 LPA, (Avanti Polar Lipids, Cat# 857120P) was set to 0%. Curves were fitted and IC₅₀ values calculated by the following model using proprietary analysis software:

$$Y = \text{Bottom} + (100 - \text{Bottom}) / (1 + 10^{((\text{LogIC}_{50} - X) * \text{HillSlope}))})$$

Where X is the logarithm of concentration. Y is the response

; Y starts at Bottom and goes to Top with a sigmoid shape.

Material

Microtiter plate: PS-Microplate, 384-hole, small volume, black Corning, Cat#3677

Protein: Recombinant autotaxin (baculoviral Hi5 expression)

Substrate: L-a-lysophosphatidyl choline (chicken egg); Avanti Polar Lipids # 830071P

Standard: C14 LPA, Avanti Polar Lipids, Cat# 857120P

Detection Reagent: Amplex Red Reagent ; Invitrogen # A12222; dissolved in 1.923 ml of DMSO peroxidase Type VI-A (horseradish), Sigma # P6782; dissolved in 7,45 ml of test buffer, Choline Oxidase ; Sigma # C5896; dissolved in 2,47 ml test buffer

Detection Reagent Mix: 1:100 dilution of Amplex Red Reagent in test buffer

Test buffer: 200 mM Tris-HCl, Merck, Cat # 1.08219, pH 7.9; 0.1 % BSA, lipid free, Roche Cat#775835

Table 3

Compound	Chemical Name	IC ₅₀ value [M] or %CTRL (1E-05)
1	3-(2-Hydroxy-phenyl)-1-[1'-(3'-trifluoromethyl-biphenyl-2-carbonyl)-[4,4']bipiperidinyl-1-yl]-propan-1-one	< 1,00E-06
2	4-[4-(1H-Benzotriazol-5-ylcarbamoyl)-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 4-chloro-benzyl ester	< 1,00E-06

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3	5-Oxo-1-{1-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-pyrrolidine-3-carboxylic acid (1H-benzotriazol-5-yl)-amide	< 1,00E-06
4	4-[4-(1H-Benzotriazol-5-ylcarbamoyl)-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-06
5	4-[5-(2-Oxo-2,3-dihydro-benzooxazol-6-ylcarbamoyl)-pyridin-3-yl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester	< 1,00E-05
6	4-[5-(1H-Benzotriazol-5-ylcarbamoyl)-pyridin-3-yl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester	< 3,00E-05
7	4-[3-[(1H-Benzotriazole-5-carbonyl)-amino]-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 4-chloro-benzyl ester	< 1,00E-05
8	1H-Benzotriazole-5-carboxylic acid (1-{1-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-pyrrolidin-3-yl)-amide	< 1,00E-05
9	4-[3-[(1H-Benzotriazole-5-carbonyl)-amino]-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-06
10	4-[3-(1H-Benzotriazol-5-ylcarbamoyl)-phenyl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester	< 3,00E-05
11	[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidinyl-1-yl]-(4'-methyl-biphenyl-2-yl)-methanone	< 1,00E-06
12	1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidinyl-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-06
13	[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidinyl-1-yl]-(4'-methyl-biphenyl-3-yl)-methanone	< 1,00E-05
15	4-[2-Oxo-4-(2-oxo-2,3-dihydro-benzooxazol-6-ylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-06
16	4-[2-Oxo-4-(2-oxo-2,3-dihydro-benzooxazol-6-ylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 4-chloro-benzyl ester	< 1,00E-05
17	4-[2-Oxo-4-(2-pyridin-2-yl-ethylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-05
19	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-06
20	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-06
21	3-(2-Hydroxy-phenyl)-1-[4-[4-(3'-trifluoromethyl-biphenyl-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl]-propan-1-one	< 1,00E-06
22	1-[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidinyl-1-yl]-3-(4-trifluoromethyl-phenyl)-propan-1-one	< 1,00E-06

23	1-[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidinyl-1-yl]-2-(4-chloro-phenyl)-ethanone	< 3,00E-05
24	1-{1-[4-Methyl-2-(4-trifluoromethyl-phenyl)-thiazole-5-carbonyl]-piperidin-4-yl}-5-oxo-pyrrolidine-3-carboxylic acid (1H-benzotriazol-5-yl)-amide	< 1,00E-05
25	4-[4-(2-Oxo-2,3-dihydro-benzooxazole-6-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-06
27	4-[4-(1H-Benzotriazol-5-ylcarbamoyl)-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid tetrahydro-furan-2-ylmethyl ester	83%
28	4-[2-Oxo-4-(2-pyridin-4-yl-ethylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-05
29	9-(1H-Benzotriazole-5-carbonyl)-3,9-diaza-spiro[5.5]undecane-3-carboxylic acid 3,5-dichloro-benzyl ester	< 3,00E-05
30	4-[2-Oxo-4-(2-pyridin-3-yl-ethylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 3,00E-05
31	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (3,5-dichloro-phenyl)-amide	< 3,00E-05
32	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-trifluoromethyl-phenyl)-propan-1-one	< 1,00E-06
33	6-(4-{3-(4-Trifluoromethyl-phenyl)-propionyl}-piperazin-1-yl)-piperidine-1-carbonyl)-3H-benzooxazol-2-one	< 1,00E-06
34	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 1,00E-06
35	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	< 1,00E-06
36	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester	< 1,00E-06
37	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzyl ester	< 1,00E-06
38	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 5-chloro-2-methoxy-benzyl ester	< 1,00E-06
39	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-chloro-benzyl ester	< 1,00E-05
40	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-methyl-benzyl ester	< 1,00E-05
41	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-benzyl ester	< 1,00E-06

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42	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester	< 1,00E-06
43	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-isopropyl-benzyl ester	< 1,00E-06
45	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,4-dichloro-benzyl ester	< 1,00E-06
46	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,4-dichloro-benzyl ester	< 1,00E-06
47	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethoxy-benzyl ester	< 1,00E-06
48	(1H-Benzotriazol-5-yl)-(4-{4-[4-methyl-2-(4-trifluoromethyl-phenyl)-thiazole-5-carbonyl]-piperazin-1-yl}-piperidin-1-yl)-methanone	< 1,00E-05
49	6-(4-{1-[3-(4-Trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-piperazine-1-carbonyl)-3H-benzooxazol-2-one	< 1,00E-06
50	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-trifluoromethyl-phenyl)-propan-1-one	< 1,00E-05
51	3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-06
52	(1H-Benzotriazol-5-yl)-{4-[4-(5-trifluoromethyl-1H-benzimidazole-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-methanone	< 1,00E-05
53	4-{4-[2-(1H-Imidazol-4-yl)-ethylcarbamoyl]-2-oxo-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-05
54	4-[2-Oxo-4-(4-[1,2,4]triazol-1-yl-phenylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-05
55	4-[2-Oxo-4-(4-pyrazol-1-yl-phenylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-05
56	4-{4-[4-(3-Methyl-5-oxo-4,5-dihydro-pyrazol-1-yl)-phenylcarbamoyl]-2-oxo-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-05
57	4-[2-Oxo-4-(4-[1,2,3]thiadiazol-4-yl-phenylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester	< 1,00E-05
58	(E)-3-(3H-Imidazol-4-yl)-1-(4-{1-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-piperazin-1-yl)-propenone	< 1,00E-05
59	N-[4-(4-{1-[3-(4-Trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-piperazine-1-carbonyl)-phenyl]-methanesulfonamide	< 1,00E-05
60	1-(4-[4-[3-(1H-Imidazol-2-yl)-benzoyl]-piperazin-1-yl]-piperidin-1-yl)-3-(4-trifluoromethyl-phenyl)-propan-1-one	< 3,00E-05

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61	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-chloro-2-fluoro-phenyl)-propan-1-one	< 1,00E-05
62	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester	< 1,00E-06
63	3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester	< 1,00E-05
64	6-(4-{4-[3-(4-Chloro-2-fluoro-phenyl)-propionyl]-piperazin-1-yl}-piperidine-1-carbonyl)-3H-benzooxazol-2-one	< 1,00E-05
65	4-[1-(2-Oxo-2,3-dihydro-benzooxazole-6-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester	< 1,00E-06
66	1-{3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidin-1-yl}-3-(4-chloro-2-fluoro-phenyl)-propan-1-one	< 1,00E-05
67	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-chloro-2-fluoro-phenyl)-propan-1-one	< 1,00E-05
68	3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 1,00E-05
69	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,5-dichloro-phenyl)-propan-1-one	< 1,00E-05
70	4-[1-(1H-Benzotriazole-5-carbonyl)-pyrrolidin-3-ylamino]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 1,00E-05
71	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 1,00E-06
72	2-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-N-(4-chloro-3-trifluoromethyl-phenyl)-2-oxo-acetamide	< 1,00E-05
73	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 2-fluoro-4-trifluoromethyl-benzyl ester	< 1,00E-06
74	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 2-fluoro-5-trifluoromethyl-benzyl ester	< 1,00E-05
75	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-fluoro-5-trifluoromethyl-benzyl ester	< 1,00E-06
76	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethoxy-benzyl ester	< 1,00E-06
77	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester	< 1,00E-06
78	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-4-fluoro-benzyl ester	< 1,00E-06

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79	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,4,5-trifluoro-benzyl ester	< 1,00E-06
80	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-bromo-benzyl ester	< 1,00E-06
81	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-nitro-phenyl)-propan-1-one	< 1,00E-06
82	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(2,4-dichloro-phenyl)-propan-1-one	< 1,00E-05
83	(E)-1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-bromo-2-fluoro-phenyl)-propenone	< 1,00E-06
84	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-isopropyl-phenyl)-propan-1-one	< 1,00E-06
85	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,4-dichloro-benzyl ester	< 1,00E-06
86	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzyl ester	< 1,00E-06
87	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-isopropyl-benzyl ester	< 1,00E-06
88	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-methanesulfonyl-phenyl)-propan-1-one	89%
89	4-[4-[(S)-2-Amino-3-(1H-imidazol-4-yl)-propionyl]-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 3,00E-05
90	(E)-1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(3,5-dichloro-phenyl)-propenone	< 1,00E-06
91	(1H-Benzotriazol-5-yl)-{4-[1-(3,5-dichloro-4-fluoro-benzoyl)-piperidin-4-yl]-piperazin-1-yl}-methanone	66%
92	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-4-trifluoromethyl-benzyl ester	< 1,00E-06
93	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-4-fluoro-benzyl ester	< 1,00E-06
94	4-(3-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-oxo-propyl)-benzonitrile	77%
95	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester	< 1,00E-06
96	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-4-trifluoromethyl-benzyl ester	< 1,00E-06

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97	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dichloro-4-fluoro-benzyl ester	< 1,00E-06
98	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dibromo-4-methyl-benzyl ester	< 1,00E-06
99	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 5-chloro-2-fluoro-3-trifluoromethyl-benzyl ester	< 1,00E-06
100	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-fluoro-3-trifluoromethyl-benzyl ester	< 1,00E-06
101	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-4-trifluoromethoxy-benzyl ester	< 1,00E-06
102	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(3-fluoro-4-trifluoromethyl-phenyl)-ethanone	24%
103	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(2-fluoro-4-trifluoromethyl-phenyl)-ethanone	< 3,00E-05
104	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,5-dichloro-4-fluoro-phenyl)-propan-1-one	< 1,00E-05
105	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3-fluoro-4-trifluoromethyl-phenyl)-propan-1-one	< 1,00E-06
106	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-fluoro-3,5-dimethyl-phenyl)-propan-1-one	< 1,00E-05
107	4-[3-(2-Oxo-2,3-dihydro-benzooxazol-6-yl)-4,5-dihydro-pyrazol-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 1,00E-05
108	4-[3-(2-Oxo-2,3-dihydro-benzooxazol-6-yl)-pyrazol-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 1,00E-05
109	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylmethyl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 1,00E-05
110	4-{5-[(1H-Benzotriazole-5-carbonyl)-amino]-pyridin-2-yl}-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 3,00E-05
111	4-[5-(1H-Benzotriazol-5-ylcarbonyl)-pyridin-2-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 1,00E-05
112	4-[4-(3-Amino-1H-indazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 1,00E-05
113	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,3-difluoro-4-trifluoromethyl-benzyl ester	< 1,00E-06
114	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,3,5,6-tetrafluoro-4-trifluoromethyl-benzyl ester	< 1,00E-06

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115	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-trifluoromethyl-benzyl ester	< 1,00E-06
116	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-4,5-difluoro-benzyl ester	< 1,00E-06
117	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-3-fluoro-benzyl ester	< 1,00E-06
118	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-fluoro-4-trifluoromethoxy-benzyl ester	< 1,00E-06
119	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-methyl-3-trifluoromethyl-benzyl ester	< 1,00E-05
120	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-methoxy-3,5-dimethyl-benzyl ester	< 1,00E-05
121	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,4,5-trimethoxy-benzyl ester	< 3,00E-05
122	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzylamide	< 1,00E-05
123	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-chloro-phenyl)-propane-1,2-dione	< 1,00E-05
124	(1H-Benzotriazol-5-yl)-{4-[4-(4'-trifluoromethyl-biphenyl-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-methanone	< 1,00E-05
125	(1H-Benzotriazol-5-yl)-{4-[4-(3'-trifluoromethyl-biphenyl-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-methanone	< 1,00E-05
126	4-[1-(2-Oxo-2,3-dihydro-benzothiazole-6-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 1,00E-05
127	4-[1-(2-Thioxo-2,3-dihydro-1H-benzoimidazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 1,00E-05
128	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-trifluoromethoxy-benzyl ester	< 1,00E-06
129	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3-chloro-4,5-difluoro-phenyl)-propan-1-one	< 1,00E-05
130	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,5-dibromo-4-methyl-phenyl)-propan-1-one	< 1,00E-05
131	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(4-chloro-phenyl)-1H-[1,2,3]triazol-4-ylmethyl ester	< 3,00E-05
132	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(4-trifluoromethyl-phenyl)-ethyl ester	< 1,00E-05

133	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(3,5-dichloro-4-fluorophenyl)-propan-1-one	< 1,00E-05
134	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-methyl-2H-indazol-3-ylmethyl ester	<3,00E-05
135	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid [3,3']bithiophenyl-5-ylmethyl ester	< 1,00E-06
136	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid benzo[1,3]dioxol-5-ylmethyl ester	< 1,00E-05
137	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,3-dihydro-benzo[1,4]dioxin-6-ylmethyl ester	< 1,00E-05
138	2-(1H-Benzotriazol-5-yl)-1-(4-{4-[2-(4-trifluoromethyl-phenyl)-acetyl]-piperazin-1-yl}-piperidin-1-yl)-ethanone	< 1,00E-05
139	4-[1-(2-1H-Benzotriazol-5-yl-acetyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	<1,00E-06
140	4-[1-(2-1H-Benzotriazol-5-yl-acetyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (3,5-dichlorophenyl)-amide	< 1,00E-05
141	2-(1H-Benzotriazol-5-yl)-1-(4-{4-[2-(3,5-bis-trifluoromethyl-phenyl)-acetyl]-piperazin-1-yl}-piperidin-1-yl)-ethanone	< 1,00E-05
142	6-(4-[4-[2-(3,5-Bis-trifluoromethyl-phenyl)-acetyl]-piperazin-1-yl]-piperidine-1-carbonyl)-3H-benzooxazol-2-one	< 1,00E-05
143	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-4-trifluoromethoxy-benzyl ester	< 1,00E-06
144	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 2,3,5,6-tetrafluoro-4-trifluoromethyl-benzyl ester	< 1,00E-06
145	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-3-trifluoromethoxy-benzyl ester	< 1,00E-05
146	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-bromo-5-chloro-benzyl ester	<1,00E-06
147	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dimethyl-benzyl ester	< 1,00E-06
148	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dibromobenzyl ester	< 1,00E-06
149	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-methylsulfanylbenzyl ester	< 1,00E-05
150	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dimethoxybenzyl ester	< 1,00E-05

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151	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-ethyl-benzyl ester	< 1,00E-06
152	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-fluoro-benzyl ester	< 1,00E-06
153	4-[1-(2-Oxo-2,3-dihydro-benzothiazole-6-sulfonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	< 1,00E-05
154	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-bromo-5-fluoro-benzyl ester	< 1,00E-06
155	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-bromo-benzyl ester	< 1,00E-06
156	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-bromo-2-fluoro-benzyl ester	< 1,00E-05
157	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-bromo-2,6-difluoro-benzyl ester	< 1,00E-05
158	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-fluoro-4-trifluoromethyl-benzyl ester	< 1,00E-06
159	(1H-Benzotriazol-5-yl)-(4-[4-(5-bromo-benzofuran-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl)-methanone	< 1,00E-05
160	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(4-trifluoromethoxy-phenoxy)-ethanone	< 1,00E-05
161	4-[1-(2-Thioxo-2,3-dihydro-1H-benzoimidazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	< 1,00E-06
162	4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	< 1,00E-06
163	4-[1-(2-Oxo-2,3-dihydro-benzothiazole-6-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	< 1,00E-06
164	1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-trifluoromethyl-phenyl)-propane-1,3-dione	< 1,00E-05
165	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,4-difluoro-5-trifluoromethyl-benzyl ester	< 1,00E-06
166	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-bromo-4-trifluoromethoxy-benzyl ester	< 1,00E-06
167	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-difluoromethoxy-benzyl ester	< 1,00E-06
168	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-difluoromethylsulfanyl-benzyl ester	< 1,00E-06

5	169	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (R)-1-(3,5-bis-trifluoromethyl-phenyl)-ethyl ester	< 1,00E-06
	170	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,2,2-trifluoro-1-p-tolyl-1-trifluoromethyl-ethyl ester	< 1,00E-05
	171	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (S)-1-(3,5-bis-trifluoromethyl-phenyl)-ethyl ester	< 1,00E-05
	172	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(3,5-bis-trifluoromethyl-phenyl)-ethyl ester	< 1,00E-05
	173	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-methyl-5-phenyl-furan-3-ylmethyl ester	< 1,00E-05
10	174	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-(4-chloro-phenyl)-4-methyl-thiazol-5-ylmethyl ester	< 1,00E-05
	175	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-(4-chloro-phenyl)-2-oxo-oxazolidin-5-ylmethyl ester	< 1,00E-05
	176	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,4-dimethyl-phenyl)-butan-1-one	< 3,00E-05
15	177	1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-trifluoromethyl-phenyl)-butan-1-one	< 1,00E-05
	178	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(3,5-bis-trifluoromethyl-phenyl)-cyclopropyl ester	< 1,00E-06
	179	(1H-Benzotriazol-5-yl)-(4-{4-[2-(4-chloro-phenyl)-cyclopropanecarbonyl]-piperazin-1-yl}-piperidin-1-yl)-methanone	< 1,00E-05
20	180	2-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-N-(3,5-bis-trifluoromethyl-phenyl)-acetamide	< 1,00E-05
	181	2-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-N-(5-chloro-2-methoxy-phenyl)-acetamide	< 1,00E-05
	182	4-[(1-(1H-Benzotriazole-5-carbonyl)-pyrrolidin-3-yl)-methyl-amino]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester	< 1,00E-05
25	183	4-[1-(2-Oxo-2,3-dihydro-benzooxazole-6-sulfonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester	< 1,00E-05
	184	4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-3-oxo-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	< 1,00E-06
	185	4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid benzyl ester	< 3,00E-05
	186	4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester	< 1,00E-06

5	187	4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester	< 1,00 E-06
	188	4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzyl ester	< 1,00 E-05
	189	4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3-fluoro-4-trifluoromethyl-benzyl ester	< 1,00 E-05

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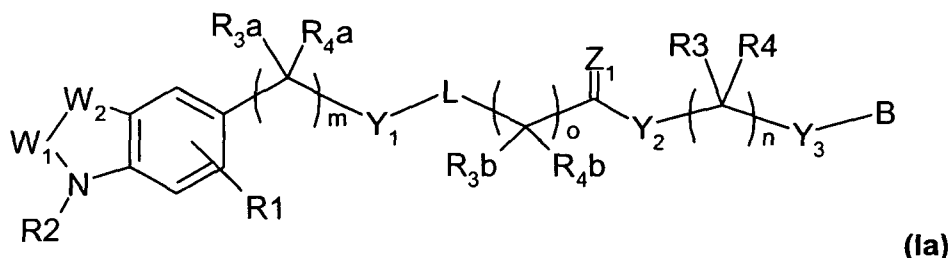
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Claims

1. Compound according to formula (I)



10 wherein:

W_1, W_2 together independently form “-N=N-, -C(O)-O-, -C(O)-S-, -C(O)-N(R5)-, -C(S)-N(R5a)-, -C(O)-C(R6)(R7)-, -N=C[N(R8)(R9)]-”;

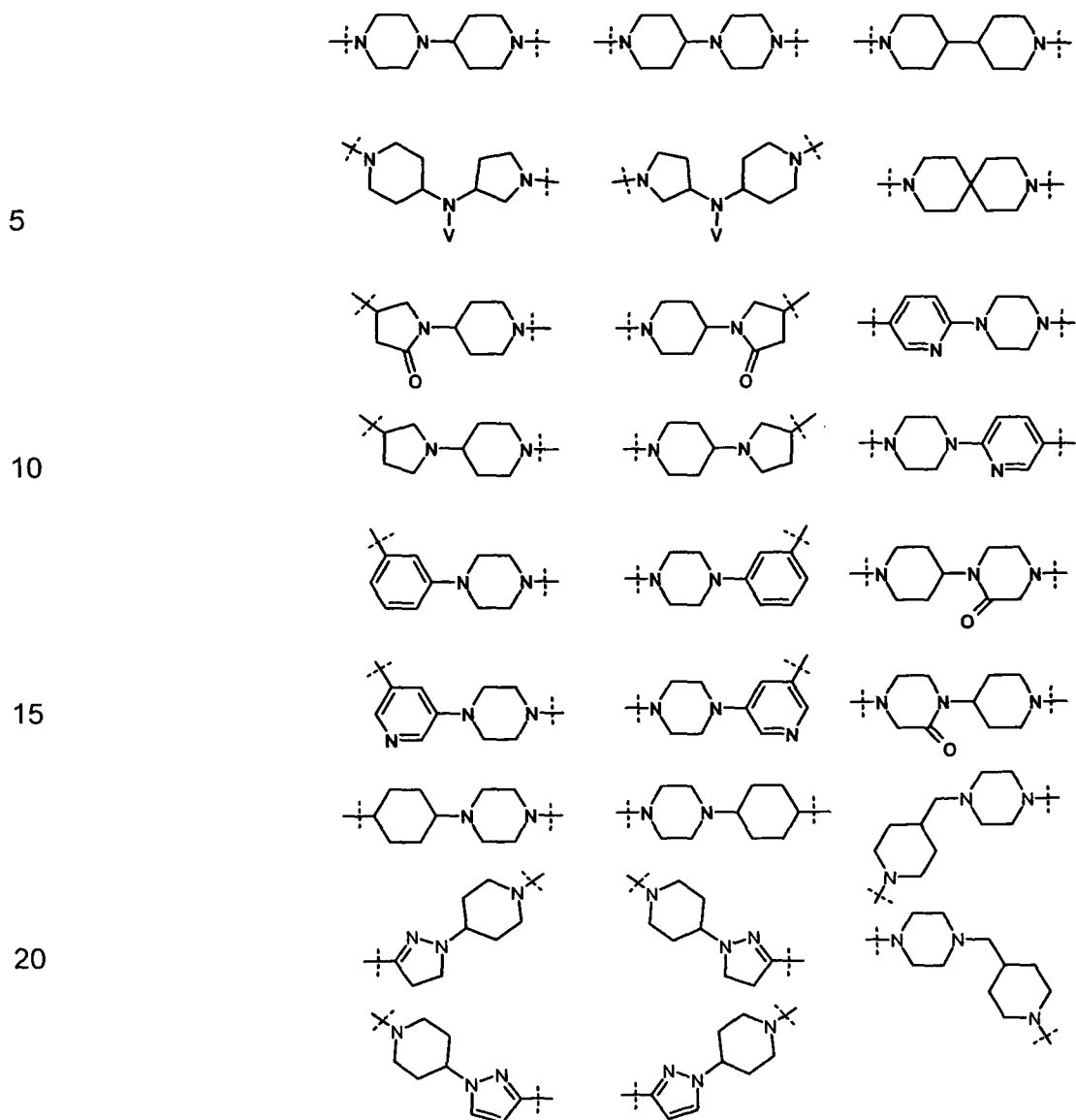
Y_1 is independently selected from the group consisting of “-C(O)-, -C(S)-, -S(O)₂-, -N(R10)-C(O)-, -C(O)-N(R11)-, -OC(O)-, single bond”;

Y_2 is independently selected from the group consisting of “-C(R12)(R13)-, -O-, -N(R14)-, -C(O)-, -C(O)-NH-, single bond”;

Y_3 is independently selected from the group consisting of “-O-, -C(O)-, single bond”;

Z_1 is independently selected from the group consisting of “O, S, N(R15)”;

L is independently selected from the group consisting of the group consisting of:



25 B is independently selected from the group consisting of "cycloalkyl, heterocyclyl, aryl, heteroaryl", wherein "cycloalkyl, heterocyclyl, aryl, heteroaryl" can be independently substituted with one or more identical or different substituents selected from the group consisting of: "(i) „hydrogen, alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl, halogen, -F, -Cl, -Br, -I, -CN, -CF₃, -SF₃, -N₃, -NH₂, -NHX₁, -NX₂X₃, -NO₂, -OH, =O, -OCF₃, -SCF₃, -OCHF₂, -SCHF₂, -SH, -O-SO₃H, -OP(O)(OH)₂, -CHO, -COOH, -C(O)NH₂, -SO₃H, -P(O)(OH)₂, -C(O)-X₄, -C(O)O-X₅, -C(O)NH-X₆, -C(O)NX₇X₈, -O-X₉, -O(-X₁₀-O)_a-H (a = 1, 2, 3, 4, 5), -O(-X₁₁-

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$O)_b-X12$ ($b = 1, 2, 3, 4, 5$), $-OC(O)-X13$, $-OC(O)-O-X14$, $-OC(O)-$
 $NHX15$, $-O-C(O)-NX16X17$, $-OP(O)(OX18)(OX19)$, $-$
 $OSi(X20)(X21)(X22)$, $-OS(O_2)-X23$, $-NHC(O)-NH_2$, $-NHC(O)-X24$,
 $-NX25C(O)-X26$, $-NH-C(O)-O-X27$, $-NH-C(O)-NH-X28$, $-NH-$
 $C(O)-NX29X30$, $-NX31-C(O)-O-X32$, $-NX33-C(O)-NH-X34$, $-$
 $NX35-C(O)-NX36X37$, $-NHS(O_2)-X38$, $-NX39S(O_2)-X40$, $-S-X41$,
 $-S(O)-X42$, $-S(O_2)-X43$, $-S(O_2)NH-X44$, $-S(O_2)NX45X46$, $-$
 $S(O_2)O-X47$, $-P(O)(OX48)(OX49)$, $-Si(X50)(X51)(X52)$, $-C(NH)-$
 NH_2 , $-C(NX53)-NH_2$, $-C(NH)-NHX54$, $-C(NH)-NX55X56$, $-$
 $C(NX57)-NHX58$, $-C(NX59)-NX60X61$, $-NH-C(O)-NH-O-X62$, $-$
 $NH-C(O)-NX63-O-X64$, $-NX65-C(O)-NX66-O-X67$, $-N(-C(O)-$
 $NH-O-X68)_2$, $-N(-C(O)-NX69-O-X70)_2$, $-N(-C(O)-NH-O-X71)(-$
 $C(O)-NX72-O-X73)$, $-C(S)-X74$, $-C(S)-O-X75$, $-C(S)-NH-X76$, $-$
 $C(S)-NX77X78$, $-C(O)-NH-O-X79$, $-C(O)-NX80-O-X81$, $-C(S)-$
 $NH-O-X82$, $-C(S)-NX83-O-X84$, $-C(O)-NH-NH-X85$, $-C(O)-NH-$
 $NX86X87$, $-C(O)-NX88-NX89X90$, $-C(S)-NH-NH-X91$, $-C(S)-$
 $NH-NX92X93$, $-C(S)-NX94-NX95X96$, $-C(O)-C(O)-O-X97$, $-$
 $C(O)-C(O)-NH_2$, $-C(O)-C(O)-NHX98$, $-C(O)-C(O)-NX99X100$, $-$
 $C(S)-C(O)-O-X101$, $-C(O)-C(S)-O-X102$, $-C(S)-C(S)-O-X103$, $-$
 $C(S)-C(O)-NH_2$, $-C(S)-C(O)-NHX104$, $-C(S)-C(O)-NX105X106$, $-$
 $C(S)-C(S)-NH_2$, $-C(S)-C(S)-NHX107$, $-C(S)-C(S)-NX108X109$, $-$
 $C(O)-C(S)-NH_2$, $-C(O)-C(S)-NHX110$, $-C(O)-C(S)-NX111X112$ ";

wherein X1, X2, X3, X4, X5, X6, X7, X8, X9, X10, X11, X12, X13,
 X14, X15, X16, X17, X18, X19, X20, X21, X22, X23, X24, X25, X26,
 X27, X28, X29, X30, X31, X32, X33, X34, X35, X36, X37, X38, X39,
 X40, X41, X42, X43, X44, X45, X46, X47, X48, X49, X50, X51, X52,
 X53, X54, X55, X56, X57, X58, X59, X60, X61, X62, X63, X64, X65,
 X66, X67, X68, X69, X70, X71, X72, X73, X74, X75, X76, X77, X78,
 X79, X80, X81, X82, X83, X84, X85, X86, X87, X88, X89, X90, X91,
 X92, X93, X94, X95, X96, X97, X98, X99, X100, X101, X102, X103,
 X104, X105, X106, X107, X108, X109, X110, X111, X112 are
 independently from each other selected from the group consisting of:
 „alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl,
 heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl“ and
 wherein alternatively X7, X8 and/or X16, X17 and/or X29, X30 and/or

X36, X37 and/or X45, X46 and/or X55, X56 and/or X60, X61 and/or X77, X78 and/or X86, X87 and/or X89, X90 and/or X92, X93 and/or X95, X96 and/or X99, X100 and/or X105, X106 and/or X108, X109 and/or X111, X112 respectively together can also form

5 „heterocyclyl“;

wherein optionally above substituents of substituents group (i) can in turn independently from each other be substituted with one or more identical or different substituents V;

10 R1, R2, R3, R3a, R3b, R4, R4a, R4b, R5, R5a, R6, R7, R8, R9, R10, R11, R12, R13, R14, R15 are independently from each other selected from the group consisting of: “V”; alternatively, R3a and R4a, R3b and R4b as well as R3 and R4 together can form “cycloalkyl” or “heterocyclyl”;

15 V is independently selected from the group consisting of: “(i)
 „hydrogen, alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl, halogen, -F, -Cl, -Br, -I, -CN, -CF₃, -SF₃, -N₃, -NH₂, -NHA1, -NA2A3, -NO₂, -OH, =O, -OCF₃, -SCF₃, -OCHF₂, -SCHF₂, -SH, -O-SO₃H, -OP(O)(OH)₂, -CHO, -COOH, -C(O)NH₂, -SO₃H, -P(O)(OH)₂, -C(O)-A4, -C(O)O-A5, -C(O)NH-A6, -C(O)NA7A8, -O-A9, -O(-A10-O)_a-H (a = 1, 2, 3, 4, 5), -O(-A11-O)_b-A12 (b = 1, 2, 3, 4, 5), -OC(O)-A13, -OC(O)-O-A14, -OC(O)-NHA15, -O-C(O)-NA16A17, -OP(O)(OA18)(OA19), -OSi(A20)(A21)(A22), -OS(O₂)-A23, -NHC(O)-NH₂, -NHC(O)-A24, -NA25C(O)-A26, -NH-C(O)-O-A27, -NH-C(O)-NH-A28, -NH-C(O)-NA29A30, -NA31-C(O)-O-A32, -NA33-C(O)-NH-A34, -NA35-C(O)-NA36A37, -NHS(O₂)-A38, -NA39S(O₂)-A40, -S-A41, -S(O)-A42, -S(O₂)-A43, -S(O₂)NH-A44, -S(O₂)NA45A46, -S(O₂)O-A47, -P(O)(OA48)(OA49), -Si(A50)(A51)(A52), -C(NH)-NH₂, -C(NA53)-NH₂, -C(NH)-NHA54, -C(NH)-NA55A56, -C(NA57)-NHA58, -C(NA59)-NA60A61, -NH-C(O)-NH-O-A62, -NH-C(O)-NA63-O-A64, -NA65-C(O)-NA66-O-A67, -N(-C(O)-NH-O-A68)₂, -N(-C(O)-NA69-O-A70)₂, -N(-C(O)-NH-O-A71)(-C(O)-NA72-O-A73), -C(S)-A74, -C(S)-O-A75, -C(S)-NH-A76, -

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C(S)-NA77A78, -C(O)-NH-O-A79, -C(O)-NA80-O-A81, -C(S)-NH-O-A82, -C(S)-NA83-O-A84, -C(O)-NH-NH-A85, -C(O)-NH-NA86A87, -C(O)-NA88-NA89A90, -C(S)-NH-NH-A91, -C(S)-NH-NA92A93, -C(S)-NA94-NA95A96, -C(O)-C(O)-O-A97, -C(O)-C(O)-NH₂, -C(O)-C(O)-NHA98, -C(O)-C(O)-NA99A100, -C(S)-C(O)-O-A101, -C(O)-C(S)-O-A102, -C(S)-C(S)-O-A103, -C(S)-C(O)-NH₂, -C(S)-C(O)-NHA104, -C(S)-C(O)-NA105A106, -C(S)-C(S)-NH₂, -C(S)-C(S)-NHA107, -C(S)-C(S)-NA108A109, -C(O)-C(S)-NH₂, -C(O)-C(S)-NHA110, -C(O)-C(S)-NA111A112";

wherein A1, A2, A3, A4, A5, A6, A7, A8, A9, A10, A11, A12, A13, A14, A15, A16, A17, A18, A19, A20, A21, A22, A23, A24, A25, A26, A27, A28, A29, A30, A31, A32, A33, A34, A35, A36, A37, A38, A39, A40, A41, A42, A43, A44, A45, A46, A47, A48, A49, A50, A51, A52, A53, A54, A55, A56, A57, A58, A59, A60, A61, A62, A63, A64, A65, A66, A67, A68, A69, A70, A71, A72, A73, A74, A75, A76, A77, A78, A79, A80, A81, A82, A83, A84, A85, A86, A87, A88, A89, A90, A91, A92, A93, A94, A95, A96, A97, A98, A99, A100, A101, A102, A103, A104, A105, A106, A107, A108, A109, A110, A111, A112 are independently from each other selected from the group consisting of: „alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl“ and wherein alternatively A7, A8 and/or A16, A17 and/or A29, A30 and/or A36, A37 and/or A45, A46 and/or A55, A56 and/or A60, A61 and/or A77, A78 and/or A86, A87 and/or A89, A90 and/or A92, A93 and/or A95, A96 and/or A99, A100 and/or A105, A106 and/or A108, A109 and/or A111, A112 respectively together can also form „heterocyclyl“;

wherein optionally above substituents of substituents group (i) can in turn independently from each other be substituted with one or more identical or different substituents V;

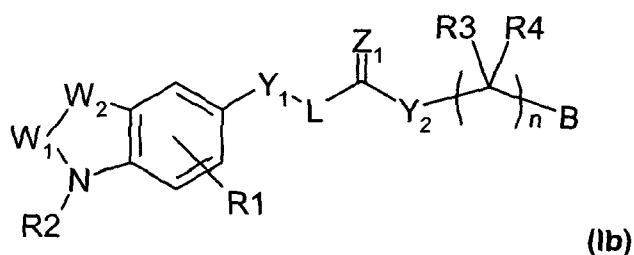
m is independently 0, 1, 2, or 3;

n is independently 0, 1, 2, or 3;

o is independently 0, 1, 2, or 3;

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

2. Compound according to formula (I)



wherein:

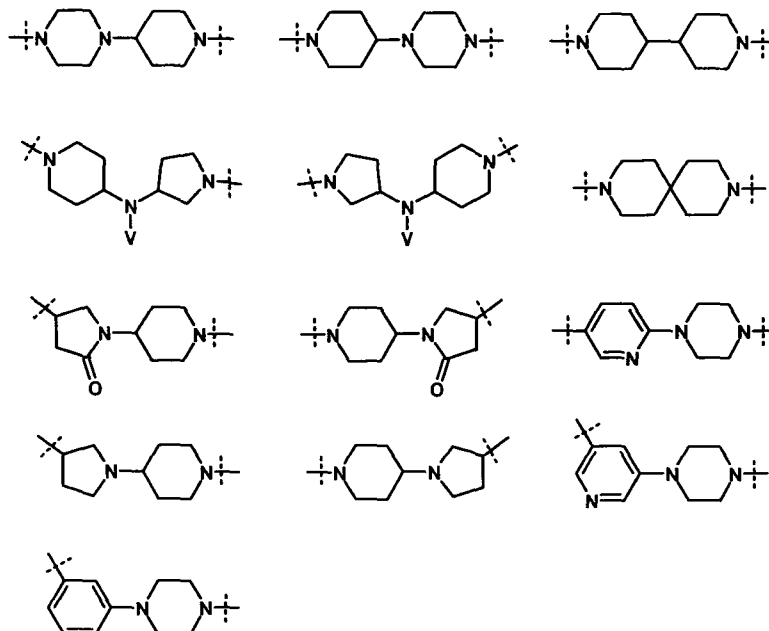
W_1, W_2 together independently form “-N=N-, -C(O)-O-, -C(O)-S-, -C(O)-N(R5)-, -C(O)-C(R6)(R7)-, -N=C[N(R8)(R9)]-”;

Y_1 is independently selected from the group consisting of “-C(O)-, -C(S)-, -N(R10)-C(O)-, -C(O)-N(R11)-, -OC(O)-, single bond”;

Y_2 is independently selected from the group consisting of “-C(R12)(R13)-, -O-, -N(R14)-, -C(O)-NH-, single bond”;

Z_1 is independently selected from the group consisting of “O, S, N(R15)”;

L is independently selected from the group consisting of the group consisting of:



B is independently selected from the group consisting of "cycloalkyl, heterocyclyl, aryl, heteroaryl", wherein "cycloalkyl, heterocyclyl, aryl, heteroaryl" can be independently substituted with one or more identical or different substituents selected from the group consisting of: "(i) „hydrogen, alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl, halogen, -F, -Cl, -Br, -I, -CN, -CF₃, -N₃, -NH₂, -NHX₁, -NX₂X₃, -NO₂, -OH, -OCF₃, -SCF₃, -SH, -O-SO₃H, -OP(O)(OH)₂, -CHO, -COOH, -C(O)NH₂, -SO₃H, -P(O)(OH)₂, -C(O)-X₄, -C(O)O-X₅, -C(O)NH-X₆, -C(O)NX₇X₈, -O-X₉, -O(-X₁₀-O)_a-H (a = 1, 2, 3, 4, 5), -O(-X₁₁-O)_b-X₁₂ (b = 1, 2, 3, 4, 5), -OC(O)-X₁₃, -OC(O)-O-X₁₄, -OC(O)-NHX₁₅, -O-C(O)-NX₁₆X₁₇, -OP(O)(OX₁₈)(OX₁₉), -OSi(X₂₀)(X₂₁)(X₂₂), -OS(O₂)-X₂₃, -NHC(O)-NH₂, -NHC(O)-X₂₄, -NX₂₅C(O)-X₂₆, -NH-C(O)-O-X₂₇, -NH-C(O)-NH-X₂₈, -NH-C(O)-NX₂₉X₃₀, -NX₃₁-C(O)-O-X₃₂, -NX₃₃-C(O)-NH-X₃₄, -NX₃₅-C(O)-NX₃₆X₃₇, -NHS(O₂)-X₃₈, -NX₃₉S(O₂)-X₄₀, -S-X₄₁, -S(O)-X₄₂, -S(O₂)-X₄₃, -S(O₂)NH-X₄₄, -S(O₂)NX₄₅X₄₆, -S(O₂)O-X₄₇, -P(O)(OX₄₈)(OX₄₉), -Si(X₅₀)(X₅₁)(X₅₂), -C(NH)-NH₂, -C(NX₅₃)-NH₂, -C(NH)-NHX₅₄, -C(NH)-NX₅₅X₅₆, -C(NX₅₇)-NHX₅₈, -C(NX₅₉)-NX₆₀X₆₁, -NH-

C(O)-NH-O-X62, -NH-C(O)-NX63-O-X64, -NX65-C(O)-NX66-O-X67, -N(-C(O)-NH-O-X68)₂, -N(-C(O)-NX69-O-X70)₂, -N(-C(O)-NH-O-X71)(-C(O)-NX72-O-X73), -C(S)-X74, -C(S)-O-X75, -C(S)-NH-X76, -C(S)-NX77X78, -C(O)-NH-O-X79, -C(O)-NX80-O-X81, -C(S)-NH-O-X82, -C(S)-NX83-O-X84, -C(O)-NH-NH-X85, -C(O)-NH-NX86X87, -C(O)-NX88-NX89X90, -C(S)-NH-NH-X91, -C(S)-NH-NX92X93, -C(S)-NX94-NX95X96, -C(O)-C(O)-O-X97, -C(O)-C(O)-NH₂, -C(O)-C(O)-NHX98, -C(O)-C(O)-NX99X100, -C(S)-C(O)-O-X101, -C(O)-C(S)-O-X102, -C(S)-C(S)-O-X103, -C(S)-C(O)-NH₂, -C(S)-C(O)-NHX104, -C(S)-C(O)-NX105X106, -C(S)-C(S)-NH₂, -C(S)-C(S)-NHX107, -C(S)-C(S)-NX108X109, -C(O)-C(S)-NH₂, -C(O)-C(S)-NHX110, -C(O)-C(S)-NX111X112";

wherein X1, X2, X3, X4, X5, X6, X7, X8, X9, X10, X11, X12, X13, X14, X15, X16, X17, X18, X19, X20, X21, X22, X23, X24, X25, X26, X27, X28, X29, X30, X31, X32, X33, X34, X35, X36, X37, X38, X39, X40, X41, X42, X43, X44, X45, X46, X47, X48, X49, X50, X51, X52, X53, X54, X55, X56, X57, X58, X59, X60, X61, X62, X63, X64, X65, X66, X67, X68, X69, X70, X71, X72, X73, X74, X75, X76, X77, X78, X79, X80, X81, X82, X83, X84, X85, X86, X87, X88, X89, X90, X91, X92, X93, X94, X95, X96, X97, X98, X99, X100, X101, X102, X103, X104, X105, X106, X107, X108, X109, X110, X111, X112 are independently from each other selected from the group consisting of: „alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl“ and wherein alternatively X7, X8 and/or X16, X17 and/or X29, X30 and/or X36, X37 and/or X45, X46 and/or X55, X56 and/or X60, X61 and/or X77, X78 and/or X86, X87 and/or X89, X90 and/or X92, X93 and/or X95, X96 and/or X99, X100 and/or X105, X106 and/or X108, X109 and/or X111, X112 respectively together can also form „heterocyclyl“;

wherein optionally above substituents of substituents group (i) can in turn independently from each other be substituted with one or more identical or different substituents V;

R1, R2, R3, R4, R5, R6, R7, R8, R9, R10, R11, R12, R13, R14, R15 are independently from each other selected from the group consisting of: "V";

V is independently selected from the group consisting of: "(i)
 „hydrogen, alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl, halogen, -F, -Cl, -Br, -I, -CN, -CF₃, -N₃, -NH₂, -NHA1, -NA2A3, -NO₂, -OH, -OCF₃, -SCF₃, -SH, -O-SO₃H, -OP(O)(OH)₂, -CHO, -COOH, -C(O)NH₂, -SO₃H, -P(O)(OH)₂, -C(O)-A4, -C(O)O-A5, -C(O)NH-A6, -C(O)NA7A8, -O-A9, -O(-A10-O)_a-H (a = 1, 2, 3, 4, 5), -O(-A11-O)_b-A12 (b = 1, 2, 3, 4, 5), -OC(O)-A13, -OC(O)-O-A14, -OC(O)-NHA15, -O-C(O)-NA16A17, -OP(O)(OA18)(OA19), -OSi(A20)(A21)(A22), -OS(O₂)-A23, -NHC(O)-NH₂, -NHC(O)-A24, -NA25C(O)-A26, -NH-C(O)-O-A27, -NH-C(O)-NH-A28, -NH-C(O)-NA29A30, -NA31-C(O)-O-A32, -NA33-C(O)-NH-A34, -NA35-C(O)-NA36A37, -NHS(O₂)-A38, -NA39S(O₂)-A40, -S-A41, -S(O)-A42, -S(O₂)-A43, -S(O₂)NH-A44, -S(O₂)NA45A46, -S(O₂)O-A47, -P(O)(OA48)(OA49), -Si(A50)(A51)(A52), -C(NH)-NH₂, -C(NA53)-NH₂, -C(NH)-NHA54, -C(NH)-NA55A56, -C(NA57)-NHA58, -C(NA59)-NA60A61, -NH-C(O)-NH-O-A62, -NH-C(O)-NA63-O-A64, -NA65-C(O)-NA66-O-A67, -N(-C(O)-NH-O-A68)₂, -N(-C(O)-NA69-O-A70)₂, -N(-C(O)-NH-O-A71)(-C(O)-NA72-O-A73), -C(S)-A74, -C(S)-O-A75, -C(S)-NH-A76, -C(S)-NA77A78, -C(O)-NH-O-A79, -C(O)-NA80-O-A81, -C(S)-NH-O-A82, -C(S)-NA83-O-A84, -C(O)-NH-NH-A85, -C(O)-NH-NA86A87, -C(O)-NA88-NA89A90, -C(S)-NH-NH-A91, -C(S)-NH-NA92A93, -C(S)-NA94-NA95A96, -C(O)-C(O)-O-A97, -C(O)-C(O)-NH₂, -C(O)-C(O)-NHA98, -C(O)-C(O)-NA99A100, -C(S)-C(O)-O-A101, -C(O)-C(S)-O-A102, -C(S)-C(S)-O-A103, -C(S)-C(O)-NH₂, -C(S)-C(O)-NHA104, -C(S)-C(O)-NA105A106, -C(S)-C(S)-NH₂, -C(S)-C(S)-NHA107, -C(S)-C(S)-NA108A109, -C(O)-C(S)-NH₂, -C(O)-C(S)-NHA110, -C(O)-C(S)-NA111A112";

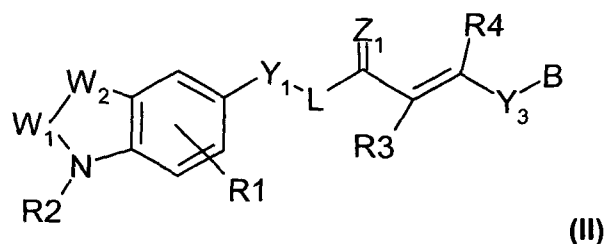
wherein A1, A2, A3, A4, A5, A6, A7, A8, A9, A10, A11, A12, A13, A14, A15, A16, A17, A18, A19, A20, A21, A22, A23, A24, A25, A26, A27, A28, A29, A30, A31, A32, A33, A34, A35, A36, A37, A38, A39, A40, A41, A42, A43, A44, A45, A46, A47, A48, A49, A50, A51, A52, A53, A54, A55, A56, A57, A58, A59, A60, A61, A62, A63, A64, A65, A66, A67, A68, A69, A70, A71, A72, A73, A74, A75, A76, A77, A78, A79, A80, A81, A82, A83, A84, A85, A86, A87, A88, A89, A90, A91, A92, A93, A94, A95, A96, A97, A98, A99, A100, A101, A102, A103, A104, A105, A106, A107, A108, A109, A110, A111, A112 are independently from each other selected from the group consisting of: „alkyl, (C₉-C₃₀)alkyl, cycloalkyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl, heteroaryl, heteroarylalkyl“ and wherein alternatively A7, A8 and/or A16, A17 and/or A29, A30 and/or A36, A37 and/or A45, A46 and/or A55, A56 and/or A60, A61 and/or A77, A78 and/or A86, A87 and/or A89, A90 and/or A92, A93 and/or A95, A96 and/or A99, A100 and/or A105, A106 and/or A108, A109 and/or A111, A112 respectively together can also form „heterocyclyl“;

wherein optionally above substituents of substituents group (i) can in turn independently from each other be substituted with one or more identical or different substituents V;

n is independently 0, 1, 2, or 3;

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

3. A compound according to formula (II)



wherein according to formulae (Ia) or (Ib) as claimed in claim 1 or 2:

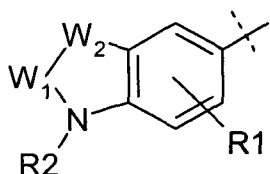
Y_2 independently is a single bond;

n independently is 2;

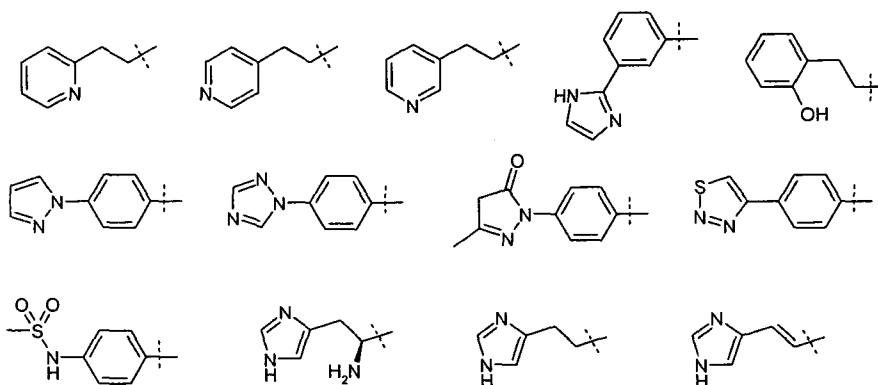
W_1 , W_2 , Y_1 , Y_3 , L , Z_1 , B , R_1 , R_2 , R_3 , R_4 have the meanings according to formulae (Ia) or (Ib) as claimed in claim 1 or 2;

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

4. The compound according to formulae (Ia), (Ib) and (II) as claimed in any of claims 1 to 3, wherein:



is independently substituted by a chemical group selected from the group consisting of:



and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

5. The compound according to formulae (Ia), (Ib) and (II) as claimed in any of claims 1 to 4, wherein:

W_1 , W_2 together independently form " $-N=N-$, $-C(O)-O-$, $-C(O)-S-$, $-C(S)-N(R_{5a})-$ ";

- Y₁ is independently selected from the group consisting of “-C(O)-, -N(R10)-C(O)-, -C(O)-N(R11)-, -OC(O)-, -S(O)₂-, single bond”;
- Z₁ is independently “O”;
- 5 B is independently selected from the group consisting of “(4-chloro-phenyl)-1H-[1,2,3]triazol-4-yl, [3,3']bithiophenyl-5-yl, 1H-benzotriazol-5-yl, 1H-imidazol-4-yl, 2-(4-chloro-phenyl)-4-methyl-thiazol-5-yl, 2-(4-chloro-phenyl)-cyclopropane-yl, 2-(4-trifluoromethyl-phenyl)-thiazol-5-yl, 2,3,5,6-tetrafluoro-4-trifluoromethyl-phenyl, 2,3-difluoro-4-trifluoromethyl-phenyl, 2,3-dihydro-benzo[1,4]dioxin-6-yl, 2,4-dichloro-phenyl, 2-chloro-phenyl, 2-fluoro-4-trifluoromethyl-phenyl, 2-fluoro-5-trifluoromethyl-phenyl, 2-methyl-2H-indazol-3-yl, 2-methyl-5-phenyl-furan-3-yl, 2-pyridin-2-yl, 3-(4-chloro-phenyl)-2-oxo-oxazolidin-5-yl, 3,4,5-trifluoro-phenyl, 3,4,5-trimethoxy-phenyl, 3,4-dichloro-phenyl, 3,4-difluoro-5-trifluoromethyl-phenyl, 3,4-dimethyl-phenyl, 3,5-bis-trifluoromethyl-phenyl, 3,5-dibromo-4-methyl-phenyl, 3,5-dibromo-phenyl, 3,5-dichloro-4-fluoro-phenyl, 3,5-dichloro-phenyl, 3,5-dimethoxy-phenyl, 3,5-dimethyl-phenyl, 3'-trifluoromethyl-biphenyl-2-yl, 3-bromo-4-trifluoromethoxy-phenyl, 3-bromo-5-chloro-phenyl, 3-bromo-5-fluoro-phenyl, 3-chloro-4,5-difluoro-phenyl, 3-chloro-4-fluoro-phenyl, 3-chloro-4-trifluoromethoxy-phenyl, 3-chloro-4-trifluoromethyl-phenyl, 3-chloro-5-fluoro-phenyl, 3-chloro-5-trifluoromethyl-phenyl, 3-chloro-phenyl, 3-fluoro-4-trifluoromethoxy-phenyl, 3-fluoro-4-trifluoromethyl-phenyl, 3-fluoro-5-trifluoromethyl-phenyl, 3-trifluoromethoxy-phenyl, 3-trifluoromethyl-phenyl, 4-(1,2,3-thiadiazol-4-yl)-phenyl, 4-(1,2,4-triazol-1-yl)-phenyl, 4-(3-methyl-5-oxo-4,5-dihydropyrazol-1-yl)-phenyl, 4'-methyl-biphenyl-2-yl, 4'-methyl-biphenyl-3-yl, 4-bromo-2,6-difluoro-phenyl, 4-bromo-2-fluoro-phenyl, 4-bromo-phenyl, 4-chloro-2-fluoro-phenyl, 4-chloro-3-fluoro-phenyl, 4-chloro-3-trifluoromethoxy-phenyl, 4-chloro-3-trifluoromethyl-phenyl, 4-chloro-phenyl, 4-cyano-phenyl, 4-difluoromethoxy-phenyl, 4-difluoromethylsulfanyl-phenyl, 4-ethyl-phenyl, 4-fluoro-3,5-dimethyl-phenyl, 4-fluoro-3-trifluoromethyl-phenyl, 4-isopropylphenyl, 4-methanesulfonyl-phenyl, 4-methoxy-3,5-dimethyl-phenyl, 4-methyl-2-(4-trifluoromethyl-phenyl)-thiazol-5-

yl, 4-methyl-3-trifluoromethyl-phenyl, 4-methyl-phenyl, 4-methylsulfanyl-phenyl, 4-nitro-phenyl, 4-trifluoromethoxy-phenyl, 4'-trifluoromethyl-biphenyl-2-yl, 4-trifluoromethyl-phenyl, 4-trifluoromethylsulfanyl-phenyl, 5-bromo-benzofuran-2-yl, 5-chloro-2-fluoro-3-trifluoromethyl-phenyl, 5-chloro-2-methoxy-phenyl, 5-trifluoromethyl-1H-benzimidazole-2-yl, benzo[1,3]dioxol-5-yl, phenyl, tetrahydrofuran-2-yl";

R1, R2, R3, R3a, R3b, R4, R4a, R4b, R5, R5a, R6, R7, R8, R9, R10, R11, R12, R13, R14, R15 are independently from each other selected from the group consisting of: "hydrogen, alkyl, methyl, ethyl, isopropyl, halogen, -F, -Br, -Cl, -CN, -CF₃, -SF₃, -OCF₃, -SCF₃, -OCHF₂, -SCHF₂, -O-alkyl, -O-methyl, -S-alkyl, -S-methyl, -NO₂, -S(O)₂-methyl"

V is independently selected from the group consisting of "hydrogen, alkyl, methyl, ethyl, isopropyl, halogen, -F, -Br, -Cl, -CN, -CF₃, -SF₃, -OCF₃, -SCF₃, -OCHF₂, -SCHF₂, =O, -O-alkyl, -O-methyl, -S-alkyl, -S-methyl, -NO₂, -S(O)₂-methyl";

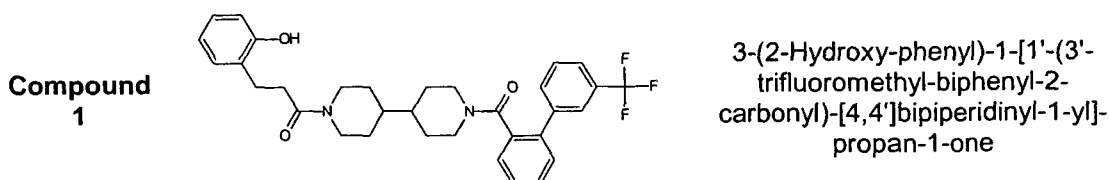
m is independently 0, 1 or 2;

n is independently 0, 1 or 2;

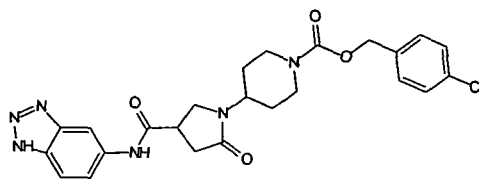
o is independently 0, 1 or 2;

and the physiologically acceptable salts, derivatives, prodrugs, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

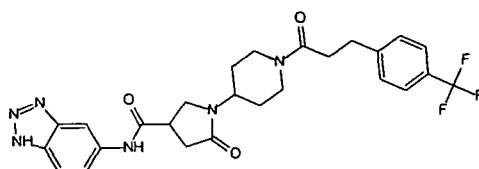
6. A compound selected from the group consisting of:



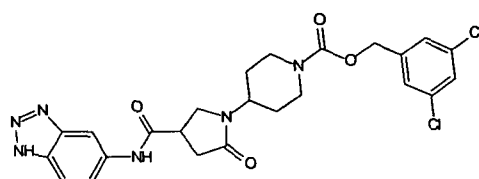
145

**Compound
2**

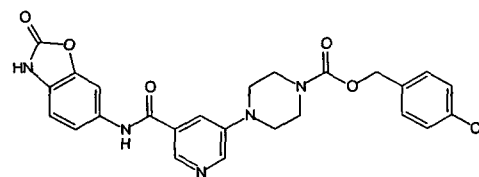
4-[4-(1H-Benzotriazol-5-ylcarbamoyl)-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 4-chloro-benzyl ester

**Compound
3**

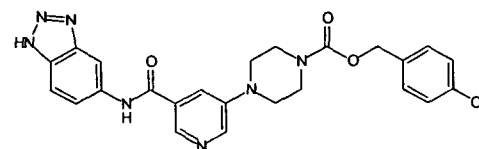
5-Oxo-1-{1-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-pyrrolidine-3-carboxylic acid (1H-benzotriazol-5-yl)-amide

**Compound
4**

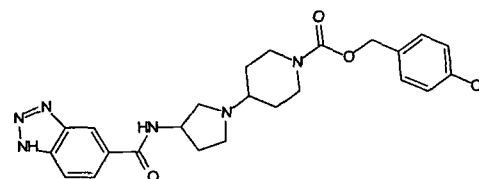
4-[4-(1H-Benzotriazol-5-ylcarbamoyl)-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

**Compound
5**

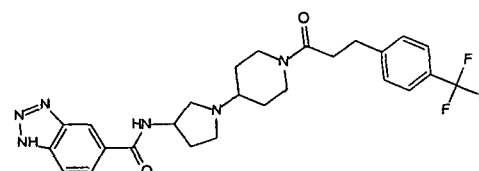
4-[5-(2-Oxo-2,3-dihydro-benzooxazol-6-ylcarbamoyl)-pyridin-3-yl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester

**Compound
6**

4-[5-(1H-Benzotriazol-5-ylcarbamoyl)-pyridin-3-yl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester

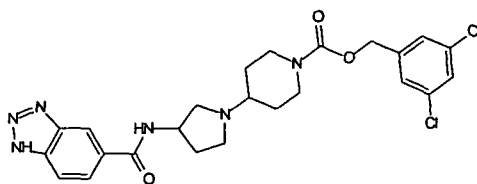
**Compound
7**

4-{3-[(1H-Benzotriazole-5-carbonyl)-amino]-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 4-chloro-benzyl ester

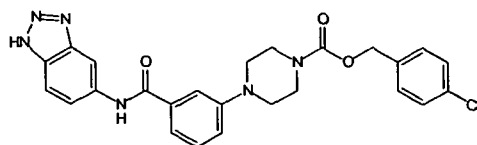
**Compound
8**

1H-Benzotriazole-5-carboxylic acid (1-{1-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-pyrrolidin-3-yl)-amide

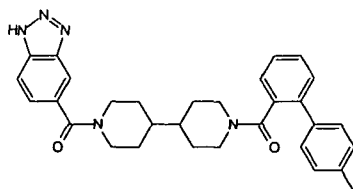
146

Compound 9

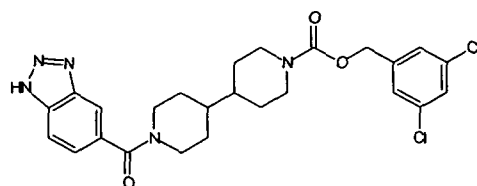
4-{3-[(1H-Benzotriazole-5-carbonyl)-amino]-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

Compound 10

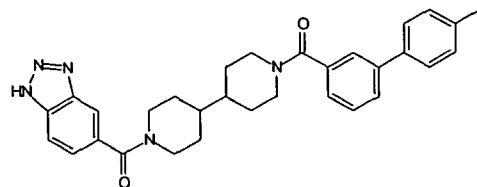
4-[3-(1H-Benzotriazol-5-ylcarbonyl)-phenyl]-piperazine-1-carboxylic acid 4-chloro-benzyl ester

Compound 11

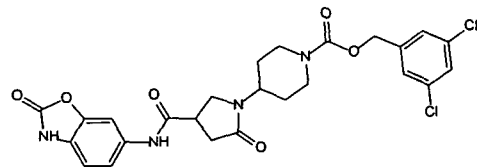
[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidiny-1-yl]-(4'-methyl-biphenyl-2-yl)-methanone

Compound 12

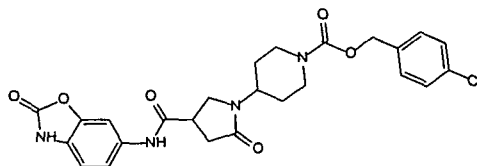
1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidiny-1-carboxylic acid 3,5-dichloro-benzyl ester

Compound 13

[1'-(1H-Benzotriazole-5-carbonyl)-[4,4']bipiperidiny-1-yl]-(4'-methyl-biphenyl-3-yl)-methanone

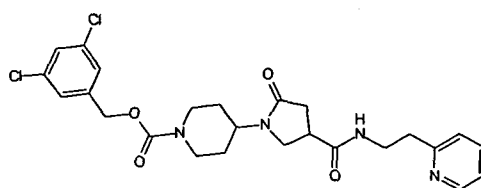
Compound 15

4-[2-Oxo-4-(2-oxo-2,3-dihydro-benzooxazol-6-ylcarbonyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

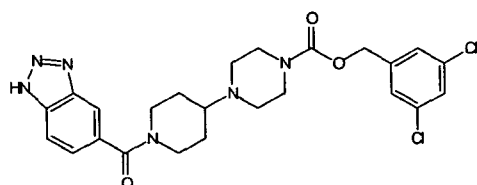
Compound 16

4-[2-Oxo-4-(2-oxo-2,3-dihydro-benzooxazol-6-ylcarbonyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 4-chloro-benzyl ester

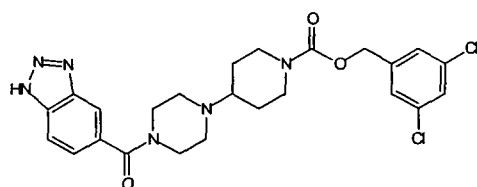
147

**Compound
17**

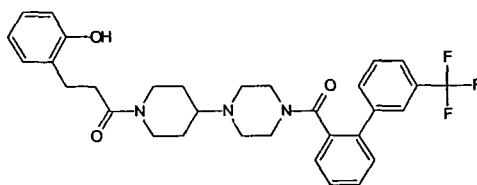
4-[2-Oxo-4-(2-pyridin-2-yl-
ethylcarbamoyl)-pyrrolidin-1-yl]-
piperidine-1-carboxylic acid 3,5-
dichloro-benzyl ester

**Compound
19**

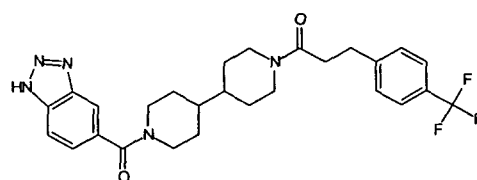
4-[1-(1H-Benzotriazole-5-
carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 3,5-
dichloro-benzyl ester

**Compound
20**

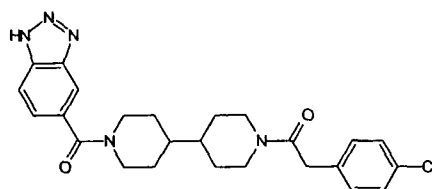
4-[4-(1H-Benzotriazole-5-
carbonyl)-piperazin-1-yl]-
piperidine-1-carboxylic acid 3,5-
dichloro-benzyl ester

**Compound
21**

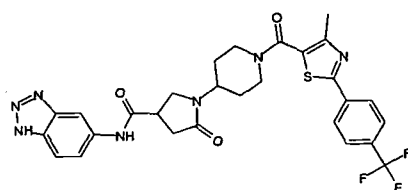
3-(2-Hydroxy-phenyl)-1-{4-[4-(3'-
trifluoromethyl-biphenyl-2-
carbonyl)-piperazin-1-yl]-piperidin-
1-yl}-propan-1-one

**Compound
22**

1-[1'-(1H-Benzotriazole-5-
carbonyl)-[4,4']bipiperidinyl-1-yl]-
3-(4-trifluoromethyl-phenyl)-
propan-1-one

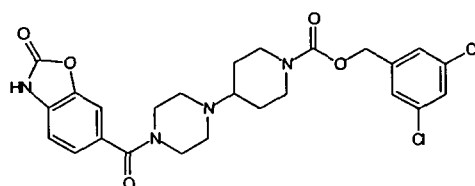
**Compound
23**

1-[1'-(1H-Benzotriazole-5-
carbonyl)-[4,4']bipiperidinyl-1-yl]-
2-(4-chloro-phenyl)-ethanone

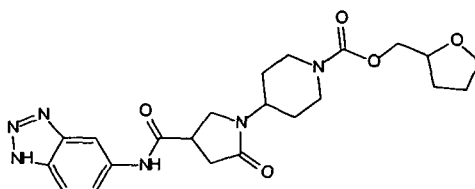
**Compound
24**

1-[1-(4-Methyl-2-(4-
trifluoromethyl-phenyl)-thiazole-5-
carbonyl)-piperidin-4-yl]-5-oxo-
pyrrolidine-3-carboxylic acid (1H-
benzotriazol-5-yl)-amide

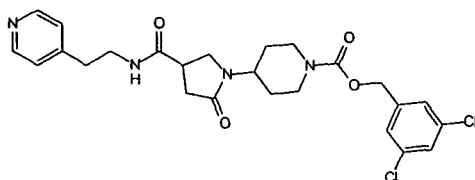
148

**Compound
25**

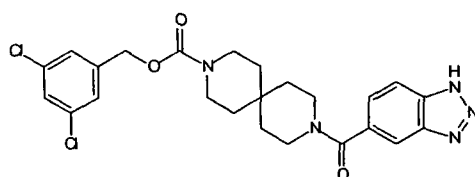
4-[4-(2-Oxo-2,3-dihydro-benzooxazole-6-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

**Compound
27**

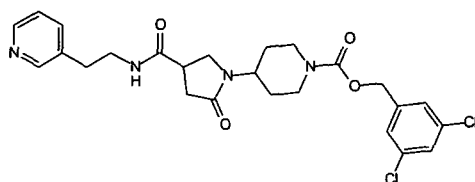
4-[4-(1H-Benzotriazol-5-ylcarbonyl)-2-oxo-pyrrolidin-1-yl]-piperidine-1-carboxylic acid tetrahydro-furan-2-ylmethyl ester

**Compound
28**

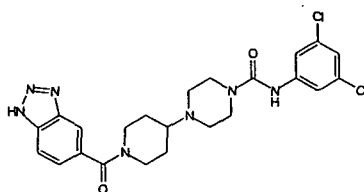
4-[2-Oxo-4-(2-pyridin-4-ylethylcarbonyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

**Compound
29**

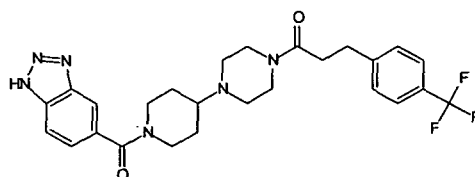
9-(1H-Benzotriazole-5-carbonyl)-3,9-diaza-spiro[5.5]undecane-3-carboxylic acid 3,5-dichloro-benzyl ester

**Compound
30**

4-[2-Oxo-4-(2-pyridin-3-ylethylcarbonyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

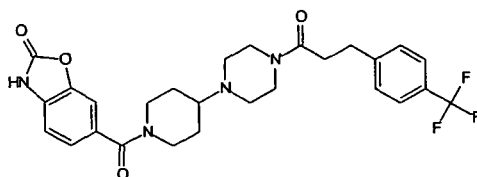
**Compound
31**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (3,5-dichloro-phenyl)-amide

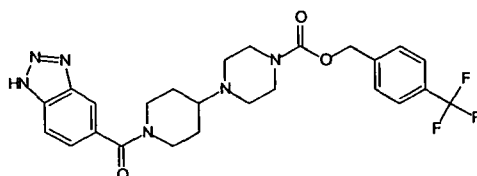
**Compound
32**

1-[4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl]-3-(4-trifluoromethyl-phenyl)-propan-1-one

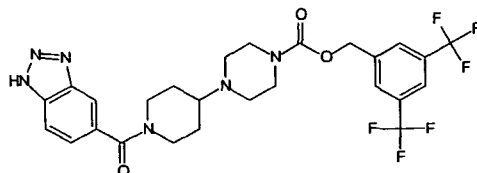
149

**Compound
33**

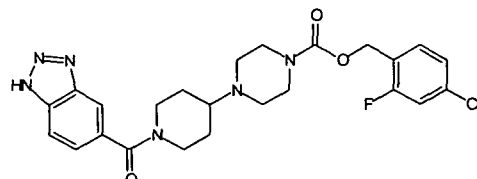
6-(4-{4-[3-(4-Trifluoromethyl-phenyl)-propionyl]-piperazin-1-yl}-piperidine-1-carbonyl)-3H-benzoxazol-2-one

**Compound
34**

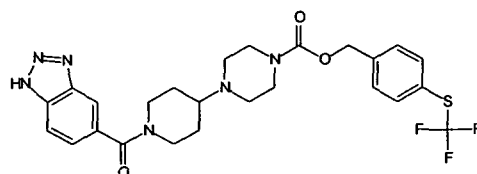
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

**Compound
35**

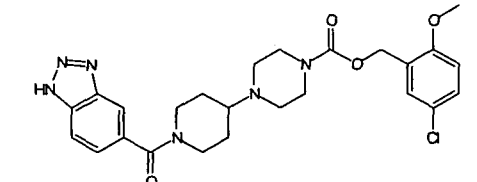
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

**Compound
36**

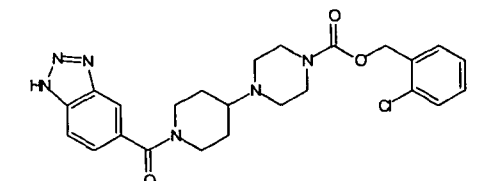
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester

**Compound
37**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzyl ester

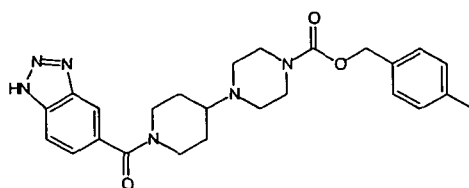
**Compound
38**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 5-chloro-2-methoxy-benzyl ester

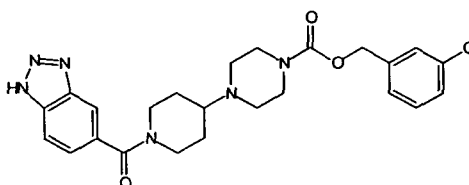
**Compound
39**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-chloro-benzyl ester

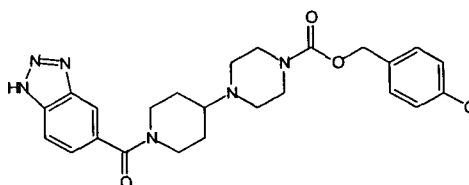
150

**Compound
40**

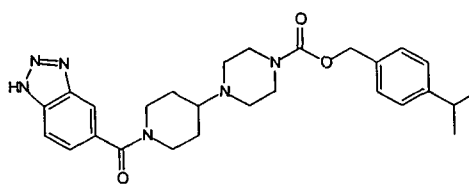
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 4-
methyl-benzyl ester

**Compound
41**

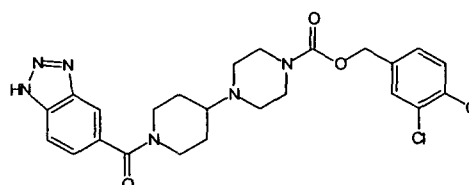
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 3-
chloro-benzyl ester

**Compound
42**

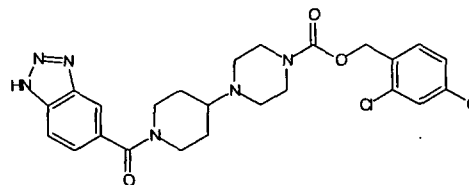
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 4-
chloro-benzyl ester

**Compound
43**

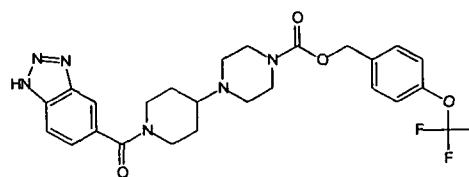
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 4-
isopropyl-benzyl ester

**Compound
45**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 3,4-
dichloro-benzyl ester

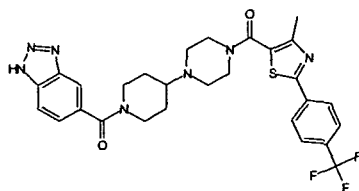
**Compound
46**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 2,4-
dichloro-benzyl ester

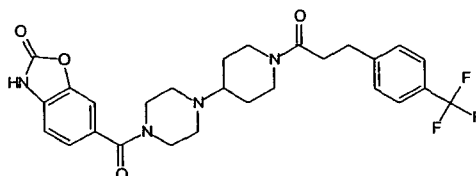
**Compound
47**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-
piperazine-1-carboxylic acid 4-
trifluoromethoxy-benzyl ester

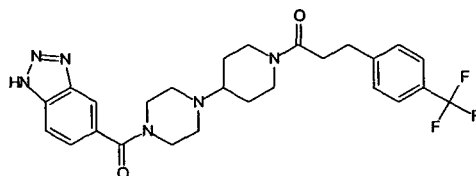
151

Compound 48

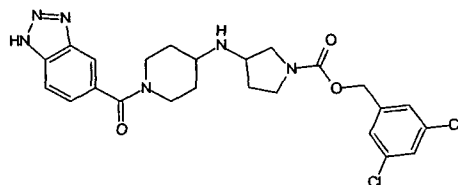
(1H-Benzotriazol-5-yl)-(4-{4-[4-methyl-2-(4-trifluoromethyl-phenyl)-thiazole-5-carbonyl]-piperazin-1-yl}-piperidin-1-yl)-methanone

Compound 49

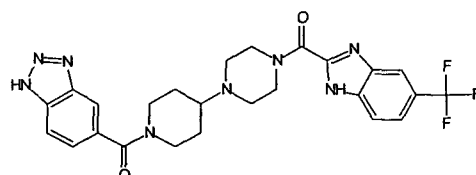
6-(4-{1-[3-(4-Trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-piperazine-1-carbonyl)-3H-benzooxazol-2-one

Compound 50

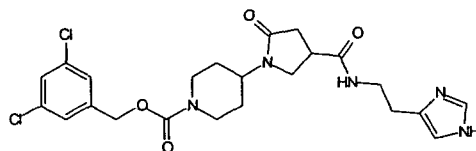
1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-trifluoromethyl-phenyl)-propan-1-one

Compound 51

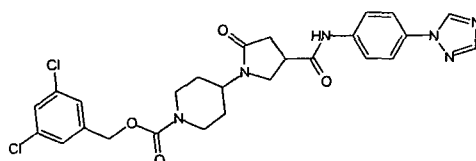
3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidine-1-carboxylic acid 3,5-dichloro-benzyl ester

Compound 52

(1H-Benzotriazol-5-yl)-(4-{4-[4-(5-trifluoromethyl-1H-benzimidazole-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-methanone

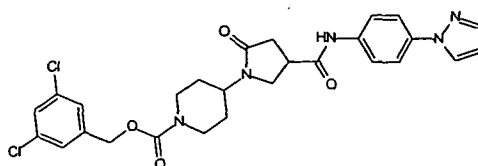
Compound 53

4-{4-[2-(1H-Imidazol-4-yl)-ethylcarbamoyl]-2-oxo-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

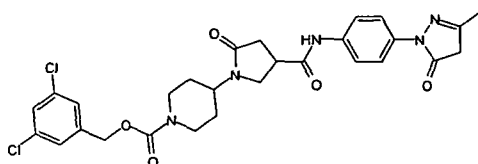
Compound 54

4-[2-Oxo-4-(4-[1,2,4]triazol-1-yl)-phenylcarbamoyl]-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

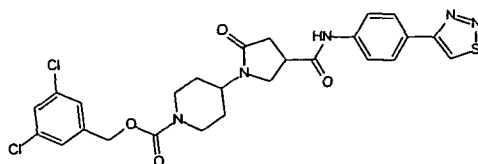
152

Compound 55

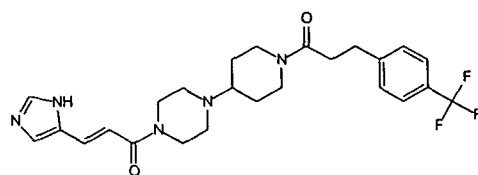
4-[2-Oxo-4-(4-pyrazol-1-yl-phenylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

Compound 56

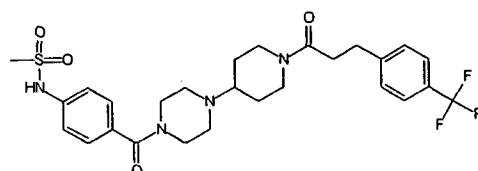
4-{4-[4-(3-Methyl-5-oxo-4,5-dihydro-pyrazol-1-yl)-phenylcarbamoyl]-2-oxo-pyrrolidin-1-yl}-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

Compound 57

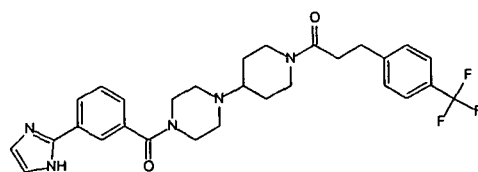
4-[2-Oxo-4-(4-[1,2,3]thiadiazol-4-yl-phenylcarbamoyl)-pyrrolidin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-benzyl ester

Compound 58

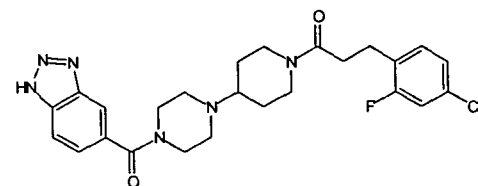
(E)-3-(3H-Imidazol-4-yl)-1-(4-{1-[3-(4-trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-piperazin-1-yl)-propanone

Compound 59

N-[4-(4-{1-[3-(4-Trifluoromethyl-phenyl)-propionyl]-piperidin-4-yl}-piperazine-1-carbonyl)-phenyl]-methanesulfonamide

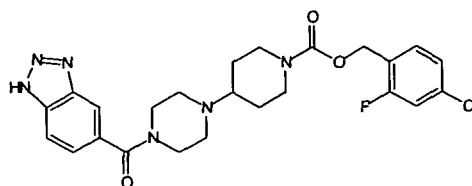
Compound 60

1-(4-{4-[3-(1H-Imidazol-2-yl)-benzoyl]-piperazin-1-yl}-piperidin-1-yl)-3-(4-trifluoromethyl-phenyl)-propan-1-one

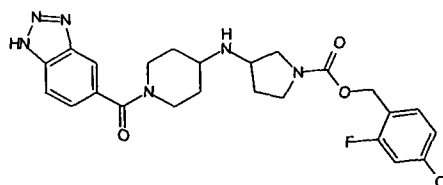
Compound 61

1-(4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl)-3-(4-chloro-2-fluoro-phenyl)-propan-1-one

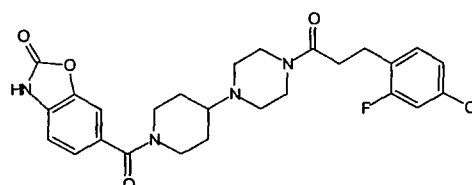
153

Compound 62

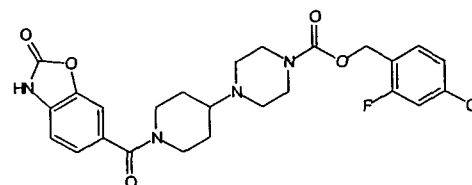
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester

Compound 63

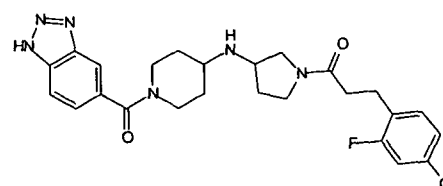
3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester

Compound 64

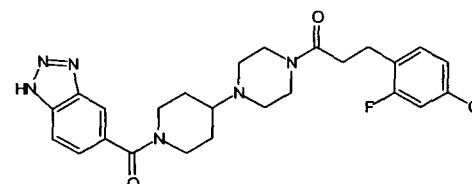
6-(4-{4-[3-(4-Chloro-2-fluorophenyl)-propionyl]-piperazin-1-yl}-piperidine-1-carbonyl)-3H-benzooxazol-2-one

Compound 65

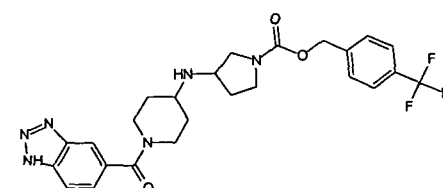
4-[1-(2-Oxo-2,3-dihydro-benzooxazole-6-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-2-fluoro-benzyl ester

Compound 66

1-{3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidin-1-yl}-3-(4-chloro-2-fluoro-phenyl)-propan-1-one

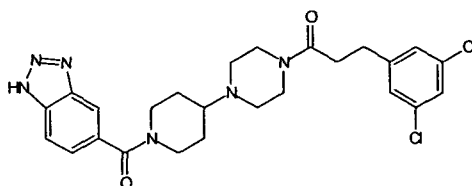
Compound 67

1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-chloro-2-fluoro-phenyl)-propan-1-one

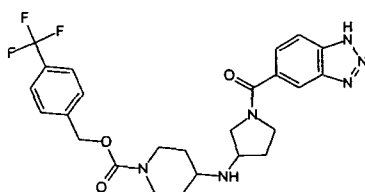
Compound 68

3-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylamino]-pyrrolidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

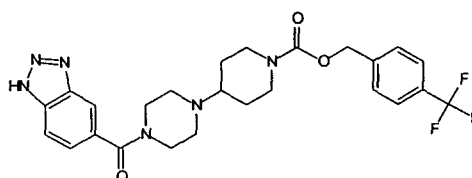
154

**Compound
69**

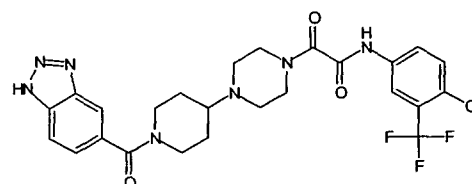
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,5-dichloro-phenyl)-propan-1-one

**Compound
70**

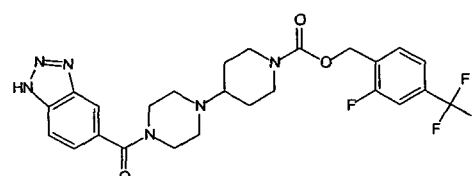
4-[1-(1H-Benzotriazole-5-carbonyl)-pyrrolidin-3-ylamino]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

**Compound
71**

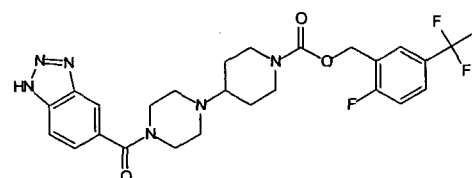
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

**Compound
72**

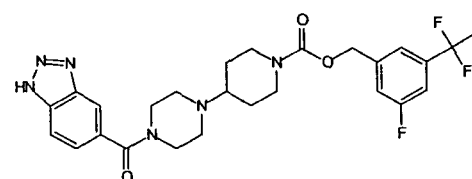
2-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-N-(4-chloro-3-trifluoromethyl-phenyl)-2-oxo-acetamide

**Compound
73**

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 2-fluoro-4-trifluoromethyl-benzyl ester

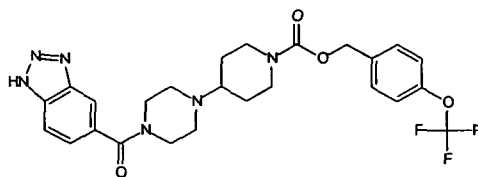
**Compound
74**

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 2-fluoro-5-trifluoromethyl-benzyl ester

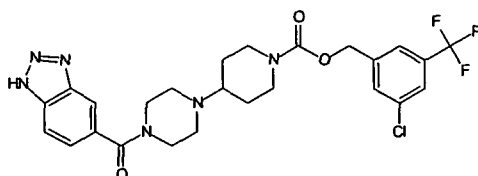
**Compound
75**

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-fluoro-5-trifluoromethyl-benzyl ester

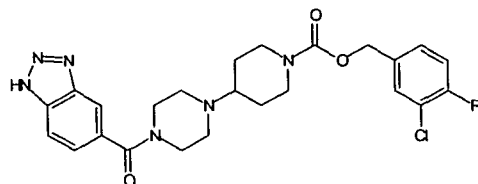
155

**Compound
76**

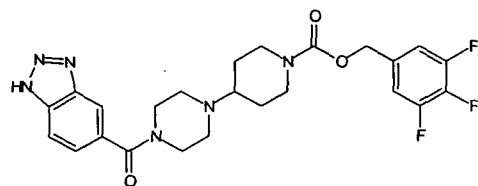
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethoxy-benzyl ester

**Compound
77**

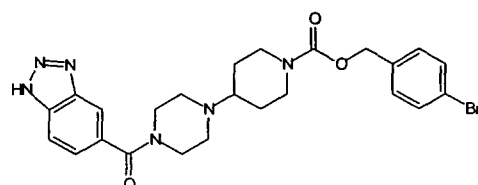
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester

**Compound
78**

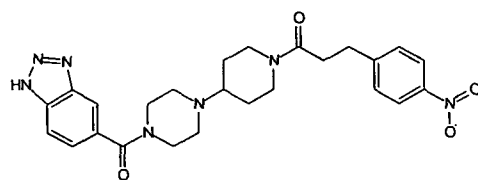
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-4-fluoro-benzyl ester

**Compound
79**

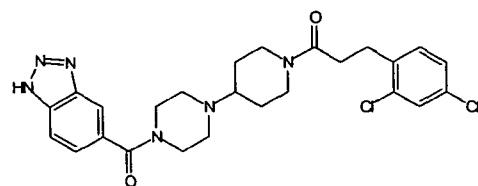
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,4,5-trifluoro-benzyl ester

**Compound
80**

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-bromo-benzyl ester

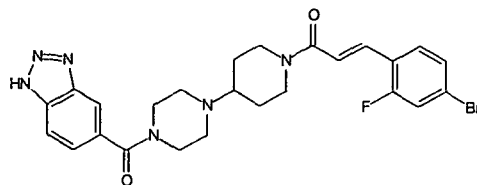
**Compound
81**

1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-nitro-phenyl)-propan-1-one

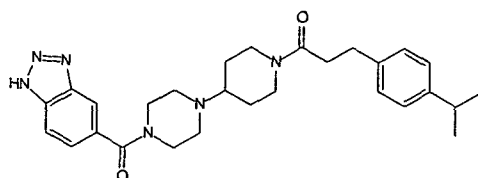
**Compound
82**

1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(2,4-dichloro-phenyl)-propan-1-one

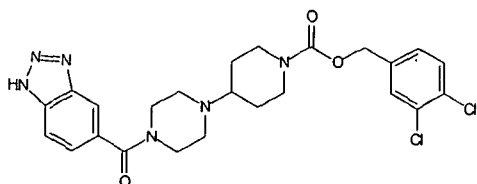
156

Compound 83

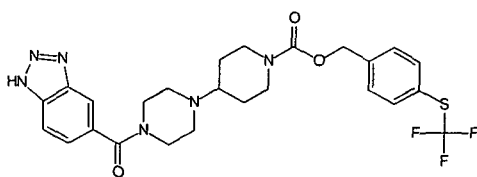
(E)-1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-bromo-2-fluoro-phenyl)propan-1-one

Compound 84

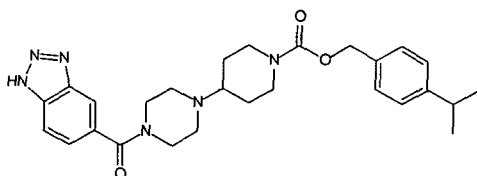
1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-isopropyl-phenyl)propan-1-one

Compound 85

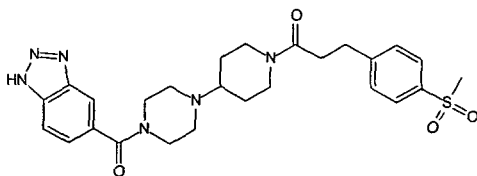
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,4-dichloro-benzyl ester

Compound 86

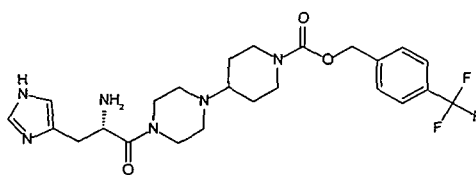
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzyl ester

Compound 87

4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-isopropyl-benzyl ester

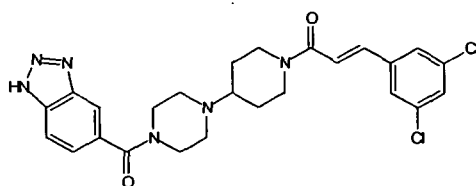
Compound 88

1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-methanesulfonyl-phenyl)propan-1-one

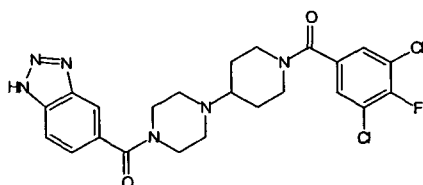
Compound 89

4-{4-[(S)-2-Amino-3-(1H-imidazol-4-yl)-propionyl]-piperazin-1-yl}-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

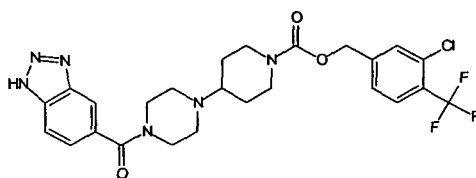
157

Compound 90

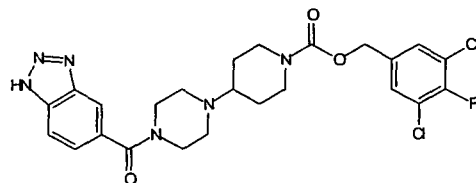
(E)-1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(3,5-dichloro-phenyl)-propenone

Compound 91

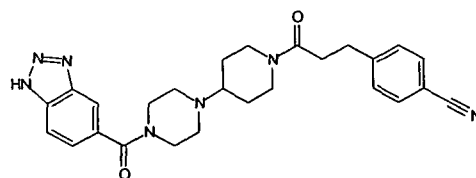
(1H-Benzotriazol-5-yl)-{4-[1-(3,5-dichloro-4-fluoro-benzoyl)-piperidin-4-yl]-piperazin-1-yl}-methanone

Compound 92

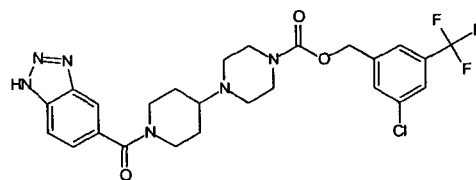
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-4-trifluoromethyl-benzyl ester

Compound 93

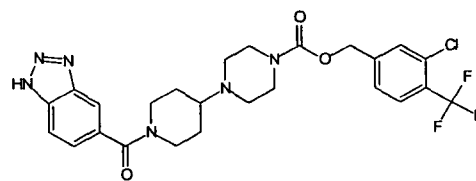
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-dichloro-4-fluoro-benzyl ester

Compound 94

4-(3-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-oxo-propyl)-benzonitrile

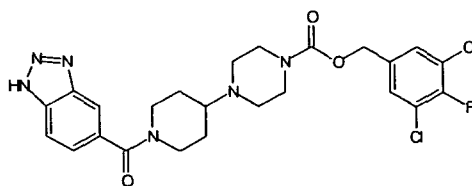
Compound 95

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester

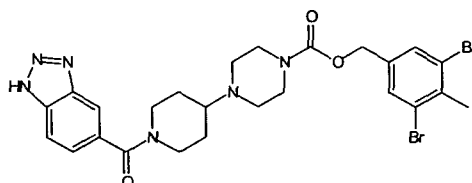
Compound 96

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-4-trifluoromethyl-benzyl ester

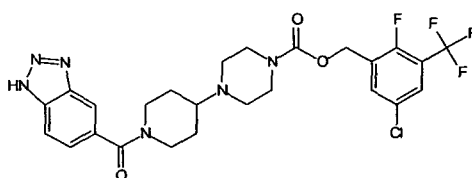
158

**Compound
97**

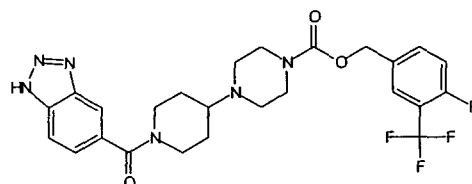
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dichloro-4-fluoro-benzyl ester

**Compound
98**

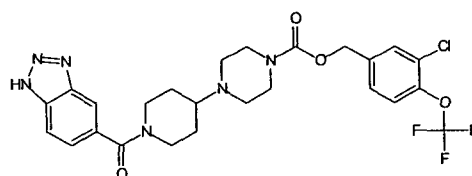
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dibromo-4-methyl-benzyl ester

**Compound
99**

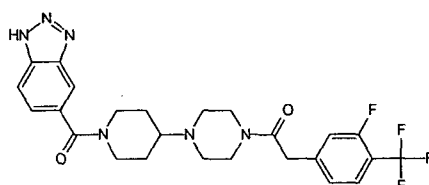
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 5-chloro-2-fluoro-3-trifluoromethyl-benzyl ester

**Compound
100**

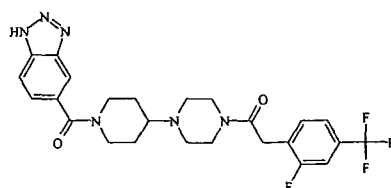
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-fluoro-3-trifluoromethyl-benzyl ester

**Compound
101**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-4-trifluoromethoxy-benzyl ester

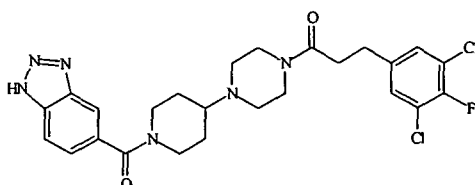
**Compound
102**

1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(3-fluoro-4-trifluoromethyl-phenyl)-ethanone

**Compound
103**

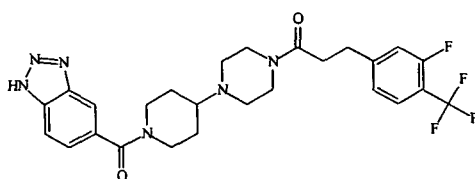
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(2-fluoro-4-trifluoromethyl-phenyl)-ethanone

5

**Compound
104**

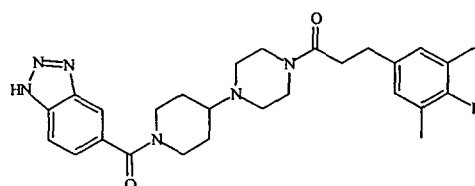
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,5-dichloro-4-fluorophenyl)-propan-1-one

10

**Compound
105**

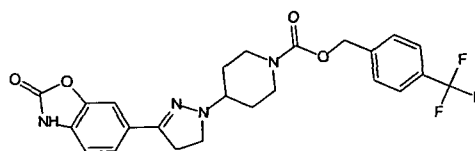
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3-fluoro-4-trifluoromethylphenyl)-propan-1-one

15

**Compound
106**

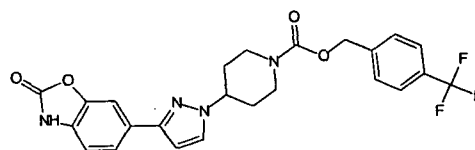
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-fluoro-3,5-dimethylphenyl)-propan-1-one

20

**Compound
107**

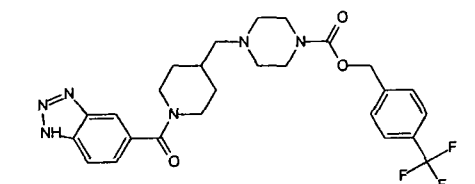
4-[3-(2-Oxo-2,3-dihydrobenzooxazol-6-yl)-4,5-dihydropyrazol-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethylbenzyl ester

25

**Compound
108**

4-[3-(2-Oxo-2,3-dihydrobenzooxazol-6-yl)-pyrazol-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethylbenzyl ester

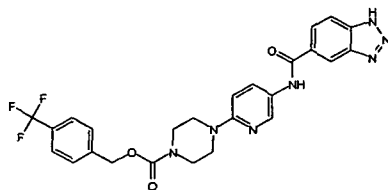
30

**Compound
109**

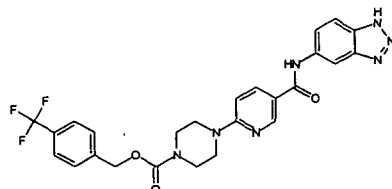
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-ylmethyl]-piperazine-1-carboxylic acid 4-trifluoromethylbenzyl ester

35

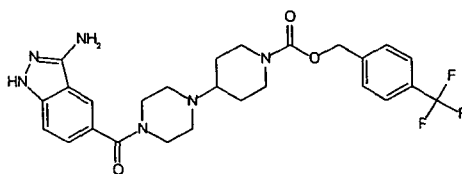
160

**Compound
110**

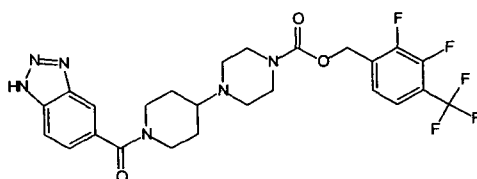
4-{5-[(1H-Benzotriazole-5-carbonyl)-amino]-pyridin-2-yl}-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

**Compound
111**

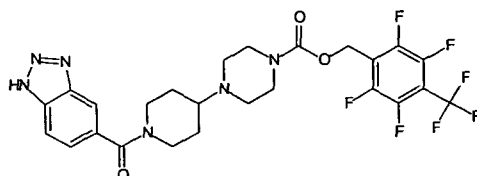
4-[5-(1H-Benzotriazol-5-ylcarbonyl)-pyridin-2-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

**Compound
112**

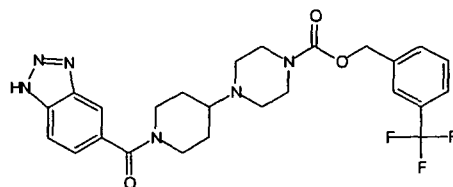
4-[4-(3-Amino-1H-indazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

**Compound
113**

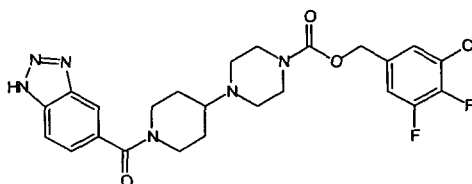
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,3-difluoro-4-trifluoromethyl-benzyl ester

**Compound
114**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,3,5,6-tetrafluoro-4-trifluoromethyl-benzyl ester

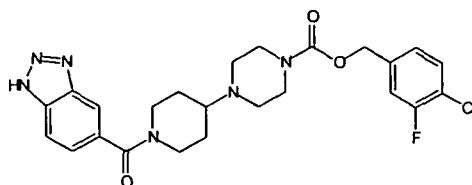
**Compound
115**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-trifluoromethyl-benzyl ester

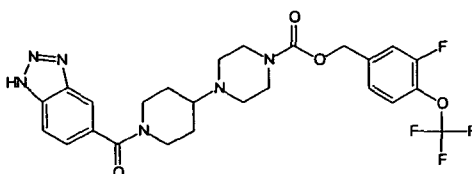
**Compound
116**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-4,5-difluoro-benzyl ester

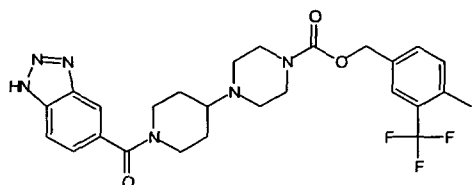
161

**Compound
117**

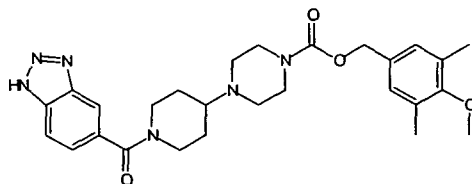
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-3-fluoro-benzyl ester

**Compound
118**

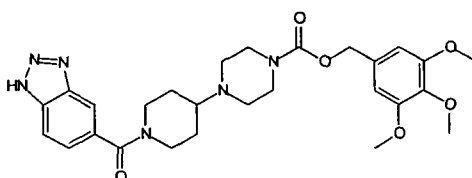
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-fluoro-4-trifluoromethoxy-benzyl ester

**Compound
119**

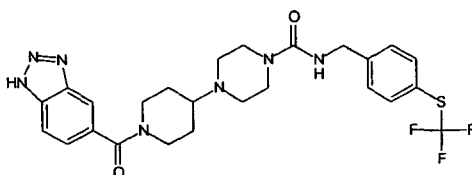
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-methyl-3-trifluoromethyl-benzyl ester

**Compound
120**

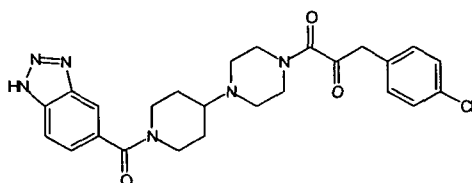
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-methoxy-3,5-dimethyl-benzyl ester

**Compound
121**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,4,5-trimethoxy-benzyl ester

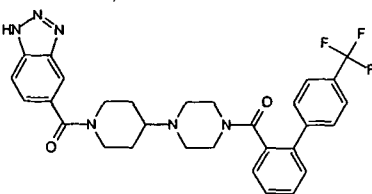
**Compound
122**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzylamide

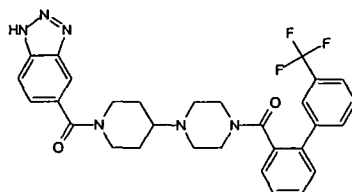
**Compound
123**

1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-chloro-phenyl)-propane-1,2-dione

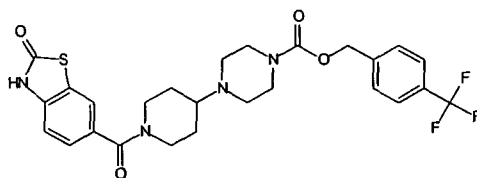
162

**Compound
124**

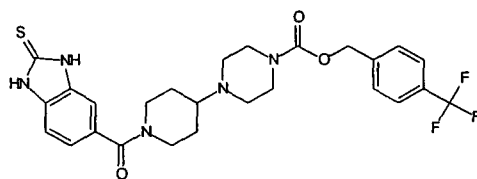
(1H-Benzotriazol-5-yl)-{4-[4-(4'-trifluoromethyl-biphenyl-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-methanone

**Compound
125**

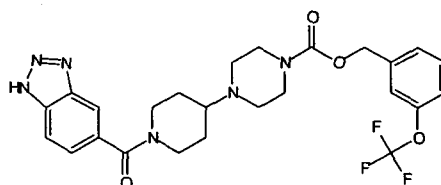
(1H-Benzotriazol-5-yl)-{4-[4-(3'-trifluoromethyl-biphenyl-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-methanone

**Compound
126**

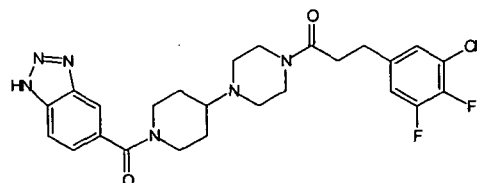
4-[1-(2-Oxo-2,3-dihydro-benzothiazole-6-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

**Compound
127**

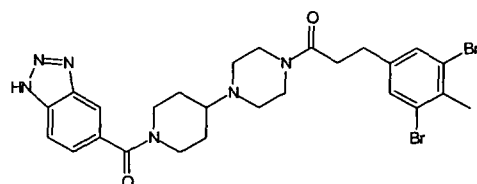
4-[1-(2-Thioxo-2,3-dihydro-1H-benzimidazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

**Compound
128**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-trifluoromethoxy-benzyl ester

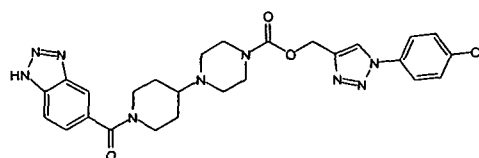
**Compound
129**

1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3-chloro-4,5-difluorophenyl)-propan-1-one

**Compound
130**

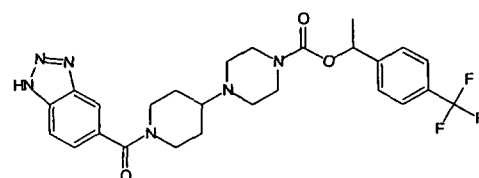
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,5-dibromo-4-methylphenyl)-propan-1-one

5

Compound 131

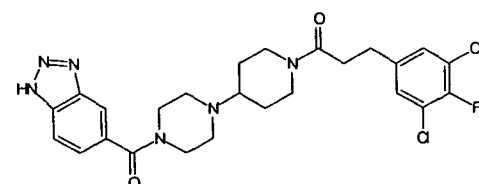
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(4-chloro-phenyl)-1H-[1,2,3]triazol-4-ylmethyl ester

10

Compound 132

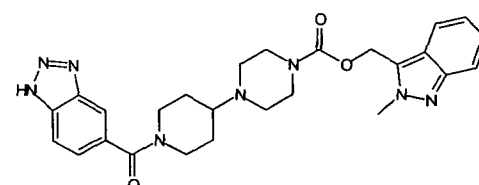
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(4-trifluoromethyl-phenyl)-ethyl ester

15

Compound 133

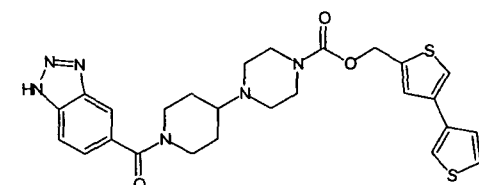
1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(3,5-dichloro-4-fluoro-phenyl)-propan-1-one

20

Compound 134

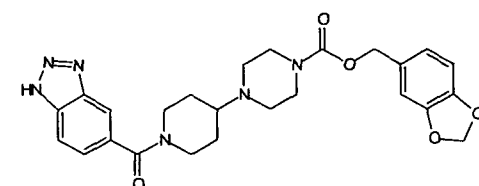
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-methyl-2H-indazol-3-ylmethyl ester

25

Compound 135

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid [3,3']bithiophenyl-5-ylmethyl ester

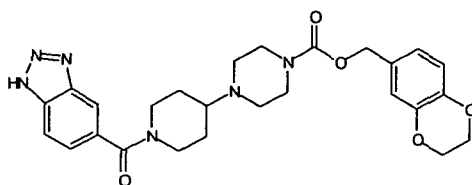
30

Compound 136

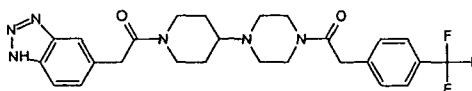
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid benzo[1,3]dioxol-5-ylmethyl ester

35

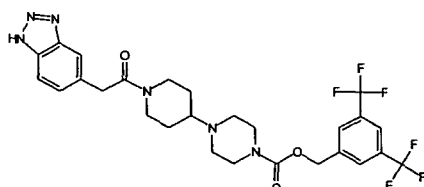
164

**Compound
137**

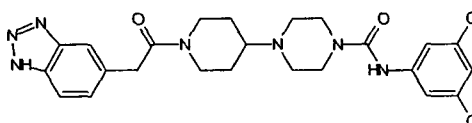
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,3-dihydro-benzo[1,4]dioxin-6-ylmethyl ester

**Compound
138**

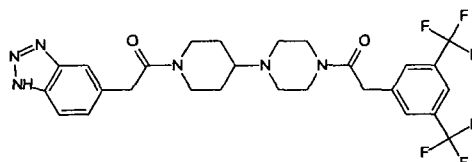
2-(1H-Benzotriazol-5-yl)-1-(4-{4-[2-(4-trifluoromethyl-phenyl)-acetyl]-piperazin-1-yl}-piperidin-1-yl)-ethanone

**Compound
139**

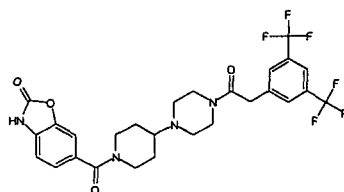
4-[1-(2-1H-Benzotriazol-5-yl)-acetyl]-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

**Compound
140**

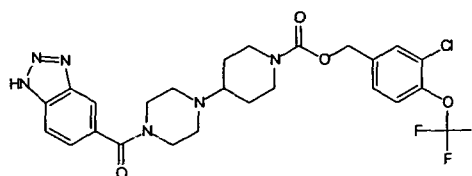
4-[1-(2-1H-Benzotriazol-5-yl)-acetyl]-piperidin-4-yl]-piperazine-1-carboxylic acid (3,5-dichloro-phenyl)-amide

**Compound
141**

2-(1H-Benzotriazol-5-yl)-1-(4-{4-[2-(3,5-bis-trifluoromethyl-phenyl)-acetyl]-piperazin-1-yl}-piperidin-1-yl)-ethanone

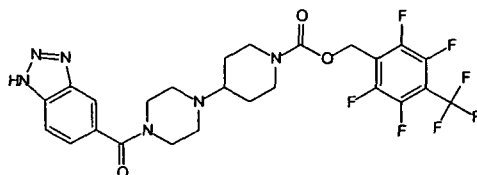
**Compound
142**

6-(4-{4-[2-(3,5-Bis-trifluoromethyl-phenyl)-acetyl]-piperazin-1-yl}-piperidine-1-carbonyl)-3H-benzooxazol-2-one

**Compound
143**

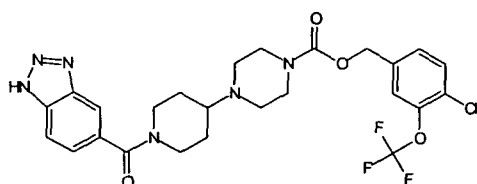
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3-chloro-4-trifluoromethoxy-benzyl ester

5

**Compound
144**

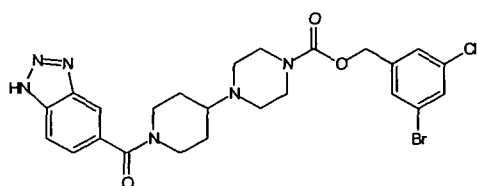
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 2,3,5,6-tetrafluoro-4-trifluoromethyl-benzyl ester

10

**Compound
145**

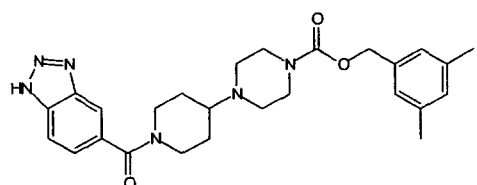
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-chloro-3-trifluoromethoxy-benzyl ester

15

**Compound
146**

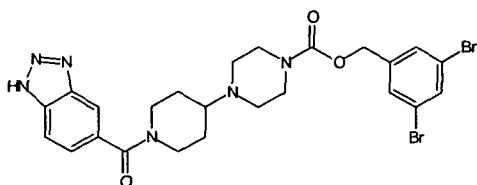
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-bromo-5-chloro-benzyl ester

20

**Compound
147**

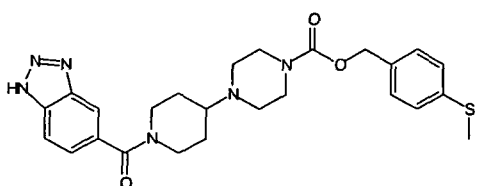
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dimethyl-benzyl ester

25

**Compound
148**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dibromo-benzyl ester

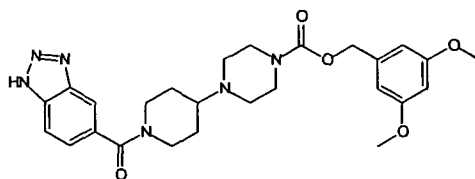
30

**Compound
149**

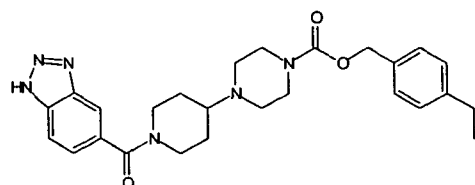
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-methylsulfanyl-benzyl ester

35

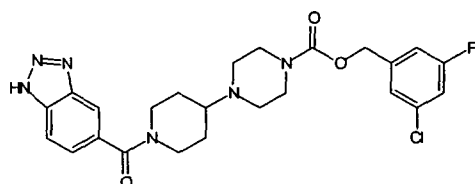
166

**Compound
150**

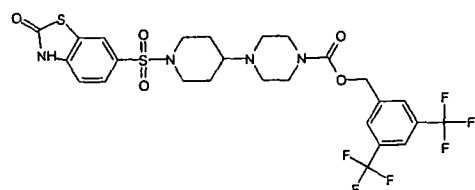
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-dimethoxy-benzyl ester

**Compound
151**

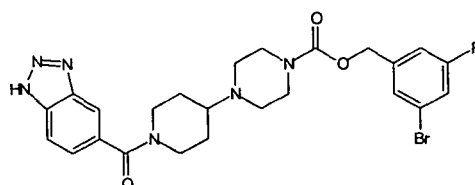
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-ethyl-benzyl ester

**Compound
152**

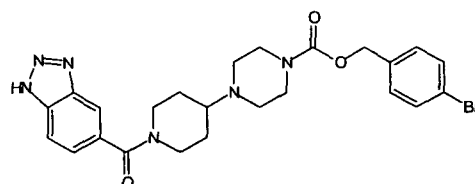
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-fluoro-benzyl ester

**Compound
153**

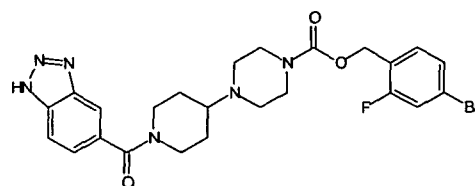
4-[1-(2-Oxo-2,3-dihydro-benzothiazole-6-sulfonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

**Compound
154**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-bromo-5-fluoro-benzyl ester

**Compound
155**

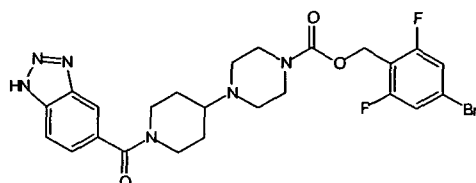
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-bromo-benzyl ester

**Compound
156**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-bromo-2-fluoro-benzyl ester

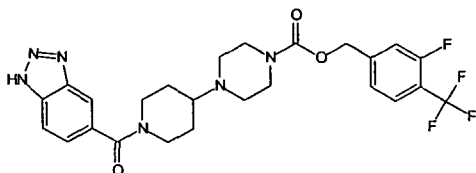
167

5

**Compound
157**

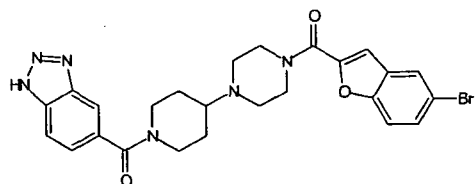
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-bromo-2,6-difluoro-benzyl ester

10

**Compound
158**

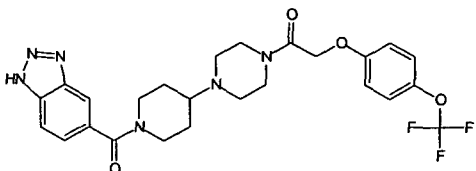
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-fluoro-4-trifluoromethyl-benzyl ester

15

**Compound
159**

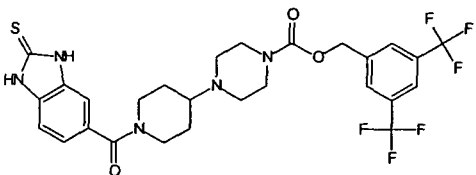
(1H-Benzotriazol-5-yl)-{4-[4-(5-bromo-benzofuran-2-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-methanone

20

**Compound
160**

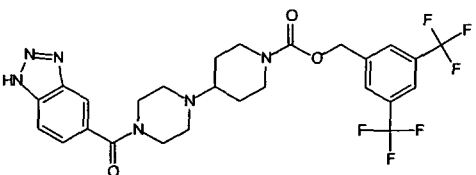
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-2-(4-trifluoromethoxy-phenoxy)-ethanone

25

**Compound
161**

4-[1-(2-Thioxo-2,3-dihydro-1H-benzimidazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

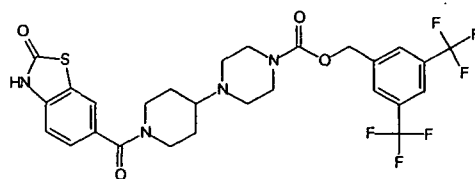
30

**Compound
162**

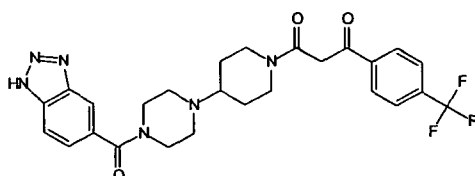
4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

35

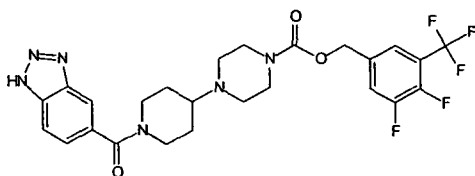
168

Compound 163

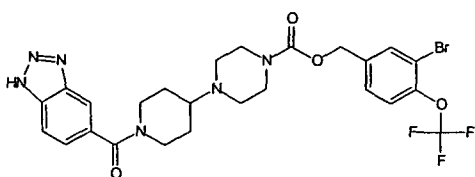
4-[1-(2-Oxo-2,3-dihydro-benzothiazole-6-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

Compound 164

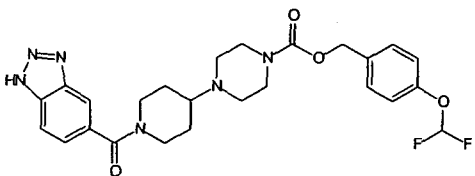
1-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-3-(4-trifluoromethyl-phenyl)-propane-1,3-dione

Compound 165

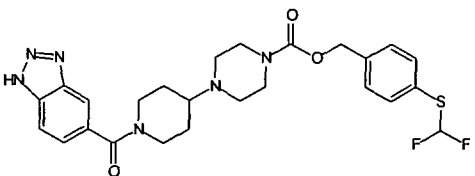
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3,4-difluoro-5-trifluoromethyl-benzyl ester

Compound 166

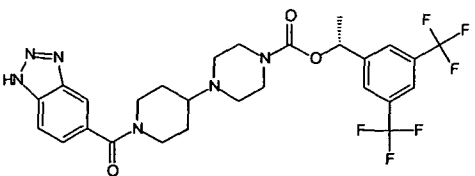
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-bromo-4-trifluoromethoxy-benzyl ester

Compound 167

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-difluoromethoxy-benzyl ester

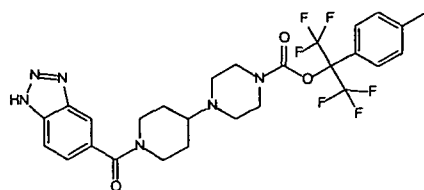
Compound 168

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 4-difluoromethylsulfanyl-benzyl ester

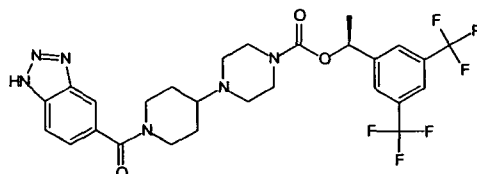
Compound 169

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (R)-1-(3,5-bis-trifluoromethyl-phenyl)-ethyl ester

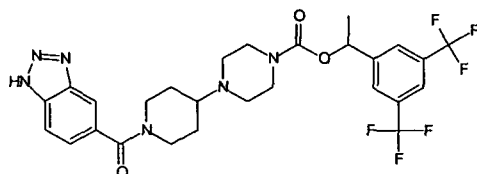
169

**Compound
170**

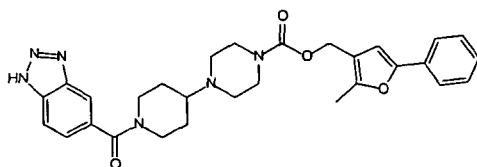
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2,2,2-trifluoro-1-p-tolyl-1-trifluoromethyl-ethyl ester

**Compound
171**

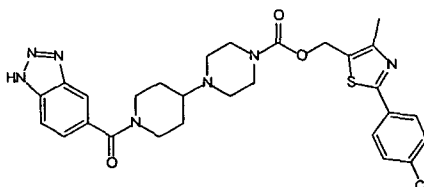
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid (S)-1-(3,5-bis-trifluoromethyl-phenyl)-ethyl ester

**Compound
172**

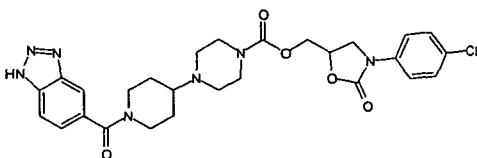
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(3,5-bis-trifluoromethyl-phenyl)-ethyl ester

**Compound
173**

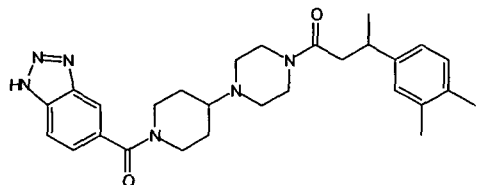
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-methyl-5-phenyl-furan-3-ylmethyl ester

**Compound
174**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 2-(4-chloro-phenyl)-4-methyl-thiazol-5-ylmethyl ester

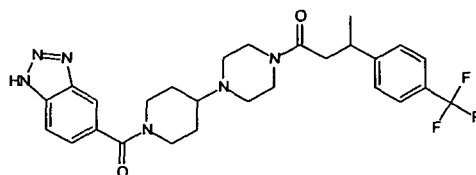
**Compound
175**

4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-(4-chloro-phenyl)-2-oxo-oxazolidin-5-ylmethyl ester

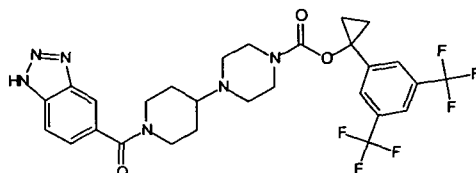
**Compound
176**

1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(3,4-dimethyl-phenyl)-butan-1-one

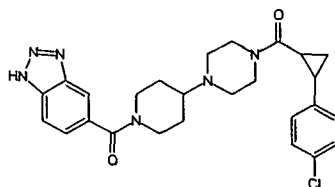
170

**Compound
177**

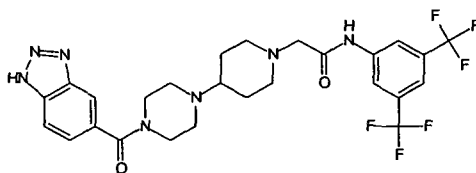
1-{4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazin-1-yl}-3-(4-trifluoromethyl-phenyl)-butan-1-one

**Compound
178**

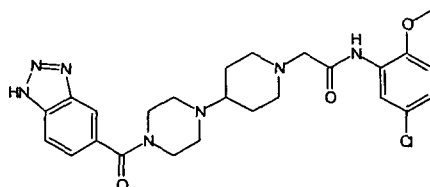
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 1-(3,5-bis-trifluoromethyl-phenyl)-cyclopropyl ester

**Compound
179**

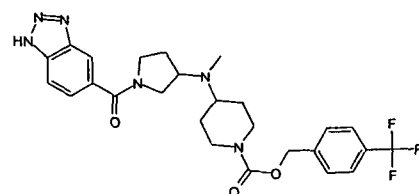
(1H-Benzotriazol-5-yl)-(4-{4-[2-(4-chloro-phenyl)-cyclopropanecarbonyl]-piperazin-1-yl}-piperidin-1-yl)-methanone

**Compound
180**

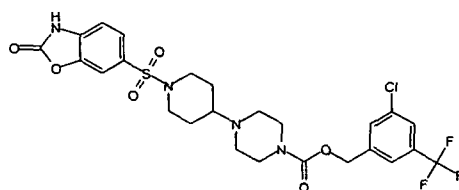
2-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-N-(3,5-bis-trifluoromethyl-phenyl)-acetamide

**Compound
181**

2-{4-[4-(1H-Benzotriazole-5-carbonyl)-piperazin-1-yl]-piperidin-1-yl}-N-(5-chloro-2-methoxy-phenyl)-acetamide

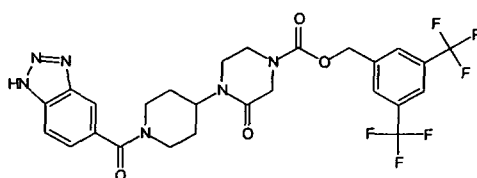
**Compound
182**

4-{[1-(1H-Benzotriazole-5-carbonyl)-pyrrolidin-3-yl]-methyl-amino}-piperidine-1-carboxylic acid 4-trifluoromethyl-benzyl ester

**Compound
183**

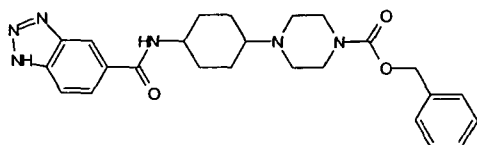
4-[1-(2-Oxo-2,3-dihydro-benzooxazole-6-sulfonyl)-piperidin-4-yl]-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester

5

**Compound
184**

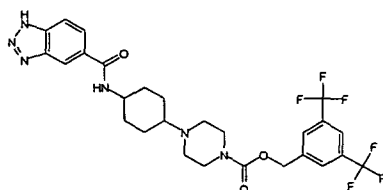
4-[1-(1H-Benzotriazole-5-carbonyl)-piperidin-4-yl]-3-oxo-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

10

**Compound
185**

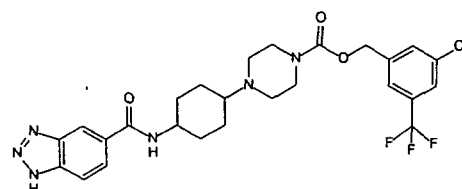
4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid benzyl ester

15

**Compound
186**

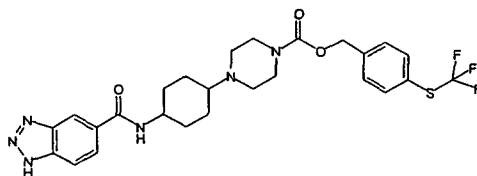
4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3,5-bis-trifluoromethyl-benzyl ester

20

**Compound
187**

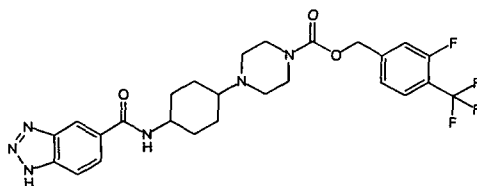
4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3-chloro-5-trifluoromethyl-benzyl ester

25

**Compound
188**

4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 4-trifluoromethylsulfanyl-benzyl ester

30

**Compound
189**

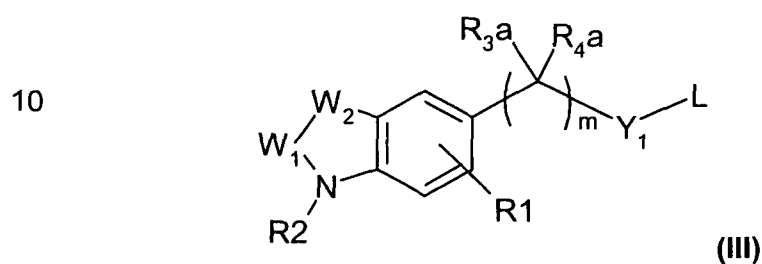
4-{4-[(1H-Benzotriazole-5-carbonyl)-amino]-cyclohexyl}-piperazine-1-carboxylic acid 3-fluoro-4-trifluoromethyl-benzyl ester

35

7. Use of a compound as claimed in any of claims 1 to 6 as autotaxin inhibitor.

5 8. Process for the preparation of a compound according to formulae (Ia), (Ib) or (II) as claimed in any of claims 1 to 6 comprising the steps:

(a) reacting a compound of the formula (III),



wherein L is selected from the group consisting of:

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173

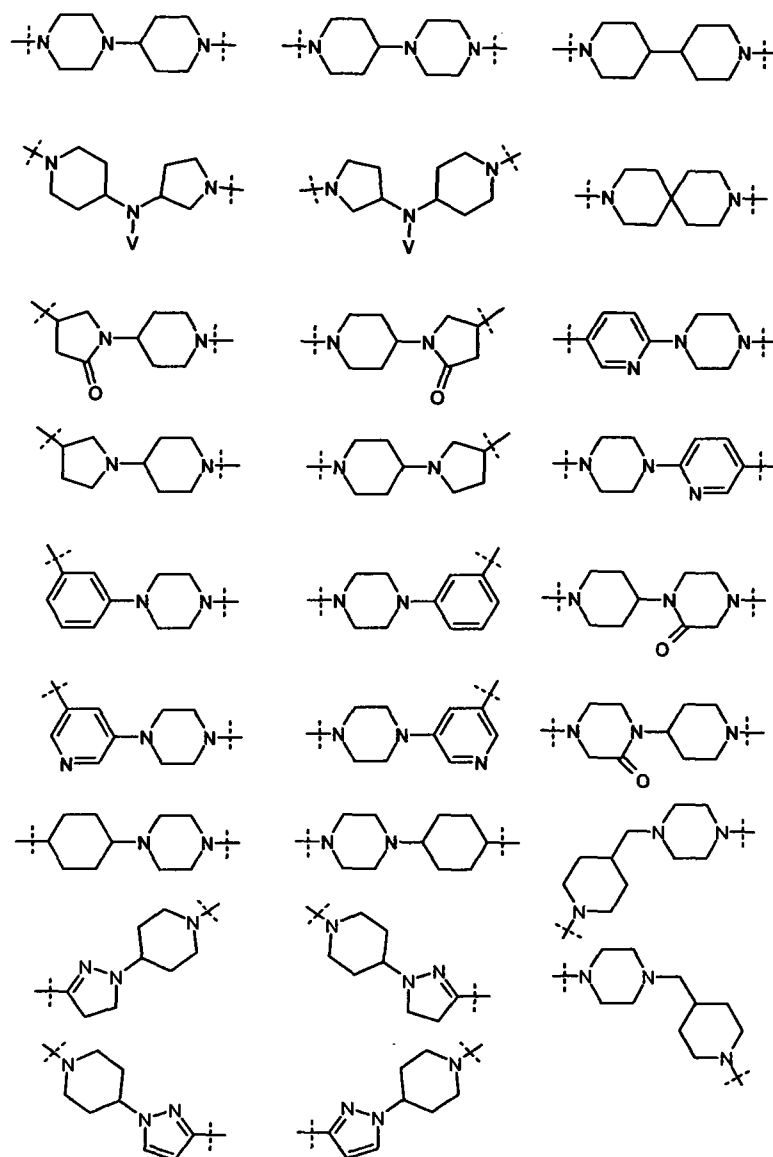
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10

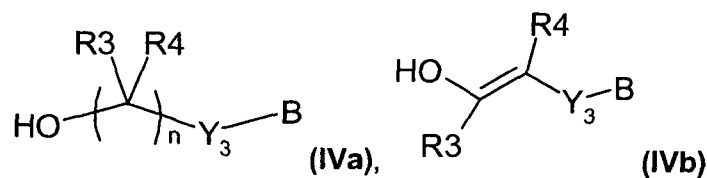
15

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25



and W_1 , W_2 , Y_1 , R_1 , R_2 , R_{3a} , R_{4a} , V , m have the meanings as indicated in any of claims 1 to 6, with a compound of the formula (IVa) or (IVb),

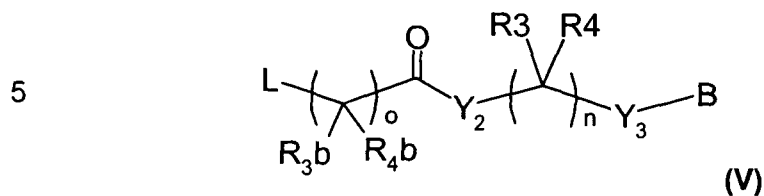


wherein B , Y_3 , R_3 , R_4 , n have the meanings as indicated in any of claims 1 to 6, and a compound selected from the group consisting of: 1,1'-carbonyldiimidazole, phosgene, diphosgene, triphosgene to yield a compound according to formulae (Ia), (Ib) or (II) as indicated in any of claims 1 to 6, in which Z_1 denotes "O" and Y_2 denotes "O";

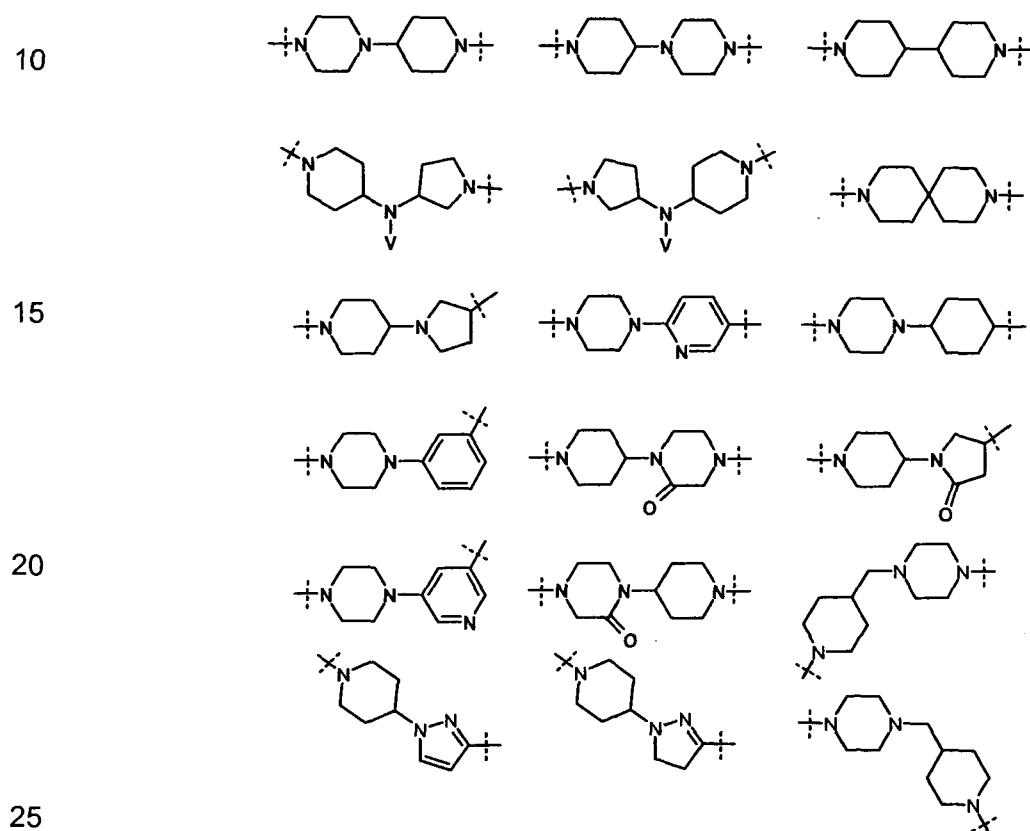
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or

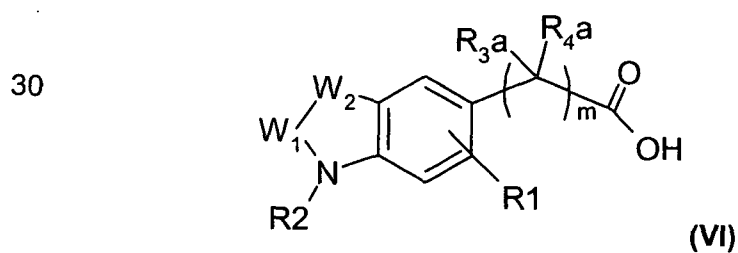
(b) reacting a compound of the formula (V)



wherein L is selected from the group consisting of:



and B, Y₂, Y₃, R₃, R₄, R_{3b}, R_{4b}, V, n, o have the meanings as indicated in any of claims 1 to 6, with a compound of the formula (VI)



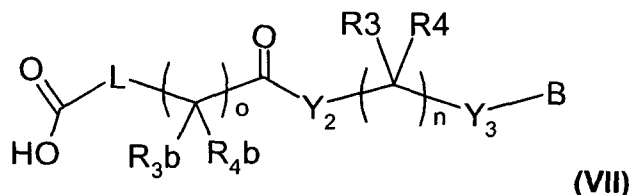
wherein W₁, W₂, R₁, R₂, R_{3a}, R_{4a}, m have the meanings as indicated in any of claims 1 to 6, to yield a compound according to formulae (Ia), (Ib) or

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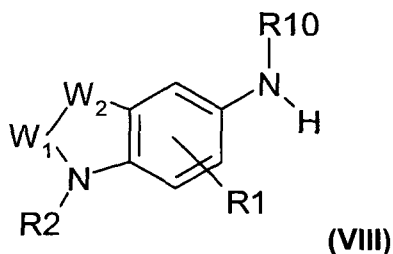
(II) as indicated in any of claims 1 to 6, in which Z_1 denotes "O", and Y_1 denotes " $-C(O)-$ ";

or

(c) reacting a compound of the formula (VII)



wherein L, Y_2 , Y_3 , B, R₃, R₄, R_{3b}, R_{4b}, n, o have the meanings as indicated in any of claims 1 to 6, with a compound of formula (VIII)



wherein W_1 , W_2 , R₁, R₂, R₁₀ have the meanings as indicated in any of claims 1 to 6, to yield a compound according to formulae (Ia), (Ib) or (II) as indicated in any of claims 1 to 6, in which Z_1 denotes "O" and Y_1 denotes " $-N(R_{10})-C(O)-$ ";

or

(d) liberating them from one of their functional derivatives (e.g. with protection groups) by treating them with an acidic, basic, solvolyzing or hydrogenolyzing agent.

and/or converting a base or an acid of a compound according to formulae (Ia), (Ib) or (II) into one of its salts.

9. Medicament comprising at least one compound as claimed in any of claims 1 to 6.

10. Medicament comprising at least one compound as claimed in any of claims 1 to 6 for use in the treatment and/or prophylaxis of physiological and/or pathophysiological conditions, which are caused, mediated and/or propagated by increased lysophosphatic acid levels and/or the activation of autotaxin.
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11. Medicament comprising at least one compound as claimed in any of claims 1 to 6 for use in the treatment and/or prophylaxis of physiological and/or pathophysiological conditions selected from the group consisting of: "cancer, tumour, malignant tumours, benign tumours, solid tumours, sarcomas, carcinomas, hyperproliferative disorders, carcinoids, Ewing sarcomas, Kaposi sarcomas, brain tumours, tumours originating from the brain and/or the nervous system and/or the meninges, gliomas, glioblastomas, neuroblastomas, stomach cancer, kidney cancer, kidney cell carcinomas, prostate cancer, prostate carcinomas, connective tissue tumours, soft tissue sarcomas, pancreas tumours, liver tumours, head tumours, neck tumours, laryngeal cancer, oesophageal cancer, thyroid cancer, osteosarcomas, retinoblastomas, thymoma, testicular cancer, lung cancer, lung adenocarcinoma, small cell lung carcinoma, bronchial carcinomas, breast cancer, mamma carcinomas, intestinal cancer, colorectal tumours, colon carcinomas, rectum carcinomas, gynaecological tumours, ovary tumours/ovarian tumours, uterine cancer, cervical cancer, cervix carcinomas, cancer of body of uterus, corpus carcinomas, endometrial carcinomas, urinary bladder cancer, urogenital tract cancer, bladder cancer, skin cancer, epithelial tumours, squamous epithelial carcinoma, basaliomas, spinaliomas, melanomas, intraocular melanomas, leukaemias, monocyte leukaemia, chronic leukaemias, chronic myelotic leukaemia, chronic lymphatic leukemia, acute leukaemias, acute myelotic leukaemia, acute lymphatic leukemia, lymphomas, angiogenesis, arteriosclerosis, ophthalmic diseases, choroidal neovascularization, diabetic retinopathy, inflammatory diseases, arthritis, neurodegeneration, restenosis, wound healing and/or transplant rejection".
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12. The medicament as claimed in any of claims 9 to 11, wherein in such medicament comprises at least one additional pharmacologically active substance.
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13. The medicament as claimed in any of claims 9 to 11, wherein the medicament is applied *before and/or during and/or after treatment* with at least one additional pharmacologically active substance.

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14. Pharmaceutical composition comprising a therapeutically effective amount of at least one compound as claimed in any of claims 1 to 6.

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15. The pharmaceutical composition as claimed in claim 14 further comprising at least one additional compound selected from the group consisting of physiologically acceptable excipients, auxiliaries, adjuvants, diluents, carriers and/or additional pharmaceutically active substance other than the compound as claimed in any of claims 1 to 5.

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16. Kit comprising a therapeutically effective amount of at least one compound as claimed in any of claims 1 to 6 and/or at least one pharmaceutical composition as claimed in any of claims 14 to 15 and a therapeutically effective amount of at least one further pharmacologically active substance other than the compound as claimed in any of claims 1 to 6.

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