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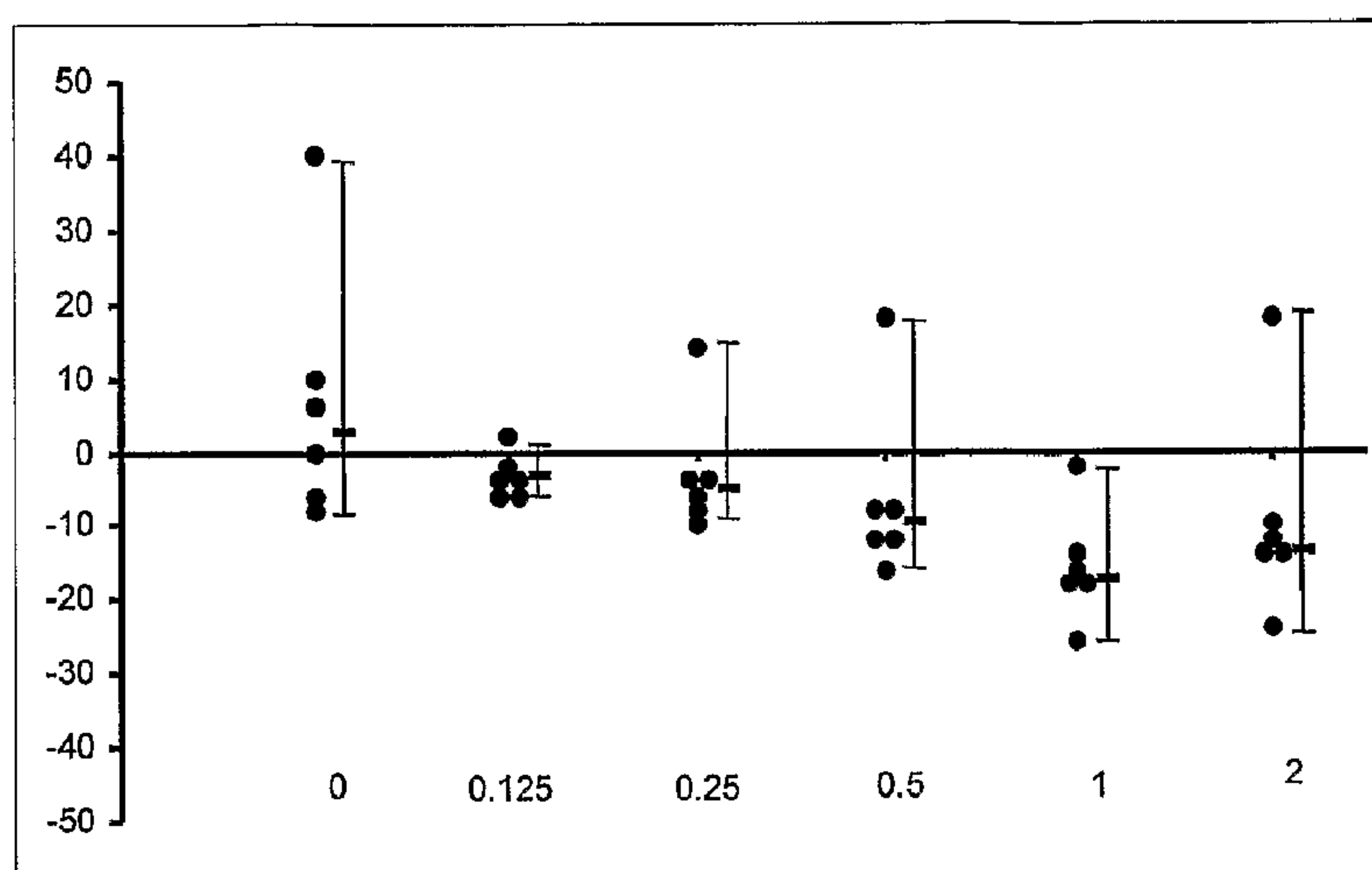
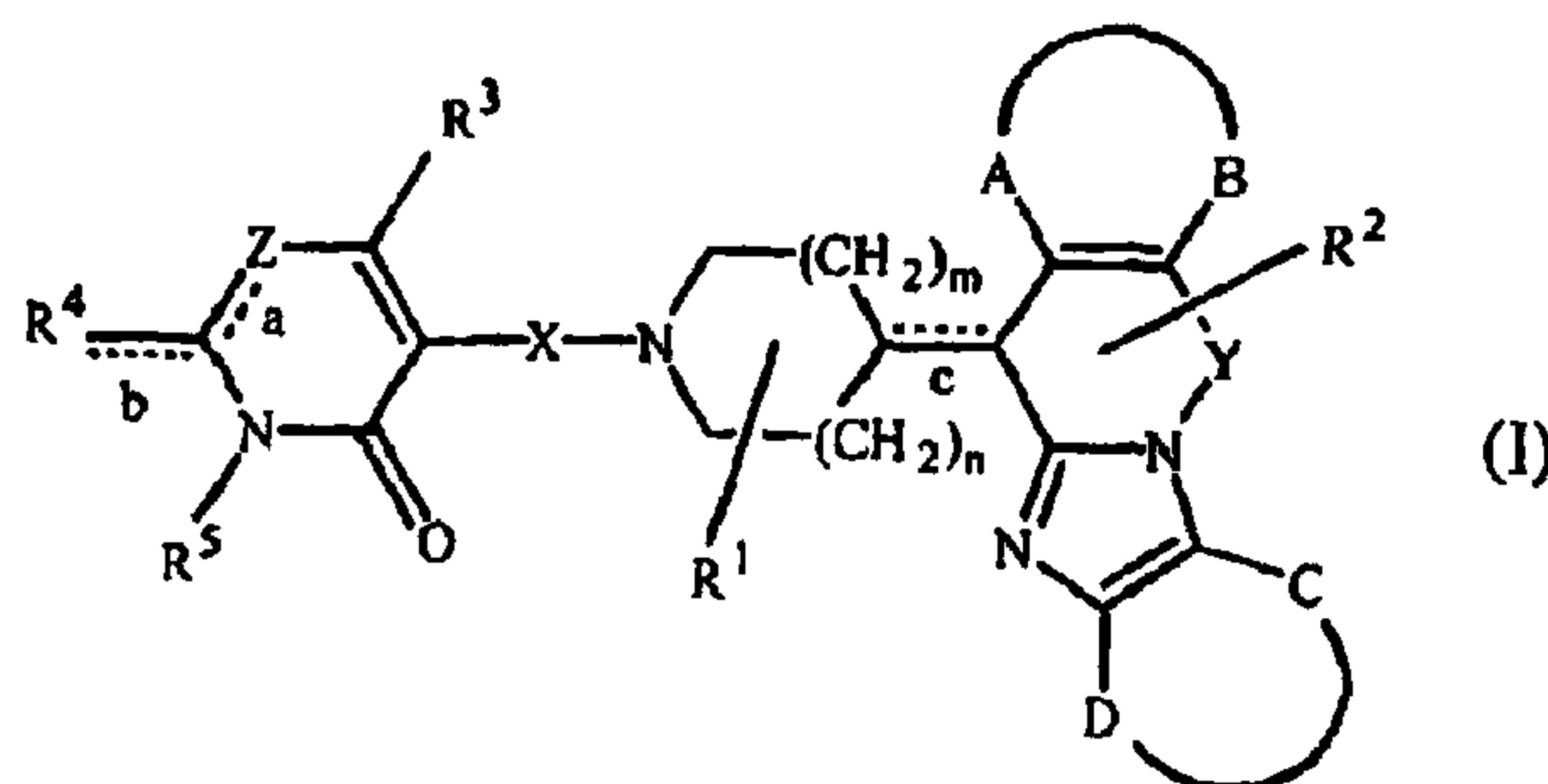
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(54) Titre : UTILISATION D'ANTI-HISTAMINIQUES POUR UNE DIMINUTION AIGUE DE LA PRESSION
INTRACRANIENNE ELEVEE

(54) Title: THE USE OF ANTI-HISTAMINICS FOR ACUTE REDUCTION OF ELEVATED INTRACRANIAL PRESSURE



(57) Abrégé/Abstract:

The invention concerns a novel histamine receptor antagonist and the use of an histamine receptor antagonist for the reduction of intracranial pressure (ICP), in particular for the prevention and treatment of elevated intracranial pressure and/or secondary



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(57) **Abrégé(suite)/Abstract(continued)**:

ischaemia, in particular caused by brain injury, more in particular caused by traumatic (TBI) and non-traumatic brain injury. The novel compounds comprise compounds according to the general Formula (I) the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the N-oxide form thereof. In particular, the preferred compound is 3-[2-[4-(11,12-dihydro-6H-benzimidazo[2,1-b][3]benzazepin-6-yl)-2-(phenylmethyl)-1-piperidinyl]ethyl]-2,10-dimethyl pyrimido[1,2-a]benzimidazol-4(10H)-one, the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the N-oxide form thereof. Also claimed is the novel use of commercially available histamine H1-and H2-receptor antagonists for the reduction of intracranial pressure (ICP).

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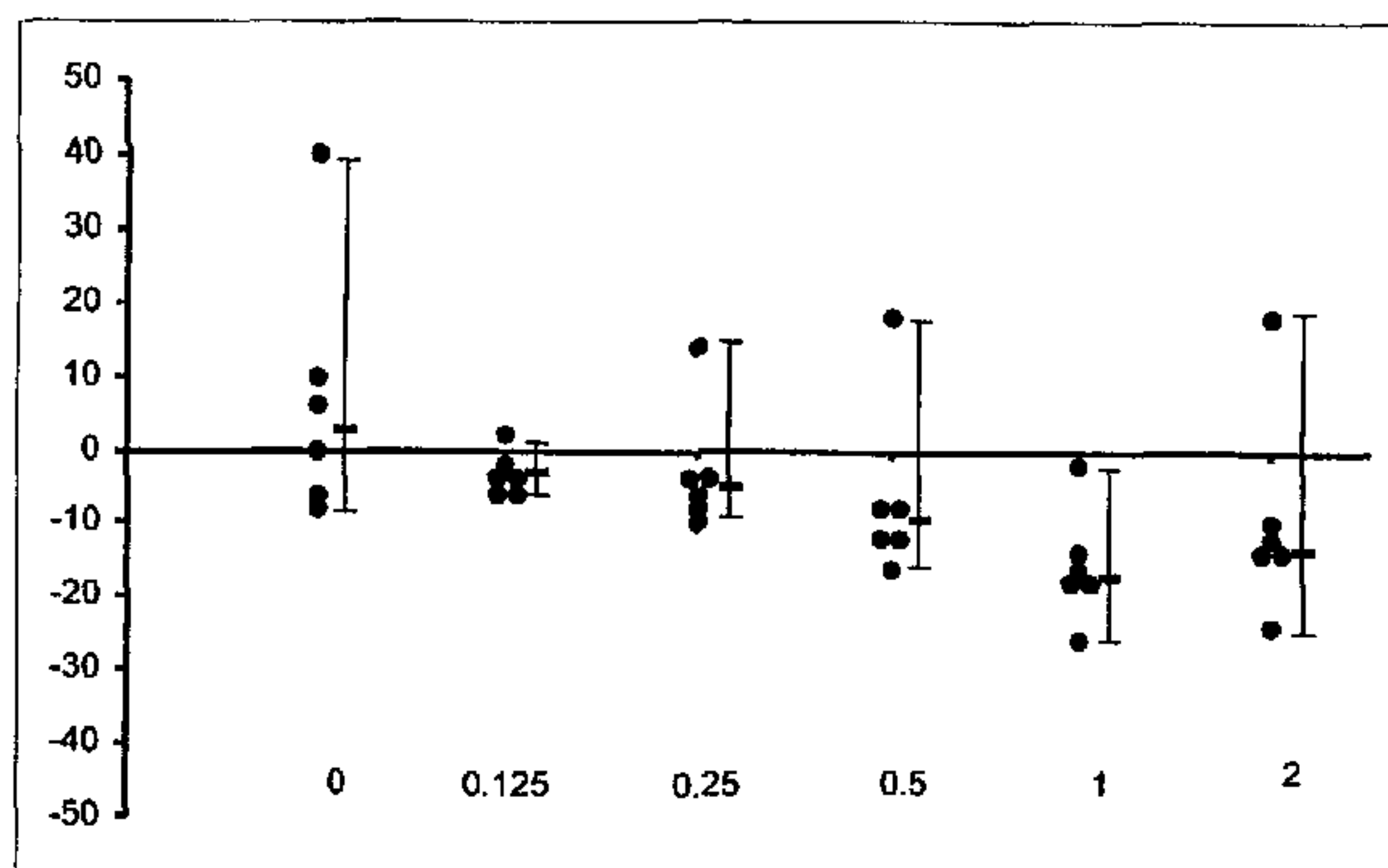
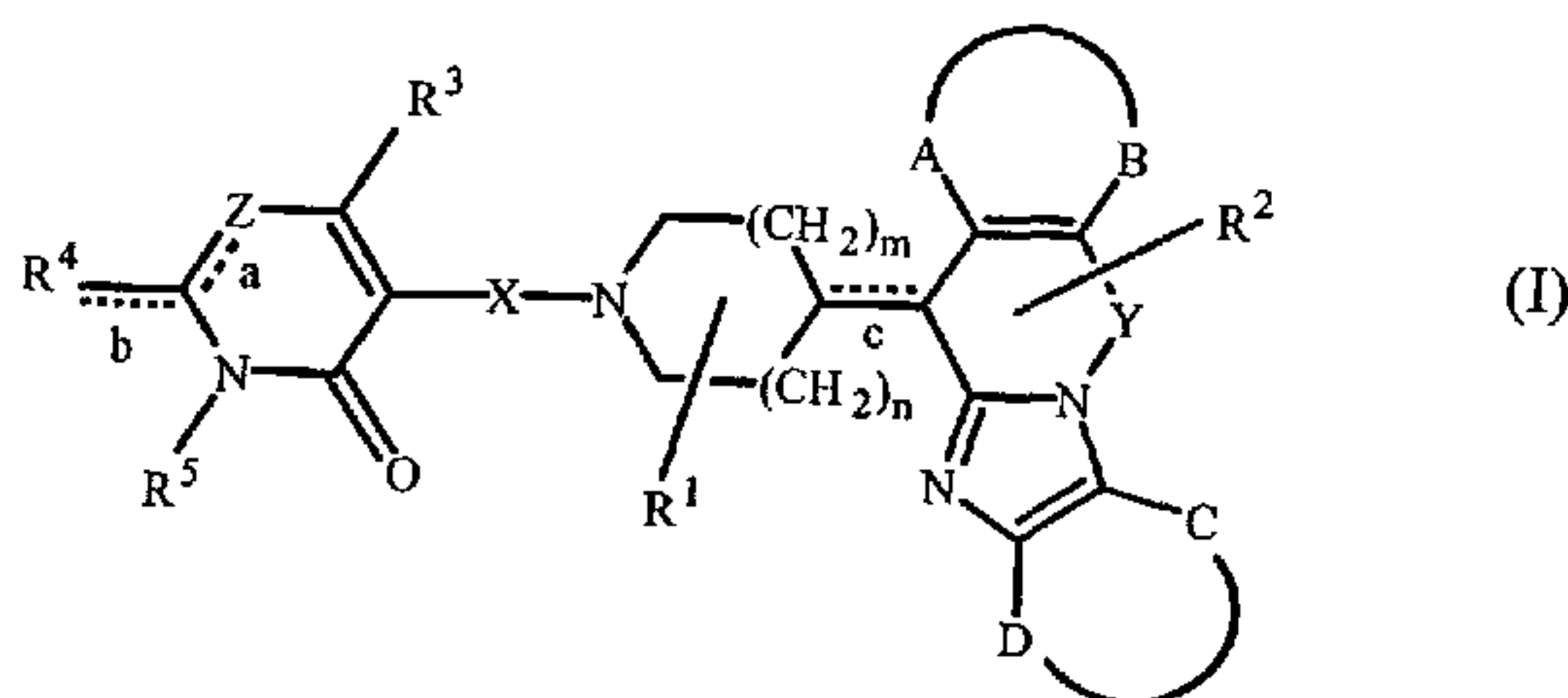
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[Continued on next page]

(54) Title: THE USE OF ANTI-HISTAMINICS FOR ACUTE REDUCTION OF ELEVATED INTRACRANIAL PRESSURE



Dose dependence of the ICP reducing effect of Compound 2 during a 10 min infusion period. X-axis : Dose (mg/kg/min) ; Y-axis : Change in ICP as percentage of initial value.

(57) Abstract: The invention concerns a novel histamine receptor antagonist and the use of an histamine receptor antagonist for the reduction of intracranial pressure (ICP), in particular for the prevention and treatment of elevated intracranial pressure and/or secondary ischaemia, in particular caused by brain injury, more in particular caused by traumatic (TBI) and non-traumatic brain injury. The novel compounds comprise compounds according to the general Formula (I) the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof. In particular, the preferred compound is 3-[2-[4-(11,12-dihydro-6*H*-benzimidazo[2,1-*b*][3]benzazepin-6-yl)-2-(phenylmethyl)-1-piperidinyl]ethyl]-2,10-dimethylpyrimido[1,2-*a*]benzimidazol-4(10*H*)-one, the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof. Also claimed is the novel use of commercially available histamine H1- and H2-receptor antagonists for the reduction of intracranial pressure (ICP).

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THE USE OF ANTI-HISTAMINICS FOR ACUTE REDUCTION OF ELEVATED INTRACRANIAL PRESSURE

5 The invention concerns a novel histamine receptor antagonist and the use of an
histamine receptor antagonist for the reduction of intracranial pressure (ICP), in
particular for the prevention and treatment of elevated intracranial pressure and/or
secondary ischaemia, in particular caused by brain injury, more in particular caused by
traumatic (TBI) and non-traumatic brain injury.

10

TBI is a significant problem in developed countries. In the USA each year about
500,000 head injuries are severe enough to require hospitalisation. Mortality is high
and approximately 80,000 of these TBI-patients face a life-long debilitating loss of
function, 5,000 develop epilepsy and 2,000 live in a persistent vegetative state. TBI is
15 the leading cause of death and disability in young adults today at an estimated cost in
1989 of over \$25 billion per year.

Primary irreversible damage after brain trauma includes hemorrhage, contusion,
neuronal necrosis and diffuse axonal injury. This damage, together with possible
20 cardiovascular and respiratory depression, can induce acute secondary features
including edema (vasogenic and/or cellular), secondary bleeding, alterations of cerebral
blood volume (CBV), disturbed autoregulation of cerebral blood flow (CBF) and
ischaemia. Edema, bleeding and an increase of CBV will increase the total brain
volume and consequently the intracranial pressure (ICP). This in turn can lead to
25 further progression of ischaemia, infarction, and, in severe cases, herniation of the brain
stem with possible acute respiratory depression and death. Therapy in TBI should
therefore be directed to the interruption of the pathologic cascade and the reduction of
the brain volume and ICP. Prevention of a life threatening secondary increase in ICP,
which often occurs e.g. in the post-acute phase after trauma or after cardiac
30 resuscitation, is also a target for pharmacological treatment.

At present, the clinical tools for ICP reduction are limited. Standard treatment
schedules include surgical drainage of the ventricles, blood pressure management,

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mannitol infusion, hyperventilation and high dose barbiturate therapy. Side effects of the non-surgical treatments include brain ischaemia, rebound effects on ICP and an increased risk for bacterial infections and sepsis. Also, various compounds with different mechanisms of actions (e.g. bradykinin antagonism, calcium antagonism, oxidative stress inhibition, glutamate receptor blockade and anti-epilepsy) have been tested in phase II and III clinical trials or are still under investigation (focus on outcome, not on ICP). Up to date no compound has been approved for the acute treatment of intracranial pressure (K.K. Jain, *Chapter 4 : Neuroprotection in Acute Trauma*, 'Neuroprotection in CNS Disorders: Commercial Opportunities'. A Jain PharmaBiotech Report: 65-73, 2000). Obviously, there is a need for pharmaceuticals and/or therapies for the treatment of elevated intracranial pressure (ICP) and/or secondary ischaemia, in particular caused by brain injury, more in particular caused by traumatic brain injury (TBI).

The inventors have now found that substituted tetracyclic imidazole derivatives according to the general Formula (I) show histamine H1- and/or H2-receptor antagonist activity. Furthermore, the compounds have been shown to be particular useful for the reduction of intracranial pressure (ICP), in particular for the prevention and treatment of elevated intracranial pressure and/or secondary ischaemia, in particular caused by brain injury, more in particular caused by traumatic (TBI) and non-traumatic brain injury.

Furthermore, the inventors have found that compounds that antagonize the histamine H1- and/or H2-receptors (commonly called anti-histaminics) are also usefull for the reduction of intracranial pressure (ICP), in particular for the prevention and treatment of elevated intracranial pressure and/or secondary ischaemia, in particular caused by brain injury, more in particular caused by traumatic (TBI) and non-traumatic brain injury.

Characteristical to all compounds is that they are able to acutely reduce the intracranial pressure when administered to the bloodstream of a mamal, in particular by intravenous administration. Advantageously and very important, said compounds

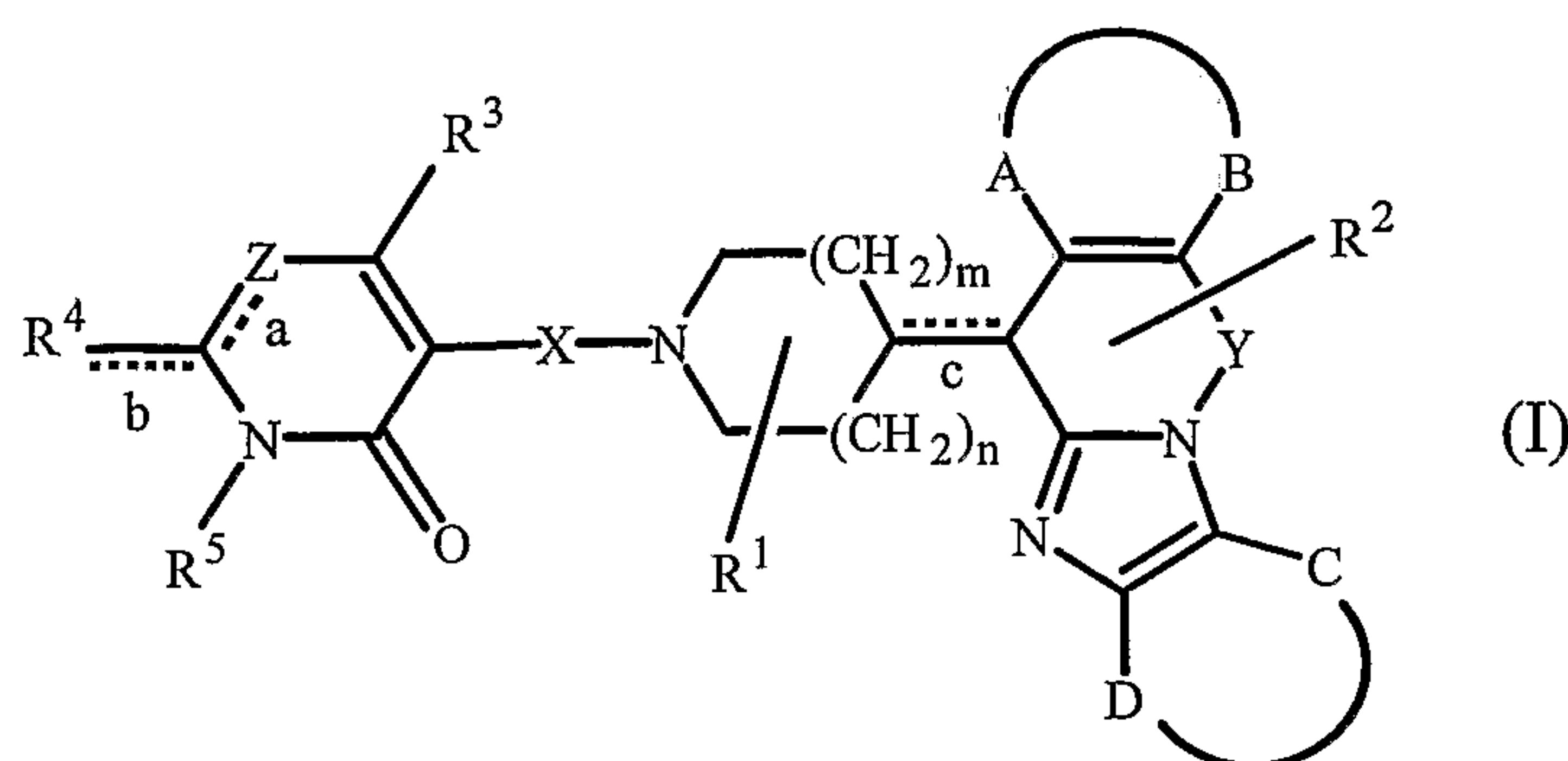
reduce the ICP while having little or no effect on the blood pressure, in particular a blood pressure-lowering effect, which is a most desired property of a potential drug.

Hitherto, histamine H1- and/or H2-receptor antagonists have not been developed for
5 lowering the ICP, in particular for post-traumatic lowering of the ICP. Mohanty et al. in Journal of the Neurological Sciences, 1989, 90:87-97 observed that histamine played a role in the forming of traumatically induced brain edema. Increased brain water content and elevated plasma and brain histamine levels were *prevented* by prior treatment with the histamine H2-receptor antagonist cimetidine. However,
10 meypyramine (an histamine H1-receptor antagonist) failed to reduce the increased brain water content and the histamine levels in the plasma and brain remained high. The effect on the ICP, in particular the action of histamine antagonists *after* a rise of the ICP for acutely reducing an increased ICP was not researched nor was the effect on the blood pressure.

15

Without being restricted theretoo, it is the opinion of the inventors that, in view of the fact that the histamine receptor antagonists show the ability to reduce a *normal* ICP in the absence of brain edema and in view of the fact that the histamine receptor antagonists do not or only marginally influence the blood pressure, an effect attributed
20 to peripheral vasodilatation, the mechanism of action is not one that acts purely on the reduction of brain edema nor via vasodilation, effects which are known in the prior art to occur for anti-histaminics.

Hence, the purpose of the present invention is to provide a substituted tetracyclic
25 imidazole derivatives for use as an histamine antagonist, in particular as an histamine H1-antagonist, more in particular as an antagonist showing both histamine H1- and H2-antagonist activity, according to the general Formula (I)



the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof, wherein :

m is 1 or 2 ;

5 *n* is 0, 1 or 2 ;

a, *b*, *c* independently are a single or a double bond ;

X is a covalent bond or a bivalent C₁₋₆alkanediyl radical wherein one or more -CH₂- groups may be optionally replaced with -O-, -S-, -CO-, or -NR⁷- ; wherein :

10 R⁷ is hydrogen, alkyl, Ar, Ar-alkyl, Het, Het-alkyl, hydroxyalkyl, alkyloxy, alkyloxyalkyl, alkyloxyalkyloxyalkyl, aminoalkyl, mono- or dialkylaminoalkyl, formyl, alkylcarbonylaminoalkyl, alkylcarbonyloxyalkyl, alkyloxycarbonyl, alkyloxycarbonylalkyl, alkylaminocarbonyl, alkylaminocarbonylalkyl, hydroxyalkyloxyalkyl, aminocarbonyl, aminocarbonylalkyl, alkyloxycarbonyl or alkylcarbonyloxyalkyloxyalkyl ;

Y is a bivalent C₁₋₄alkanediyl or C₂₋₄ alkenediyl radical ;

Z is N, in which case *a* is a double bond and *b* is a single bond or N-R⁷ in which case *a* is a single bond, *b* is a double bond and R⁷ is defined as above ;

20 R¹, R² independently are hydrogen, hydroxy, alkyl, alkyloxy, Ar, Ar-alkyl, di(Ar-)alkyl, Het or Het-alkyl ;

-A-B- independently is a bivalent radical of formula

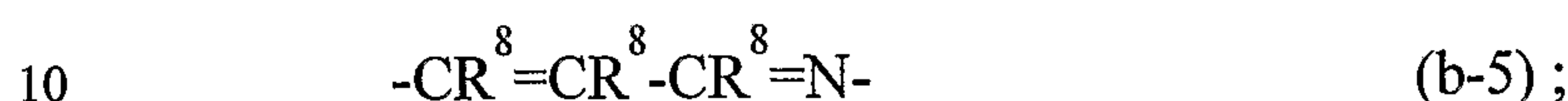


25 wherein

R^8 each independently is hydrogen, halo, hydroxy, alkyl or alkyloxy ;

E is a bivalent radical of formula -O-, -S- or -NR⁷- wherein R⁷ is defined as above ;

5 -C-D- independently is a bivalent radical of formula



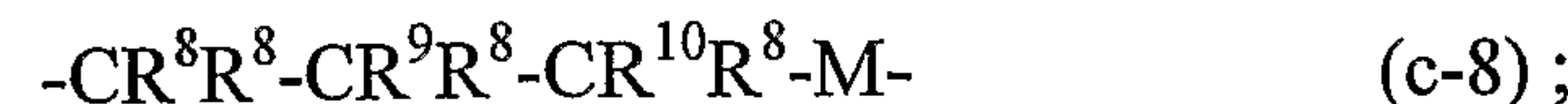
wherein R⁸ is defined as above ;

R³ is hydrogen, halo, hydroxy, alkyl, alkyloxy, Ar, Ar-alkyl, di(Ar-)alkyl, Het or Het-alkyl ;

R⁴ is hydrogen, alkyl, amino, alkylamino, Ar-amino, Het-amino,
15 alkylcarbonylamino, Ar-carbonylamino, Het-carbonylamino,
alkylaminocarbonylamino, Ar-aminocarbonylamino, Het-
aminocarbonylamino, alkyloxyalkylamino, Ar-oxyalkylamino or Het-
oxyalkylamino ;

R⁵ is hydrogen or alkyl ;

20 or R⁴ and R⁵ together may form a bivalent radical of Formula



wherein

R^7 and R^8 are defined as above ;

R^9 , R^{10} independently are hydrogen, alkyl, halo, haloalkyl ;

or R^9 and R^{10} together may form a bivalent radical of formula

5 $-CR^8=CR^8-CR^8=CR^8-$; and

M is a bivalent radical of formula $-CH_2-$, $-O-$, $-S-$ or $-NR^7-$ wherein R^7 is defined as above.

In the framework of this application, Ar is a homocycle selected from the group of
 10 naphthyl and phenyl, each optionally substituted with 1, 2 or 3 substituents, each
 substituent independently selected from the group of hydroxy, halo, cyano, nitro,
 amino, mono- or dialkylamino, alkyl, haloalkyl, alkyloxy, haloalkyloxy, carboxyl,
 alkyloxycarbonyl, aminocarbonyl and mono- or dialkylaminocarbonyl. Preferably, Ar
 is a naphthyl or phenyl, each optionally substituted with 1 substituent, each substituent
 15 independently selected from the group of halo or alkyl.

In the framework of this application, Het is a monocyclic heterocycle selected from the
 group of pyrrolyl, pyrazolyl, imidazolyl, furanyl, thienyl, oxazolyl, isoxazolyl,
 thiazolyl, isothiazolyl, pyridinyl, pyrimidinyl, pyrazinyl and pyridazinyl; or a bicyclic
 20 heterocycle selected from the group of quinolinyl, quinoxalinyl, indolyl,
 benzimidazolyl, benzoxazolyl, benzisoxazolyl, benzothiazolyl, benzisothiazolyl,
 benzofuranyl and benzothienyl ; each monocyclic and bicyclic heterocycle may
 optionally be substituted on a carbon atom with halo, hydroxy, alkyl or alkyloxy.
 Preferably, Het is pyridinyl, pyrazinyl or indolyl.

25 In the framework of this application, alkyl is a straight or branched saturated
 hydrocarbon radical having from 1 to 6 carbon atoms ; or is a cyclic saturated
 hydrocarbon radical having from 3 to 6 carbon atoms ; or is a a cyclic saturated
 hydrocarbon radical having from 3 to 6 carbon atoms attached to a straight or branched
 30 saturated hydrocarbon radical having from 1 to 6 carbon atoms ; wherein each carbon
 atom can be optionally substituted with halo, hydroxy, alkyloxy or oxo. Preferably,
 alkyl is methyl, ethyl or cyclohexylmethyl.

In the framework of this application, halo is a substituent selected from the group of fluoro, chloro, bromo and iodo and haloalkyl is a straight or branched saturated hydrocarbon radical having from 1 to 6 carbon atoms or a cyclic saturated hydrocarbon radical having from 3 to 6 carbon atoms, wherein one or more carbon atoms are substituted with one or more halo-atoms. Preferably, halo is fluoro or chloro and preferably, haloalkyl is trifluoromethyl.

A preferred group of compounds are those compounds according to Formula (I), the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof, in which -A-B- is a bivalent radical of formula (a-1) or (a-3), wherein E is a bivalent radical of formula -O-, -S- or -NR⁷- wherein R⁷ is hydrogen, R⁸ is hydrogen, -C-D- is a bivalent radical of formula (b-1) or (b-2), wherein R⁸ is hydrogen and Y is a bivalent radical of formula -CH₂-, -CH₂-CH₂- or -CH=CH-.

Another group of preferred compounds of Formula (I) are those compounds according to Formula (I), the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof, in which m and n are both 1.

Another group of preferred compounds of Formula (I) are those compounds according to Formula (I), the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof, in which R¹ and R², each independently are hydrogen, alkyl, Ar-alkyl, Het or Het-alkyl.

Yet another group of preferred compounds of Formula (I) are those compounds according to Formula (I), the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof, in which X is a bivalent radical of formula -CH₂CH₂- or -CH₂CH₂CH₂-.

Yet another group of preferred compounds of Formula (I) are those compounds according to Formula (I), the pharmaceutically acceptable acid or base addition salts

thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof, in which R^3 is hydrogen or alkyl, Z is $N-R^7$ wherein R^7 is hydrogen or alkyl, a is a single bond and b is a double bond, and R^4 and R^5 together form a bivalent radical of Formula (c-1), (c-3), (c-5), (c-7), (c-8) or (c-10) wherein R^7 and R^8 are hydrogen.

5

Yet another group of preferred compounds of Formula (I) are those compounds according to Formula (I), the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof, in which R^3 is hydrogen or alkyl, Z is $N-R^7$ wherein R^7 is hydrogen or alkyl, a is a single bond and b is a double bond, R^4 and R^5 together form a bivalent radical of Formula (c-1), (c-3), (c-5), (c-7), (c-8) or (c-10) wherein R^7 and R^8 are hydrogen and R^9 and R^{10} together form a bivalent radical of formula $-CR^8=CR^8-CR^8=CR^8-$ wherein R^8 is hydrogen.

15 More specifically, the compound 3-[2-[4-(11,12-dihydro-6*H*-benzimidazo[2,1-*b*][3]benzazepin-6-yl)-2-(phenylmethyl)-1-piperidinyl]ethyl]-2,10-dimethyl pyrimido[1,2-*a*]benzimidazol-4(10*H*)-one, the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof, are most preferred.

20

The pharmaceutically acceptable acid addition salts are defined to comprise the therapeutically active non-toxic acid addition salt forms which the compounds according to Formula (I) are able to form. Said acid addition salts can be obtained by treating the base form of the compounds according to Formula (I) with appropriate acids, for example inorganic acids, for example hydrohalic acid, in particular hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid and phosphoric acid ; organic acids, for example acetic acid, hydroxyacetic acid, propanoic acid, lactic acid, pyruvic acid, oxalic acid, malonic acid, succinic acid, maleic acid, fumaric acid, malic acid, tartaric acid, citric acid, methanesulfonic acid, ethanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, cyclamic acid, salicylic acid, p-aminosalicylic acid and pamoic acid.

30

The compounds according to Formula (I) containing acidic protons may also be converted into their therapeutically active non-toxic base addition salt forms by treatment with appropriate organic and inorganic bases. Appropriate base salts forms comprise, for example, the ammonium salts, the alkaline and earth alkaline metal salts, in particular lithium, sodium, potassium, magnesium and calcium salts, salts with organic bases, e.g. the benzathine, *N*-methyl-D-glucamine, hybramine salts, and salts with amino acids, for example arginine and lysine.

Conversely, said acid or base addition salt forms can be converted into the free forms by treatment with an appropriate base or acid.

The term addition salt as used in the framework of this application also comprises the solvates which the compounds according to Formula (I) as well as the salts thereof, are able to form. Such solvates are, for example, hydrates and alcoholates.

Among the acid addition salts, the compound 3-[2-[4-(11,12-dihydro-6*H*-benzimidazo[2,1-*b*][3]benzazepin-6-yl)-2-(phenylmethyl)-1-piperidinyl]ethyl]-2,10-dimethyl pyrimido[1,2-*a*]benzimidazol-4(10*H*)-one (E)-2-butenedioate (2:3) hydrate (1:1) including all stereoisomeric forms thereof is the most preferred compound.

Particular preferred compounds are the (A)[(2 α , 4 β)(A)] enantiomer, the (B)[(2 α , 4 β)(A)] enantiomer and a mixture thereof, of the compounds 3-[2-[4-(11,12-dihydro-6*H*-benzimidazo[2,1-*b*][3]benzazepin-6-yl)-2-(phenylmethyl)-1-piperidinyl]ethyl]-2,10-dimethyl pyrimido[1,2-*a*]benzimidazol-4(10*H*)-one and 3-[2-[4-(11,12-dihydro-6*H*-benzimidazo[2,1-*b*][3]benzazepin-6-yl)-2-(phenylmethyl)-1-piperidinyl]ethyl]-2,10-dimethyl pyrimido[1,2-*a*]benzimidazol-4(10*H*)-one (E)-2-butenedioate (2:3) hydrate (1:1).

The *N*-oxide forms of the compounds according to Formula (I) are meant to comprise those compounds of Formula (I) wherein one or several nitrogen atoms are oxidized to the so-called *N*-oxide, particularly those *N*-oxides wherein one or more nitrogens of the piperidinyl radical in Formula (1) are *N*-oxidized.

The term "stereochemically isomeric forms" as used herein defines all possible isomeric forms which the compounds of Formula (I) may possess. Unless otherwise mentioned or indicated, the chemical designation of compounds denotes the mixture of all possible stereochemically isomeric forms, said mixtures containing all
5 diastereomers and enantiomers of the basic molecular structure. More in particular, stereogenic centers may have the R- or S-configuration; substituents on bivalent cyclic (partially) saturated radicals may have either the cis- or trans-configuration. Compounds encompassing double bonds can have an E or Z-stereochemistry at said double bond. Stereochemically isomeric forms of the compounds of Formula (I) are
10 obviously intended to be embraced within the scope of this invention.

Following CAS nomenclature conventions, when two stereogenic centers of known absolute configuration are present in a molecule, an *R* or *S* descriptor is assigned (based on Cahn-Ingold-Prelog sequence rule) to the lowest-numbered chiral center, the
15 reference center. The configuration of the second stereogenic center is indicated using relative descriptors [*R**,*R**] or [*R**,*S**], where *R** is always specified as the reference center and [*R**,*R**] indicates centers with the same chirality and [*R**,*S**] indicates centers of unlike chirality. For example, if the lowest-numbered chiral center in the molecule has an *S* configuration and the second center is *R*, the stereo descriptor would
20 be specified as *S*-[*R**,*S**]. If "α" and "β" are used : the position of the highest priority substituent on the asymmetric carbon atom in the ring system having the lowest ring number, is arbitrarily always in the "α" position of the mean plane determined by the ring system. The position of the highest priority substituent on the other asymmetric carbon atom in the ring system relative to the position of the highest priority substituent
25 on the reference atom is denominated "α", if it is on the same side of the mean plane determined by the ring system, or "β", if it is on the other side of the mean plane determined by the ring system.

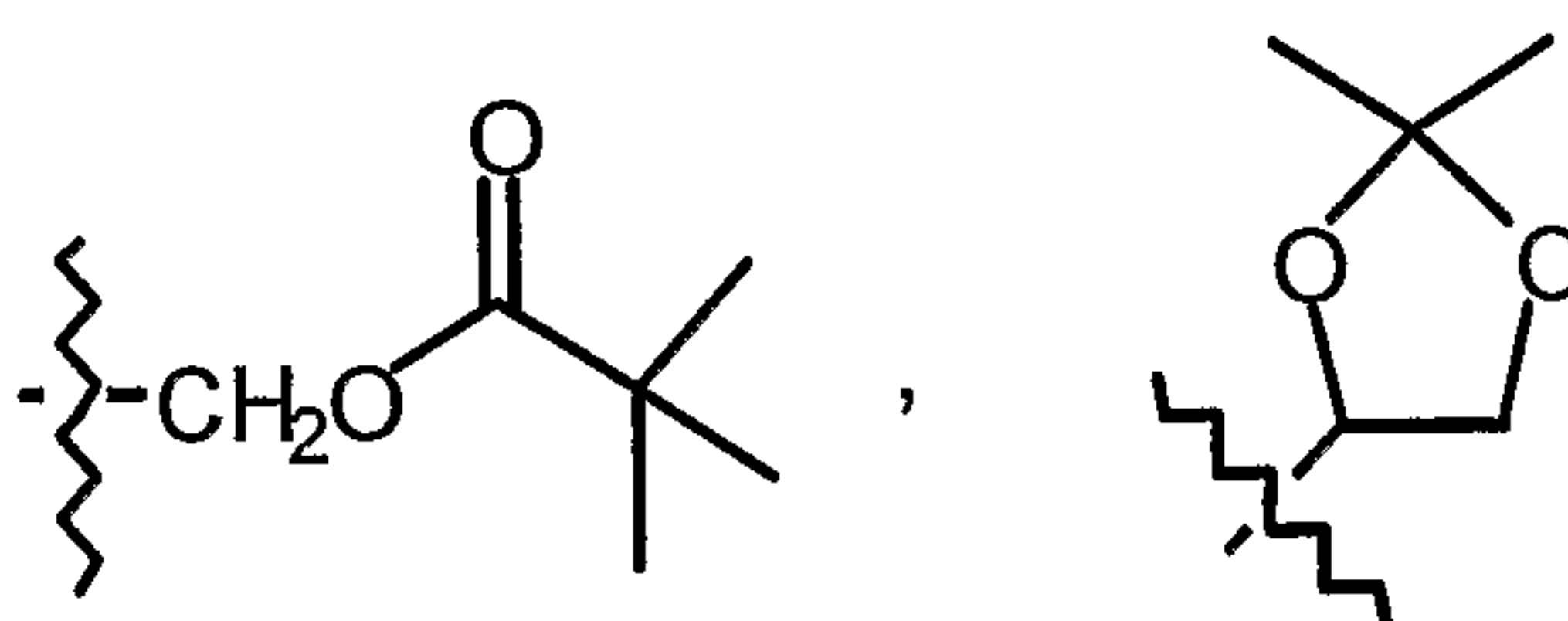
When the bond at c is a single bond, compounds of Formula (I) and some of the
30 intermediate compounds have at least two stereogenic centers in their structure. When *R*¹ is other than hydrogen, the monocyclic N-ring in Formula (I) has a further stereogenic center. This may lead to 8 stereochemically different structures.

The compounds of Formula (I) as prepared in the processes described below may be synthesized in the form of racemic mixtures of enantiomers which can be separated from one another following art-known resolution procedures. The racemic compounds
5 of Formula (I) may be converted into the corresponding diastereomeric salt forms by reaction with a suitable chiral acid. Said diastereomeric salt forms are subsequently separated, for example, by selective or fractional crystallization and the enantiomers are liberated therefrom by alkali. An alternative manner of separating the enantiomeric forms of the compounds of Formula (I) involves liquid chromatography using a chiral
10 stationary phase. Said pure stereochemically isomeric forms may also be derived from the corresponding pure stereochemically isomeric forms of the appropriate starting materials, provided that the reaction occurs stereospecifically. Preferably if a specific stereoisomer is desired, said compound will be synthesized by stereospecific methods of preparation. These methods will advantageously employ enantiomerically pure
15 starting materials.

Some of the compounds of Formula (I) may also exist in their tautomeric form. Such forms although not explicitly indicated in the above formula are intended to be included within the scope of the present invention. For instance, compounds of Formula (I)
20 wherein R^5 is H may exist in their corresponding tautomeric form.

The invention also comprises derivative compounds (usually called "pro-drugs") of the pharmacologically-active compounds according to the invention, which are degraded *in vivo* to yield the compounds according to the invention. Pro-drugs are usually (but not
25 always) of lower potency at the target receptor than the compounds to which they are degraded. Pro-drugs are particularly useful when the desired compound has chemical or physical properties that make its administration difficult or inefficient. For example, the desired compound may be only poorly soluble, it may be poorly transported across the mucosal epithelium, or it may have an undesirably short plasma half-life. Further
30 discussion on pro-drugs may be found in Stella, V. J. *et al.*, "Prodrugs", *Drug Delivery Systems*, 1985, pp. 112-176, and *Drugs*, 1985, **29**, pp. 455-473.

Pro-drugs forms of the pharmacologically-active compounds according to the invention will generally be compounds according to Formula (I), the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof, having an acid group which is esterified or amidated. Included
 5 in such esterified acid groups are groups of the formula $-\text{COOR}^x$, where R^x is a C_{1-6} alkyl, phenyl, benzyl or one of the following groups :



Amidated groups include groups of the formula $-\text{CONR}^y\text{R}^z$, wherein R^y is H,
 10 C_{1-6} alkyl, phenyl or benzyl and R^z is $-\text{OH}$, H, C_{1-6} alkyl, phenyl or benzyl.

Compounds according to the invention having an amino group may be derivatised with a ketone or an aldehyde such as formaldehyde to form a Mannich base. This base will hydrolyze with first order kinetics in aqueous solution.

15

The compounds according to Formula (I) can generally be prepared by a succession of steps, each of which is known to the skilled person. The preparation of said compounds is disclosed in a co-pending application, which is included herein by reference.

20

Apart from their use for reducing the ICP, the compounds according to Formula (I) and derivatives thereof are also useful for the treatment of other histamine H1- and H2-mediated diseases, in particular for immunomodulation in a mammal, for the suppression of hypersensitivity and/or inflammatory reactions, for the treatment and prevention of
 25 allergic diseases such as rhinitis, urticaria, asthma, anaphylaxis and the like and for the treatment of gastrointestinal conditions such as ulcers, dyspepsia, various reflux indications and the like. The invention is therefore also concerned with the use of an histamine receptor antagonist according to Formula (I) and derivatives thereof for the manufacture of a medicament for immunomodulation in a mammal, for the suppression

of hypersensitivity and/or inflammatory reactions and for the treatment and prevention of allergic diseases and gastrointestinal conditions.

5 A further aspect of the invention is to provide for a new use for histamine H1- and/or H2-receptor antagonists, in particular for acutely lowering the intracranial pressure (ICP), in particular an elevated ICP, more in particular a critically elevated ICP and/or preventing an elevated ICP and secondary ischaemia caused by brain injury. Most advantageously, the histamine H1- and/or H2-receptor antagonists do not or to a minimum extend lower or raise the blood pressure.

10

According to the invention, the histamine H1- and/or H2-receptor antagonists are either compounds and derivatives thereof according to Formula (I) or known histamine H1- and/or H2-receptor antagonists, being a discrete and limited group of medications readily recognized in the art.

15

Hitherto, histamine H1-receptor antagonists are commonly used for immunomodulation in a mammal and for the suppression of hypersensitivity and/or inflammatory reactions. In particular, the histamine H1-receptor antagonist is selected from the group of acrivastine, alimemazine, antazoline, astemizole, azatadine, azelastine, brompheniramine, buclizine, carbinoxamine, carebastine, cetirizine, chlorcyclizine, chlorpheniramine, cinnarizine, clemastine, clemizole, clocinizine, clonidine, cyclizine, cyproheptadine, descarboethoxyloratidine, dexchlorpheniramine, dimenhydrinate, dimethindene, dimethothiazine, diphenhydramine, diphenylpyraline, doxylamine, ebastine, efletirizine, epinastine, fexofenadine, hydroxyzine, ketotifen, levocabastine, loratidine, meclizine, mequitazine, methdilazine, mianserin, mizolastine, niaprazine, noberastine, norastemizole, oxatomide, oxomemazine, phenbenzamine, pheniramine, picumast, promethazine, pyrillamine, temelastine, terfenadine, trimeprazine, tripeleminamine and triprolidine, derivatives thereof and mixtures of any two or more of the foregoing.

30

Hitherto, histamine H2-receptor antagonists are commonly used for mammals suffering from certain gastrointestinal conditions such as ulcers, dyspepsia, various reflux indications and the like. In particular, the histamine H2-receptor antagonist is selected

from the group of ranitidine, cimetidine, famotidine, nizatidine, tiotidine, zolantidine, derivatives thereof and mixtures of any two or more of the foregoing.

Also, histamine receptor antagonists may exhibit both histaminic H1- and/or H2-
5 receptor antagonist activity, such as ritanserine or the compounds according to Formula (I), the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof.

While all compounds show a marked depression of the ICP, the following compounds
10 have been shown also to have no or little effect on the lowering of the blood pressure : ketotifen, chlorcyclizine, promethazine, pyrilamine, diphenylhydramine, chlorpheniramine and zolantadine.

In vitro studies can be used to evaluate the histamine antagonist activity of the present
15 compounds using appropriate receptor modelling studies.

In vivo studies can be used to evaluate the biological activity of the present compounds. To this extent, a clinically relevant rat model for traumatic brain injury (Closed Head Injury-model) was developed and used to test the compounds according to the
20 invention (K. Engelborghs et al., *Temporal changes in intracranial pressure in a modified experimental model of closed head injury*, J. Neurosurg. 89: 796-806, 1998 ; K. van Rossem et al., *Brain oxygenation after experimental closed head injury*, Adv. Exp. Med. Biol. 471: 209-215, 1999 ; K. Engelborghs et al., *Impaired autoregulation of cerebral blood flow in an experimental model of traumatic brain injury*, J. Neurotrauma, 17(8): 667-677, 2000). In one study intracranial hypertension was
25 induced by a cortical cold lesion in rabbits.

The histamine receptor antagonist according to the invention, including the compounds according to Formula (I) and the currently known histamine H1-, H2- and H1/H2-
30 receptor antagonists may be formulated into various pharmaceutical forms for administration purposes. As appropriate compositions there may be cited all compositions usually employed for systemically administering drugs. To prepare the pharmaceutical compositions of this invention, an effective amount of the particular

compound, optionally in addition salt form, as the active ingredient is combined in intimate admixture with a pharmaceutically acceptable carrier, which carrier may take a wide variety of forms depending on the form of preparation desired for administration. These pharmaceutical compositions are desirable in unitary dosage form suitable, in particular, for administration orally or by parenteral injection. For example, in preparing the compositions in oral dosage form, any of the usual pharmaceutical media may be employed such as, for example, water, glycols, oils, alcohols and the like in the case of oral liquid preparations such as suspensions, syrups, elixirs, emulsions and solutions; or solid carriers such as starches, sugars, kaolin, diluents, lubricants, binders, disintegrating agents and the like in the case of powders, pills, capsules and tablets. Because of their ease in administration, tablets and capsules represent the most advantageous oral dosage unit forms in which case solid pharmaceutical carriers are obviously employed. For parenteral compositions, the carrier will usually comprise sterile water, at least in large part, though other ingredients, for example, to aid solubility, may be included. Injectable solutions, for example, may be prepared in which the carrier comprises saline solution, glucose solution or a mixture of saline and glucose solution. Injectable suspensions may also be prepared in which case appropriate liquid carriers, suspending agents and the like may be employed. Also included are solid form preparations which are intended to be converted, shortly before use, to liquid form preparations.

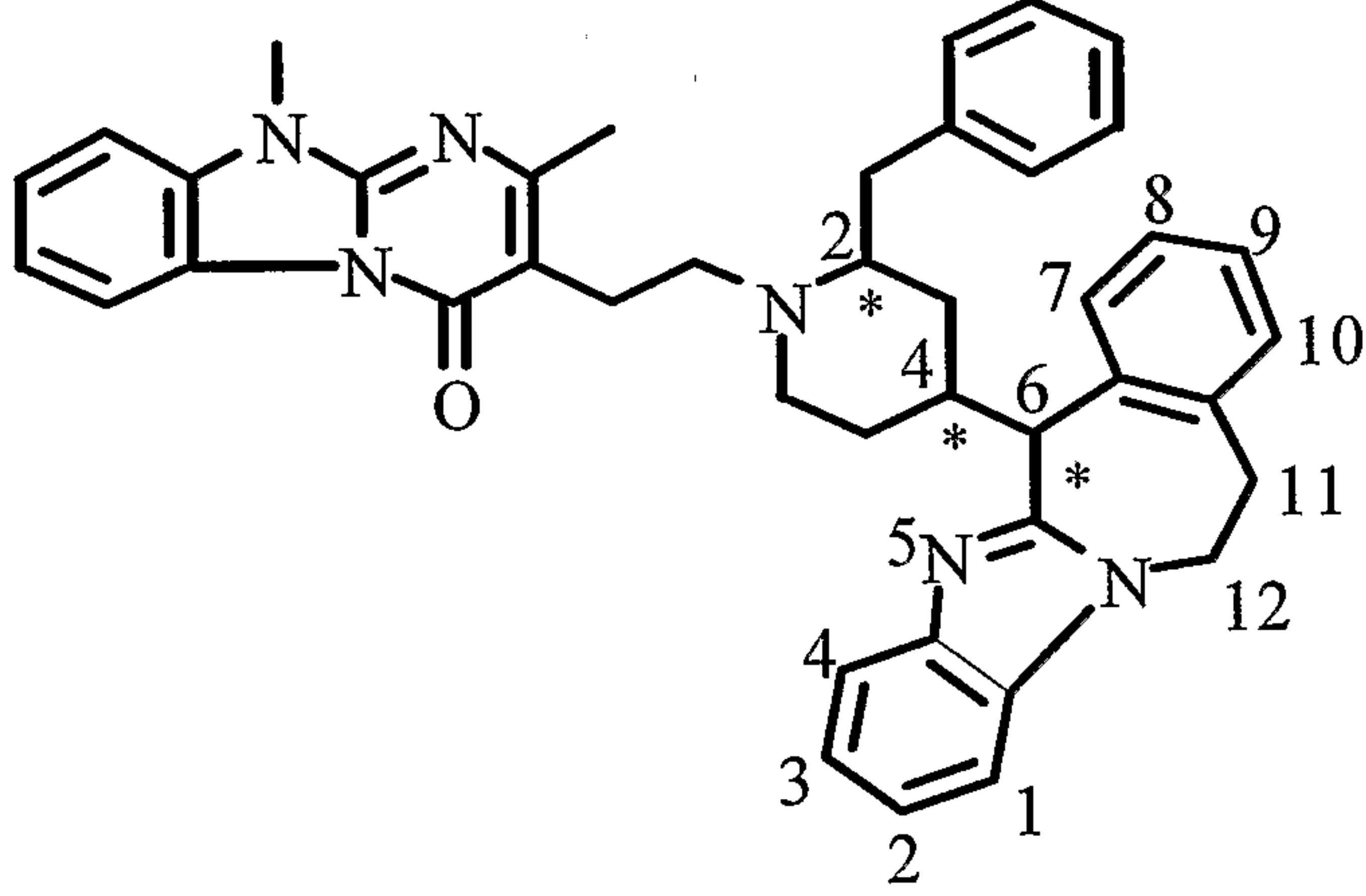
It is especially advantageous to formulate the aforementioned pharmaceutical compositions in unit dosage form for ease of administration and uniformity of dosage. Unit dosage form as used herein refers to physically discrete units suitable as unitary dosages, each unit containing a predetermined quantity of active ingredient calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. Examples of such unit dosage forms are tablets (including scored or coated tablets), capsules, pills, powder packets, wafers, suppositories, injectable solutions or suspensions and the like, and segregated multiples thereof. Most preferably, -for ease of quick administration - the aforementioned pharmaceutical composition is formulated as an injectable or perfusable solution or suspension.

The following examples illustrate the present invention without being limited thereto.

Experimental part

Of some compounds the absolute stereochemical configuration of the stereogenic
 5 carbon atom(s) therein was not experimentally determined. In those cases the
 stereochemically isomeric form which was first isolated is designated as "A" and the
 second as "B", without further reference to the actual stereochemical configuration.
 However, said "A" and "B" isomeric forms can be unambiguously characterized by a
 person skilled in the art, using art-known methods such as, for example, X-ray
 10 diffraction.

For example, for the compound pyrimido[1,2-*a*]benzimidazol-4(10*H*)-one, 3-[2-[4-
 (11,12-dihydro-6*H*-benzimidazo[2,1-*b*][3]benzazepin-6-yl)-2-(phenylmethyl)-1-
 piperidinyl]ethyl]-2,10-dimethyl, the 8 possible stereochemical isomeric forms are
 15 defined as follows :

		
CIS-forms	(2 α ,4 α)(A)	(A)[(2 α ,4 α)(A)]
		(B)[(2 α ,4 α)(A)]
	(2 α ,4 α)(B)	(A)[(2 α ,4 α)(B)]
		(B)[(2 α ,4 α)(B)]
TRANS-forms	(2 α ,4 β)(A)	(A)[(2 α ,4 β)(A)]
		(B)[(2 α ,4 β)(A)]
	(2 α ,4 β)(B)	(A)[(2 α ,4 β)(B)]
		(B)[(2 α ,4 β)(B)]

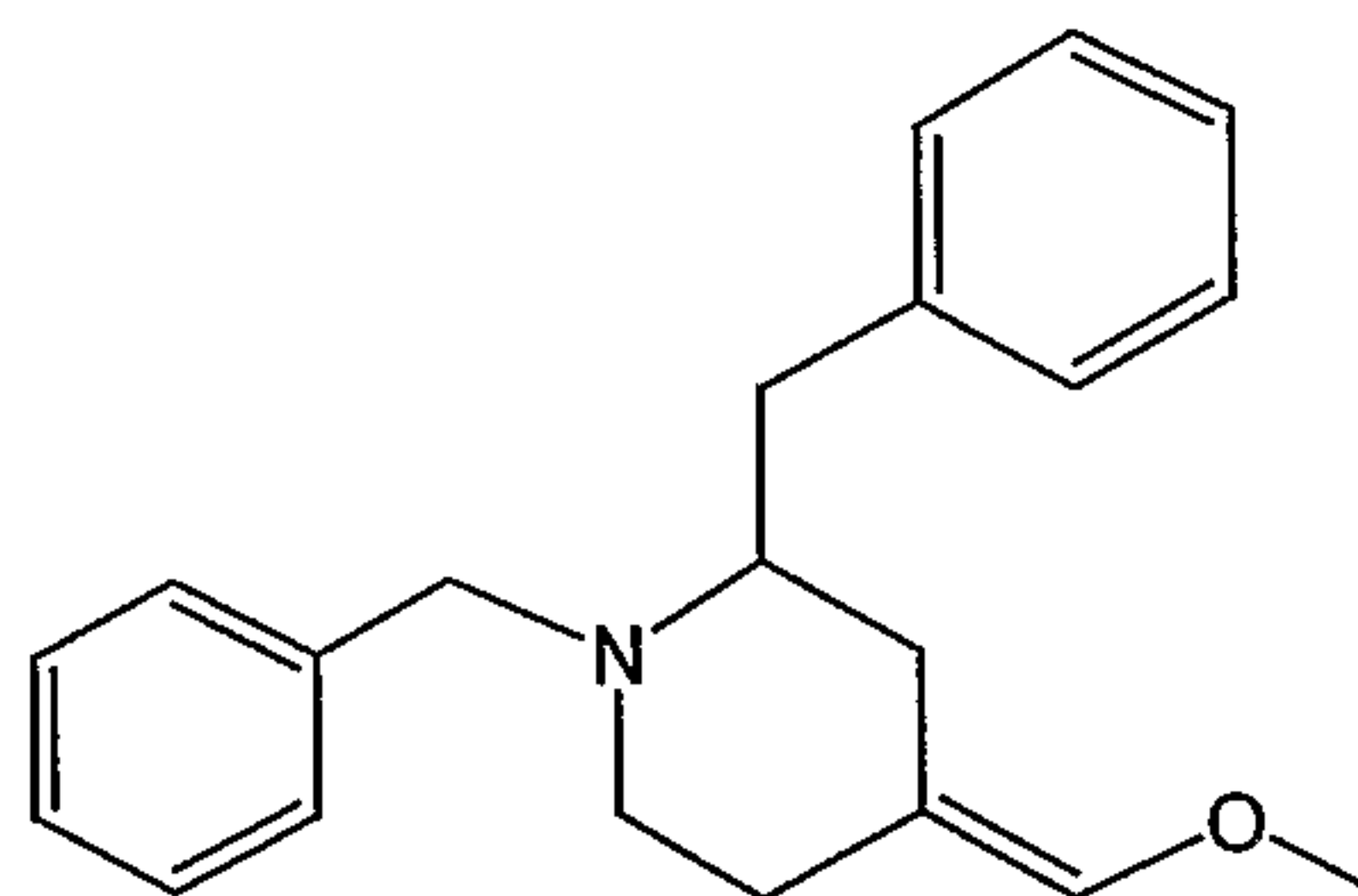
Hereinabove and hereinafter, "DMF" is defined as *N,N*-dimethylformamide, "DIPE" is defined as diisopropyl ether, "THF" is defined as tetrahydrofuran, "MIBK" is defined as methyl isobutylketone, "DIPA" is defined as diisopropylamine.

5

A. Preparation of the intermediate compounds

Example A1

a) Preparation of intermediate 1

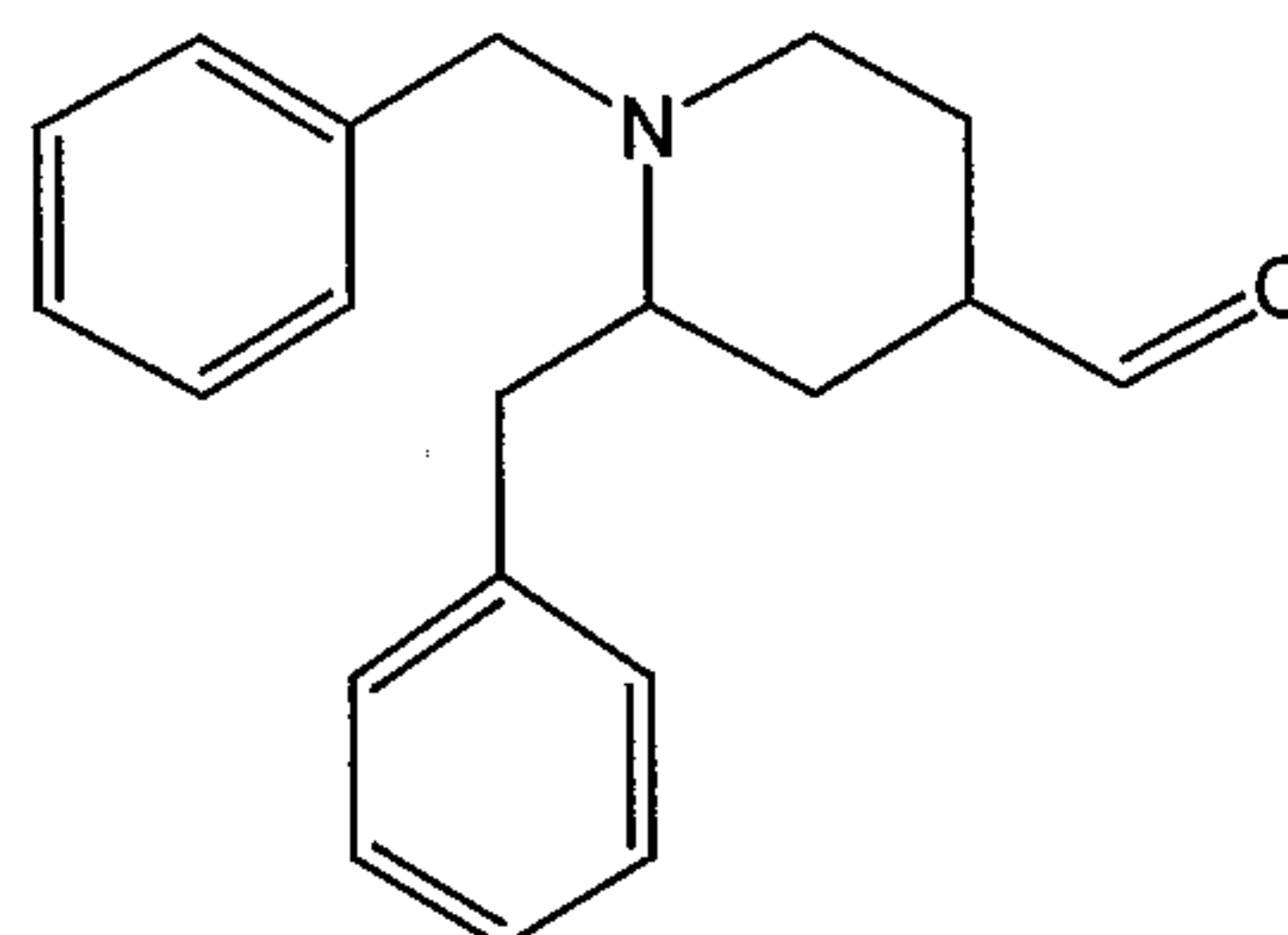


10

Use dry glassware. A mixture of (methoxymethyl)triphenylphosphoniumchloride (0.35 mol) in THF p.a.(mol. sieves) (2 l) was stirred at -50°C under N₂ flow. BuLi, 2.5M/hexane (0.35 mol) was added dropwise and the mixture was stirred at -25°C for 30 min. A solution of 1,2-bis(phenylmethyl)-4-piperidinone (0.35 mol) in THF was added dropwise at -25°C. The mixture was allowed to warm to room temperature, then stirred at room temperature overnight and decomposed with water. The organic solvent was evaporated. The aqueous concentrate was extracted with CH₂Cl₂. The organic layer was separated, dried (MgSO₄), filtered and the solvent was evaporated. The residue was purified by column chromatography over silica gel (eluent: CH₂Cl₂/CH₃OH 97.5/2.5). The pure fractions were collected and the solvent was evaporated. Yielding: 121g of 4-(methoxymethylene)-1,2-bis(phenylmethyl)piperidine enantiomeric mixture (intermediate 1) (100%).

20

b) Preparation of intermediate 2



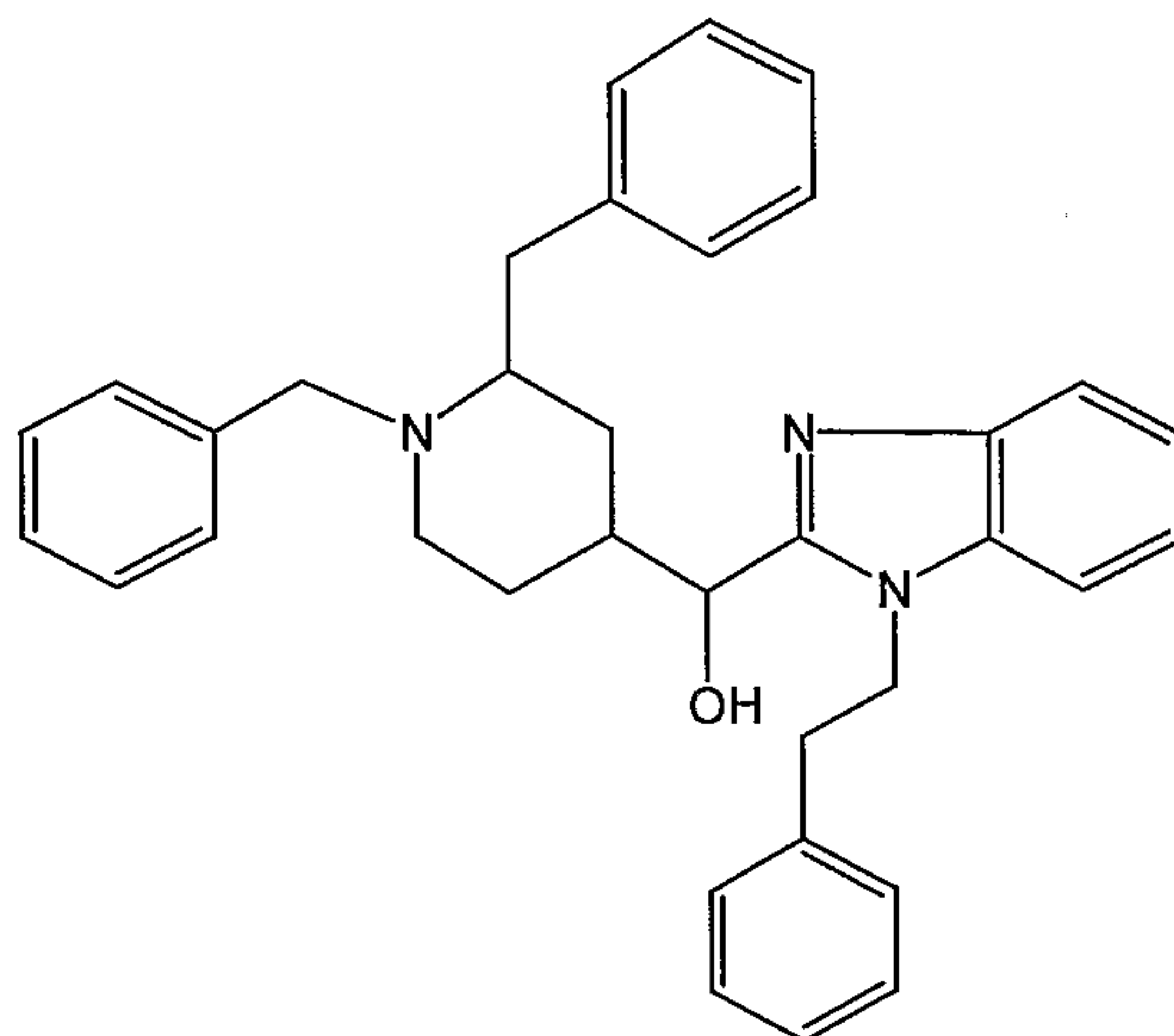
25

A mixture of intermediate 1 (0.35 mol) in THF (500ml) was stirred till complete dissolution. H₂O (900ml) and then HCl p.a. 38% (100ml) were added. The mixture was stirred and refluxed for 3 hours. The organic solvent was evaporated. The aqueous concentrate was alkalized with K₂CO₃ and extracted with CH₂Cl₂. The organic layer was separated, dried (MgSO₄), filtered and the solvent was evaporated. The residue was

purified by column chromatography over silica gel (eluent: CH₂Cl₂/CH₃OH 97/3). The pure fractions were collected and the solvent was evaporated. Yielding: 81g of 1,2-bis(phenylmethyl)-4-piperidinecarboxaldehyde enantiomeric mixture (intermediate 2) (79%).

5

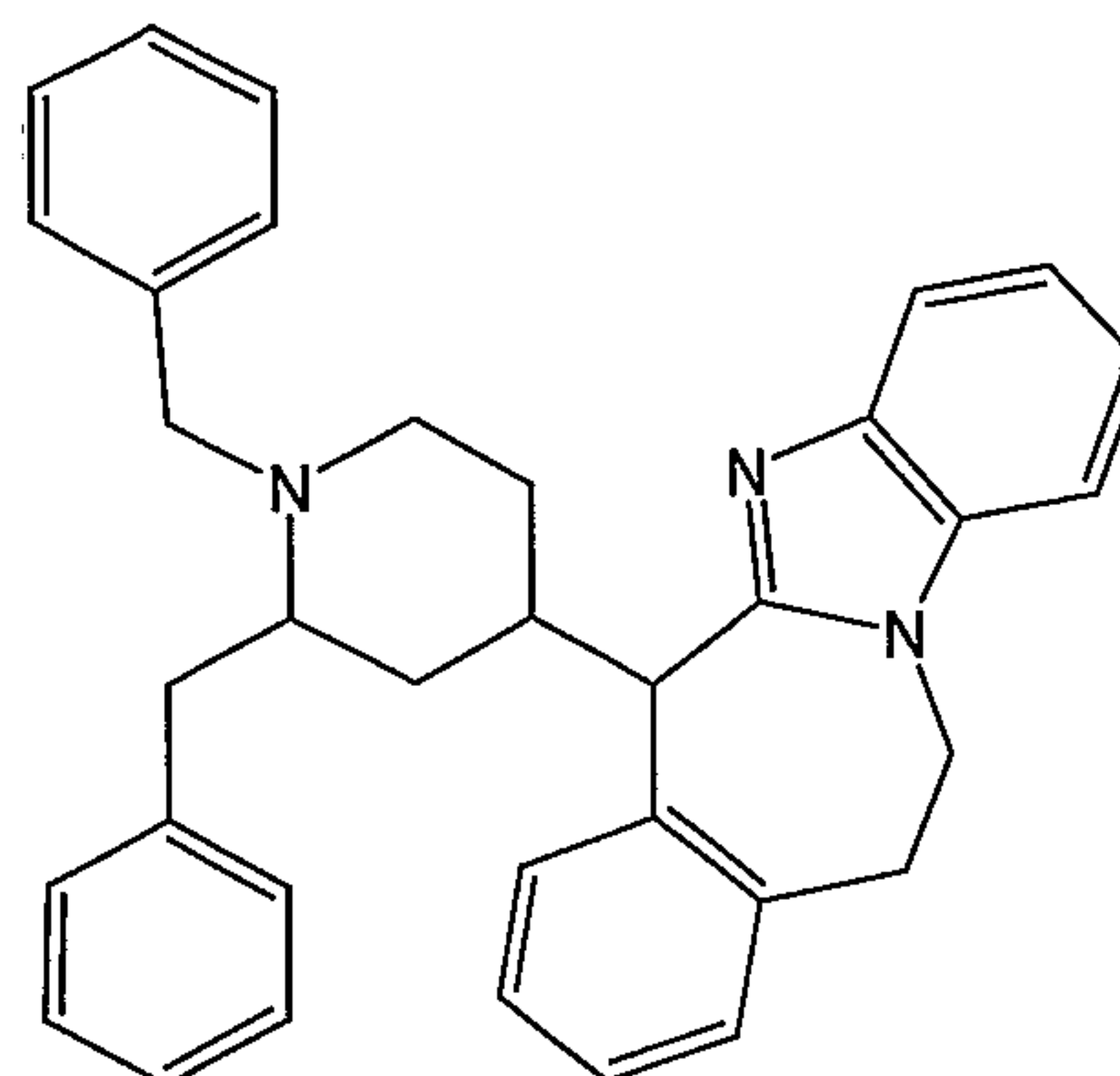
c) Preparation of intermediate 3



A mixture of DIPA (0.33 mol) in THF p.a. (previously dried on mol. sieves) (2l) was stirred at -78°C under N₂ flow. BuLi, 2.5M/hexane (0.276 mol) was added dropwise. The mixture was stirred at -78°C for 15 min. A solution of 1-(2-phenylethyl)-1H-benzimidazole

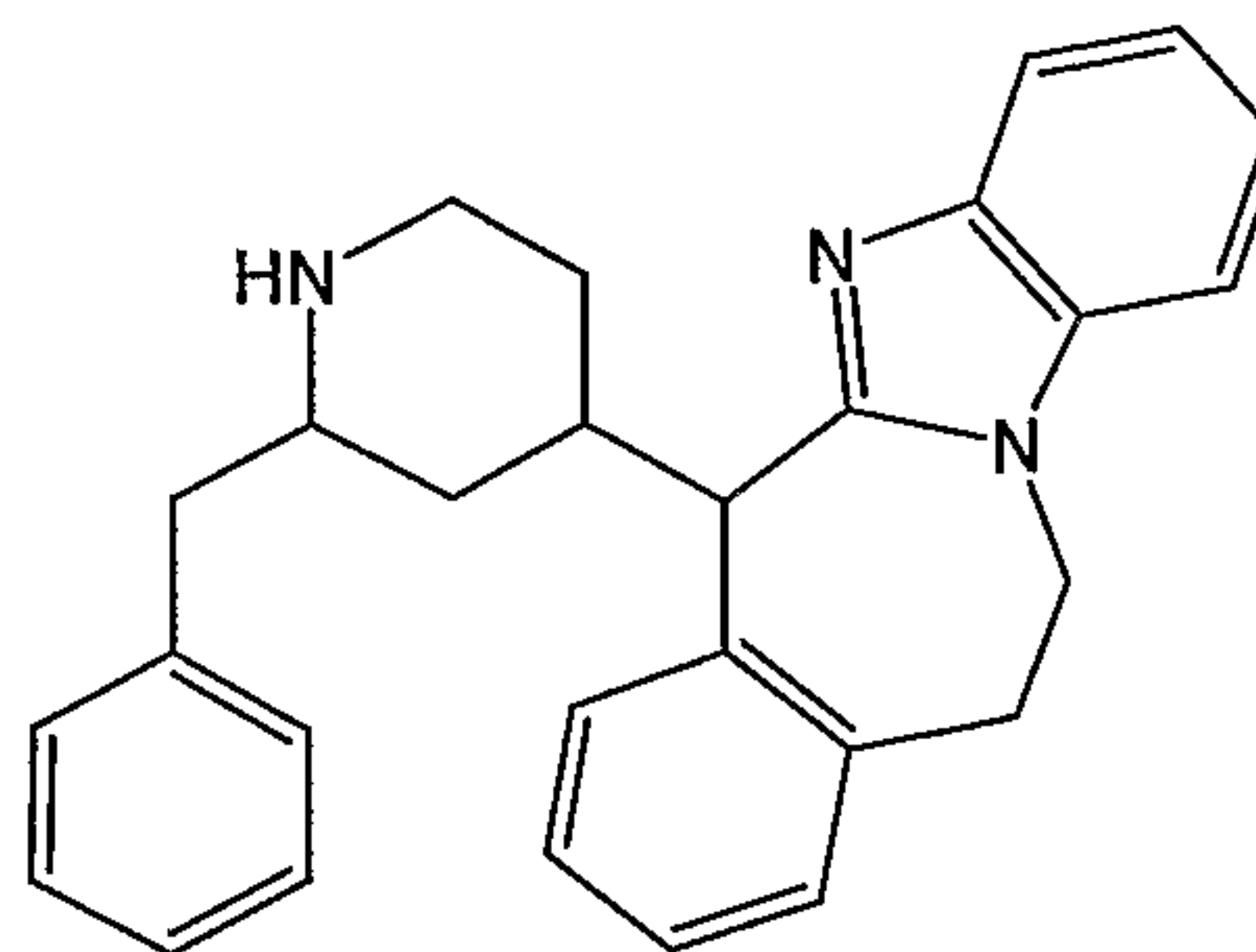
(0.276 mol) in THF was added dropwise. The mixture was stirred at -78°C for 1 hour. A solution of intermediate 2 (0.276 mol) in THF was added dropwise. The mixture was stirred at -78°C for 1 hour, then allowed to warm to room temperature, stirred at room temperature overnight and then decomposed with water. The organic solvent was evaporated. The aqueous concentrate was extracted with CH₂Cl₂. The organic layer was separated, dried (MgSO₄), filtered and the solvent was evaporated. The residue was purified by column chromatography over silica gel (eluent: CH₂Cl₂/CH₃OH 95/5 to 90/10). The pure fractions were collected and the solvent was evaporated. Yielding: 113g of α-[1,2-bis(phenylmethyl)-4-piperidinyl]-1-(2-phenylethyl)-1H-benzimidazole-2-methanol (intermediate 3)(79%).

20

d) Preparation of intermediate 4[(2 α ,4 β)(A)]

A mixture of intermediate 3 (0.22 mol) in trifluoromethanesulfonic acid (750ml) was stirred at 110°C for 7 hours. The mixture was cooled, poured out on ice, alkalized with NaOH 50% and extracted with CH₂Cl₂. The organic layer was separated, dried (MgSO₄), filtered and the solvent was evaporated. The residue was crystallized from CH₃CN. The mixture was filtered. The precipitate and the filtrate was purified separately by column chromatography over silica gel (eluent: CH₂Cl₂/CH₃OH 98.5/1.5 to 95/5). Four pure fractions were collected and their solvents were evaporated. The residues were crystallized from CH₃CN. The precipitates were filtered off and dried.

Yielding: 16g of fraction 1 [(2 α , 4 β)(A)]-6-[1,2-bis(phenylmethyl)-4-piperidiny]-11,12-dihydro-6H-benzimidazo[2,1-b][3]benzazepine (intermediate 4) (14.6%), 19.5 g of fraction 2 [(2 α , 4 β)(B)]-6-[1,2-bis(phenylmethyl)-4-piperidiny]-11,12-dihydro-6H-benzimidazo[2,1-b][3]benzazepine (17.8%), 8.66 g fraction 3 [(2 α , 4 α)(A)]-6-[1,2-bis(phenylmethyl)-4-piperidiny]-11,12-dihydro-6H-benzimidazo[2,1-b][3]benzazepine (7.9%) and 7.74 g of fraction 4 [(2 α , 4 α)(B)]-6-[1,2-bis(phenylmethyl)-4-piperidiny]-11,12-dihydro-6H-benzimidazo[2,1-b][3]benzazepine (8.9%).

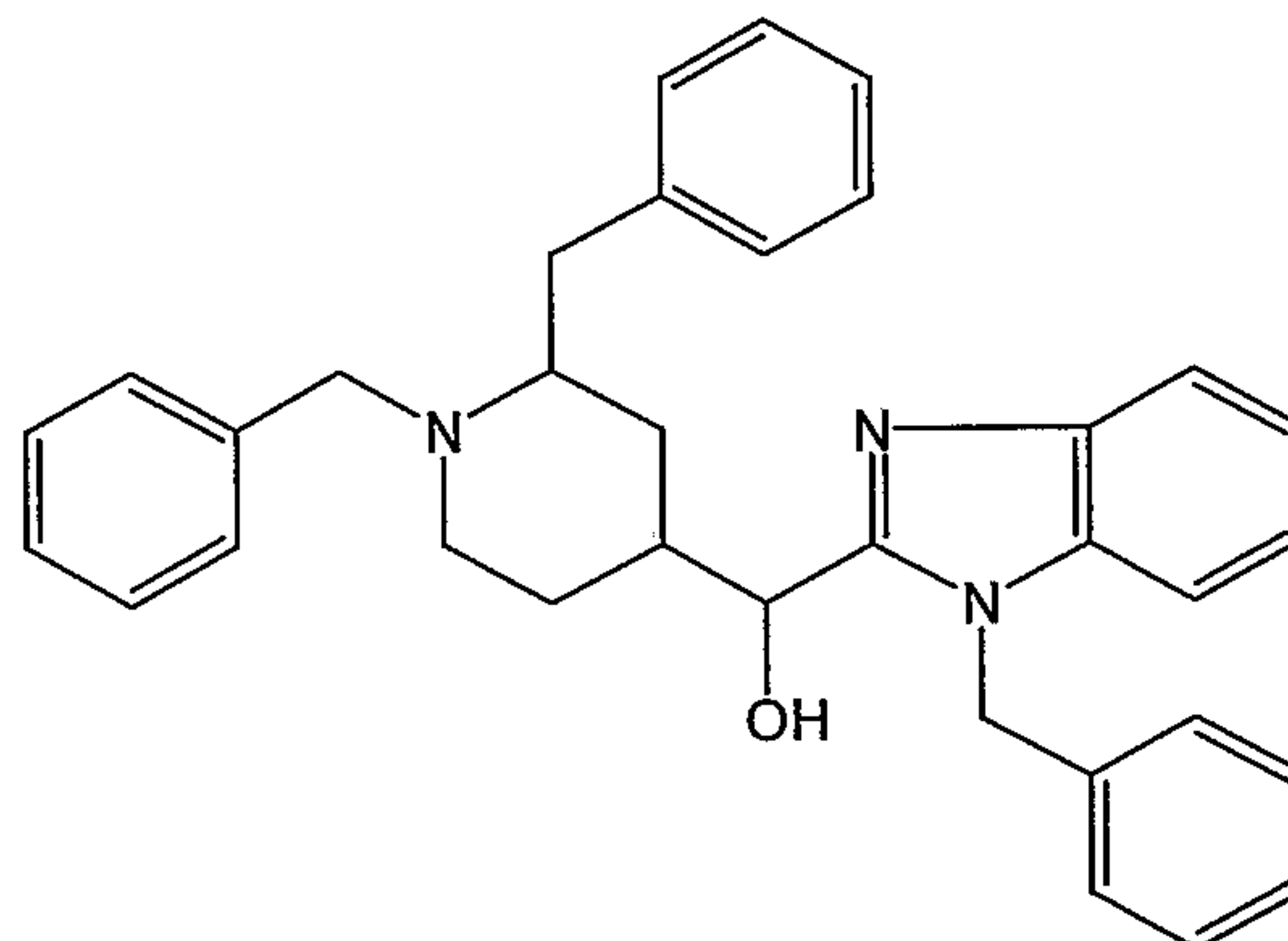
e) Preparation of intermediate 5

A mixture of intermediate 4 (0.0305 mol) in methanol (150ml) was hydrogenated at 50°C overnight with Pd/C 10% (1g) as a catalyst. After uptake of H₂ (1 equiv), the catalyst was filtered off and the filtrate was evaporated. The residue was crystallized from CH₃CN. The precipitate was filtered off and dried. Yielding: 11.66g of [(2 α ,

4β)(A)]-11,12-dihydro-6-[2-(phenylmethyl)-4-piperidiny]-6*H*-benzimidazo
[2,1-*b*][3]benzazepine (intermediate 5) (94%).

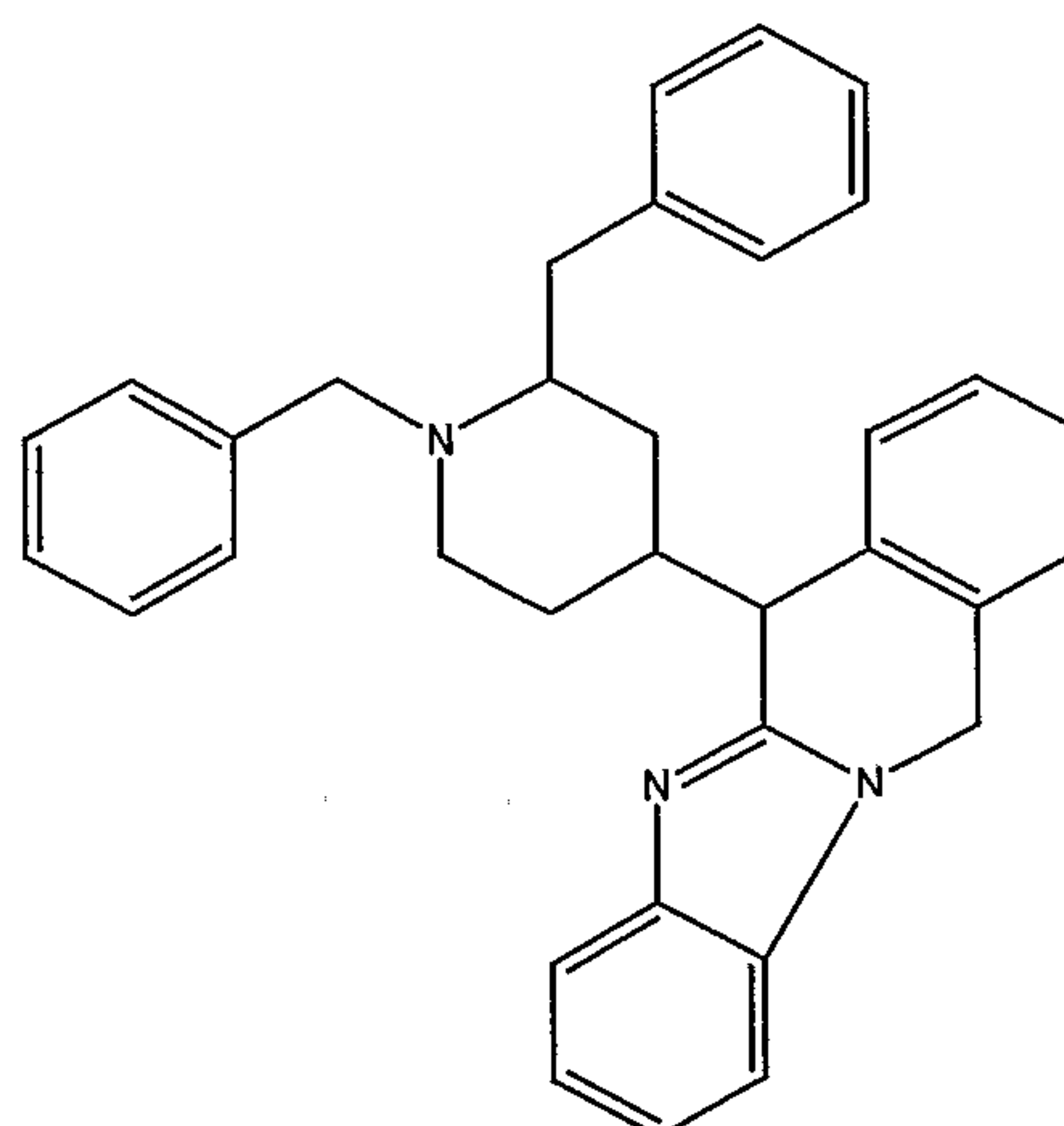
Example A2

a) Preparation of intermediate 6



- 5 Use dry glassware. A mixture of DIPA (0.22 mol) in THF p.a.(previously dried on mol.
sieves) (1400ml) was stirred at -70°C under N₂ flow. BuLi 2.5M (0.185 mol) was
added dropwise and the mixture was stirred at -70°C for 15 min. 1-(phenylmethyl)-1*H*-
benzimidazole (0.185 mol) dissolved in THF was added dropwise at -70°C and the
mixture was stirred at -70°C for 1 hour. Intermediate 2 (0.185 mol) dissolved in THF
10 was added dropwise at -70°C. The mixture was stirred at -70°C for 1 hour, then
brought slowly to room temperature, stirred at room temperature overnight and
decomposed with H₂O. The organic solvent was evaporated. The aqueous concentrate
was extracted with CH₂Cl₂. The organic layer was separated, dried (MgSO₄), filtered
and the solvent was evaporated. The residue was purified by column chromatography
15 over silica gel (eluent: CH₂Cl₂/CH₃OH 95/5). The pure fractions were collected and the
solvent was evaporated. Yielding: 91 g of intermediate 6 (98%).

b) Preparation of intermediate 7

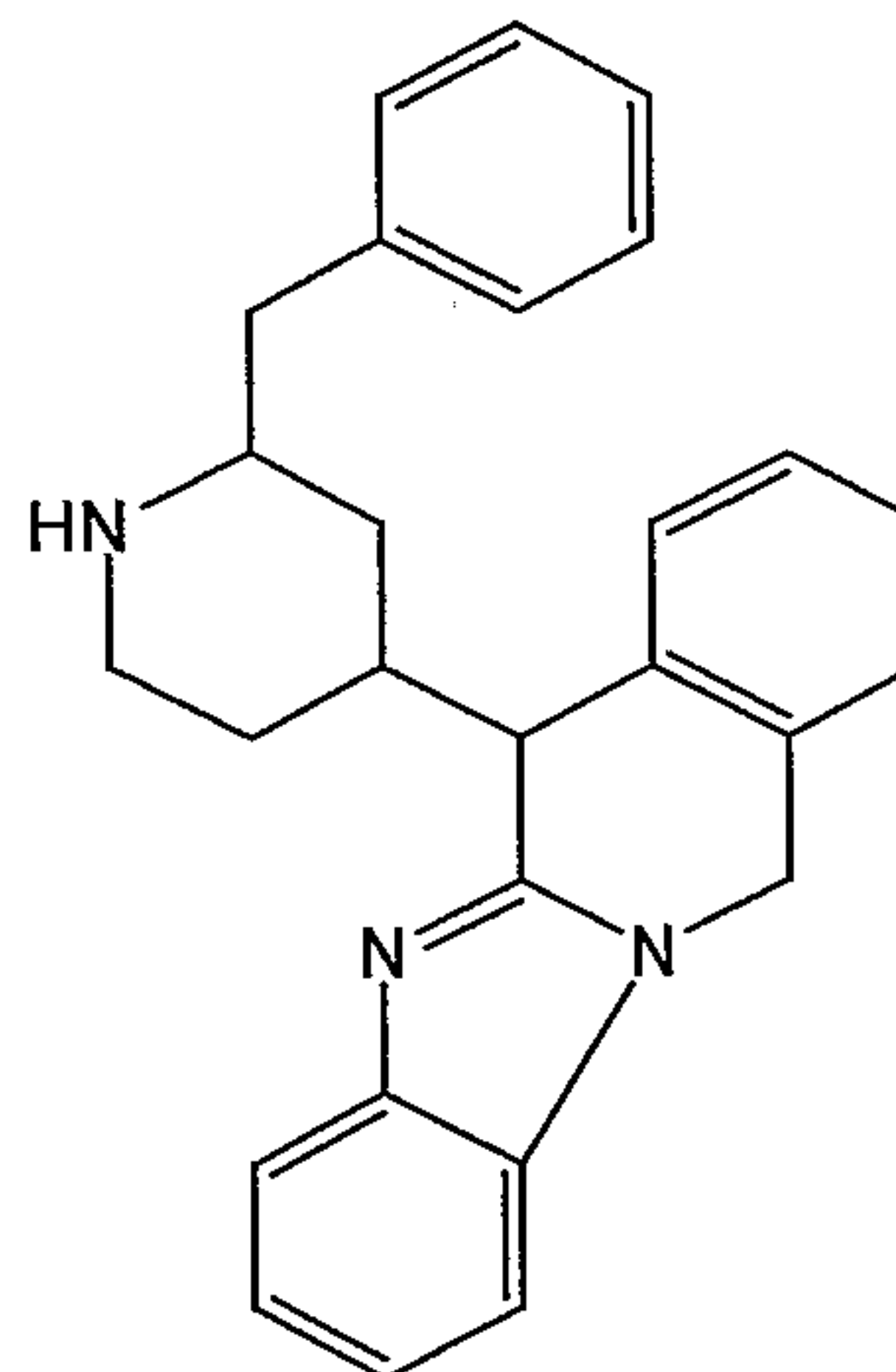


- A mixture of intermediate 6 (0.18 mol) in trifluoromethanesulfonic acid (700ml) was
stirred at 120°C under N₂ flow for 18 hours. The mixture was cooled, poured out on
20 ice, alkalized with NaOH 50% and extracted with CH₂Cl₂. The organic layer was

separated, dried (MgSO_4), filtered and the solvent was evaporated. The residue was purified by column chromatography over silica gel (eluent: $\text{CH}_2\text{Cl}_2/(\text{CH}_3\text{OH}/\text{NH}_3)$ 99/1). The pure fractions were collected and the solvent was evaporated. Yielding: 40g of intermediate 7 (46%).

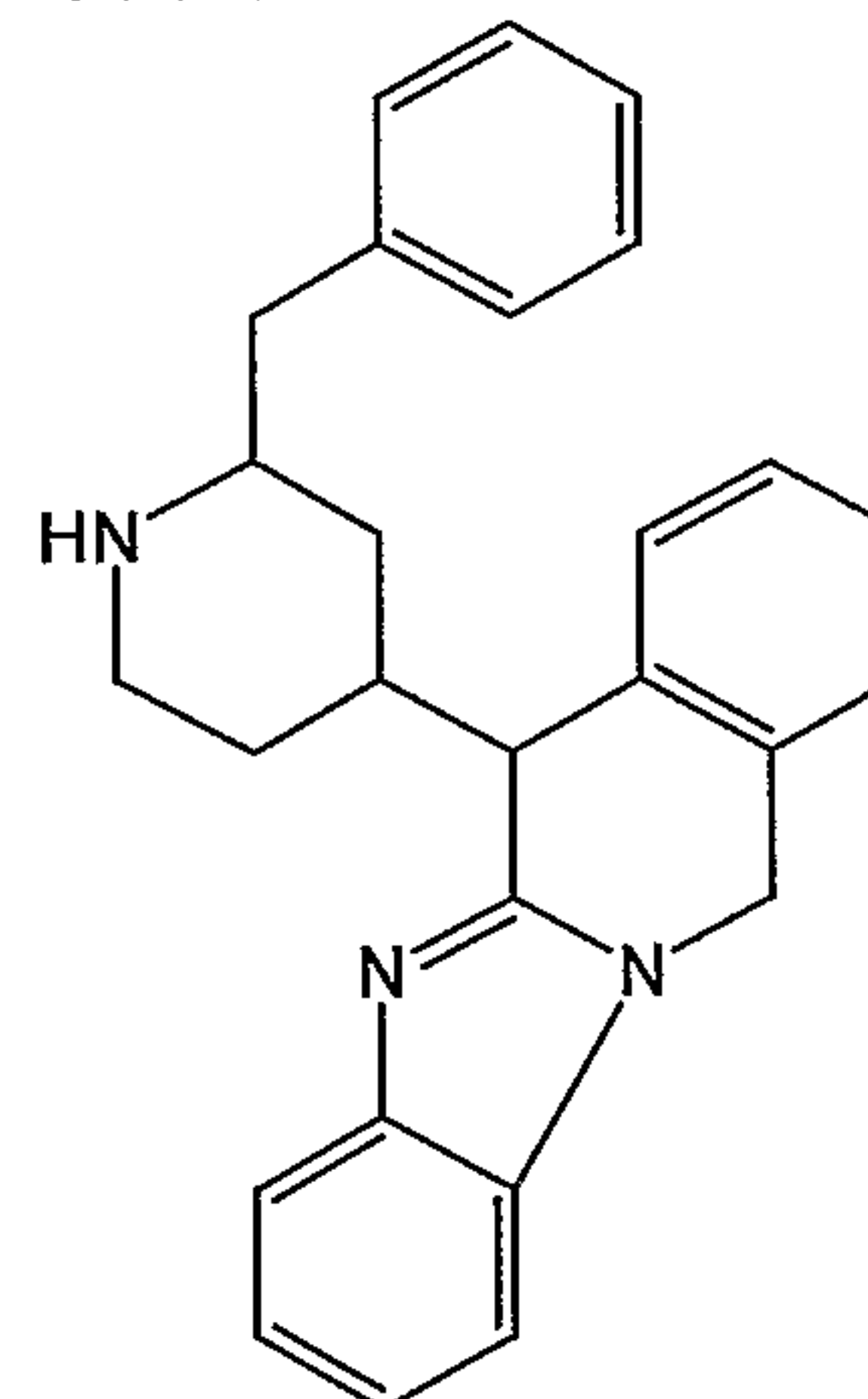
5

c) Preparation of intermediate 8



[(2 α ,4 β)(A)]

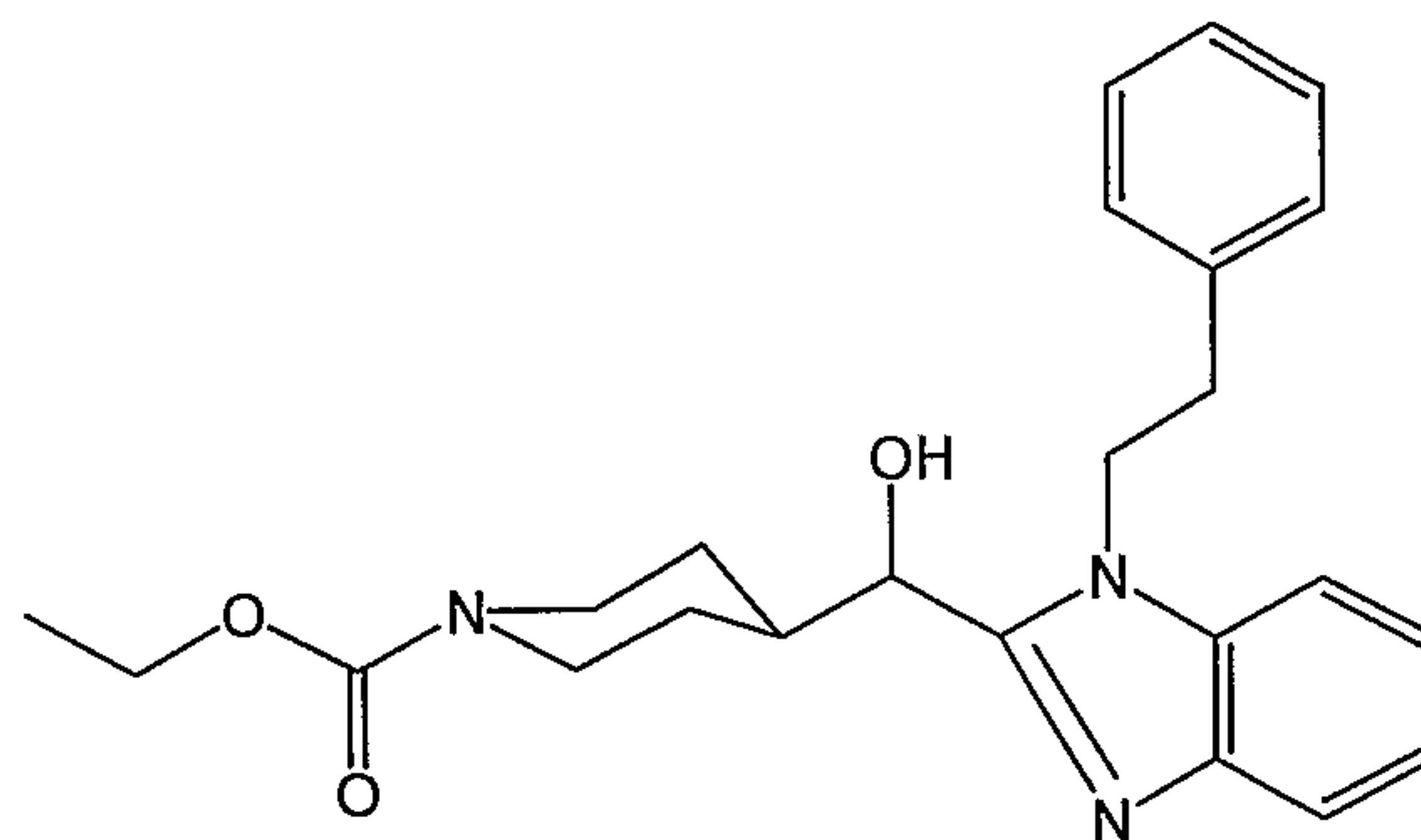
and preparation of intermediate 9



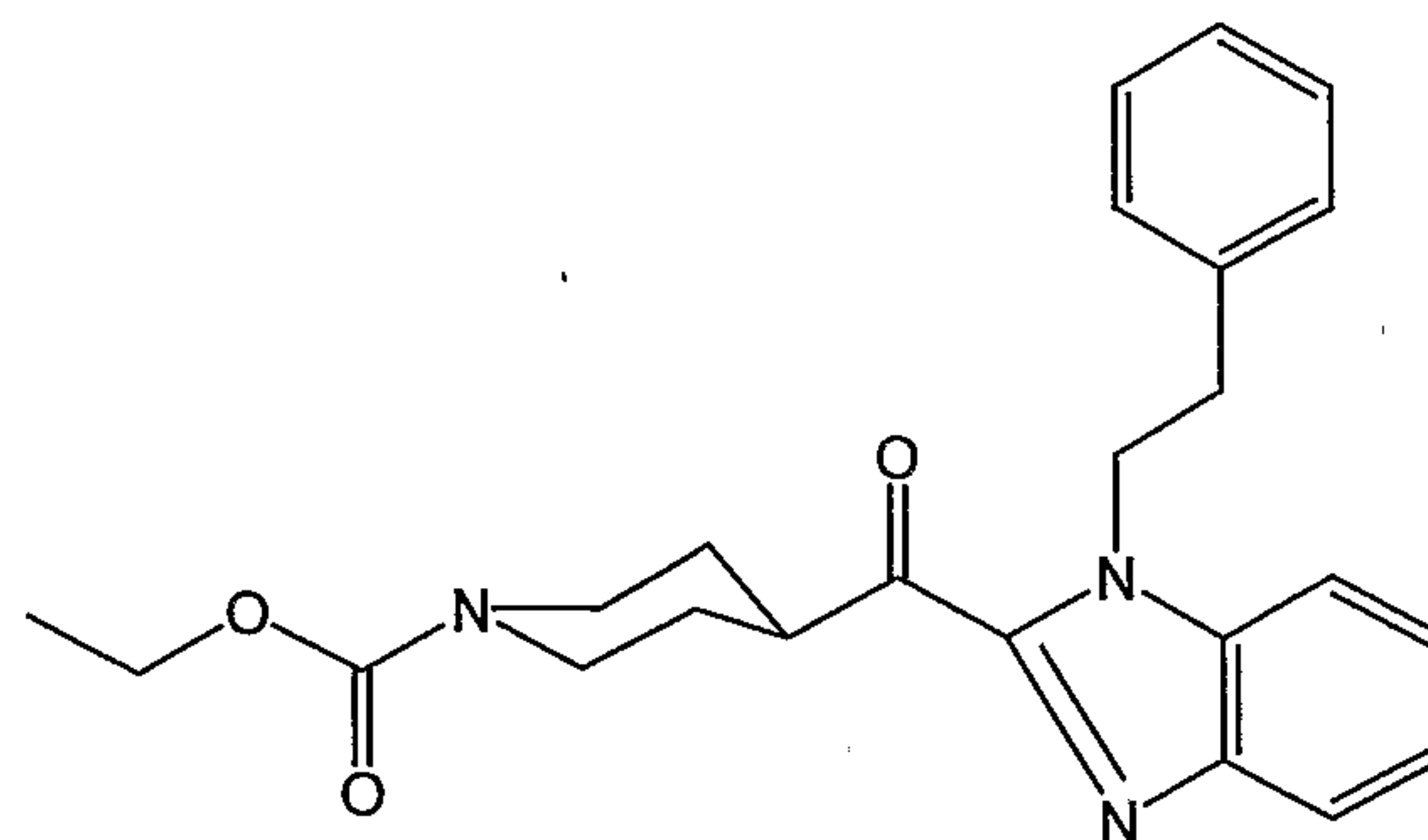
CIS

A mixture of intermediate 7 (0.081 mol) in methanol (200ml) was hydrogenated at 50°C with Pd/C 10% (2g) as a catalyst. After uptake of H_2 (1 equiv), the catalyst was filtered off and the filtrate was evaporated. This fraction was purified by column chromatography over silica gel (eluent: $\text{CH}_2\text{Cl}_2/(\text{CH}_3\text{OH}/\text{NH}_3)$ 97/3). Two pure fractions were collected and their solvents were evaporated. Yielding: Fraction 1 and 12.5g of intermediate 9 (cis isomers) (36%). Fraction 1 was crystallized from CH_3CN . The precipitate was filtered off and dried. Yielding: 4.44g of intermediate 8 (14%) ([2 α , 4 β)(A)]-racemate.

10

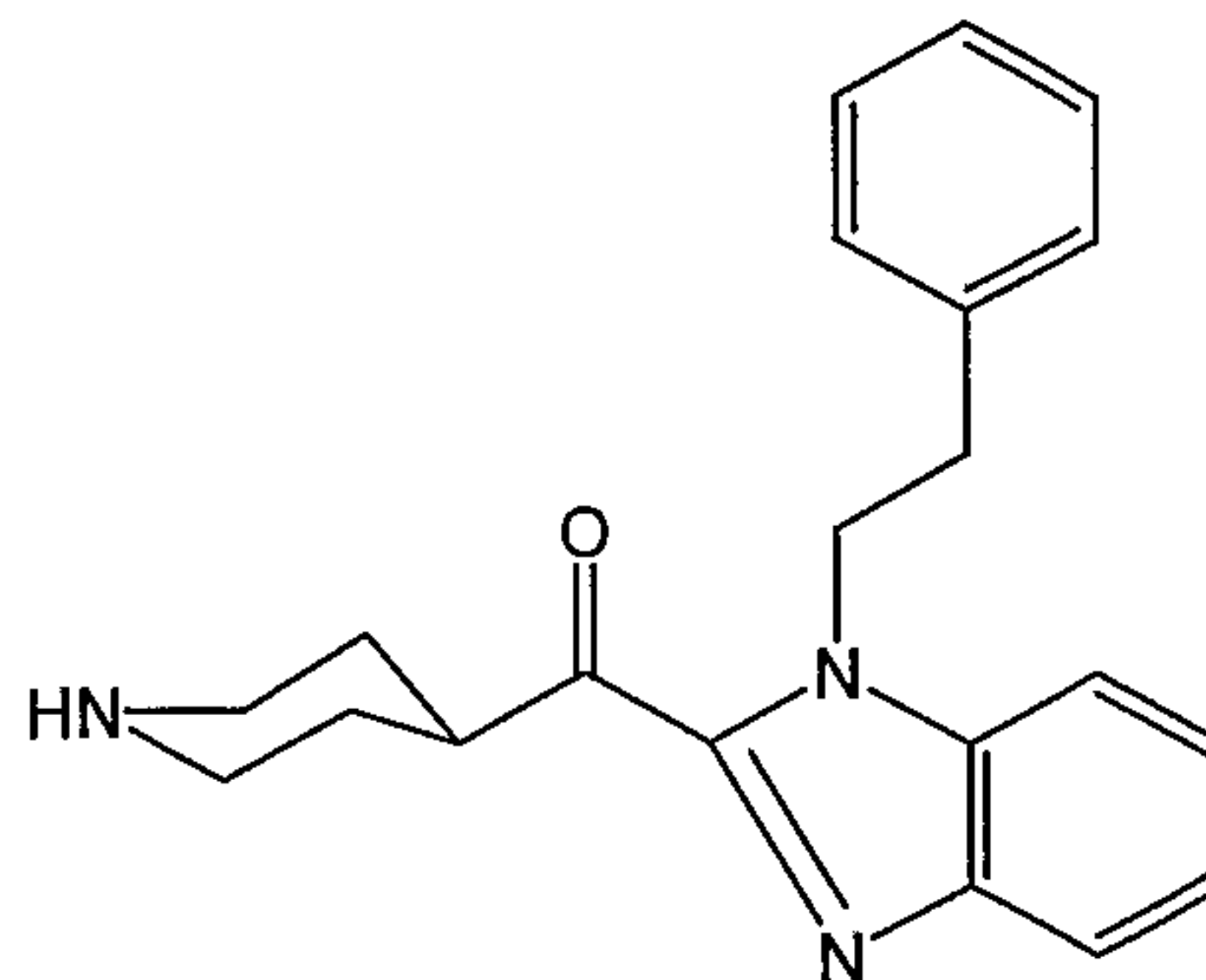
Example A3a) Preparation of intermediate 10

A mixture of DIPA (0.1mol) in THF (100ml) was stirred under N₂ flow. The mixture was cooled to -70°C and BuLi, 2.5M/hexane (40ml) was added portionwise. The temperature was allowed to reach -30°C, while stirring for 10 min. The mixture was cooled to -70°C. A solution of 1-(phenylethyl)-1H-benzimidazole (0.1mol) in THF (50ml) was added dropwise at this temperature and the mixture was stirred for 2h at -70°C. Ethyl 4-formyl-1-piperidinecarboxylate (0.1mol) was added dropwise and the mixture was stirred for 30 min at -70°C. The mixture was allowed to reach room temperature and stirring was continued for 30 min. The mixture was decomposed with water, then evaporated. The residue was stirred in water, and this mixture was extracted with CH₂Cl₂. The organic layer was separated, dried, filtered and the solvent was evaporated. The residue was purified by column chromatography over silica gel (eluent : CH₂Cl₂/CH₃OH 98/2). The pure fractions were collected and the solvent was evaporated. Yielding : 38g of ethyl 4-[[hydroxy[1-(2-phenylethyl)-1H-benzimidazol-2-yl]methyl]-1-piperidinecarboxylate (intermediate 10).

b) Preparation of intermediate 11

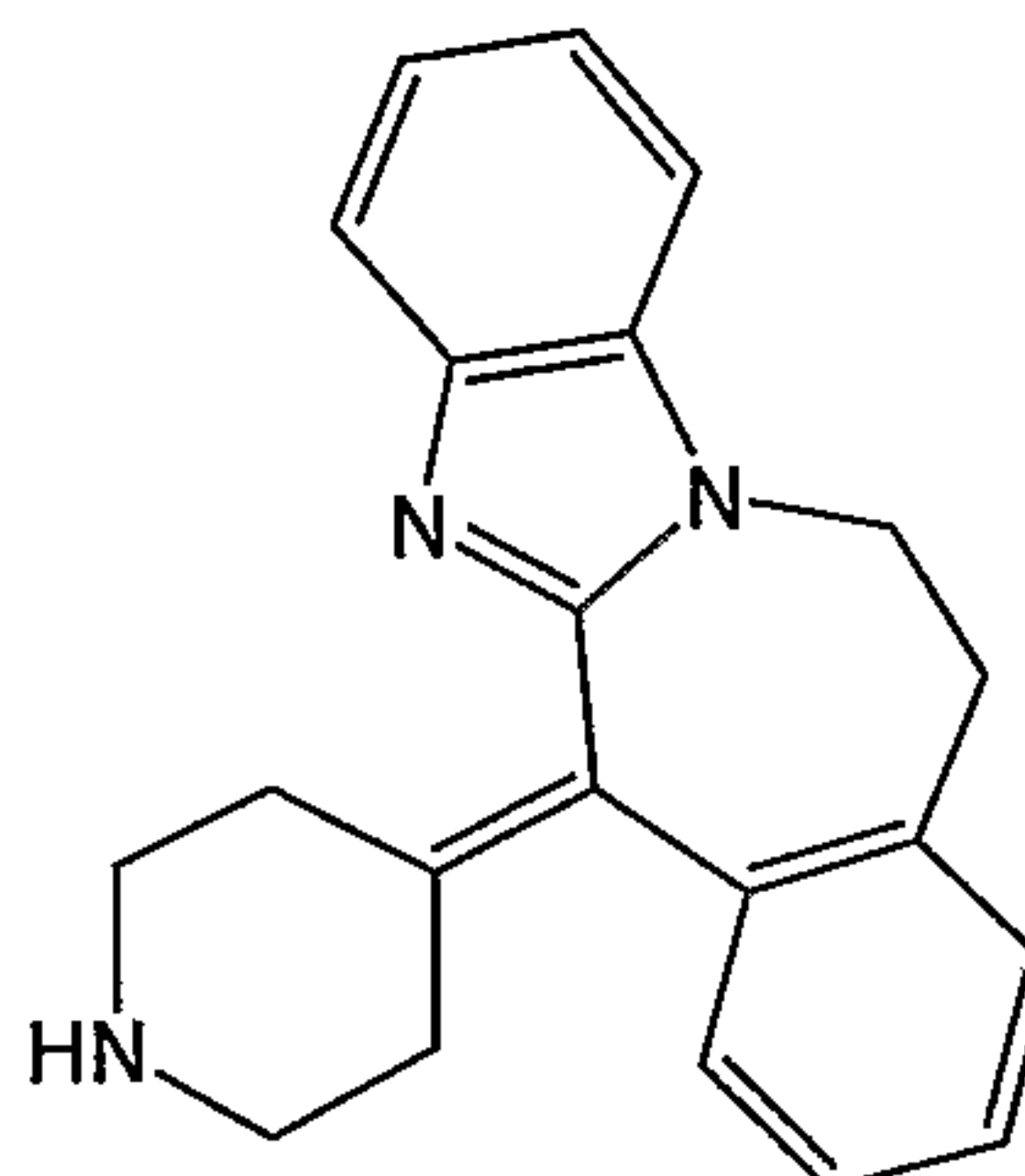
A mixture of intermediate 10 (0.011 mol) and MnO₂ (15g) in CH₂Cl₂ (150ml) was stirred overnight at room temperature. MnO₂ was filtered off over dicalite. The reaction was performed a second time with identical quantities. The mixture was stirred overnight. MnO₂ was filtered off over dicalite. The filtrate was evaporated. Yielding : 4.5g ethyl 4-[[[1-(2-phenylethyl)-1H-benzimidazol-2-yl]carbonyl]-1-piperidinecarboxylate (intermediate 11).

c) Preparation of intermediate 12

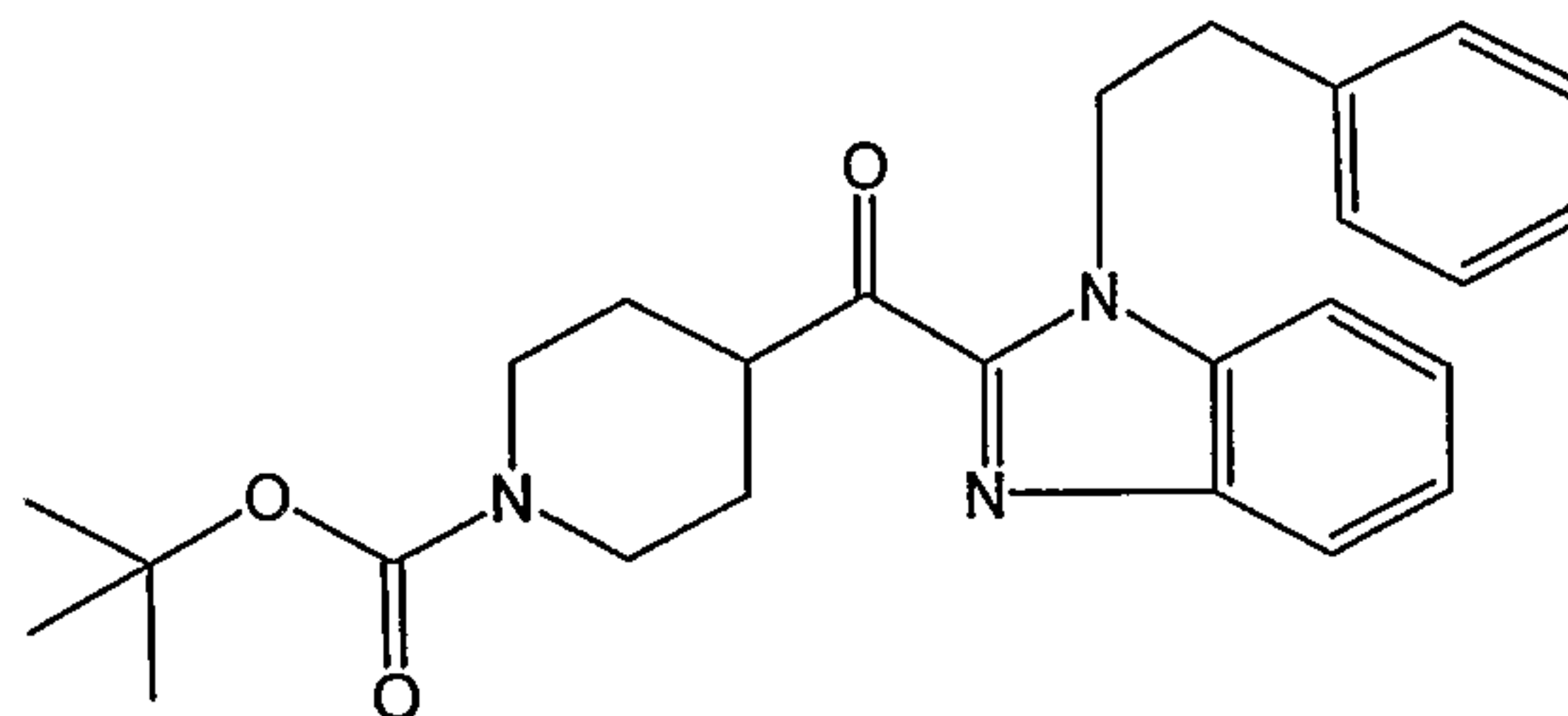


A mixture of intermediate 11 (0.011 mol) and HBr, 48%aq. (25ml) was stirred for 10h at 80°C. The solvent was evaporated. The residue was stirred in boiling 2-propanol, cooled and the resulting precipitate was filtered off and dried. A sample (1g) was recrystallized from ethanol. The crystals were filtered off and dried. Yielding: 0.5g of [1-(2-phenylethyl)-1*H*-benzimidazol-2-yl] (4-piperidinyl)methanone dihydrobromide (intermediate 12) (mp. 261.9°C).

d) Preparation of intermediate 13

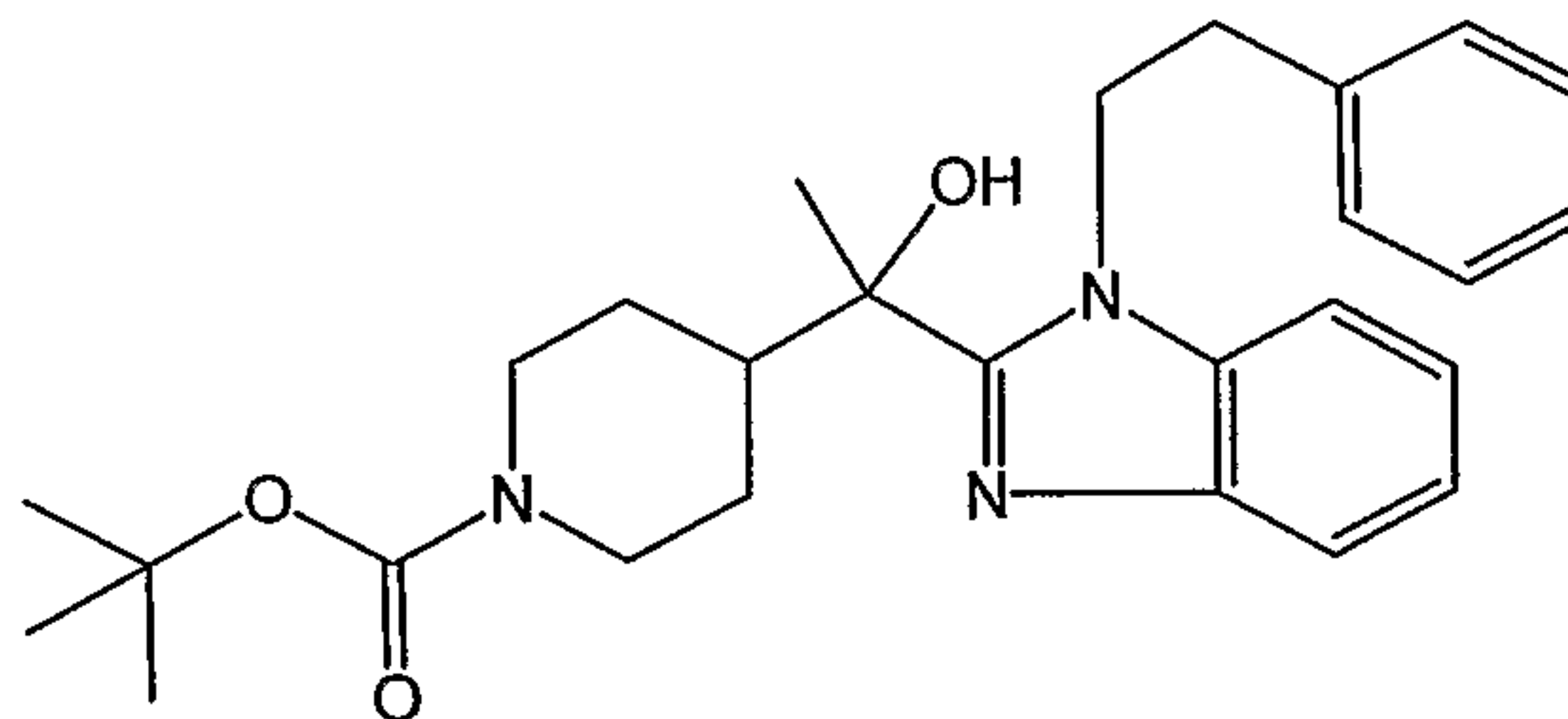


Trifluoromethanesulfonic acid (150ml) was stirred under N₂ flow. Intermediate 12 (0.1mol) was added portionwise and the resulting reaction mixture was stirred for 20h at 100°C (N₂ flow). The reaction mixture was cooled, poured out into ice (1kg) and the resulting mixture was neutralized with NaOH 50%, while stirring and cooling. This mixture was extracted with CH₂Cl₂. Precipitation resulted. The organic layer was separated. The precipitate was filtered off and recrystallized from CH₃CN. The crystals were filtered off and recrystallized again from CH₃CN. The crystals were filtered off and dried. Yielding: 3.0 g of 11,12-dihydro-6-(4-piperidinylidene)-6*H*-benzimidazo[2,1-*b*][3]benzazepine .trifluoromethanesulfonate (2:3). The separated organic liquor was combined with the mother layers, dried, filtered and the solvent was evaporated. The residue (37g) was dissolved in water/ethanol, alkalized with 50% NaOH and extracted with CH₂Cl₂. The separated organic layer was dried (MgSO₄), filtered and the solvent was evaporated. The residue was stirred in 2-propanone/DIPE, then filtered off and dried. Yielding : 16.2g of 11,12-dihydro-6-(4-piperidinylidene)-6*H*-benzimidazo[2,1-*b*][3]benzazepine (intermediate 13) (mp. 180.3°C).

Example A4a) Preparation of intermediate 14

Use dry glassware. A mixture of DIPA (1.1 mol) in THF p.a. (previously dried on mol. sieves) (3000ml) was stirred at -78°C under N₂ flow. BuLi 1.5M in hexane (1.05 mol) was added dropwise at -70°C and the mixture was stirred at -70°C for 20 min. 1-(phenylethyl)-1*H*-benzimidazole (1 mol) dissolved in THF was added dropwise at -78°C and the mixture was stirred at -78°C for 1 hour. 4-ethyl 1-(1,1-dimethyl)1,4-piperidinedicarboxylate (1.1 mol) dissolved in THF was added dropwise at -70°C. The mixture was stirred at -78°C for 1 hour, then brought to room temperature, stirred at room temperature overnight and decomposed with H₂O. The organic solvent was evaporated. The aqueous concentrate was extracted with CH₂Cl₂. The organic layer was separated, dried (MgSO₄), filtered and the solvent was evaporated. The residue was crystallized from CH₃CN. The precipitate was filtered off and dried. Yielding: 350g of intermediate 14 (81%).

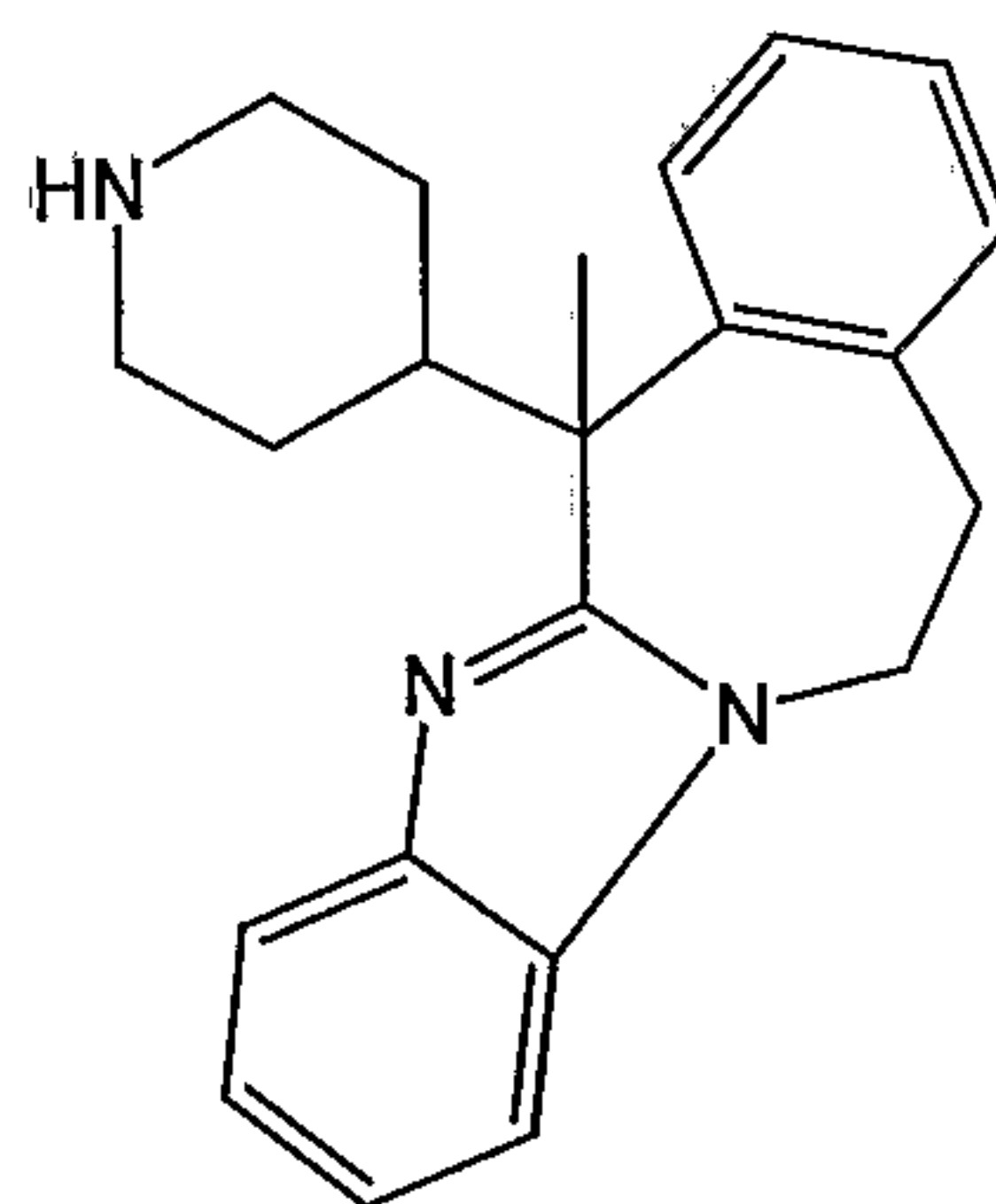
15

b) Preparation of intermediate 15

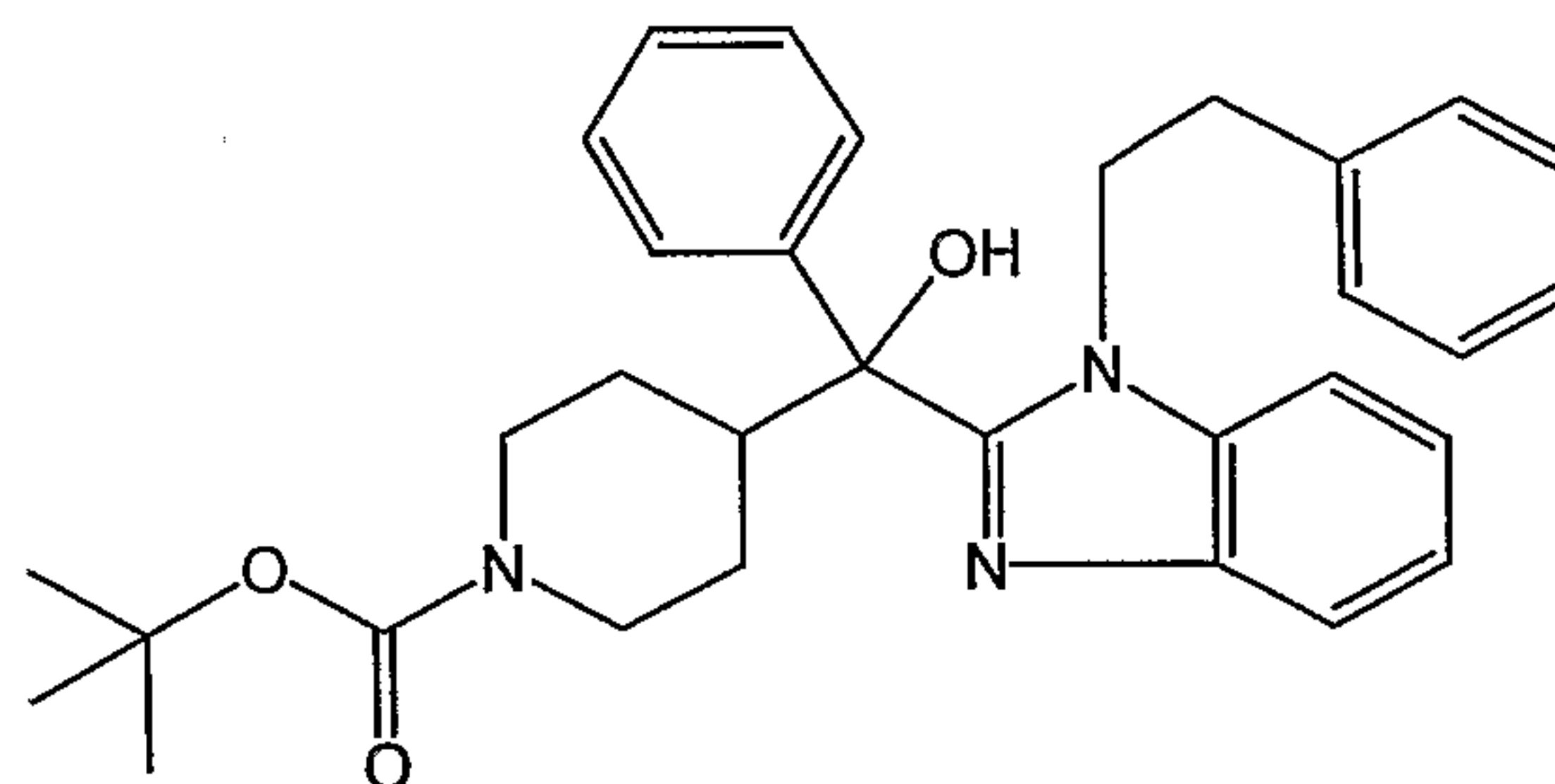
Reaction under N₂ atmosphere. Methylmagnesium chloride (0.0165 mol; 8.2 ml, 2.0 M/THF) was added dropwise to a solution of intermediate 14 (0.0150 mol) in THF (90 ml), stirred at room temperature. The resulting reaction mixture was stirred for 2 hours. Water was added. The organic solvent was evaporated and the aqueous concentrate was extracted with CH₂Cl₂. The separated organic layer was dried, filtered and the solvent evaporated. The residue (6 g) was crystallized from CH₃CN. The precipitate was filtered off and dried. Yielding: 4.3 g of intermediate 15 (64%).

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-25-

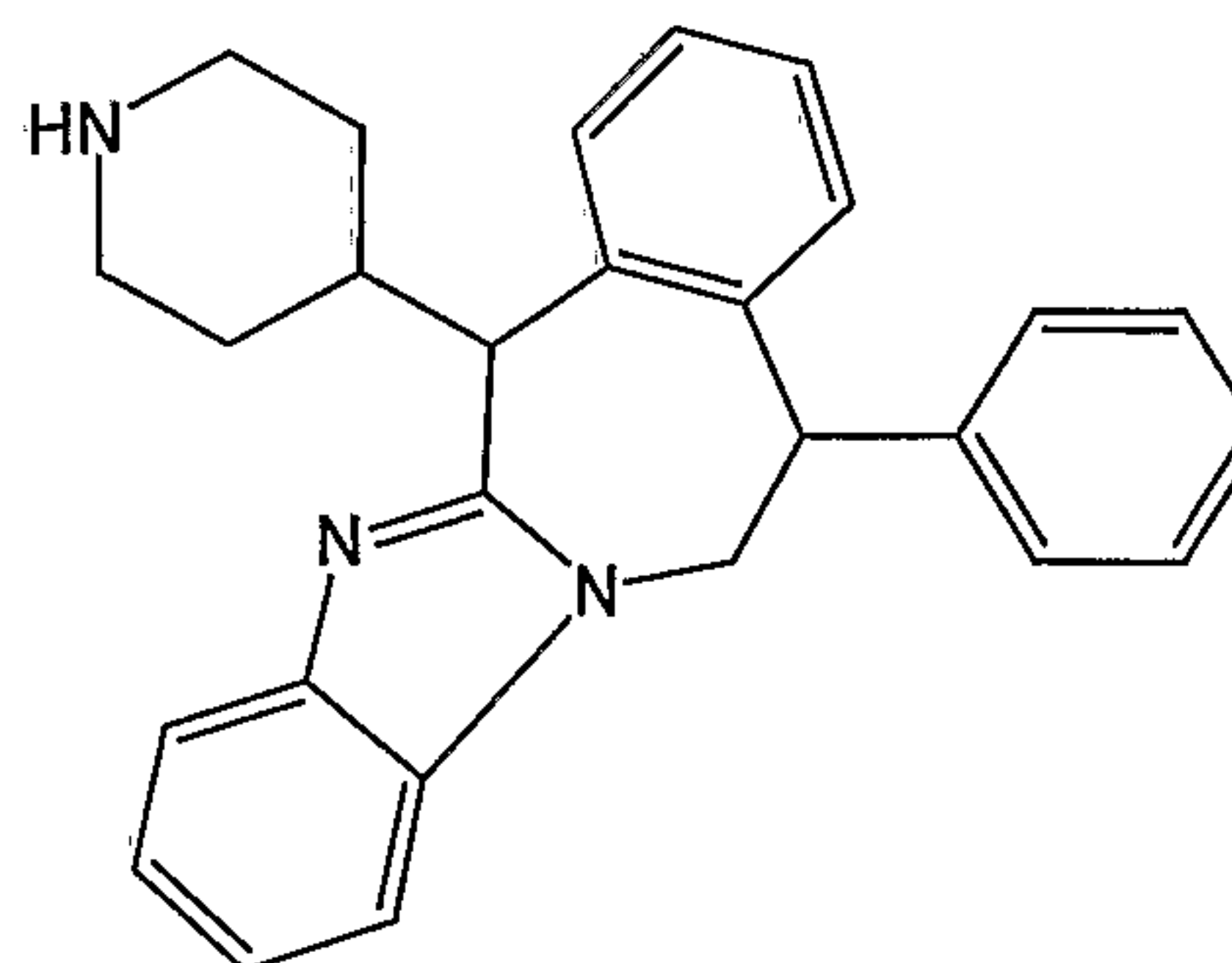
c) Preparation of intermediate 16

A mixture of intermediate 15 (0.0076 mol) in trifluoromethanesulfonic acid (29 ml) was stirred for 48 hours at room temperature. The reaction mixture was poured out into water. This mixture was alkalized with K_2CO_3 . The aqueous layer was extracted with
5 CH_2Cl_2 . The separated organic layer was dried, filtered and the solvent evaporated. The residue was purified by short open column chromatography over silica gel (eluent: $CH_2Cl_2/(CH_3OH/NH_3)$ 90/10). The pure fractions were collected and the solvent was evaporated. Yielding: 2 g of intermediate 16 (79%).

10 Example A5a) Preparation of intermediate 17

Reaction under N_2 atmosphere. Phenylmagnesium chloride (0.0440 mol) was added to a solution of intermediate 14 (0.0400 mol) in THF (200 ml), stirred at room temperature. The resulting reaction mixture was stirred for one hour. Water was added. The organic solvent was evaporated and the aqueous concentrate was extracted with
15 CH_2Cl_2 . The separated organic layer was dried, filtered and the solvent evaporated. This residue was combined with analogously obtained material and the whole (20 g) was crystallized from CH_3CN . The precipitate was filtered off and dried. Yielding: 20 g of intermediate 17 (98%).

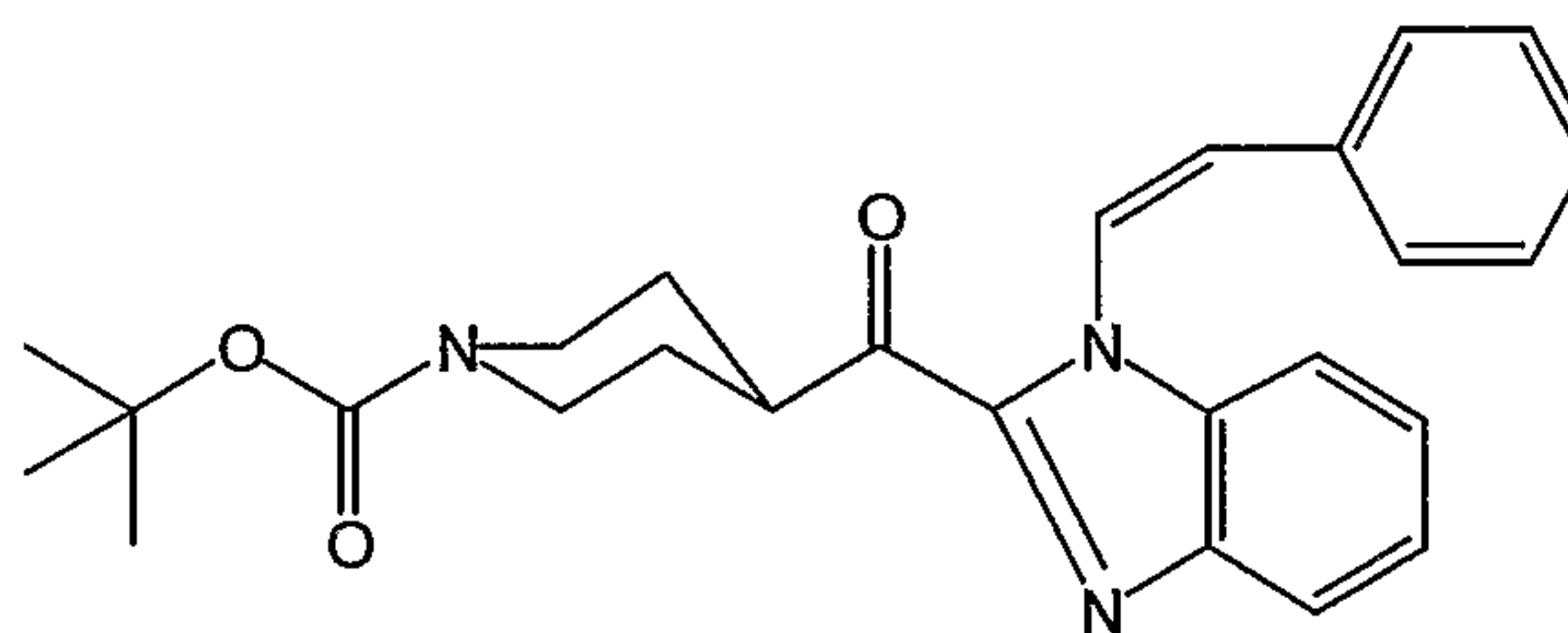
b) Preparation of intermediate 18



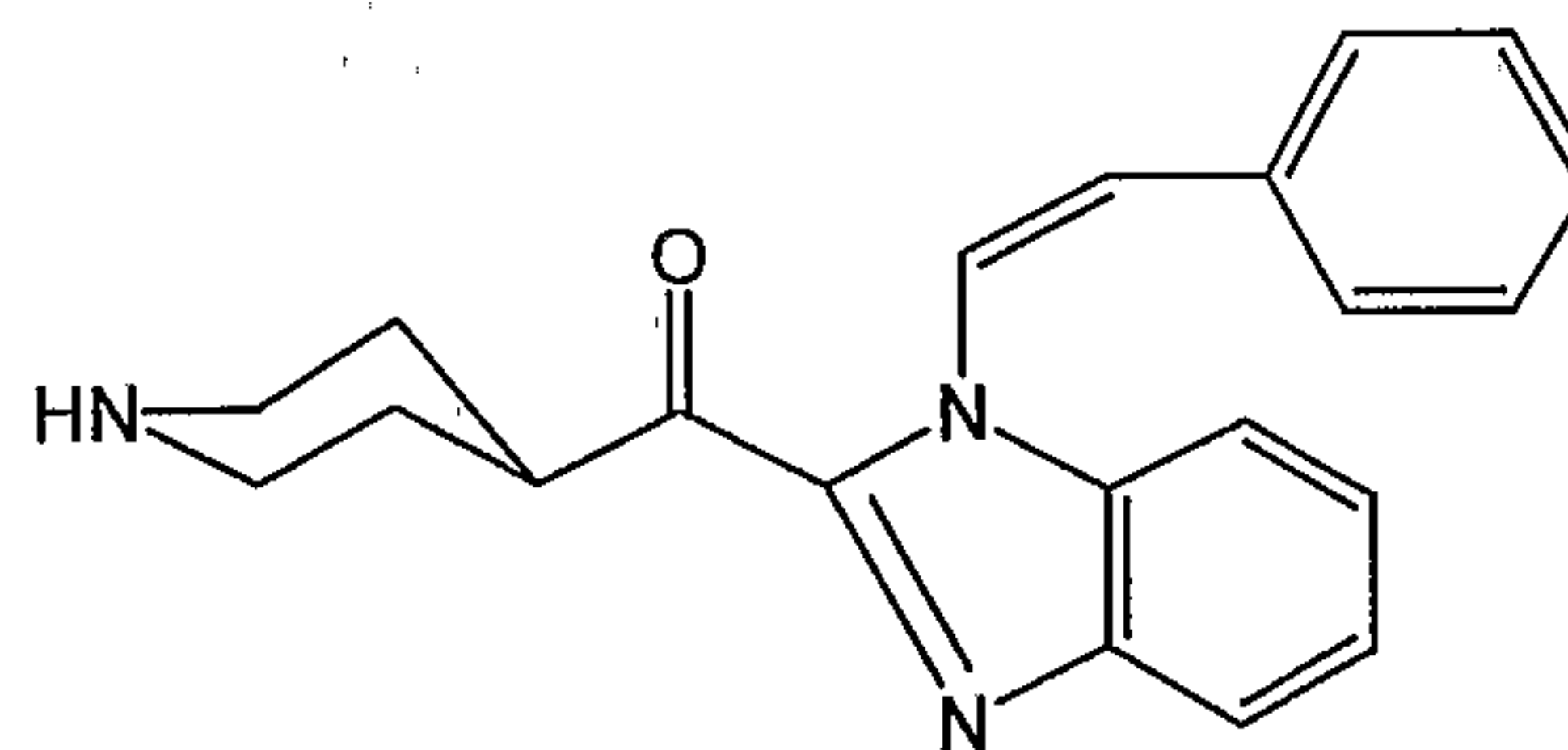
A mixture of intermediate 17 (0.0360 mol) in trifluoromethanesulfonic acid (120 ml) was stirred for 24 hours, going from 0 °C to room temperature. The reaction mixture was poured out into water. This mixture was alkalized with NaOH 50%, then extracted with CH₂Cl₂. The separated organic layer was dried, filtered and the solvent
 5 evaporated. The residue was crystallized from CH₃CN, filtered off, then purified by short open column chromatography over silica gel (eluent: CH₂Cl₂/(CH₃OH/NH₃) 90/10). The pure fractions were collected and the solvent was evaporated. Yielding: 11 g of intermediate 18 (78%).(mp. 270.7°C)

10 Example A6

a) Preparation of intermediate 19

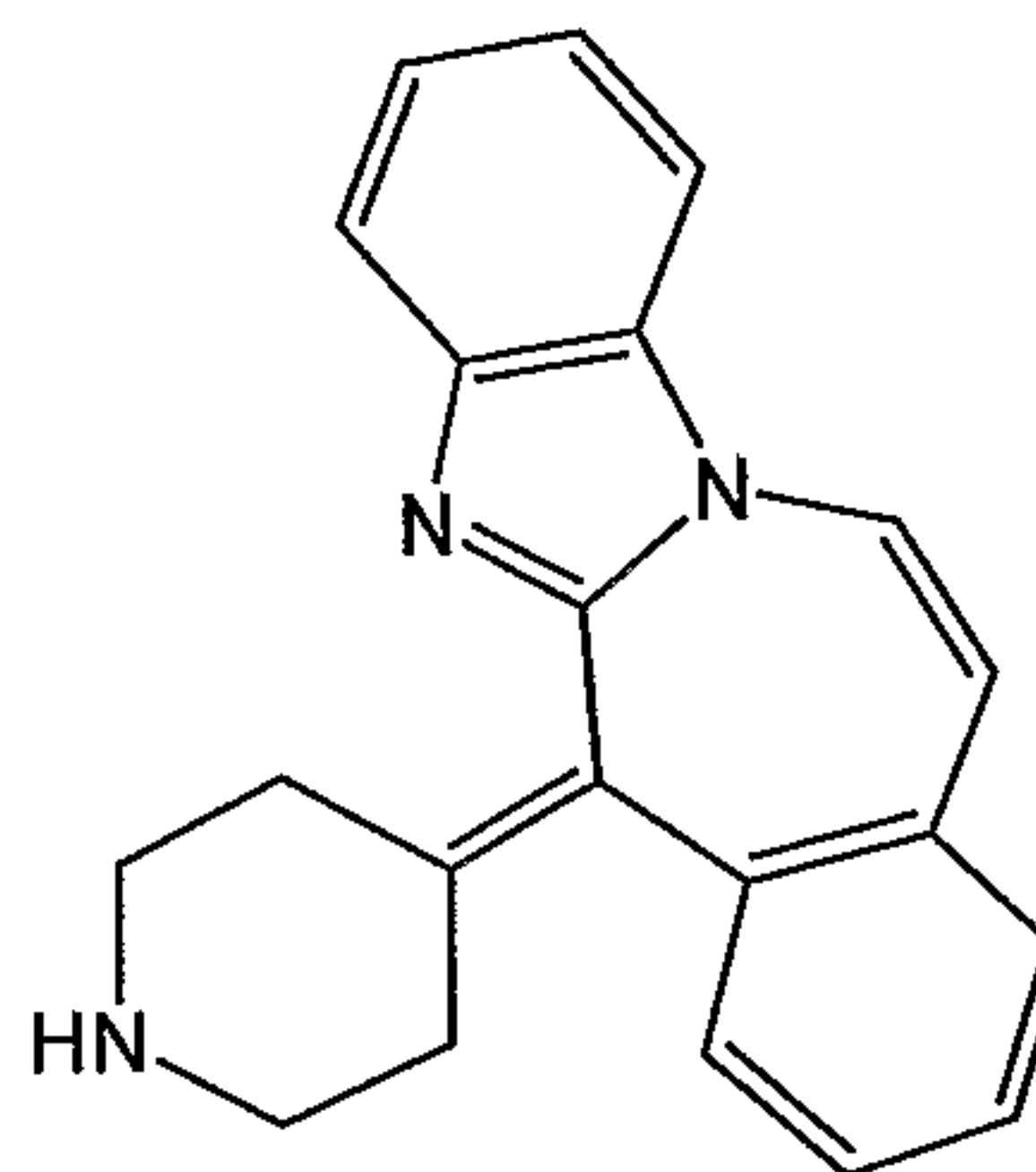


A mixture of 1-(2-phenylethenyl)-1H-benzimidazole (0.04mol) in THF (100ml) was stirred under N₂ flow and cooled to -70°C. BuLi, 2.5 M/hexane (0.04mol) was added dropwise at -70°C and stirring was continued for 30 min at -70°C. A solution of 4-ethyl 1-(1,1-dimethylethyl)-1,4-piperidinedicarboxylate (0.04mol) in THF was added
 15 dropwise and the mixture was stirred for 1h at -70°C. The temperature was allowed to reach room temperature and the mixture was decomposed with water, then extracted with CH₂Cl₂. The separated organic layer was dried (MgSO₄), filtered and the solvent was evaporated. The residue was purified by column chromatography over silica gel (eluent : CH₂Cl₂/CH₃CN 97/3 upgrading to 94/6). Two fractions were collected and the
 20 solvent was evaporated. The second fraction's residue was crystallized from DIPE/CH₃CN. The crystals were filtered off and dried. Yielding: 7.0g of (1,1-dimethylethyl) (Z)-4-[[1-(2-phenylethenyl)-1H-benzimidazol-2-yl]carbonyl]-1-piperidinecarboxylate (41%) (intermediate 19). (mp. 155.8°C)

b) Preparation of intermediate 20

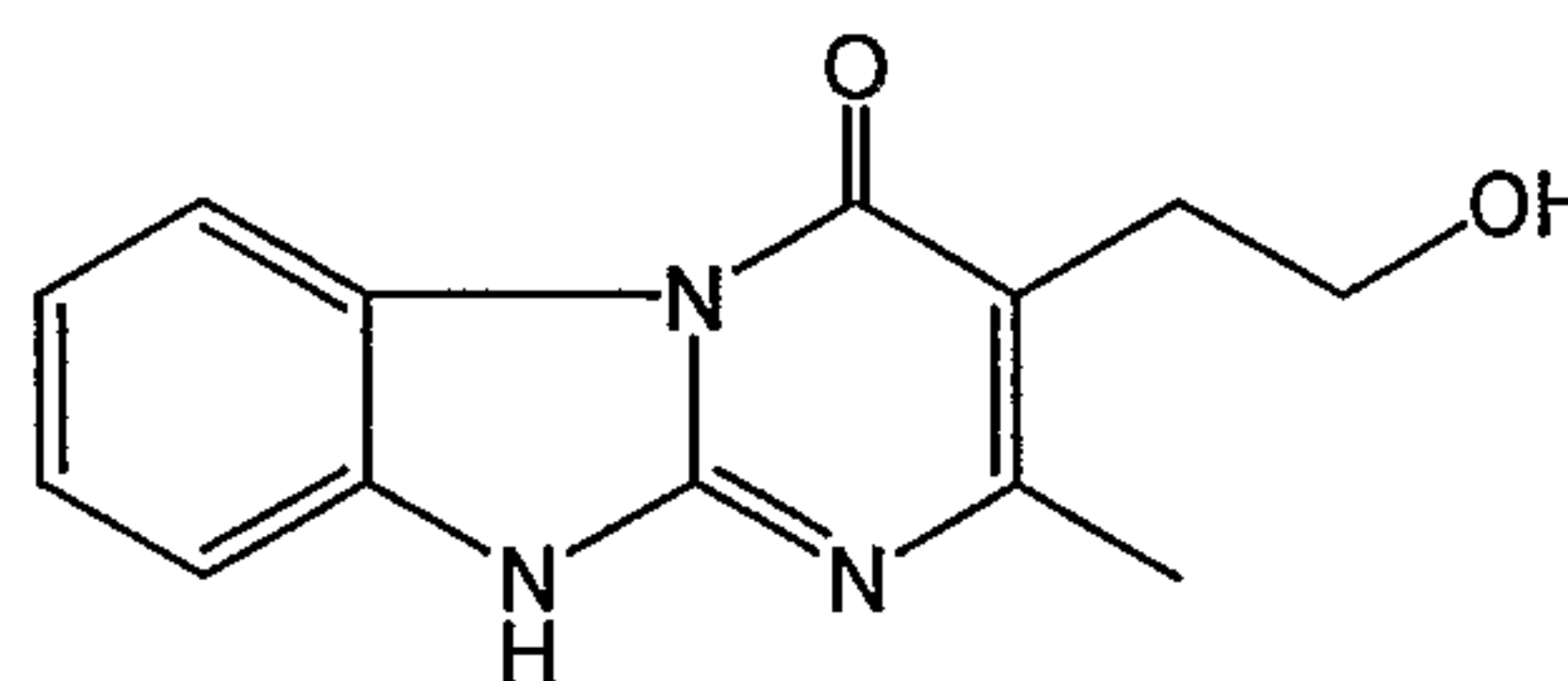
A mixture of intermediate 19 (0.043mol) in trifluoroacetic acid (130ml) was stirred for 1/2 hour at room temperature. The reaction mixture was poured out into diethylether. The precipitate was filtered off, washed with diethylether and dried. Yielding : 18g of (Z)-[1-(2-phenylethenyl)-1H-benzimidazol-2-yl] (4-piperidinyloxy)methanone

5 .trifluoroacetate (1:1) (intermediate 20) (94.0%). (mp. 202.2°C)

c) Preparation of intermediate 21

A mixture of intermediate 20 (0.0276mol), AlCl₃ (0.187mol) and NaCl (0.187mol) was stirred for 1 hour at 150°C (melt). The reaction mixture was decomposed in a mixture of ice, water and NaOH 50%. The mixture was extracted with dichloromethane and the organic layer was separated, dried, filtered and evaporated. The residue (4.3g) was purified on a glass filter over silica gel (eluent: CH₂Cl₂/(CH₃OH/NH₃) 90/10). The pure fractions were collected and the solvent was evaporated. The residue was converted into the (E)-2-butenedioic acid salt (2:3) in ethanol. The salt was filtered off and dried. Yielding : 1.8g of 6-(4-piperidinyloxy)-6H-benzimidazo[2,1-b][3]benzazepine .(E)-2-butenedioate (2:3) (13.4%) (intermediate 21). (mp. 229.4°C)

15

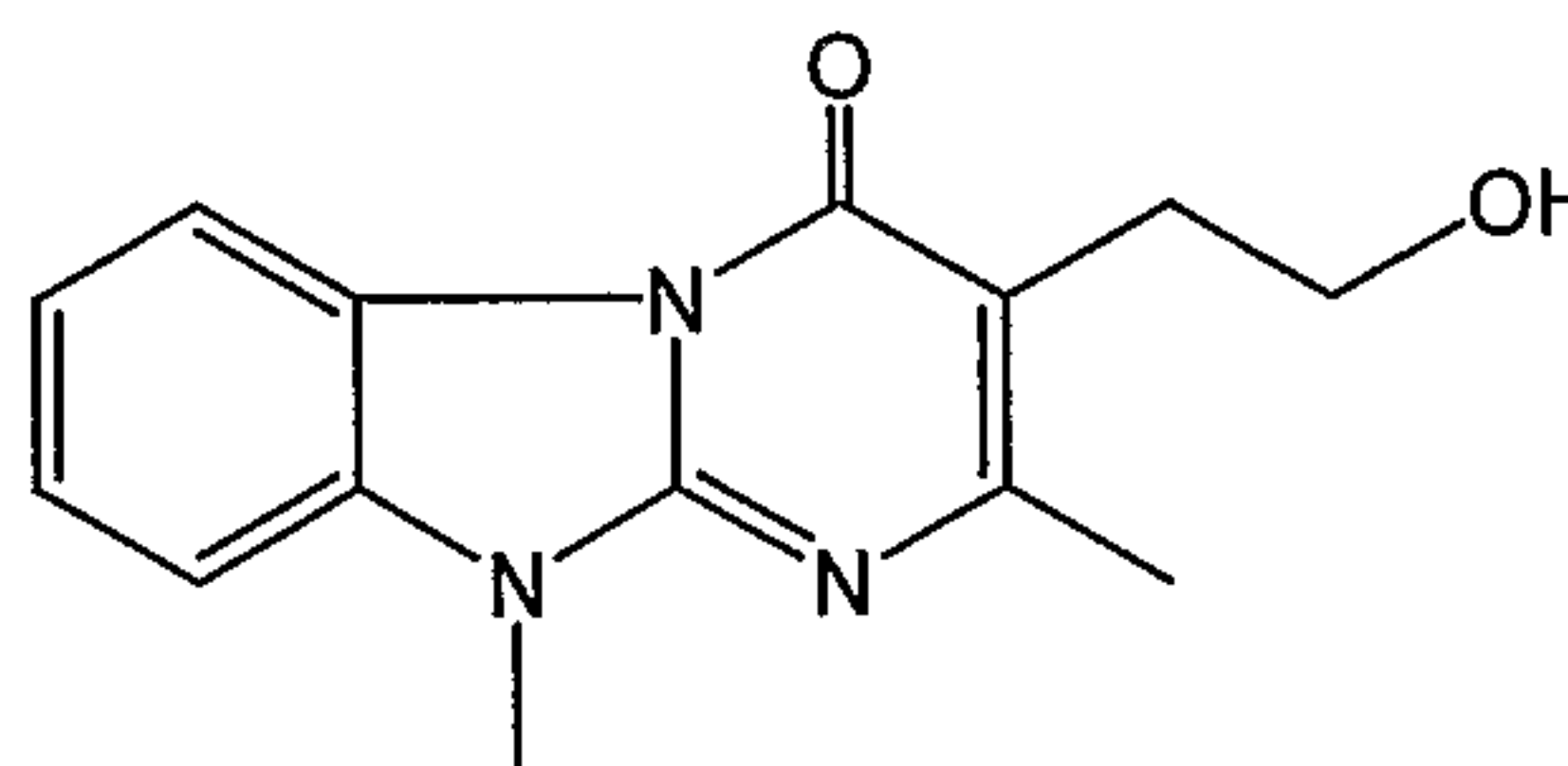
Example A7a) Preparation of intermediate 22

A mixture of 2-amine-1H-benzimidazole (0.04mol), 3-acetyldihydro-2(3H)-furanone (0.53 mol) and 4-methylbenzenesulfonic acid (4 g) in xylene (930ml) was stirred and refluxed overnight and then cooled. The precipitate was filtered off and stirred in H₂O (200ml), Na₂CO₃ (5g) and CH₂Cl₂ (500ml). The precipitate was filtered off, boiled in

20

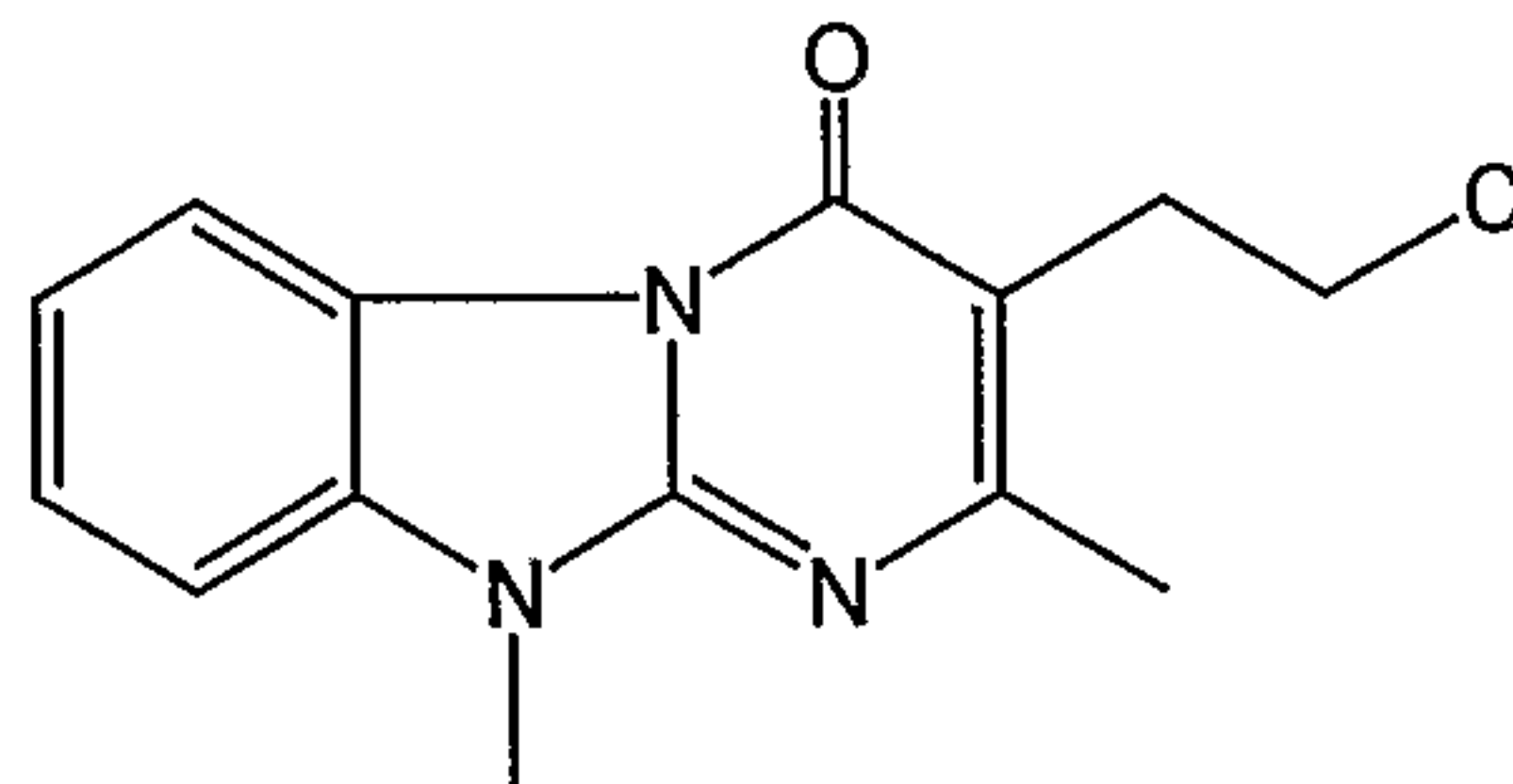
CH₃OH, filtered off and dried. Yielding: 47.4 g of 3-(2-hydroxyethyl)-2-methyl-pyrimido[1,2-a]benzimidazol-4(10*H*)-one (intermediate 22).

b) Preparation of intermediate 23

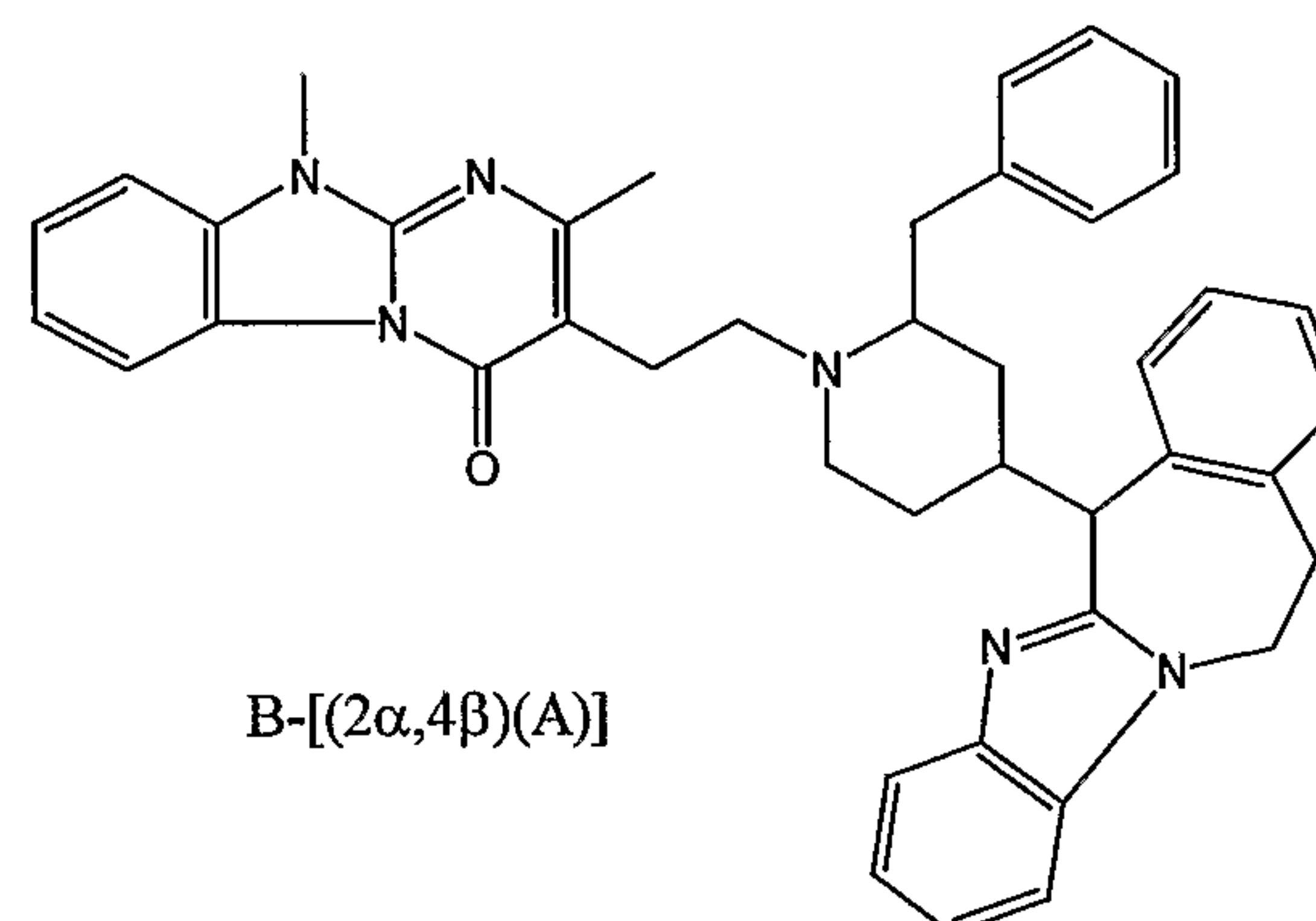


- 5 A mixture of intermediate 22 (0.025 mol) and K₂CO₃ p.a. (0.03mol) in DMF (70 ml) was stirred at 50°C. Methyl iodide (0.03mol) was added dropwise. The mixture was stirred at 50°C for 4 hours and cooled. The solvent was evaporated. The residue was boiled in CH₃OH. The precipitate was filtered off and dried. The residue was purified by HPLC over silica gel (eluent: CH₂Cl₂/(CH₃OH/NH₃) 97/3). Two pure fractions
- 10 were collected and their solvents were evaporated. Yielding: 2.08 g of 3-(2-hydroxyethyl)-2,10-dimethyl-pyrimido[1,2-a]benzimidazol-4(10*H*)-one (intermediate 23).

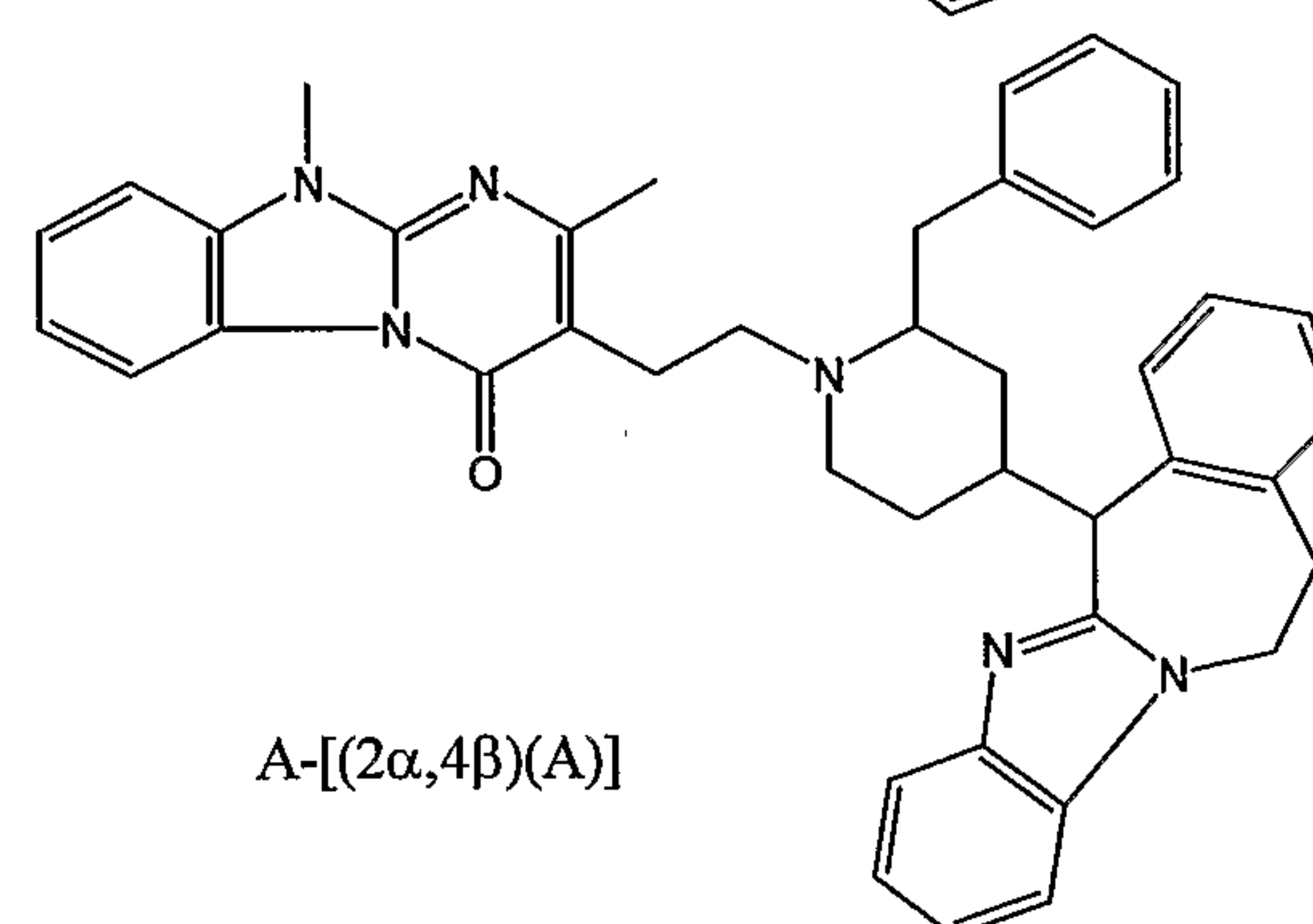
c) Preparation of intermediate 24



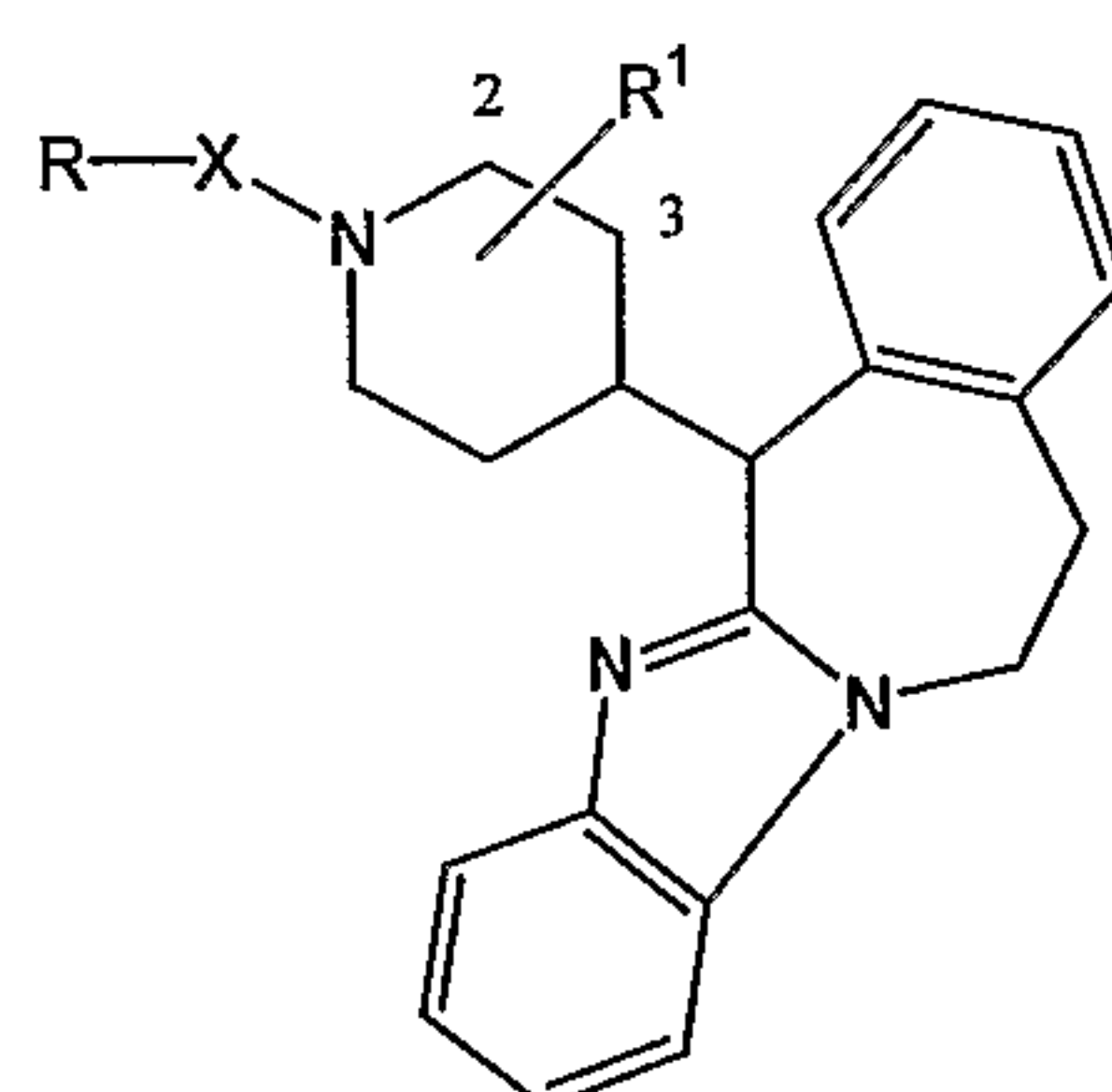
- 15 A mixture of intermediate 23 (0.02 mol) and SOCl₂ (0.06 mol) in CHCl₃ (50 ml) was stirred and refluxed for 4 hours and then cooled. H₂O was added. The mixture was alkalized with K₂CO₃ and separated into its layers. The aqueous layer was extracted with CH₂Cl₂. The combined organic layer was dried (MgSO₄), filtered and the solvent was evaporated. The residue was crystallized from CH₃CN. The precipitate was filtered off and dried. Yielding: 3.44 g of intermediate 24.

B. Preparation of the final compoundsExample B1Preparation of compound 1

B-[(2α,4β)(A)]

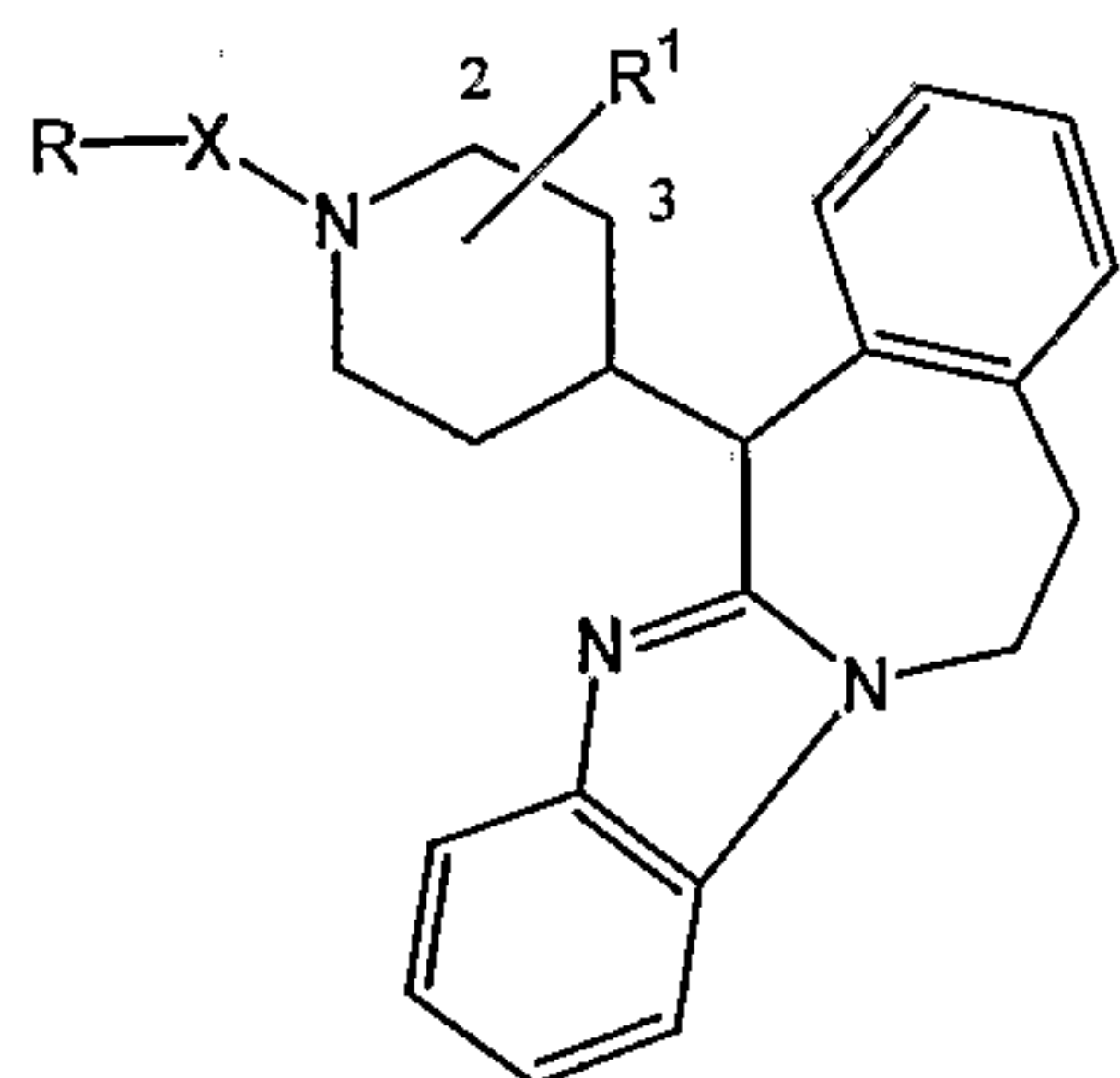
and preparation of compound 2

A-[(2α,4β)(A)]

Table 1 :

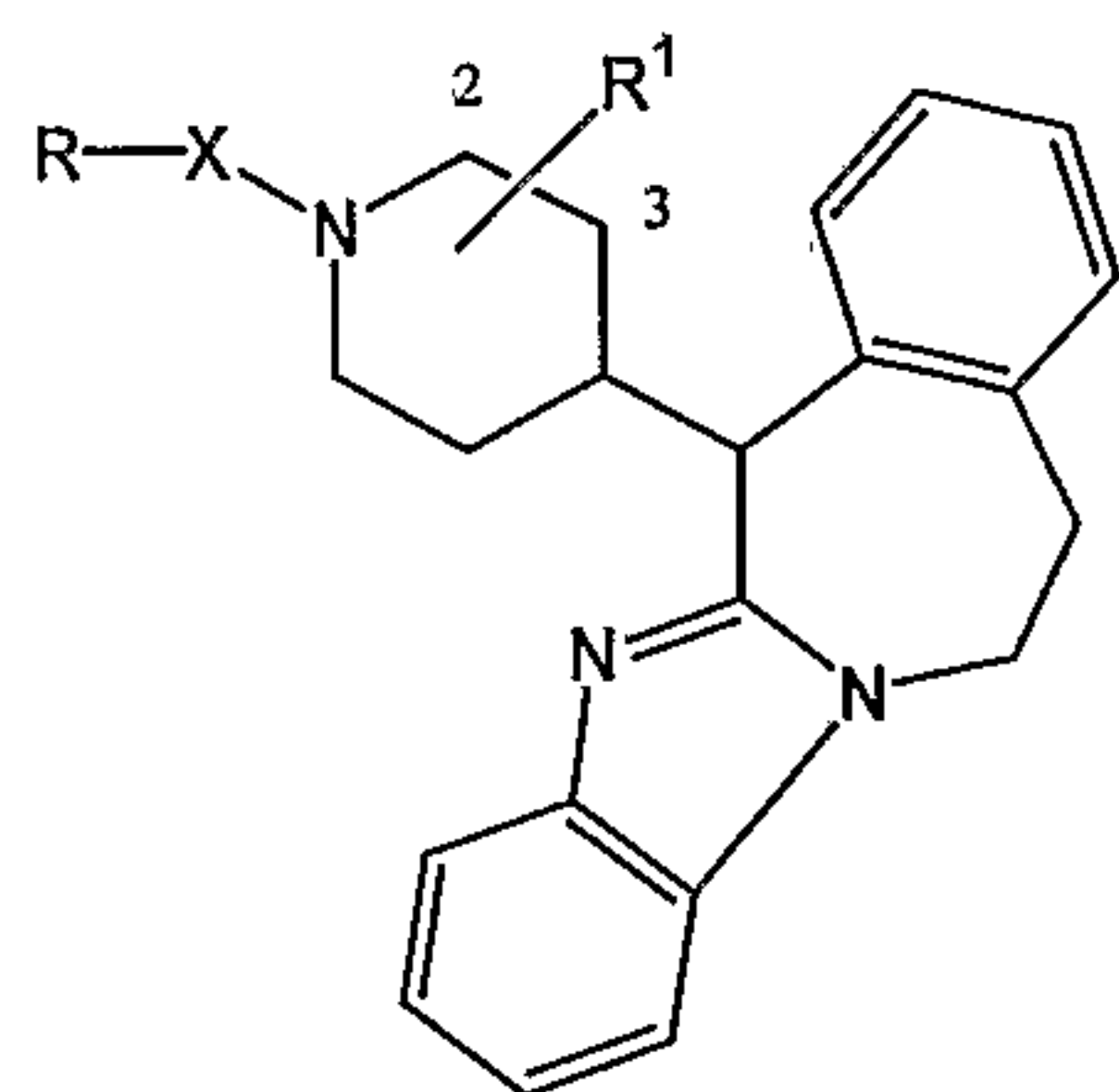
Co. nr.	Ex. nr.	R ¹	R	X	Phys.data and stereochemistry
9	B1	2-benzyl	Chemical structure of R for entry 9: 2-methyl-4-((2-phenyl-1H-imidazo[1,2-a]pyridin-3-yl)propyl)-6-oxo-1,2,3,4-tetrahydropyrimidin-5(1H)-one.	-CH ₂ -CH ₂ -	[(2a,4a)(B)]
10	B1	2-benzyl	Chemical structure of R for entry 10: 2-methyl-4-((2-phenyl-1H-imidazo[1,2-a]pyridin-3-yl)propyl)-6-oxo-1,2,3,4-tetrahydropyrimidin-5(1H)-one, with a sulfur atom at the 2-position of the imidazo[1,2-a]pyridine moiety.	-CH ₂ -CH ₂ -	[(2a,4a)(B)]

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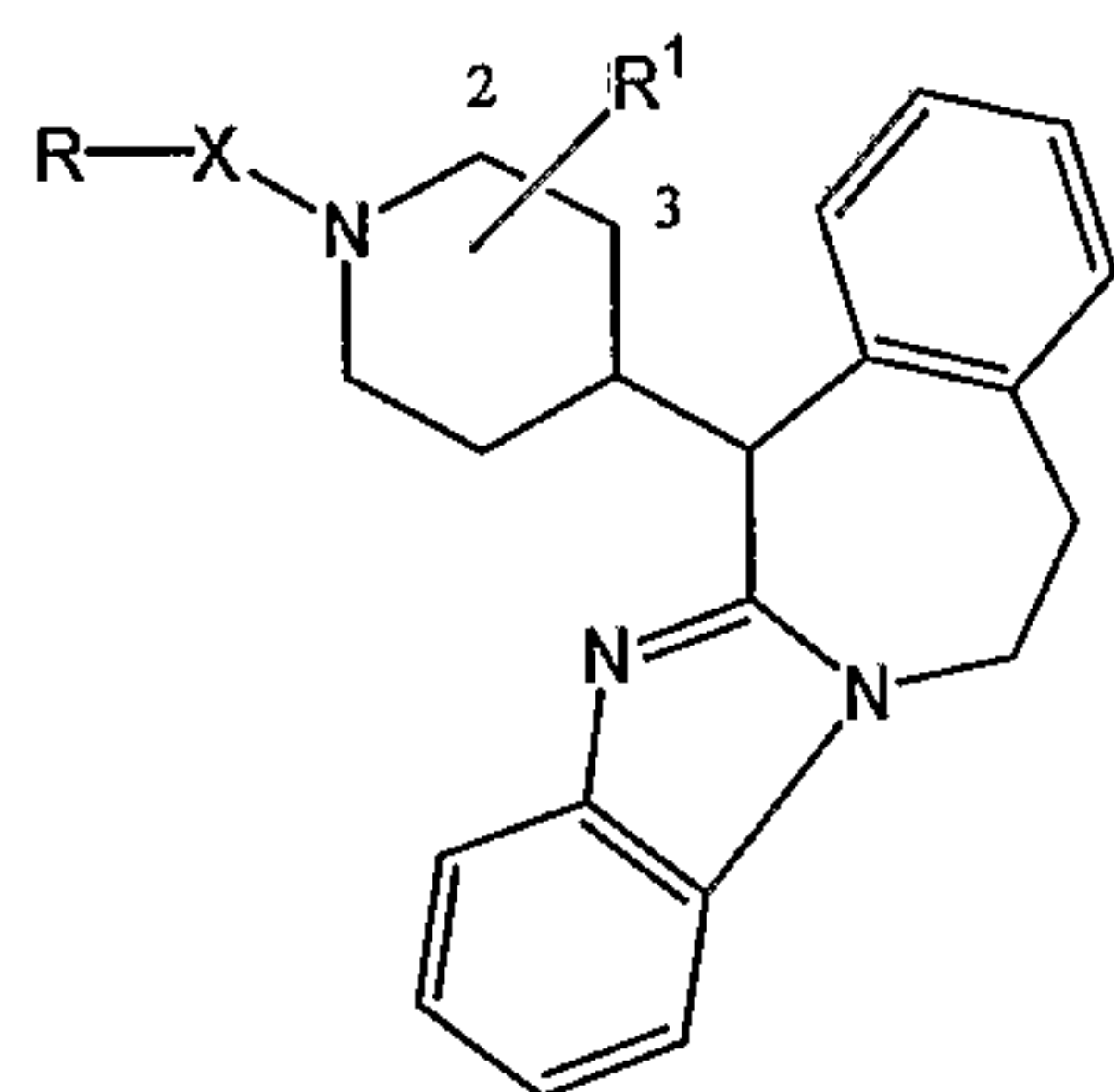


Co. nr.	Ex. nr.	R ¹	R	X	Phys.data and stereochemistry
11	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4a)(B)]; .H ₂ O(1:2)
12	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4a)(B)]
13	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4β)(B)]
14	B1	H		-CH ₂ -CH ₂ -	
15	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4β)(A)]; (E)- 2-butenedioate (2:3)
16	B1	2-benzyl		-CH ₂ -CH ₂ -	[A(2a,4α)(B)]
17	B1	2-benzyl		-CH ₂ -CH ₂ -	[B(2a,4α)(B)]

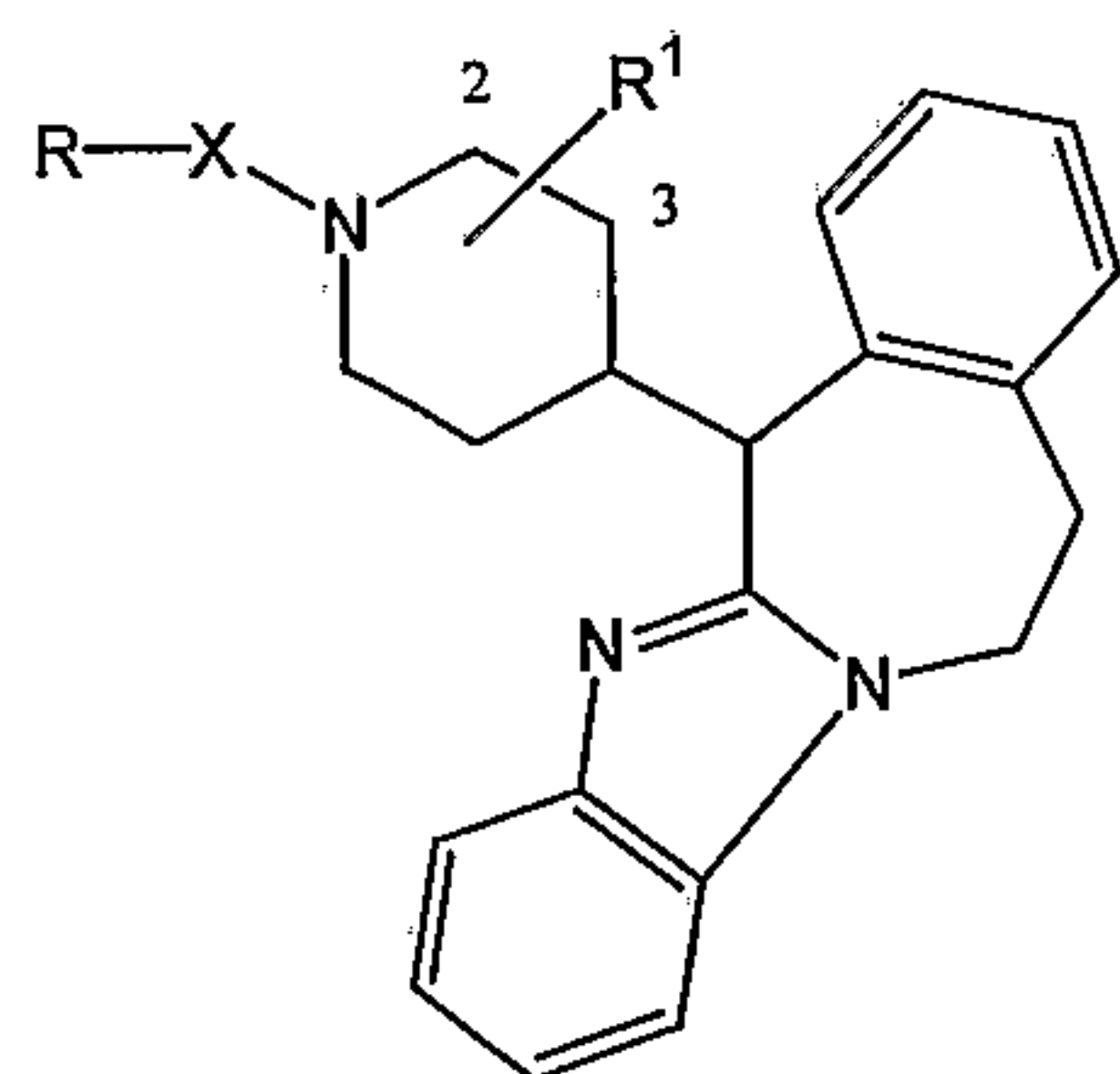
-31-



Co. nr.	Ex. nr.	R ¹	R	X	Phys.data and stereochemistry
18	B1	2-methyl cyclohexy		-CH ₂ -CH ₂ -	
19	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
20	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
21	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
22	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
23	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]

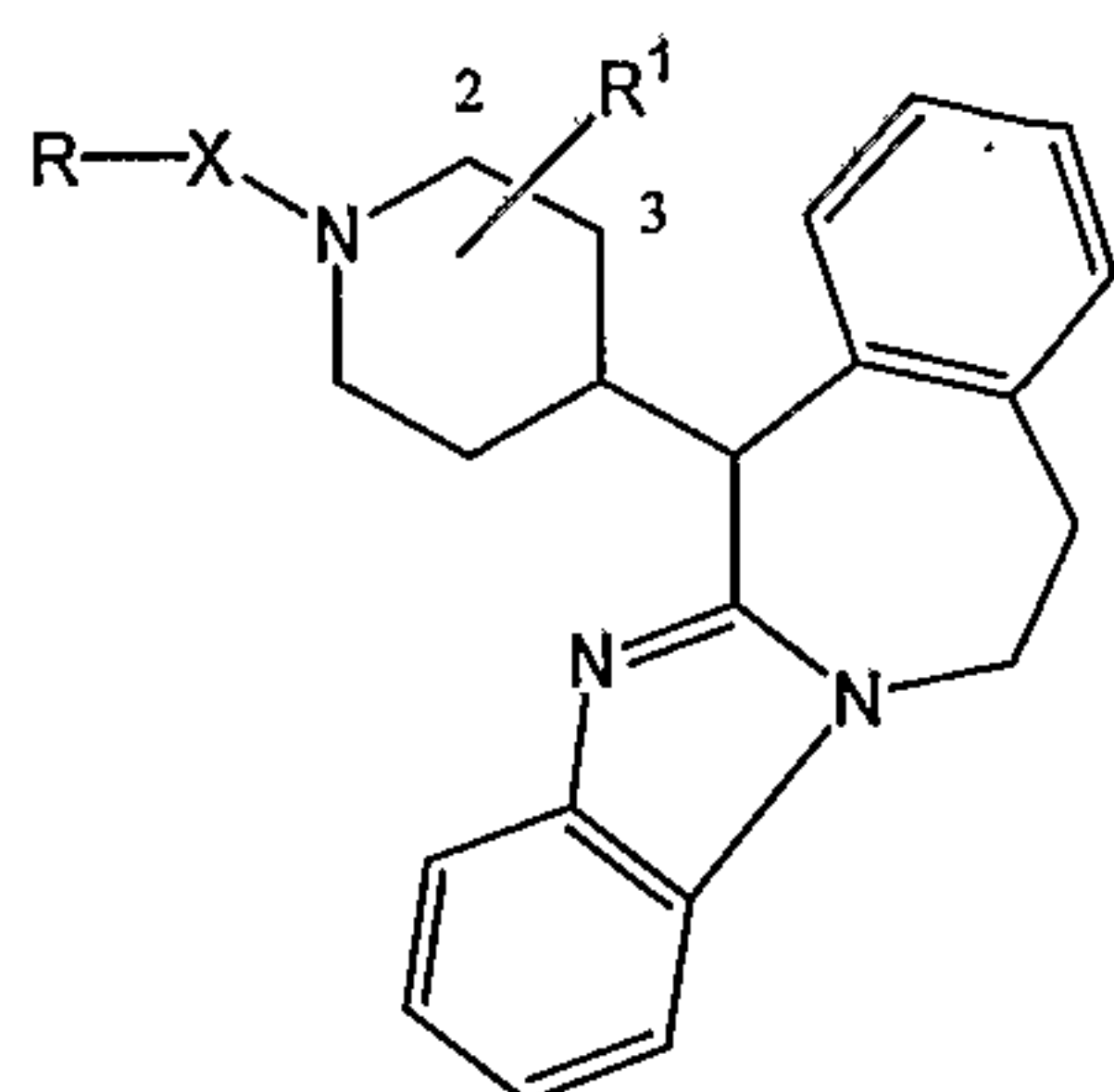


Co. nr.	Ex. nr.	R ¹	R	X	Phys.data and stereochemistry
24	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
25	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
26	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]; .H ₂ O(1:1)
27	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
28	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]



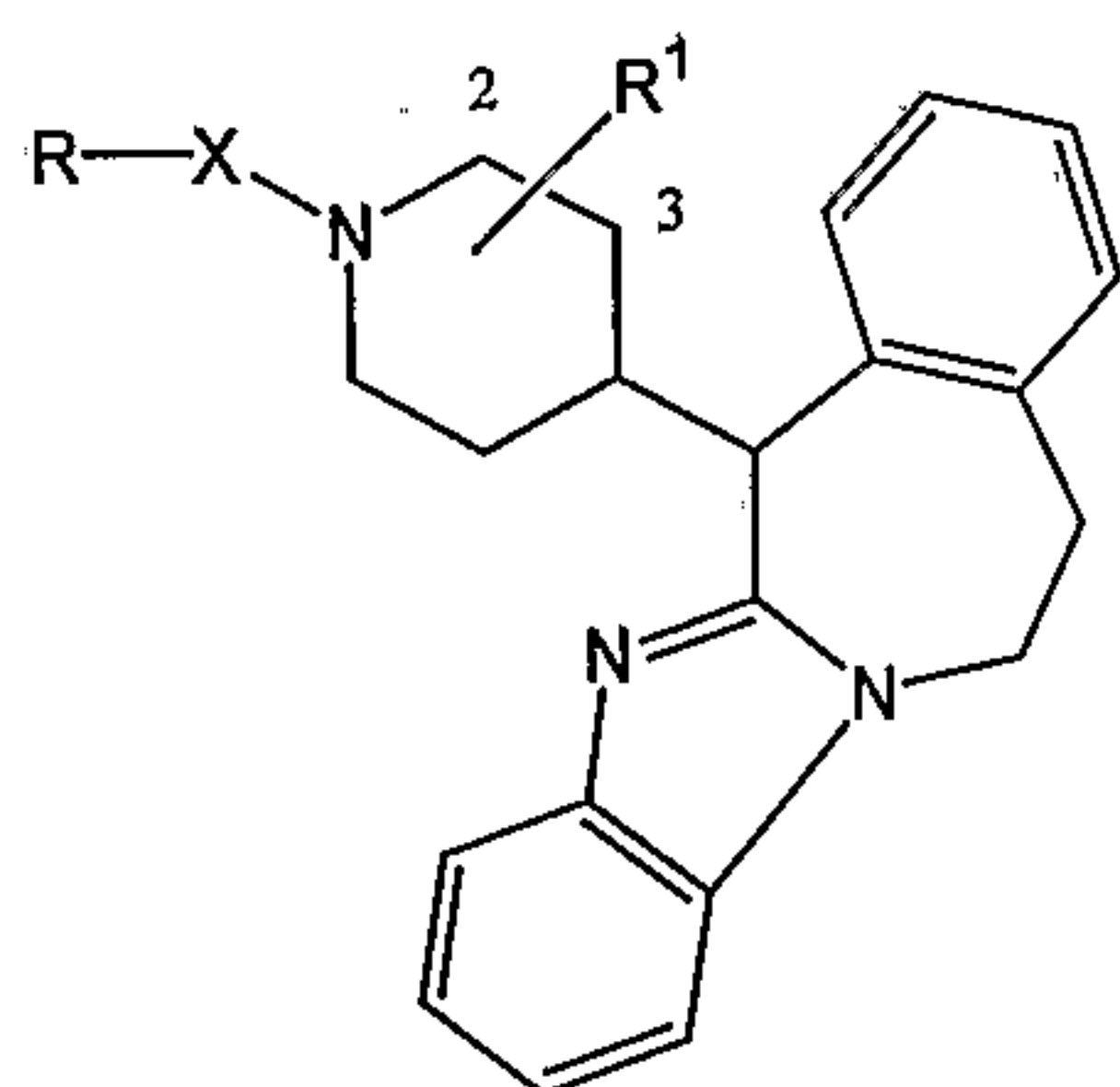
Co. nr.	Ex. nr.	R ¹	R	X	Phys.data and stereochemistry
29	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
30	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(A)]
31	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
32	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
33	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]

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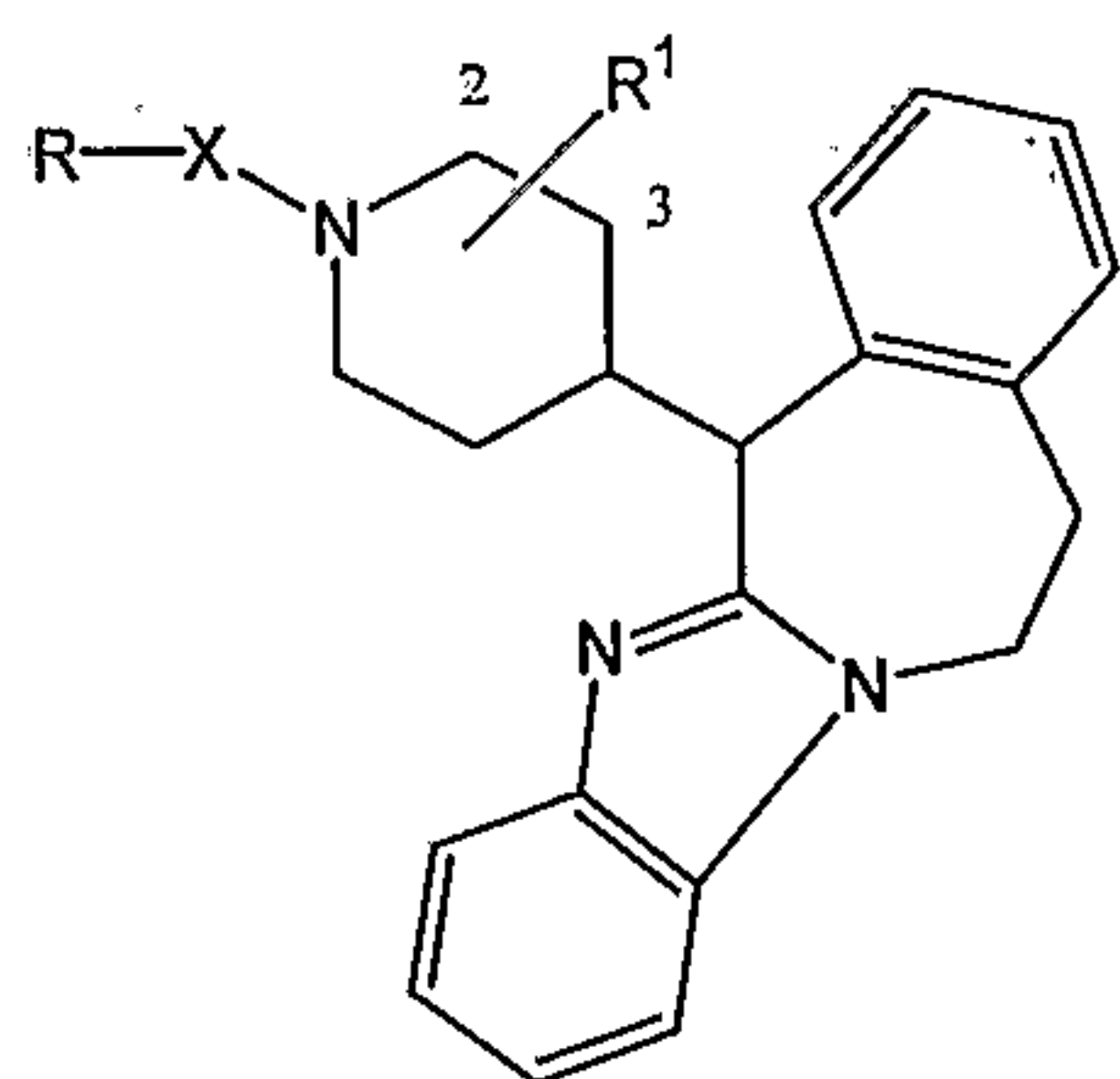
Co. nr.	Ex. nr.	R ¹	R	X	Phys.data and stereochemistry
34	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]; .H ₂ O(1:1)
35	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]; .H ₂ O(1:1)
36	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]; .H ₂ O(1:1)
37	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
38	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]; .H ₂ O(1:1)
39	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]; .H ₂ O(1:1)

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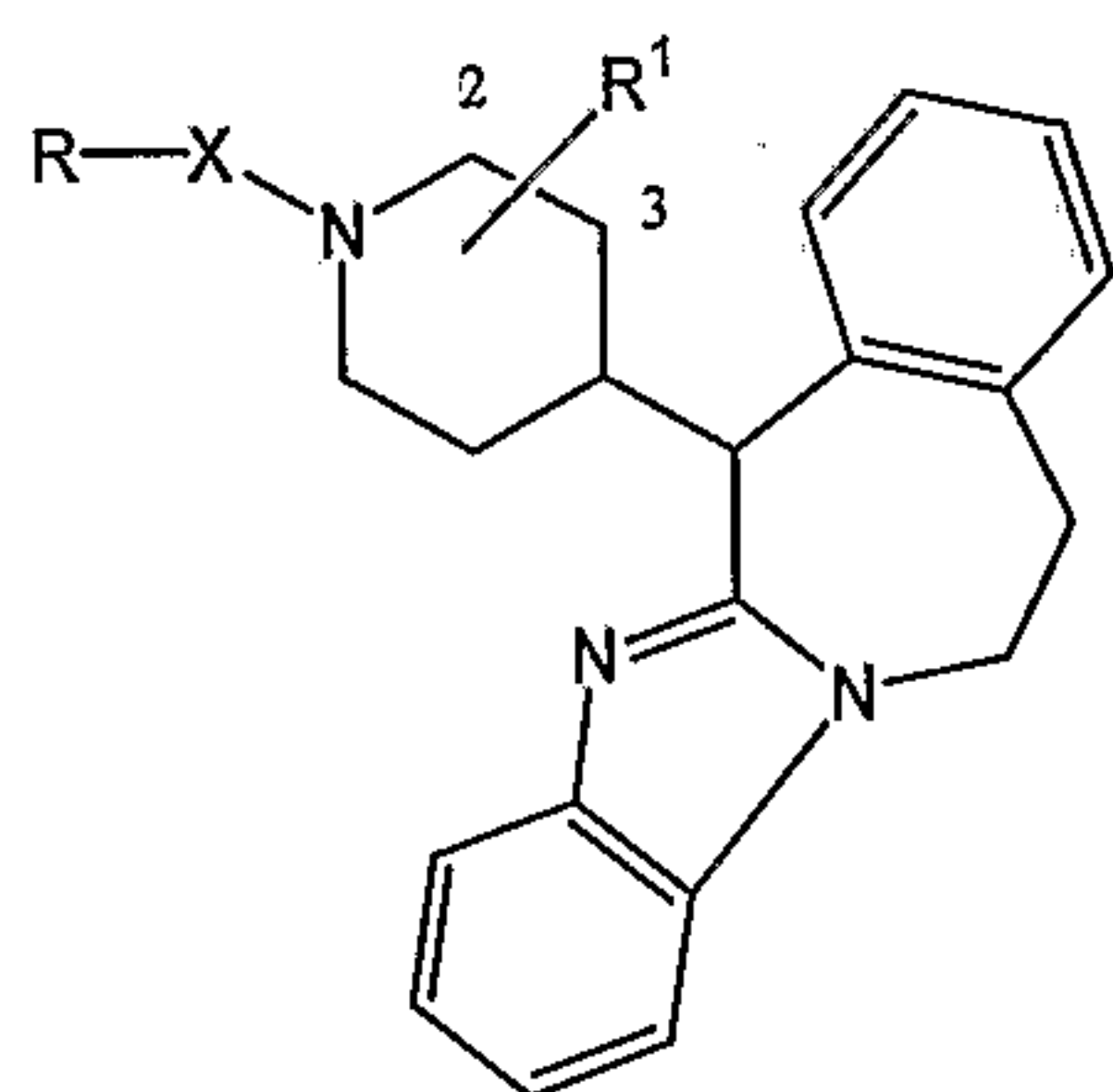


Co. nr.	Ex. nr.	R ¹	R	X	Phys.data and stereochemistry
40	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
41	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
42	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
43	B1	2-benzyl		[CH ₂] ₂ -C(=O)	[(2a,4α)(A)]
44	B1	2-benzyl		-CH ₂ -CH ₂ -	[A(2a,4β)(A)]
45	B1	2-benzyl		-CH ₂ -CH ₂ -	[B(2a,4β)(A)]

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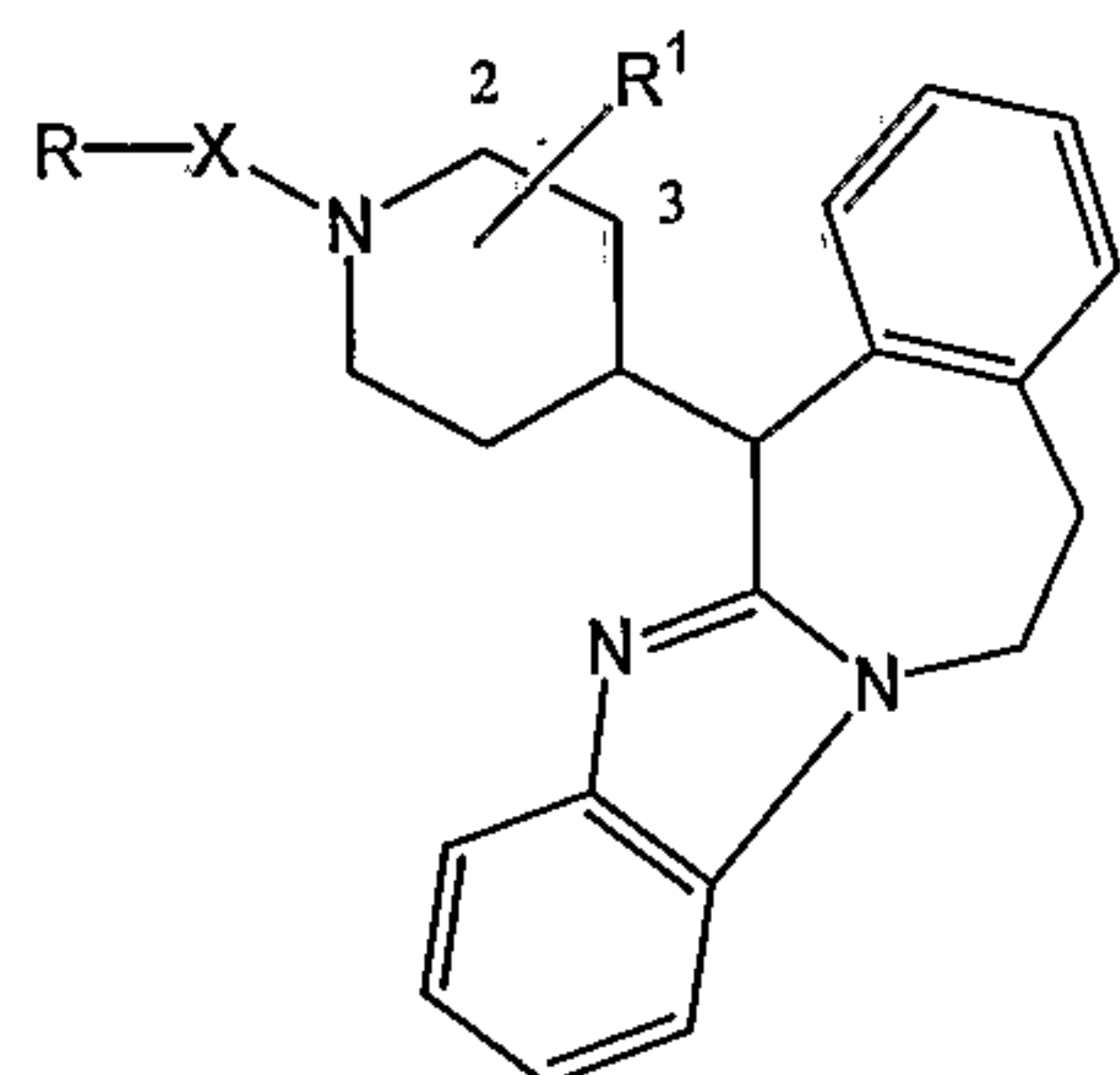


Co. nr.	Ex. nr.	R ¹	R	X	Phys.data and stereochemistry
46	B1	2-benzyl		-CH ₂ -CH ₂ -	[A(2a,4β)(B)]; Tri-fluoroacetate(1:1)
47	B1	2-benzyl		-CH ₂ -CH ₂ -	[B(2a,4β)(B)]; Tri-fluoroacetate(1:1)
48	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4β)(A)]
49	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4β)(A)]; (-)-[S(R*,R*)]- 2,3-dihydroxy butanedioate (1:2)
50	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4β)(A)]; .HCl(1:3).H ₂ O (1:2)
51	B1	2-benzyl		-CH ₂ -CH ₂ -CH ₃	[(2a,4β)(A)]; .H ₂ O(1:2)
52	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4β)(A)]



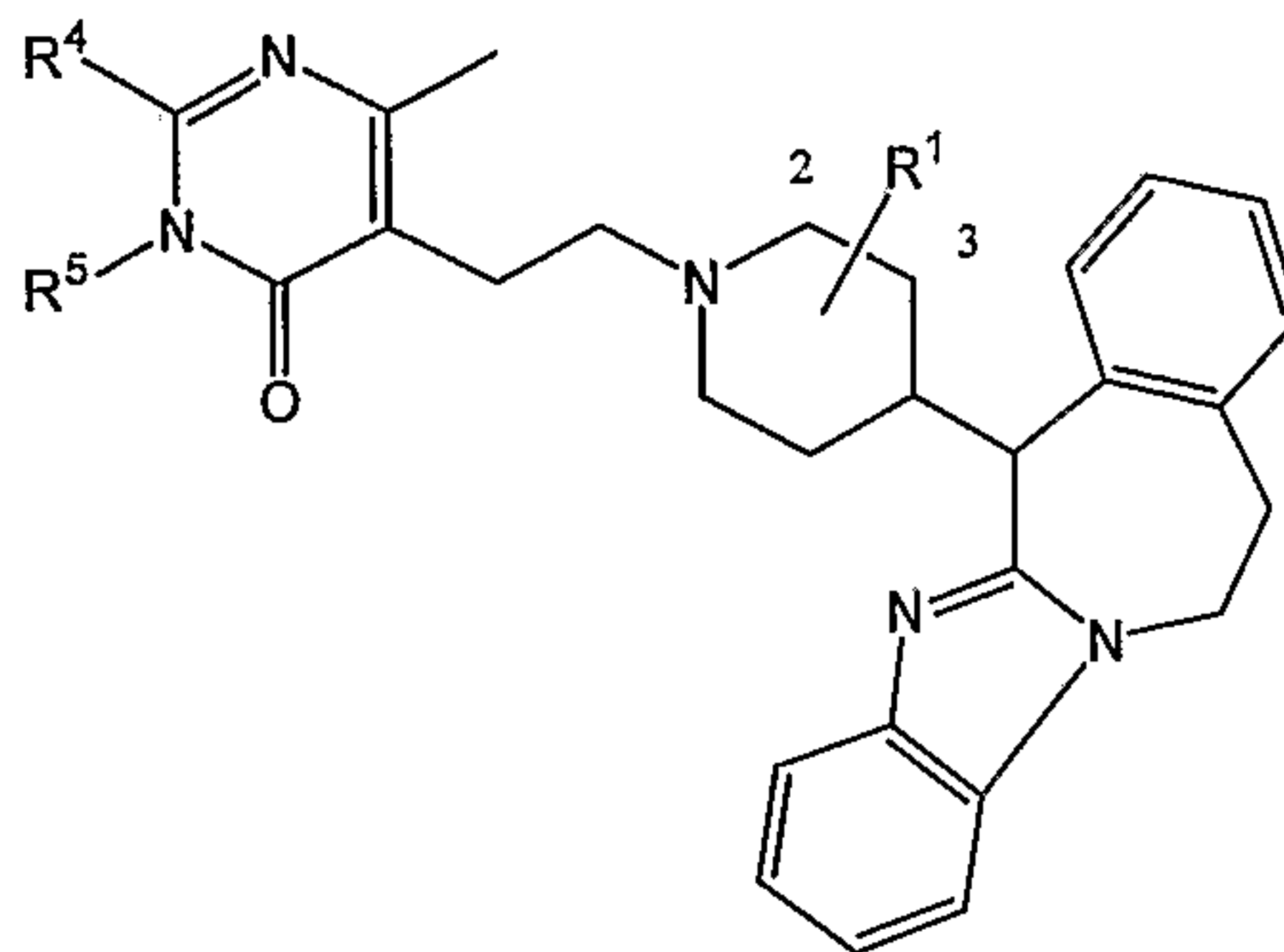
Co. nr.	Ex. nr.	R ¹	R	X	Phys.data and stereochemistry
53	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4β)(A)]
54	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4β)(A)]; (E)-2-butenedioate (1:1)
55	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4β)(A)]; (E)-2-butenedioate (1:1).H ₂ O(1:2)
2	B1	2-benzyl		-CH ₂ -CH ₂ -	[A(2a,4β)(A)]; (E)-2-butenedioate (2:3).H ₂ O(1:1)
1	B1	2-benzyl		-CH ₂ -CH ₂ -	[B(2a,4β)(A)]; (E)-2-butenedioate (2:3).H ₂ O(1:1)
56	B1	2-benzyl		CH ₂ -CH ₂ -CH	[(2a,4α)(B)]
57	B1	2-benzyl		-CH ₂ -CH ₂ -	[A(2a,4α)(A)]

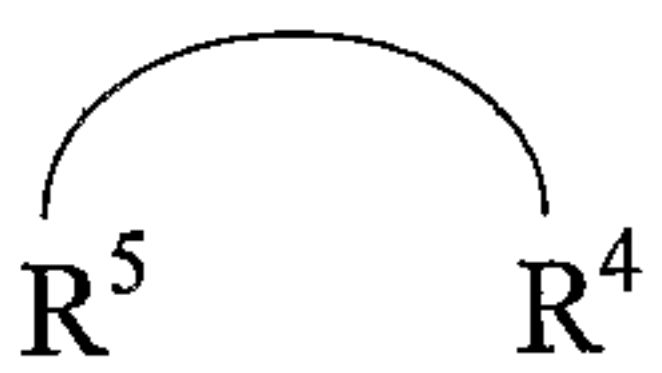
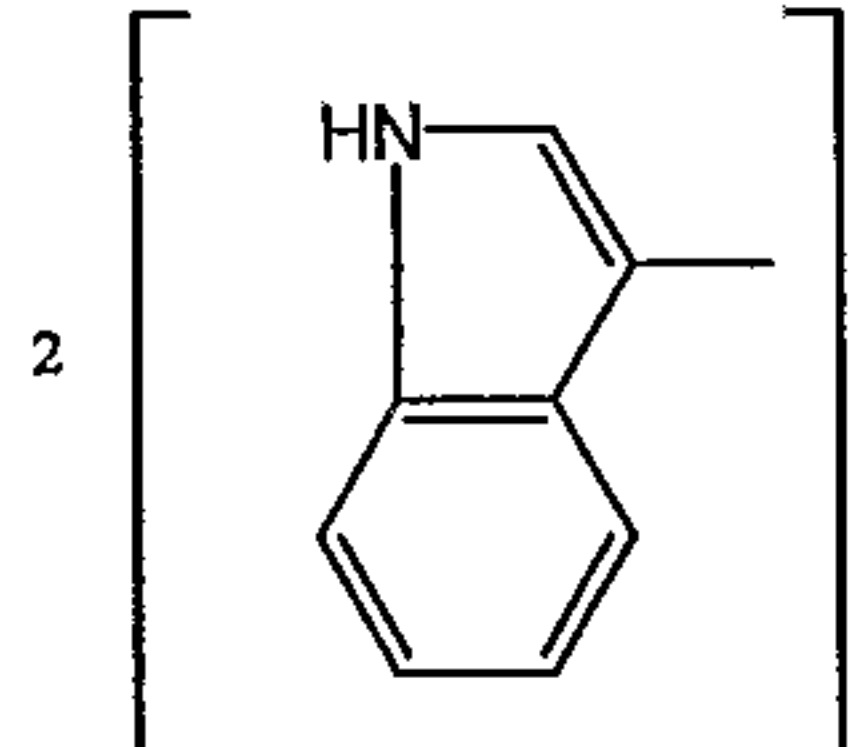
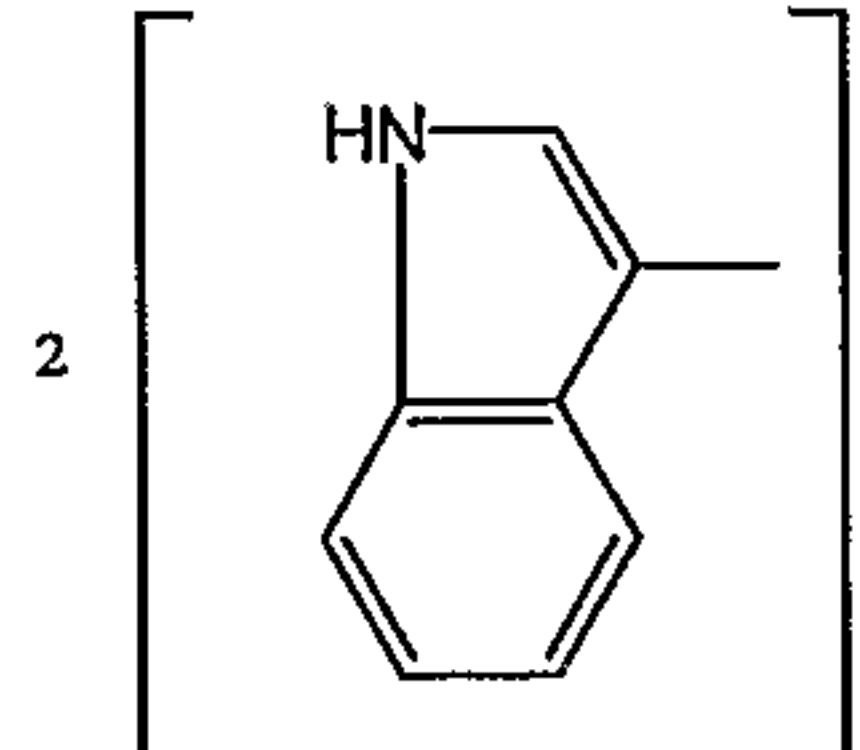
-38-



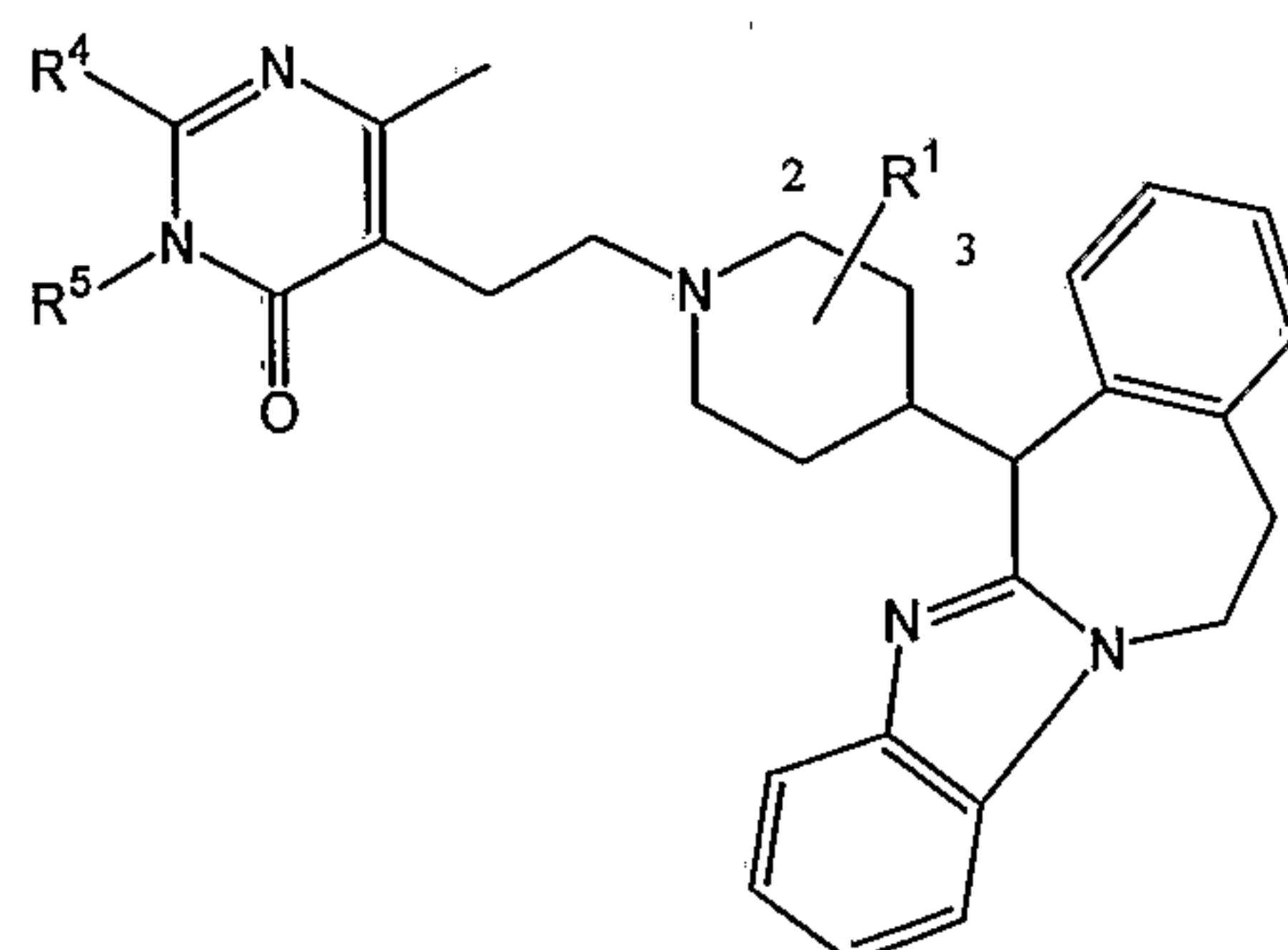
Co. nr.	Ex. nr.	R ¹	R	X	Phys.data and stereochemistry
58	B1	2-benzyl		-CH ₂ -CH ₂ -	[B(2a,4α)(A)]
59	B1	2-benzyl			[(2a,4α)(A)]
60	B1	2-benzyl		-CH ₂ -CH ₂ -	[(2a,4α)(B)]
109	B1	2-benzyl		-CH ₂ -	[(2α,4α)(B)]
152	B1	H		-CH ₂ -CH ₂ -	
153	B1	H		CH ₂ -CH ₂ -CH	(E)-2- Butenedioate (2:5)

Table 2 :



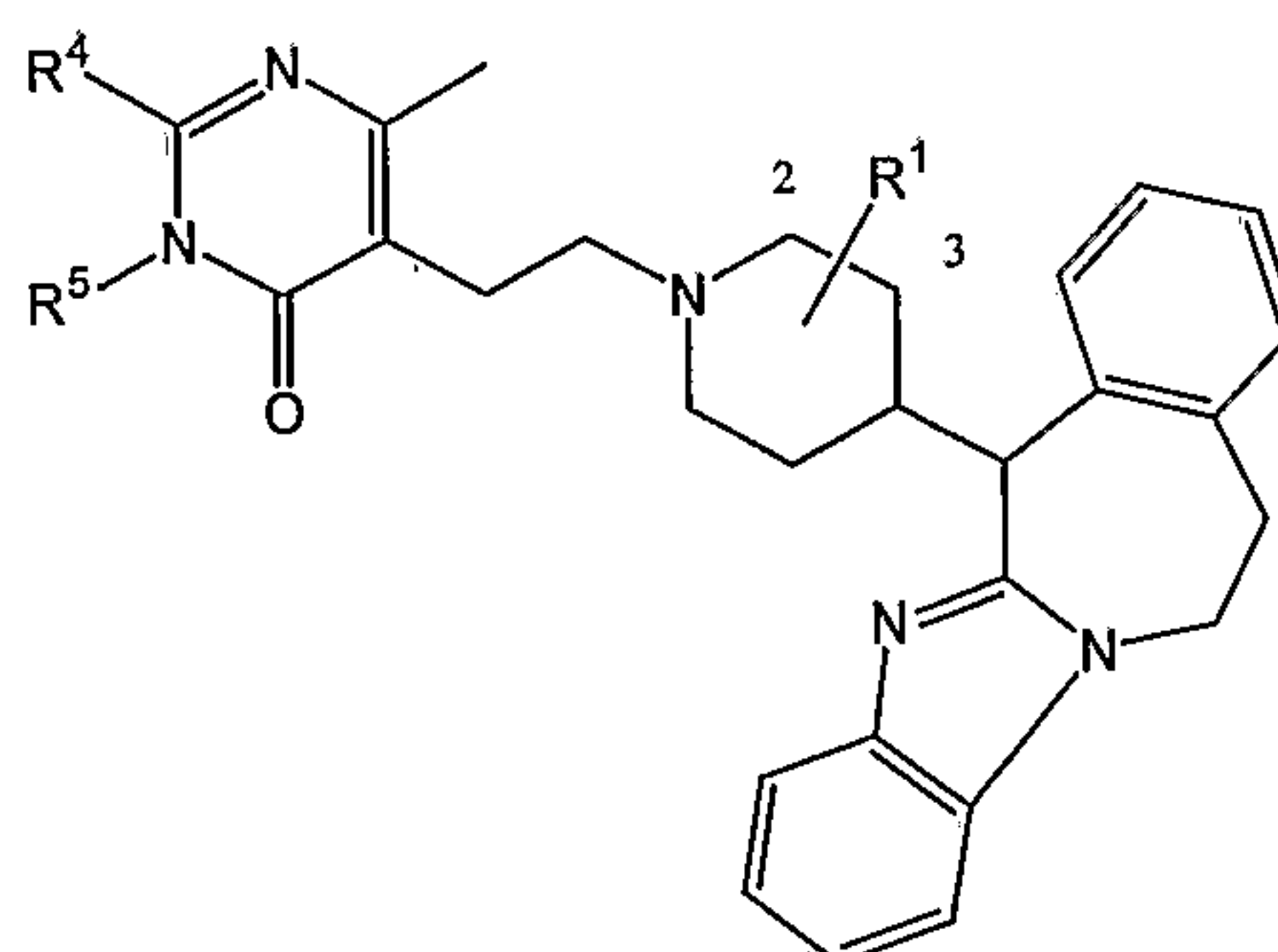
Comp.nr	Ex. nr.	R ¹		Phys.data and stereochemistry
61	B2	H	-CH=CH-S-	
62	B2	H	-CH ₂ -CH ₂ -S-	
63	B2	H	-CH ₂ -CH ₂ -CH ₂ -S-	
64	B2	H	-CH=CH-CH=CH-	
65	B2	H	-CH ₂ -C(CH ₃)=N-N(CH ₃)-	
66	B2	H	-C(CH ₃)=N-N(CH ₃)-	
67	B2	H	-CH=CH-N(CH ₃)-	
68	B2	H	-O-C(CH ₃)=CH-	
69	B2	H	-CH=C(CH ₃)-N(CH ₃)-	
70	B2	H	-CH=C(CH ₃)-CH=CH-	
71	B2	H	-C(CH ₃)=CH-S-	
72	B2	H	-CH=CH-CH=C(CH ₃)-	
73	B2	2-benzyl	-CH=CH-S-	[(2 α ,4 β)(B)]
74	B2	2-benzyl	-CH=CH-S-	[(2 α ,4 α)(A)]
75	B2	2-benzyl	-CH=CH-S-	[(2 α ,4 α)(B)]
76	B2	2-benzyl	-CH=CH-S-	[(2 α ,4 β)(A)]; (E)-2-butenedioate (1:2)ethanolate(1:1)
77	B2		-CH=CH-S-	[(2 α ,4 α)(A)]
78	B2		-CH=CH-S-	[(2 α ,4 β)(B)]

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Comp.nr	Ex. nr.	R ¹	R ⁵ R ⁴	Phys.data and stereochemistry
79	B2		-CH=CH-S-	[(2 α ,4 α)(B)]
80	B2	2-benzyl	-CH=CH-CH=CH-	[(2 α ,4 α)(B)]
81	B2	2-benzyl	-CH=CH-CH=CH-	[(2 α ,4 β)(A)]
82	B2	2-benzyl	-CH=CH-CH=CH-	[(2 α ,4 β)(B)]
83	B2	2-methylnaphthyl	-CH=CH-S-	[(2 α ,4 β)(A)]
84	B2	2-methylnaphthyl	-CH=CH-S-	[(2 α ,4 β)(B)]
85	B2	2-methylnaphthyl	-CH=CH-S-	[(2 α ,4 α)(B)]
86	B2	2-methylnaphthyl	-CH=CH-S-	[(2 α ,4 α)(A)];.H ₂ O (1:1)ethanolate(1:1)
87	B2	3-methyl	-CH=CH-S-	A-trans
88	B2	3-methyl	-CH=CH-S-	B-trans
89	B2	3-methyl	-CH=CH-CH=CH-	[(3 α ,4 β)(B)]
90	B2	3-methyl-(4-fluorophenyl)	-CH=CH-S-	[(2 α ,4 β)(A)]
91	B2	3-methyl-(4-fluorophenyl)	-CH=CH-S-	[(2 α ,4 β)(B)]
92	B2	3-methyl-(4-fluorophenyl)	-CH=CH-S-	[(2 α ,4 α)(A)]
93	B2	3-methyl-(4-fluorophenyl)	-CH=CH-S-	[(2 α ,4 α)(B)]
94	B2	3-methyl	-CH=CH-CH=CH-	[(3 α ,4 β)(A)]
95	B2	2-benzyl	-CH=C(CH ₃)-N(CH ₃)-	[(2 α ,4 α)(B)]
96	B2	2-benzyl	-CH=CH-N(CH ₃)-	[(2 α ,4 α)(B)]
97	B2	2-benzyl	-CH=CH-CH=C(CH ₃)-	[(2 α ,4 α)(B)]
98	B2	2-benzyl	-CH ₂ -CH ₂ -S-	[(2 α ,4 α)(B)]

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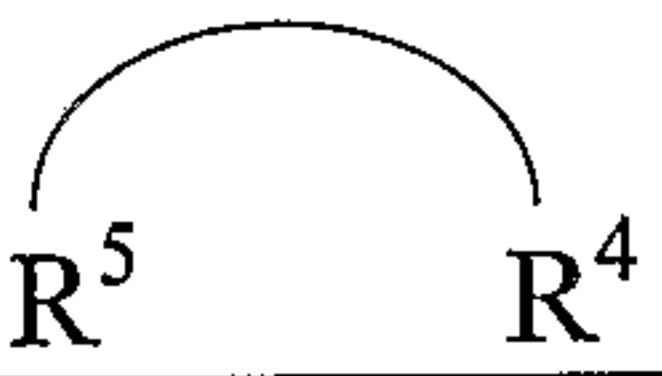
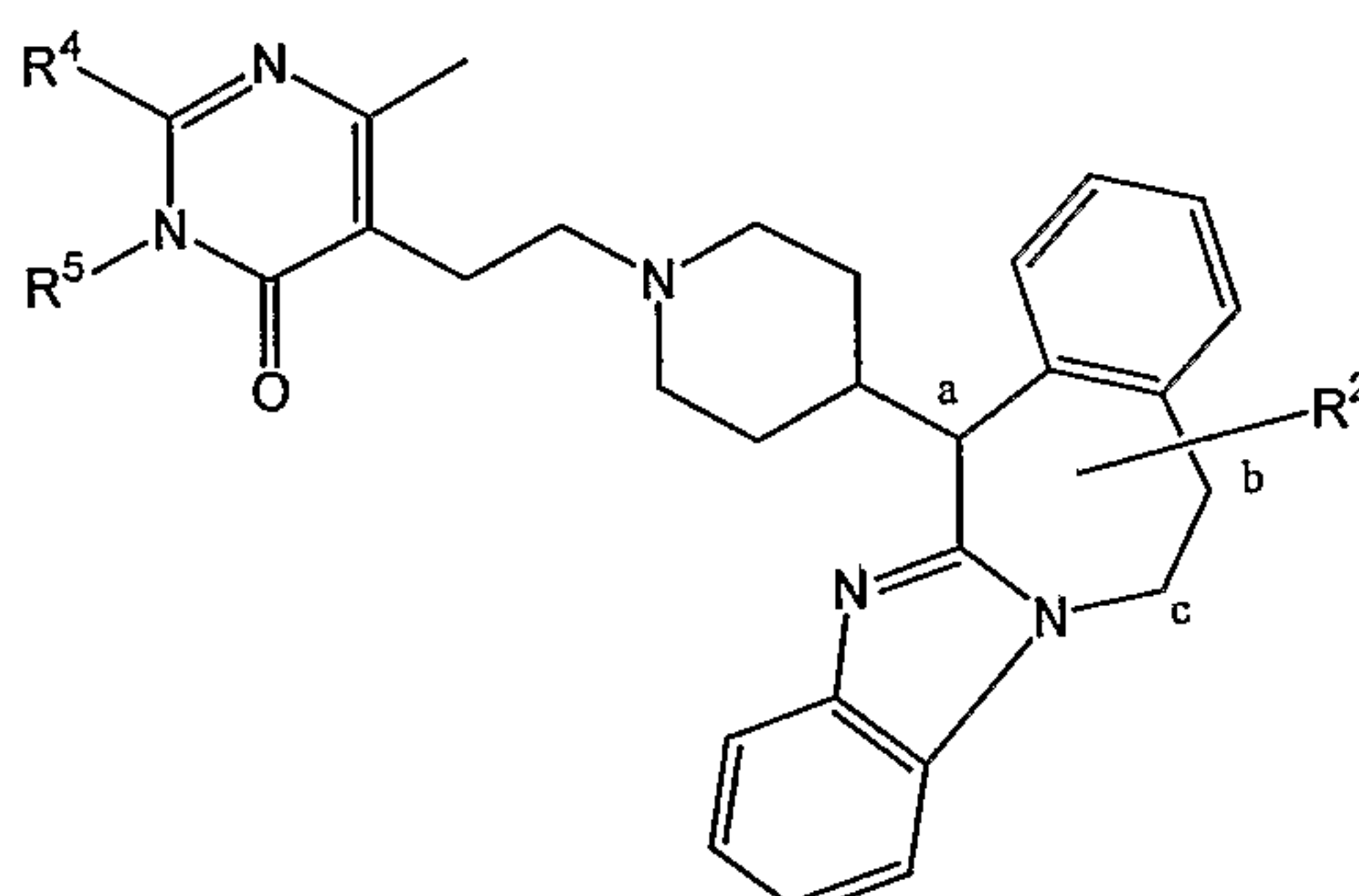
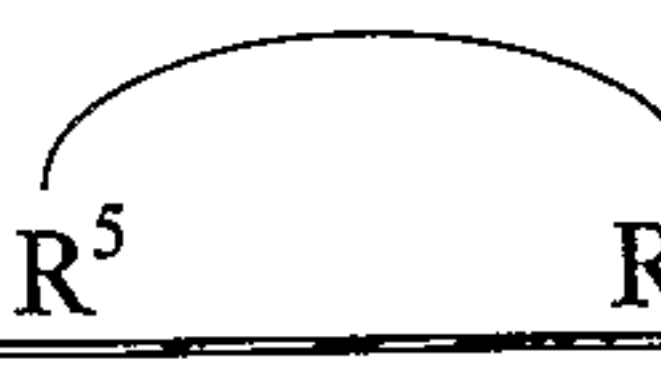
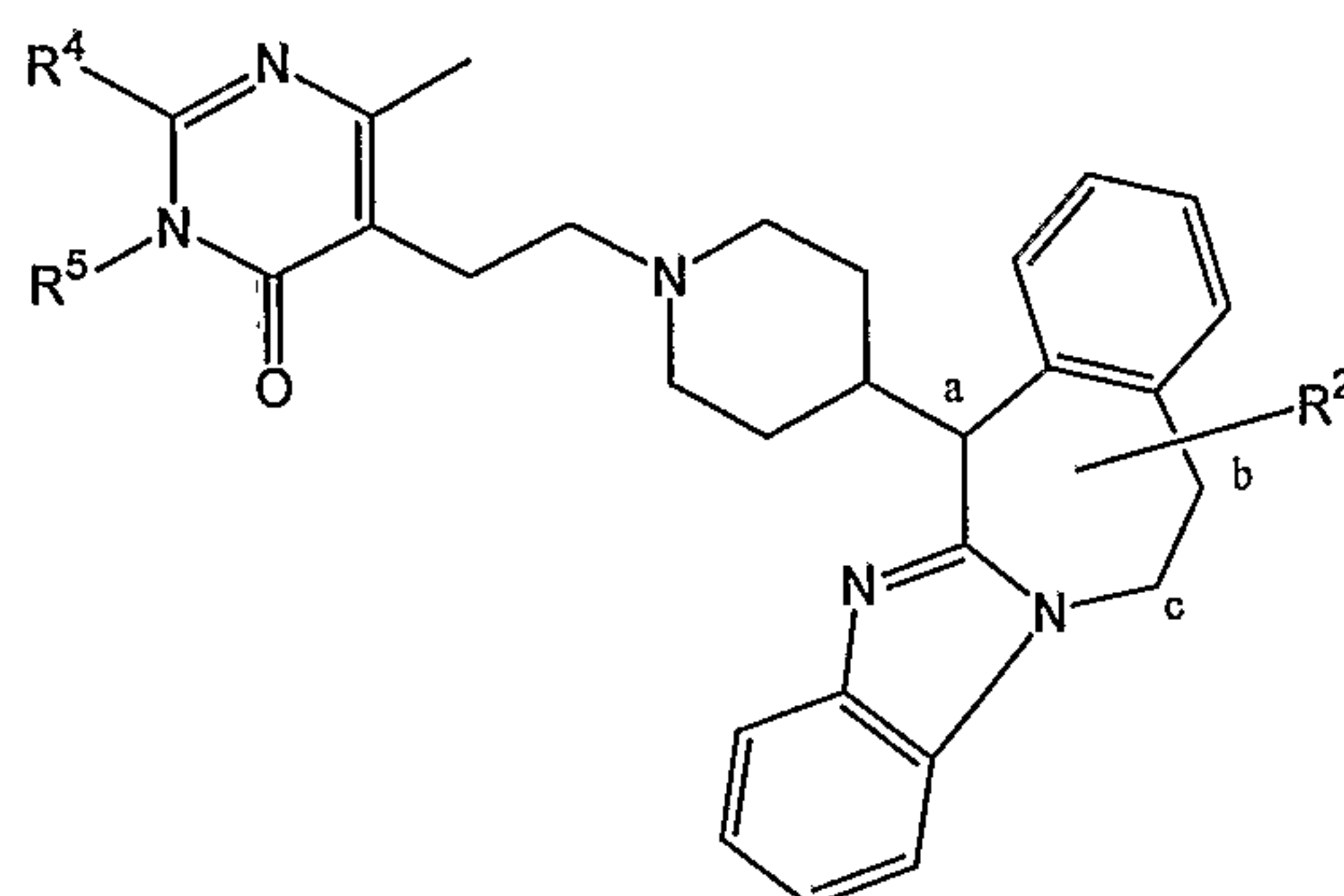
Comp.nr	Ex. nr.	R ¹		Phys.data and stereochemistry
99	B2	2-benzyl	-CH ₂ -C(CH ₃)=N-N(CH ₃)-	[(2 α ,4 α)(B)]
100	B2	2-benzyl	-CH=C(CH ₃)-CH=CH-	[(2 α ,4 α)(B)]
101	B2	2-benzyl	-C(CH ₃)=CH-C(CH ₃)=CH-	[(2 α ,4 α)(B)]
102	B2	2-benzyl	-CH=C(Cl)-CH=C(Cl)-	[(2 α ,4 α)(B)]
103	B2	2-benzyl	-CH=C(CF ₃)-CH=C(Cl)-	[(2 α ,4 α)(B)]
104	B2	4-methyl	-CH=CH-S-	
105	B2	2-methylcyclohexyl	-CH=CH-CH=CH-	
106	B2	2-benzyl	-CH=CH-S-	[A(2 α ,4 α)(B)]
107	B2	2-benzyl	-CH=CH-S-	[B(2 α ,4 α)(B)]
108	B2	2-methylcyclohexyl	-CH=CH-S-	

Table 3 :



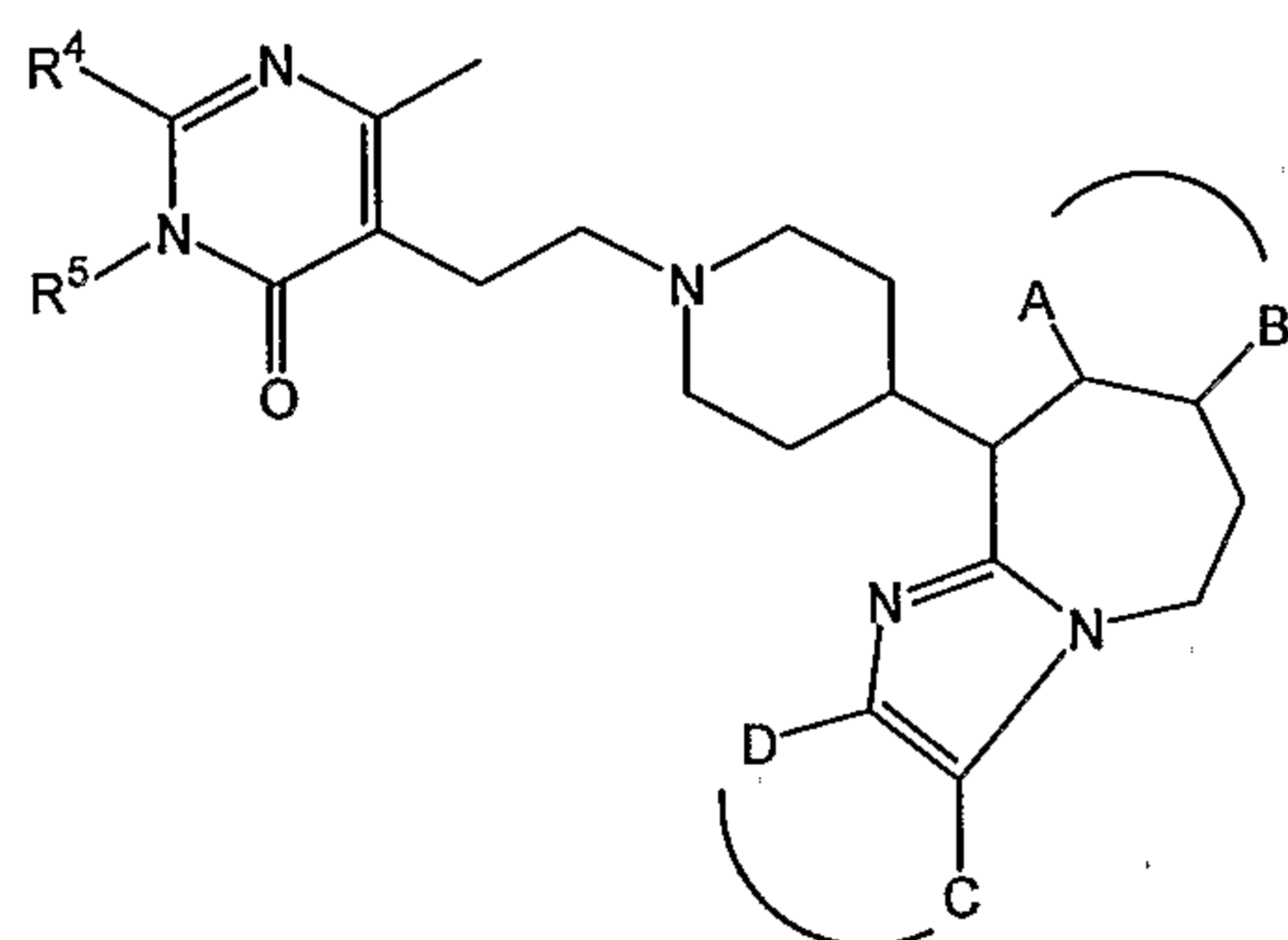
Comp.nr	Ex. nr.	R ²		Phys.data and stereochemistry
6	B4	a) -Me	-CH=CH-CH=CH-	hydrate (1:1)
110	B4	a) -Me	-CH=CH-S-	
111	B4	a) -CH ₂ -Phe	-CH=CH-CH=CH-	
112	B5	b) -CH ₂ -Phe	-CH=CH-CH=CH-	
7	B5	b) -CH ₂ -Phe	-CH=CH-S-	

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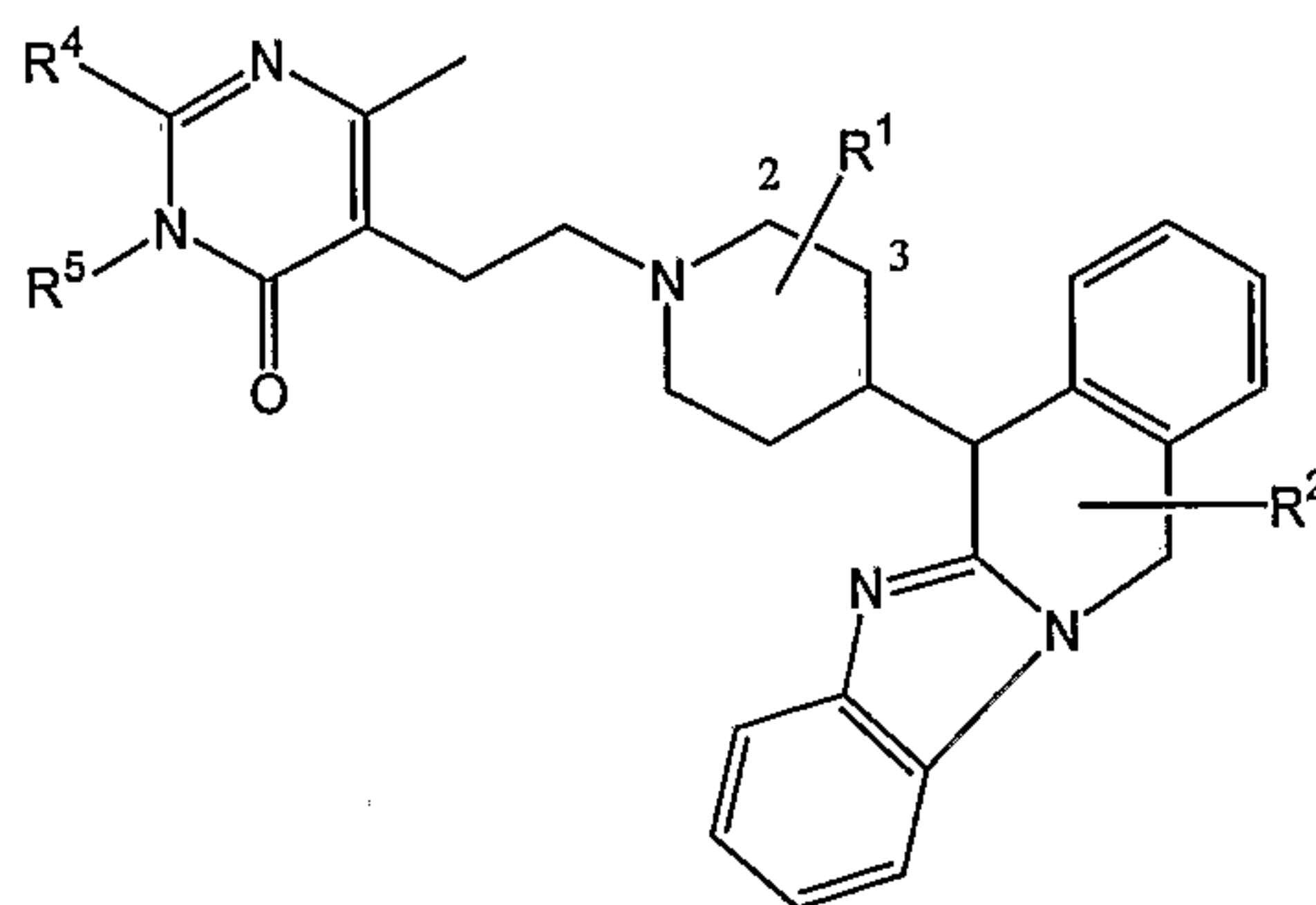
Comp.nr	Ex. nr.	R ²	R ⁵ R ⁴	Phys.data and stereochemistry
113	B4	a) -Phe	-CH=CH-S-	

Table 4:



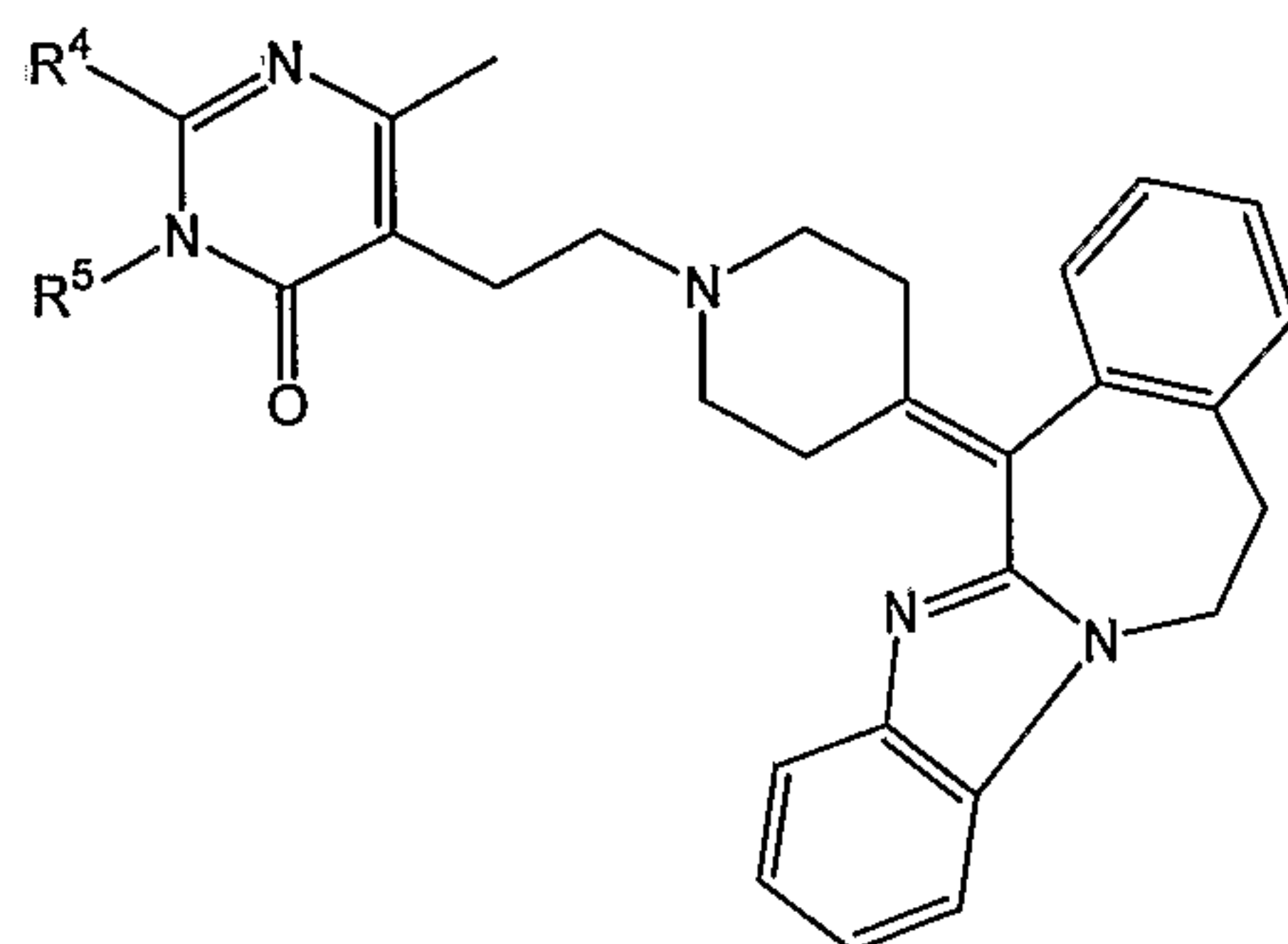
Co.nr.	Ex. nr.	R ⁵ R ⁴	A B	C D	Phys.data and stereochemistry
114	B1	-CH=CH-CH=CH-	-CH=CH-S-	-CH=CH-CH=CH-	H ₂ O(1:1) (E)-2-butenedioate(1:1)
115	B1	-CH=CH-S-	-CH=CH-S-	-CH=CH-CH=CH-	H ₂ O(2:1) (E)-2-butenedioate(2:3)
116	B1	-CH=CH-CH=CH-	-CH=CH-CH=CH-	N=CH-CH=CH-	
117	B1	-CH=CH-S-	-CH=CH-CH=CH-	N=CH-CH=CH-	
118	B1	-CH=CH-S-	-CH=CH-N(CH ₃)-	N=CH-CH=CH-	

Table 5 :



Co. nr.	Ex. nr.	R ¹	R ²	R ⁵ R ⁴	Phys.data and stereochemistry
119	B2	2-benzyl	H	-CH=CH-S-	cis
3	B2	2-benzyl	H	-CH=CH-S-	[(2 α ,4 β)(B)]
4	B2	2-benzyl	H	-CH=CH-S-	trans
120	B2	2-benzyl	H	-CH=CH-CH=CH-	[(2 α ,4 β)(B)]
121	B2	2-benzyl	H	-CH=CH-CH=CH-	[(2 α ,4 β)(A)]
122	B2	H	H	-CH ₂ -CH ₂ -CH ₂ -CH ₂ -	
123	B2	H	H	-CH ₂ -CH ₂ -CH ₂ -S-	
124	B2	H	H	-CH=CH-CH=CH-	
125	B2	H	H	-CH ₂ -CH ₂ -S-	
126	B2	H	H	-C(CH ₃)=CH-S-	
127	B2	H	H	-CH=C(CH ₃)-CH=CH-	
128	B2	H	H	-CH=CH-CH=C(CH ₃)-	
129	B2	H	H	-CH ₂ -C(CH ₃)=N-N(CH ₃)-	
130	B2	H	H	-CH=CH-N(CH ₃)-	
131	B2	H	H	-CH=C(CH ₃)-N(CH ₃)-	
132	B2	H	H	-O-C(CH ₃)=CH-	(E)-2-butenedioate (1:2)
133	B2	H	H	-C(CH ₃)=N-N(CH ₃)-	.H ₂ O(1:1)
134	B2	2-benzyl	H	-CH=CH-CH=CH-	[(2 α ,4 α)(B)]
135	B2	2-benzyl	H	-CH=CH-CH=CH-	[(2 α ,4 α)(A)]
136	B2	H	H	-CH=CH-S-	.ethanedioate(2:5) .H ₂ O (2:1)

Table 6 :



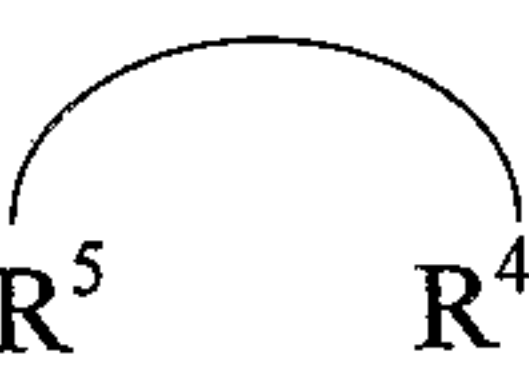
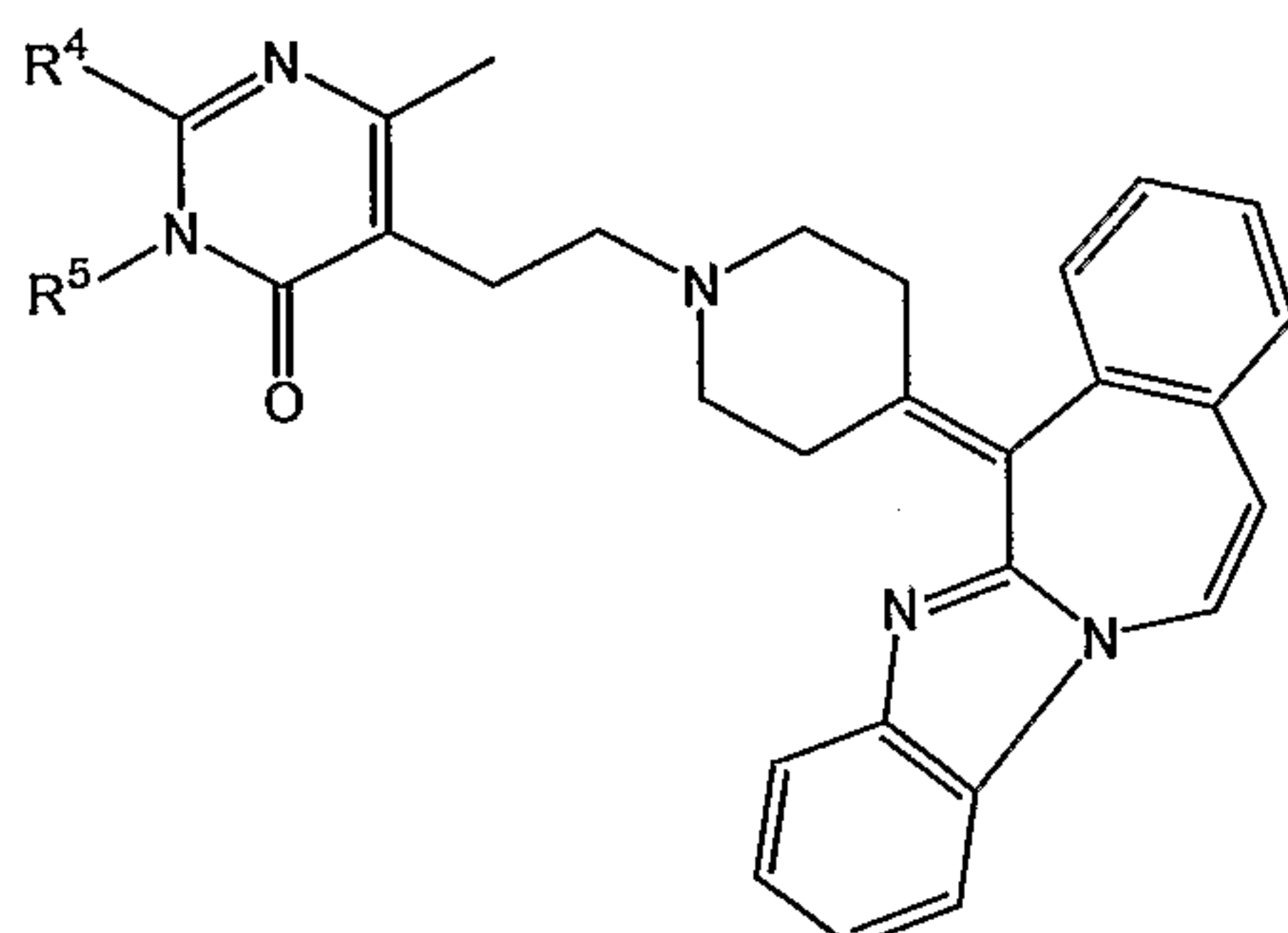
Co. nr.	Ex. nr.		Phys.data and stereochemistry
5	B3	-CH-CH-S-	
137	B3	-CH ₂ -CH ₂ -S-	
138	B3	-CH ₂ -CH ₂ -CH ₂ -S-	
139	B3	-CH=CH-CH=CH-	

Table 7 :



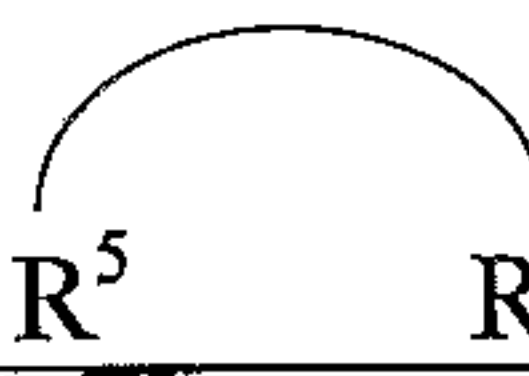
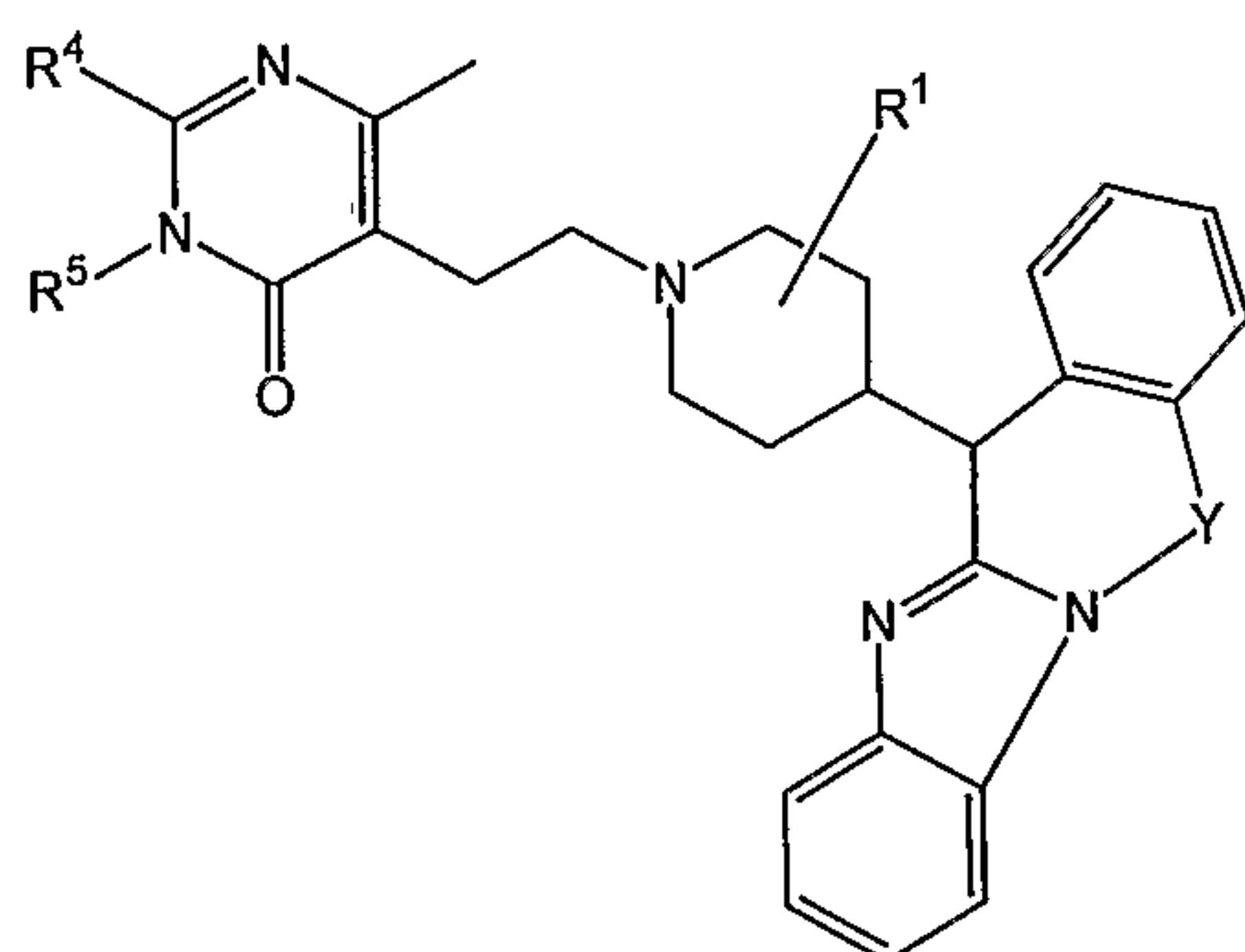
Co. nr.	Ex. nr.		Phys.data and stereochemistry
140	B6	-CH=CH-CH=CH-	
141	B6	-CH ₂ -CH ₂ -CH ₂ -S-	
8	B6	-CH=CH-S-	
142	B6	-CH ₂ -CH ₂ -S	

Table 8 :



Co. nr	Ex.m	Y	R ¹	R ⁴	R ⁵	Phys.data and stereochemistry
143	B2	-CH ₂ -CH ₂ -	H		-CH ₃	
144	B1	-CH ₂ -CH ₂ -	H		-CH ₃	
145	B1	-CH ₂ -CH ₂ -	H		-CH ₃	.H ₂ O(1:1)
146	B1	-CH ₂ -CH ₂ -	H		-CH ₃	
150	B1	-CH ₂ -CH ₂ -	2-benzyl	-NH ₂	-CH ₃	[(2 α ,4 α)(B)]
147	B2	-CH ₂ -	H	-NH ₂	-CH ₃	(Z)-2-Butenedioate (1:3).H ₂ O(1:1)
148	B2	-CH ₂ -	H		-CH ₃	
149	B2	-CH ₂ -	H		-CH ₃	
151	B2	-CH ₂ -	H		-CH ₃	.HCl(1:3).H ₂ O (1:2) .2-propanolate (2:1)

5 C. Pharmacological examplesC1. *In vitro* determination of the histaminic H₁-and H₂-antagonist activity.

Radioligand receptor binding studies were performed *in vitro* for radioligand binding of the selected compounds using a preparation of a tissue which was enriched in a particular receptor, i.e. the histamine H1- or H2-receptor. For the histamine H1-receptor, the tissue used were CHO-cells, permanently transfected with the human histamine H1-receptor. Only diphenhydramine was tested against guinea pig cells from the cerebral cortex. Competitive inhibition of [³H] Pyrilamine by the tested compounds was conducted by incubating a low (nM) concentration of the radioligand with a small sample of the tissue preparation (0.2-5 ml; 1-5 mg tissue) in a buffered medium and various concentrations of the compounds, dissolved in DMSO, spanning at least 4 orders of magnitude around the pIC₅₀ value, derived from the inhibition curve. The histaminic H2-antagonist activity was tested in much the same way as the histaminic H1-antagonist activity, using guinea pig striatum cells and [¹²⁵I]APT as the radioligand in a concentration of 0.1 nM. Incubation was done during 150 min at 22°C.

All compounds according to our invention showed a pIC₅₀ value of 5 or more for histaminic H1-antagonist activity. Several compounds showed a pIC₅₀ value of 6 or more for histaminic H1-antagonist activity. These compounds are listed in Table 9. Furthermore is observed that a commercially available typical histamine H1-antagonist (diphenhydramine) exhibits only a slightly higher histaminic H1-antagonist activity as the bulk of the compounds according to our invention. Furthermore is shown that the commercially available H2-antagonists (ranitidine and cimetidine) exhibit histamine H2-activities in the range of the (moderately high) H2-activities of the compounds according to our invention. A selection of the compounds in Table 9, including commercially available compounds, was also tested in *in vivo* experiments for their ability to reduce ICP.

Table 9 : Results of the histamine H1- and H2-antagonist activity receptor model screening.

Comp.nr	H1-antagonist activity (pIC ₅₀)	H2-antagonist activity (pIC ₅₀)
14	7.6	
94	7.0	
104	7.0	
46	6.9	
110	6.9	
1 (also tested <i>in vivo</i>)	6.7	6.0
6	6.7	
23	6.7	
78	6.7	
81	6.7	
82	6.7	
50	6.6	
55	6.6	
87	6.6	
12	6.5	
13	6.5	
15 (also tested <i>in vivo</i>)	6.5	
45 (also tested <i>in vivo</i>)	6.5	
48	6.5	
49	6.5	
53	6.5	
54	6.5	
83	6.5	
88	6.5	
20	6.4	
32 (also tested <i>in vivo</i>)	6.4	

Comp.nr	H1-antagonist activity (pIC ₅₀)	H2-antagonist activity (pIC ₅₀)
47 (also tested <i>in vivo</i>)	6.4	
57	6.4	
58	6.4	
105	6.4	
29	6.3	
51	6.3	
84	6.3	
17 (also tested <i>in vivo</i>)	6.2	
27	6.2	
37	6.2	
2 (also tested <i>in vivo</i>)	6.1	
30 (also tested <i>in vivo</i>)	6.1	
35	6.1	
56	6.1	
89	6.1	
90	6.1	
9 (also tested <i>in vivo</i>)	6.0	
31	6.0	
41	6.0	
44 (also tested <i>in vivo</i>)	6.0	
102	6.0	
Ranitidine (also tested <i>in vivo</i>)	-	5.5
Cimetidine	-	5.9
Diphenhydramine (also tested <i>in vivo</i>)	7.2	-

C.2. In vivo pharmacology

Closed Head Injury (CHI) Model

A clinically relevant rat model for traumatic brain injury was used to test the compounds according to the invention and the commercially available compounds.

5 This model mimics several clinical features of traumatic brain injury, such as increased ICP, decreased cerebral perfusion pressure, morphologic alterations including diffuse axonal injury, neuronal necrosis and contusion, impairment of autoregulation of cerebral blood flow and reduction of brain oxygenation and was applied for screening drugs with ICP-lowering effects. Trauma was induced in intubated, isoflurane
10 anesthetized (1.5 % isoflurane in a mixture of 30 % O₂ and 70 % N₂O) Sprague-Dawley rats (380-400 g) stereotaxically positioned on a table mounted on 4 springs. A 400 g steel cylinder, protected with a 9 mm diameter silicon disc, was dropped on the unprotected skull from a height of either 70 cm or 50 cm (respectively 'severe' and 'moderate' head injury). The impact area was centered between bregma and lamda.
15 ICP was recorded using a Codman microsensor probe inserted in the parietal cortex. In both severe and moderate head injuries the ICP increased immediately after trauma and remained elevated for several days. The severe head injury mode was used for the evaluation of pharmacological effects immediately after trauma (screening procedure). When survival and recovery from anesthesia was envisaged, the moderate head injury
20 mode was applied. In pharmacological studies, animals with a pathological ICP between 12.5 and 35 mm Hg were included. The changes in ICP, mean arterial blood pressure (MABP) and cerebral perfusion pressure CPP (=MABP-ICP) were expressed as percentage of the initial value at onset of the treatment.

Screening procedure for the compounds according to the invention

On a weekly base, 4 treated groups of 3 rats were compared with 3 saline treated animals. Since conventional statistical methods require a larger amount of animals, a sequential procedure was used. Sequential methods operate in different stages. At each stage, a group of animals was selected as homogeneous as possible. Animals
30 were randomly allocated to either drugs or saline. The procedure allowed to make the decision of rejecting the drug, accepting the drug as active or to continue with a new group of animals in a next stage. Given the biological relevant level of activity that must be detected, the expected fraction of false positive and negative results was

known and fixed. A sequential two-sample grouped rank test was used. A three stage sequential design with a relatively small number of animals at each stage showed to be optimal. Despite the variability in the individual data, the procedure consistently accepted reference treatments such as mannitol as active, while controls were rejected.

- 5 Clinically relevant i.v. doses of mannitol (3 g over 45 min) consistently reduced the ICP (mean reduction about 20 %).

Table 10 : Results of the screening procedure.

<i>Treatment</i> ⁽¹⁾	<i>Delta %</i> ⁽²⁾	<i>Decision</i> ⁽³⁾
Compound 9	-12.4	active
Compound 15	-23.3	active
Compound 17	-8.9	active
Compound 30	-9.3	active
Compound 32	-13.9	active
Compound 44	-14.8	active
Compound 45	-13.1	active
Compound 47	-12.0	active
CD10%	5.1	not active
CD10%+3H2T	10.0	not active
CD20%	19.1	not active
CD20%+HCl	2.4	not active
Mannitol ¹	-21.7	active
Mannitol ²	-22.1	active
Mannitol ³	-13.0	active
Mannitol ⁴	-19.3	active
Mannitol ⁵	-19.9	active

- 10 (1) Experimental compounds administered as a bolus of 1mg/kg given in 1 min, followed by an infusion of 0.5 mg/kg/min for 44 min; solvents administered as a 0.4 ml bolus in 1 min followed by an infusion of 0.2 ml/min for 44 min; mannitol given as an infusion of 67 mg/kg/min for 45 min.
- (2) Delta%: average change of the relative ICP from baseline over the treatment period.
- 15 (3) Decision: based upon sequential statistical evaluation.

CD = hydroxypropyl- β -cyclodextrin solvent

H2T = tartaric acid solvent

Mannitol¹⁻⁵: Mannitol was evaluated 5 times in separate tests (positive controls). The result of each test is mentioned.

5

Further studies

Table 11 shows the changes in some relevant physiological variables recorded during treatment after severe CHI in rats. Treatment was started at 20 min after severe head injury and involved administering a dose of 0.5 mg/kg/min during 10 minutes, followed by 0.1 mg/kg/min during 50 min.

Table 11 : Changes in relevant physiological variables during treatment after severe CHI in rats.

	Solvent (n = 10)	Compound 2 (n = 10)	Compound 1 (n = 10)	Racemate (comp. 1 and comp.2) (n = 10)
ICP (%)	1.6 (-9.4;11.1)	-15.3 (-20.0;-9.5)*	-15.4 (-22.6; -11.5)*	-19.1 (-24.9; -10.8)*
MABP (%)	-1.2 (-2.7; 3.7)	18.8 (-2.0; 31.0)*	-3.6 (-11.9; -1.5)	0.6 (-5.1; 8.5)
CPP (%)	-1.3 (-8.0; 5.8)	24.2 (0.9; 43.6)*	-1.9 (-8.9; 0.4)	7.5 (-2.4; 15.5)
ETCO ₂ (%)	8.0 (-1.2; 12.9)	-4.4 (-8.9; 2.3)*	2.2 (-0.8; 8.4)	2.4 (-7.8; 3.8)
Heart rate(%)	-2.7 (-5.4; 3.9)	-9.6 (-21.8; 0.7)	-4.1 (-11.4; 1.9)	5.6 (-11.7; 0.4)
Resp. rate(%)	3.6 (-4.3; 11.8)	6.6 (-1.3; 14.6)	5.3 (-3.3; 13.6)	9.6 (3.0; 14.8)

15 Average change over the entire treatment period, expressed as % change of initial value.

Values are medians (95% C.I.).

* = Significantly different from solvent group (p < 0.05, Dunnett's test)

Solvent : 10% hydroxypropyl-beta-cyclodextrine, tartaric acid, NaOH and mannitol in pyrogen free water ; pH = 4 ;osmolarity 312- 314 mOsm/ kg ; compound concentration 2 mg/ml.

20

Compound : pyrimido[1,2-*a*]benzimidazol-4(10*H*)-one, 3-[2-[4-(11,12-dihydro-6*H*-benzimidazo[2,1-*b*][3]benzazepin-6-yl)-2-(phenylmethyl)-1-piperidinyl]ethyl]-2,10-dimethyl (E)-2-butenedioate (2:3) hydrate (1:1)

Compound 1 : (B)[(2 α , 4 β)(A)]

Compound 2 : (A)[(2 α , 4 β)(A)]

Racemate (comp. 1 and comp. 2): (2 α , 4 β)(A), i.e. the racemic mixture of Compound 1 and Compound 2

5 ICP : Intracranial pressure

MABP : Mean arterial blood pressure

CPP : Cerebral perfusion pressure

ETCO₂ : End tidal CO₂

10 The significant effect of compound 2 on MABP is much less pronounced when the compound is given at a continuous infusion of 0.1 mg/kg.min. In this case a blood pressure peak is not present and increases in MABP larger than 20 % are not observed (median MABP increase at the end of the infusion is 9 %, n=6). The maximal reduction of ICP at this dose is comparable to the one observed when the infusion is preceded by
15 the 'loading dose' of 5 mg/kg over 10 min, but the time required to obtain this effect is longer (median: 30 min).

Effect of ranitidine and diphenhydramine on the ICP

Ranitidine was infused for 6 min at a dose of 2 mg/kg/min in the rat CHI-model after
20 inflicting severe head injury. Solvent (NaCl + H₂T) was given at the same rate. In each group, 6 rats were treated. Ranitidine was observed to yield a statistically significant larger reduction in ICP than in the solvent-treated group (7.7 % versus 0.5% reduction, which was significant at p=0.013). The percentage of reduction are calculated as % change of the ICP recorded at the onset of the treatment and at the end of the infusion.
25 No significant change in blood pressure was observed.

Diphenhydramine was infused for 10 min at a dose of 1 mg/kg/min in the rat CHI-model after inflicting severe head injury. Three rats were treated. Diphenhydramine was observed to yield a 34 % reduction in ICP without any significant effect on the
30 blood pressure.

Comparative experiments with agonists.

For comparison, two commercially available H₂-agonists (dimaprit and impromidine) were also tested by infusion into non-traumatized rats, using a dose of 0.5 mg/kg/min for 10 minutes for dimaprit and increasing doses up to 3.75 mg/kg/hour for

5 impromidine. No effects were observed. When dimaprit was dosed at the high dose of 2 mg/kg/min for 10 minutes, and impromidine was doses as a bolus of 0.5 mg/kg blood pressure and ICP were observed to drop, but recovered after the treatment.

10 It was therefor concluded that histamine H₁-and/or H₂-receptor antagonists exhibited the effect of lowering the ICP and having in the meantime hardly any significant effect on the blood pressure.

Experiments with commercially available H₁- and H₂-antagonists

15 A number of commercially available H₁- and H₂-antagonists was infused for 10 min at a dose of 0.5 mg/kg/min in the rat CHI-model after inflicting severe head injury. Solvent (NaCl + H₂T) was given at the same rate. In each group, 6 rats were treated. The results of the behavior of the ICP and BP in the first 15 minutes are summarized in Table 12.

20

Table 12 : Effect of commercially available H₁- and H₂-antagonists.

Compound	Effect on ICP	Effect on blood pressure
solvent	0	0
Cyclodextrine	-	+
Alimemazine	-	-
Antazoline	--	--
Brompheniramine	-	--
Chlorcyclizine	-	0
Chlorpheniramine	-	+

Clemastine	-	--
Clemizole	-	-
Cyproheptadine	--	--
Dimethindene	-	+
Diphenhydramine	--	0
Diphenylpyraline	-	-/0
Hydroxyzine	-	0
Ketotifen	--	0
Loratidine	-	+
Niaprazine	--	--
Oxatomide	-	-
Pheniramine	--	-
Promethazine	-	+
Pyrilamine (see Figure 5)	--	0
Ritanserine	--	-
Tiotidine	--	-/0
Zolantidine	-	0

0 : no effect ; - : decrease ; -- : strong decrease ; + : increase

Dose response for compound 1

Results of a blinded, completely randomized study of the effect of a 10 min infusion of Compound 1 at different doses (0.125, 0.25, 0.5, 1 and 2 mg/kg/min) in the rat CHI model indicate that during treatment Compound 1 invokes a sustained dose-dependent decrease of ICP (Figure 1). Starting at 1 mg/kg/min Compound 1 yields a statistically significant larger reduction in ICP than in the solvent-treated group. In the 10 min

period following the infusion a highly significant dose-dependent effect on ICP remains present (Figure 2).

5 Effects of Compound 2, Compound 1 and Racemate (Comp. 1 and Comp. 2) on brain hemoglobin concentration and oxygenation.

Near-infrared spectroscopy (NIRS) of the rat brain 'in vivo' allows to quantify non-invasively saturation of brain haemoglobin with oxygen (HbSat) and total brain haemoglobin concentration ([HbTot]). The latter is a measure for cerebral blood
10 volume (CBV). Changes in the redox state of the mitochondrial enzyme cytochrome oxidase (CytOx), an indicator for tissue oxygenation, can also be monitored.

All compounds 2, 1 and the racemate (Comp. 1 and Comp. 2) do not have a significant effect on [HbTot] when given 24 h after moderate head injury at a i.v. dose of 0.5
15 mg/kg.min during 10 min, followed by 0.1 mg/kg.min during 45 min. Only compound 2 induces a small but statistically significant reduction of HbSat. HbSat is not affected by compound 1 and the racemate (Comp. 1 and Comp. 2). At the applied dose all compounds do not have an effect on the redox state of CytOx. These results indicate that in the applied experimental conditions a vasoconstrictive effect on cerebral blood
20 vessels, if present, is limited and tissue oxygenation is not jeopardised.

Influence of anaesthesia on the effects of Compound 2

The effects of treatment with Compound 2 (i.v. infusion at a dose of 0.1 mg/kg.min during 30 min) at 24 h after moderate trauma were studied using different anesthetics
25 (isoflurane, chloralhydrate, pentobarbital). When chloralhydrate (400 mg/kg i.p.) is used as anesthetic, ICP decreases to 75% of initial value and MABP gradually increases to 110% of initial value (medians, n=6). These effects are comparable with those observed under isoflurane anesthesia. When pentobarbital (60 mg/kg i.p.) is used, Compound 2 induces a significant gradual increase in MABP up to 141 % of initial value at the end
30 of the infusion, whereas ICP decreases to 64% of initial value (medians, n=6). These results indicate that the same pattern of effects on ICP and MABP are observed under various types of anesthesia. The fact that the compound reduces the ICP significantly under pentobarbital anesthesia is important, as barbiturates are often applied in

traumatic brain injury patients. Barbiturates also reduce the ICP and an important additional effect can be obtained with the compound.

5 The effect of repeated application of Compound 1 and of mannitol on elevated ICP in traumatized rats.

Compound 1 was given 2 times with intermittent periods of 20 min at a i.v. dose of 1 mg/kg/min during 10 min, starting a first time 20 min after induction of severe head injury.

10 Mannitol was given i.v. in the same time windows as Compound 1 at a dose of 0.125 g/kg/min. The control animals received the solvent (containing 10 % HP- β -CD, pH 4) only.

Infusion with Compound 1 results in rapid reduction of ICP (Figure 3). This effect is amplified after termination of each infusion period. Blood pressure drops during Compound 1 treatment but is restored again after this episode. This is in contrast with
15 mannitol, that induces a lowering of ICP and an increase in blood pressure during each infusion followed by a decrease in blood pressure after termination of each treatment. Only in the Compound 1- treated animals a clear dissociation between the changes in blood pressure and ICP can be observed. In contrast, the mannitol treated animals exhibit more or less parallel changes in blood pressure and intracranial pressure. This
20 indicates that the pharmacological effect of Compound 1 is different from that of mannitol.

Effect of Compound 1 on cold lesion-induced rise of ICP in rabbits

Cryo-lesions were induced in adult rabbits to obtain a pathological ICP that is caused
25 by tissue oedema. A 8 mm stainless steel rod was placed at predetermined coördinates on the exposed skull of deeply anaesthetised rabbits and cooled for 10 min with liquid nitrogen. One day later the animals were re-anaesthetised and ICP and blood pressure continuously recorded as described for the rat. After a stabilisation period of 15 min, Compound 1 was infused for 10 min at a dose of 2 mg/kg/min. Solvent (preclinical
30 formulation containing 10% HP- β -CD, pH 4) was given for 10 min at a rate of 2 ml/min.

During infusion of the Compound 1, the blood pressure drops and although there is no immediate decrease in ICP, the ICP rise that is observed in the solvent-treated animals

tends to be antagonised (Figure 4). When drug infusion is terminated, blood pressure comes back to the initial value and a significant ICP reduction is seen that persists during the entire recording period.

These results indicate that the compound reduces the ICP also in non-rodent species
5 and in pathologic conditions different from closed head injury.

The effect of Compound 1 and on ICP in non-traumatized animals.

Rats

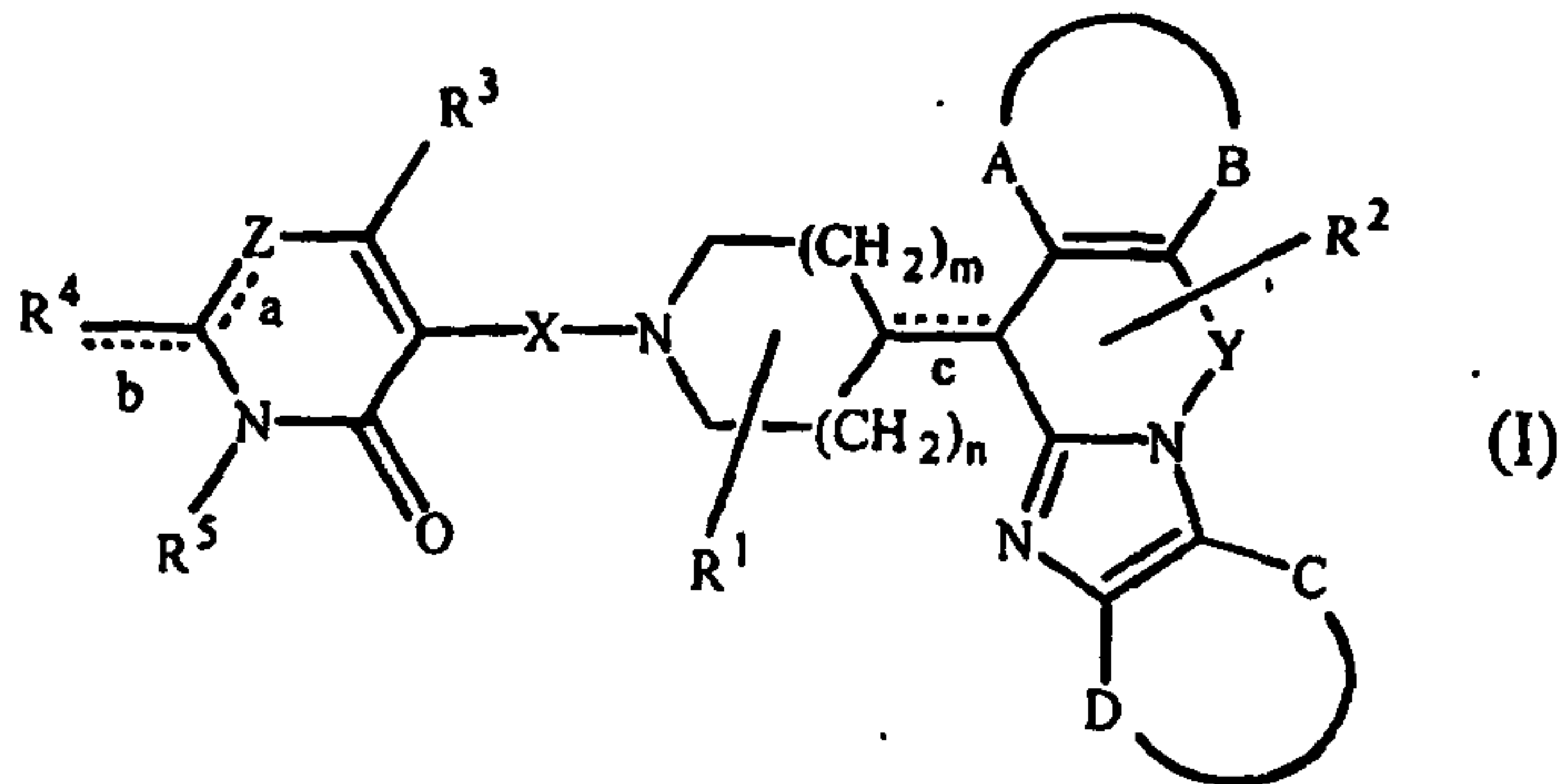
The effect of Compound 2, Compound 1 and Racemate (comp. 1 and comp. 2) on ICP,
10 MABP and CPP was tested in anaesthetised non-traumatized rats. The compounds were administered i.v. and the same dose was given as in traumatized rats (0.5 mg/kg/min during 10 minutes, followed by 0.1 mg/kg/min during 50 min). The results were comparable with those obtained in the traumatized animals.

15 Conclusion

The results obtained in traumatized animals, animals with cold lesion, and non-traumatized animals indicate that the compounds are active in various conditions, even in normal conditions. Their field of application probably includes various pathological conditions in which intracranial hypertension is present.

CLAIMS:

1. Use of a compound for the manufacture of a medicament for immunomodulation in a mammal, for the suppression of hypersensitivity or inflammatory reactions and for the treatment and prevention of allergic diseases and gastrointestinal conditions, wherein the compound is a compound according to the general Formula (I)



the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the N-oxide form thereof, wherein

m is 1 or 2;

n is 0, 1 or 2;

a,b,c independently are a single or a double bond;

X is a covalent bond or a bivalent radical of a C₁₋₆alkanediyl wherein one or more -CH₂-groups may be optionally replaced with -O-, -S-, -CO-, or NR⁷- wherein:

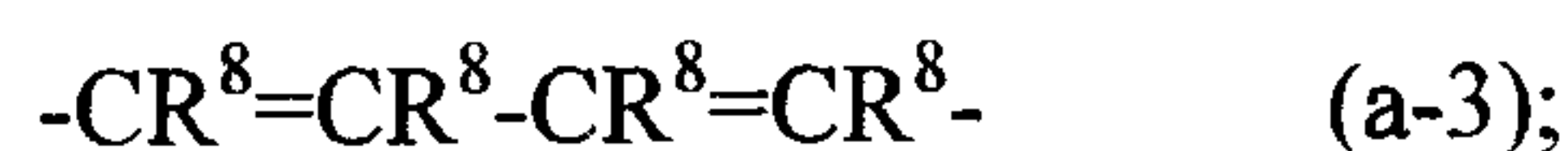
R⁷ is hydrogen, alkyl, Ar, Ar-alkyl, Het, Het-alkyl, hydroxyalkyl, alkyloxy, alkyloxyalkyl, alkyloxyalkyloxyalkyl, aminoalkyl, mono- or dialkylaminoalkyl, formyl, alkylcarbonylaminoalkyl, alkylcarbonyloxyalkyl, alkyloxycarbonyl, alkyloxycarbonylalkyl, alkylaminocarbonyl, alkylaminocarbonylalkyl, hydroxyalkyloxyalkyl, aminocarbonyl, aminocarbonylalkyl, alkyloxycarbonyl or alkylcarbonyloxyalkyloxyalkyl;

Y is bivalent C₁₋₄alkanediyl or C₂₋₄ alkenediyl radical;

Z is N, in which case a is a double bond and b is a single bond or N-R⁷ in which case a is a single bond, b is a double bond and R⁷ is defined as above;

R¹, R² independently are hydrogen, hydroxyl, alkyl, alkyloxy, Ar, Ar-alkyl, di(Ar) alkyl, Het or Het-alkyl;

-A-B- independently is a bivalent radical of formula

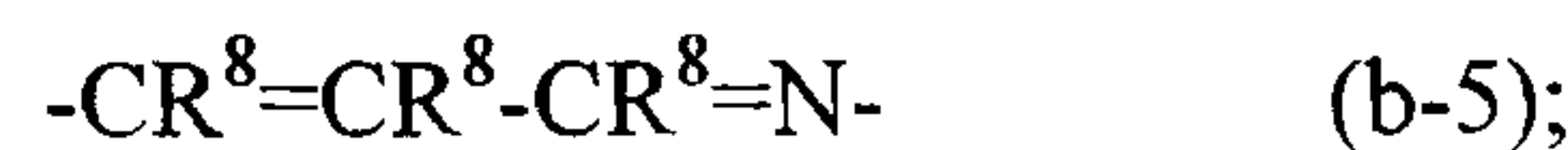
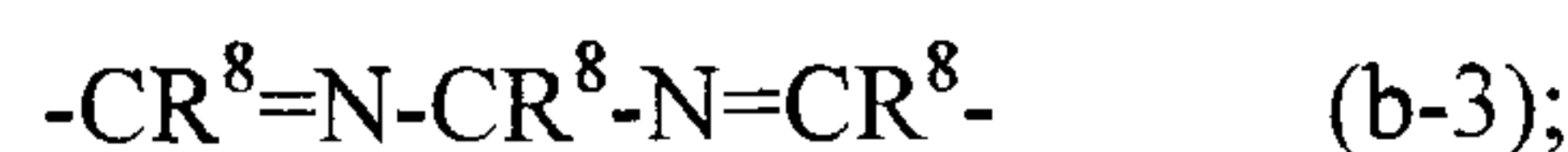
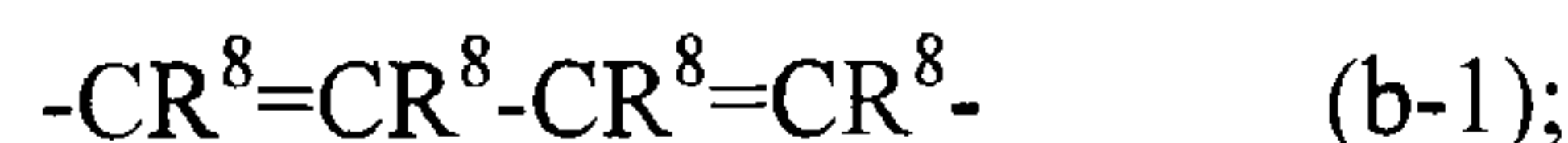


wherein

R^8 each independently is hydrogen, halo, hydroxyl, alkyl or alkyloxy;

E is a bivalent radical of formula -O- , -S- or $\text{-NR}^7\text{-}$ wherein R^7 is defined as above;

-C-D- independently is a bivalent radical of formula



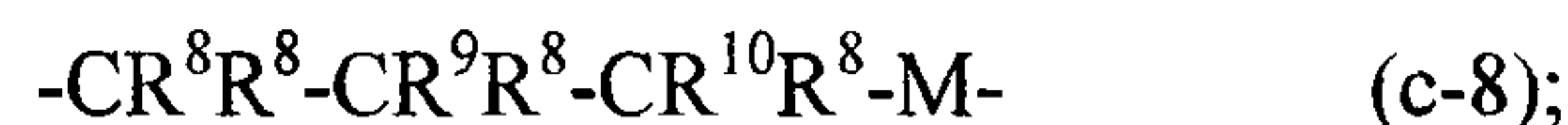
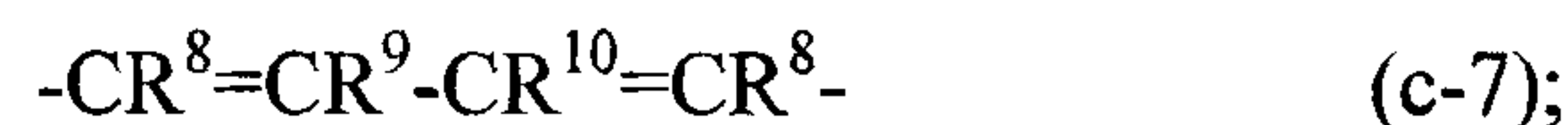
wherein R^8 is defined as above;

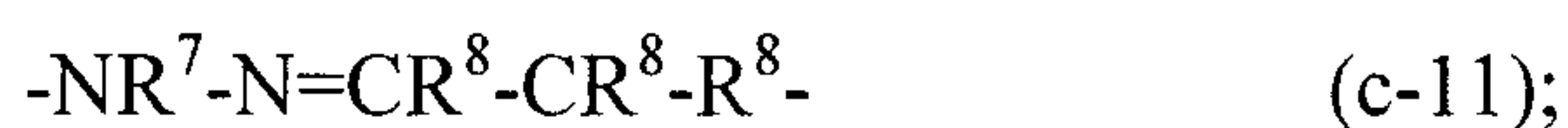
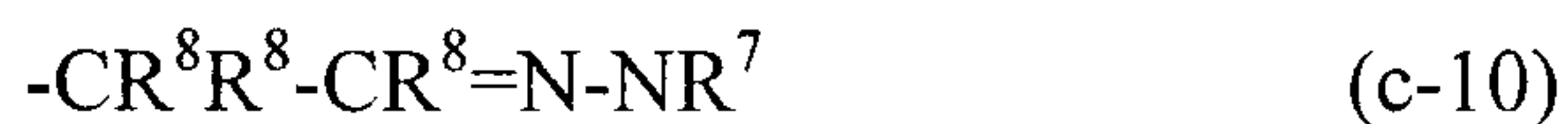
R^3 is hydrogen, halo, hydroxyl, alkyl, alkyloxy, Ar, Ar-alkyl, di(Ar-)alkyl, Het or Het-alkyl;

R^4 is hydrogen, alkyl, amino, alkylamino, Ar-amino, Het-amino, alkylcarbonylamino, Ar-aminocarbonylamino, Het-aminocarbonylamino, alkyloxyalkylamino, Ar-oxyalkylamino or Het-oxyalkylamino;

R^5 is hydrogen or alkyl;

or R^4 and R^5 together may form a bivalent radical of Formula





wherein

R^7 and R^8 are defined as above;

R^9 and R^{10} independently are hydrogen, alkyl, halo, haloalkyl;

or R^9 and R^{10} together may form a bivalent radical of formula $-\text{CR}^8=\text{CR}^8-$
 $\text{CR}^8=\text{CR}^8-$; and

M is a bivalent radical of formula $-\text{CH}_2-$, $-\text{O}-$, $-\text{S}-$ or $-\text{NR}^7$ wherein R^7 is defined as above;

Ar is a homocycle selected from the group of naphthyl and phenyl, each optionally substituted with 1, 2 or 3 substituents, each substituent independently selected from the group of hydroxyl, halo, cyano, nitro, amino, mono- or dialkylamino, alkyl, haloalkyl, alkyloxy, haloalkyloxy, carboxyl, alkyloxycarbonyl, aminocarbonyl and mono- or dialkyl-aminocarbonyl;

Het is a monocyclic heterocycle selected from the group of pyrrolyl, pyrazolyl, imidazolyl, furanyl, thienyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, pyridinyl, pyrimidinyl, pyrazinyl and pyridazinyl; or a bicyclic heterocycle selected from the group of quinolinyl, quinoxalinyl, indolyl, benzimidazolyl, benzoxazolyl, benzisoxazolyl, benzothiazolyl, benzisothiazolyl, benzofuranyl and benzothienyl; each monocyclic and bicyclic heterocycle may optionally be substituted on a carbon atom with halo, hydroxyl, alkyl or alkyloxy;

alkyl is a straight or branched saturated hydrocarbon radical having from 1 to 6 carbon atoms; or is a cyclic saturated hydrocarbon radical having from 3 to 6 carbon atoms; or is a cyclic saturated hydrocarbon radical having from 3 to 6 carbon atoms attached to a straight or branched saturated hydrocarbon radical having from 1 to 6 carbon atoms; wherein each carbon atom can be optionally substituted with halo, hydroxy, alkyloxy or oxo; halo is a substituent selected from the group of fluoro, chloro, bromo and iodo;

haloalkyl is a straight or branched saturated hydrocarbon radical having from 1 to 6 carbon atoms or a cyclic saturated hydrocarbon radical having from 3

to 6 carbon atoms, wherein one or more carbon atoms are substituted with one or more halo atoms.

2. The use according to claim 1, wherein, independently of each other, Ar is a naphthyl or phenyl, each optionally substituted with 1 substituent, each substituent independently selected from the group of halo and alkyl, Het is pyridinyl, pyrazinyl or indolyl, alkyl is methyl, ethyl or cyclohexylmethyl, halo is fluoro or chloro and haloalkyl is trifluoromethyl.
3. The use according to claim 1 to 2, wherein -A-B- is a bivalent radical of formula (a-1) or (a-3), wherein E is a bivalent radical of formula -O-, -S- or -NR⁷- wherein R⁷ is hydrogen, R⁸ is hydrogen, -C-D- is a bivalent radical of formula (b-1) or (b-2), wherein R⁸ is hydrogen and Y is a bivalent radical of formula -CH₂-, -CH₂-CH₂- or -CH=CH-.
4. The use according to any one of claims 1 to 3 wherein m and n are both 1.
5. The use according to any one of claims 1 to 4 wherein R¹ and R², each independently are hydrogen, alkyl, Ar-alkyl, Het or Het-alkyl.
6. The use according to any one of claims 1 to 5 wherein X is a bivalent radical of formula -CH₂CH₂- or -CH₂CH₂CH₂-.
7. The use according to any one of claims 1 to 6 wherein R³ is hydrogen or alkyl, Z is N-R⁷ wherein R⁷ is hydrogen or alkyl, a is a single bond and b is a double bond, and R⁴ and R⁵ together form a bivalent radical of Formula (c-1), (c-3), (c-5), (c-7), (c-8) or (c-10) wherein R⁷ and R⁸ are hydrogen.
8. The use according to claim 7, wherein R⁹ and R¹⁰ together form a bivalent radical of formula -CR⁸=CR⁸-CR⁸=CR⁸- wherein R⁸ is hydrogen.

9. The use according to claim 1, wherein the compound is 3-[2-[4-(11,12-dihydro-6*H*-benzimidazo[2,1-*b*][3]benzazepin-6-yl)-2-(phenylmethyl)-1-piperidinyl]ethyl]-2,10-dimethyl pyrimido[1.2-*a*]benzimidazol-4(10*H*)-one, the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof and the *N*-oxide form thereof.
10. The use according to claim 9, wherein the compound is 3-[2-[4-(11,12-dihydro-6*H*-benzimidazo[2,1-*b*][3]benzazepin-6-yl)-2-(phenylmethyl)-1-piperidinyl]ethyl]-2,10-dimethyl pyrimido[1.2-*a*]benzimidazol-4(10*H*)-one (E)-2-butenedioate (2-3) hydrate (1:1) and stereoisomeric forms thereof.
11. The use according to claim 9 or 10, wherein the compound is the (A)[(2 α , 4 β)(A)] enantiomer, the (B)[(2 α , 4 β)(A)] enantiomer or a mixture thereof.

-1/4-

Figures

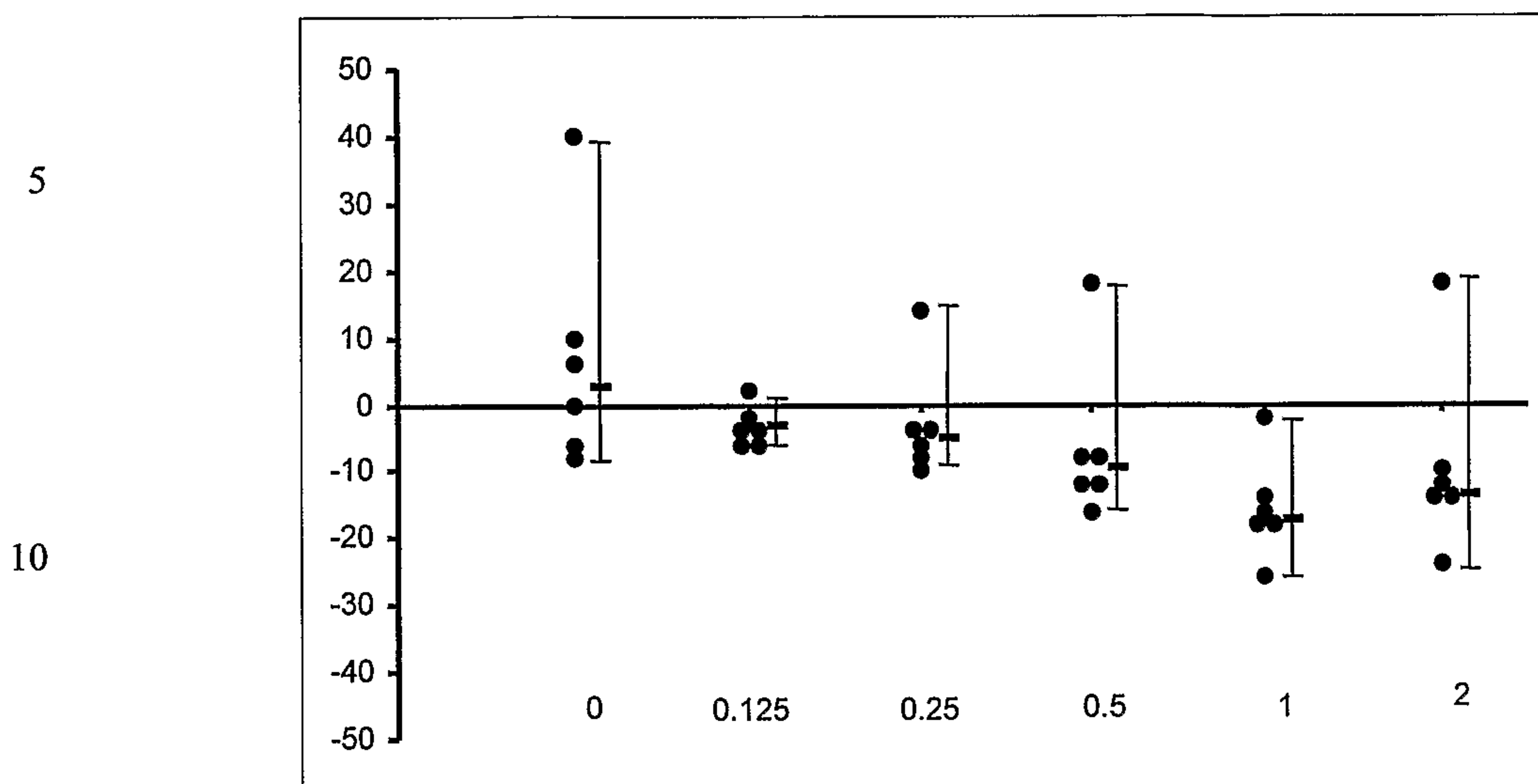


Figure 1. Dose dependence of the ICP reducing effect of Compound 2 during a 10 min infusion period. **X-axis :** Dose (mg/kg/min) ; **Y-axis :** Change in ICP as percentage of initial value.

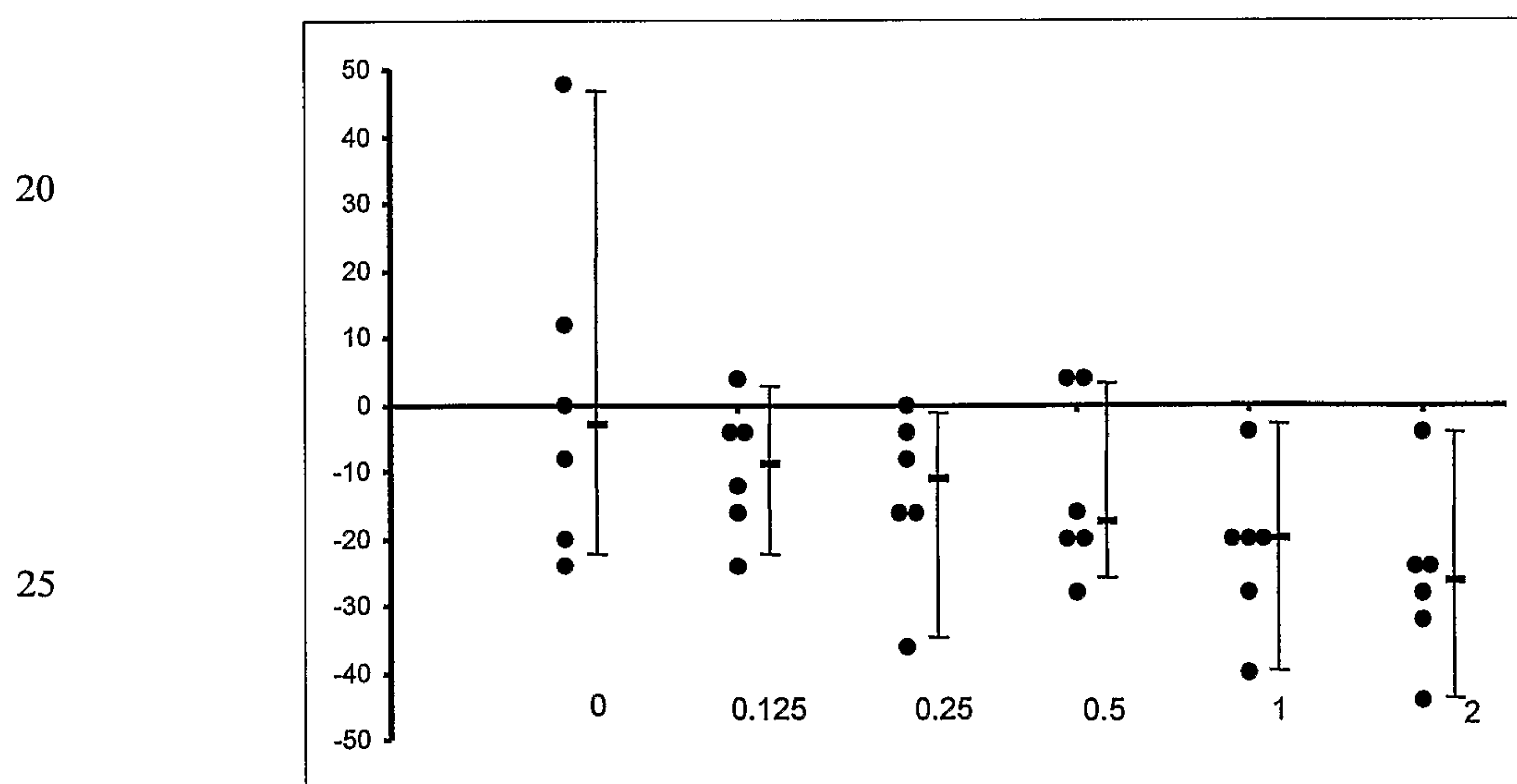


Figure 2. Dose dependence of the ICP reducing effect of Compound 2 during the 10 min post-treatment period following a 10 min infusion .

X-axis : Dose (mg/kg/min) ; **Y-axis :** Change in ICP as percentage of initial value.

-2/4-

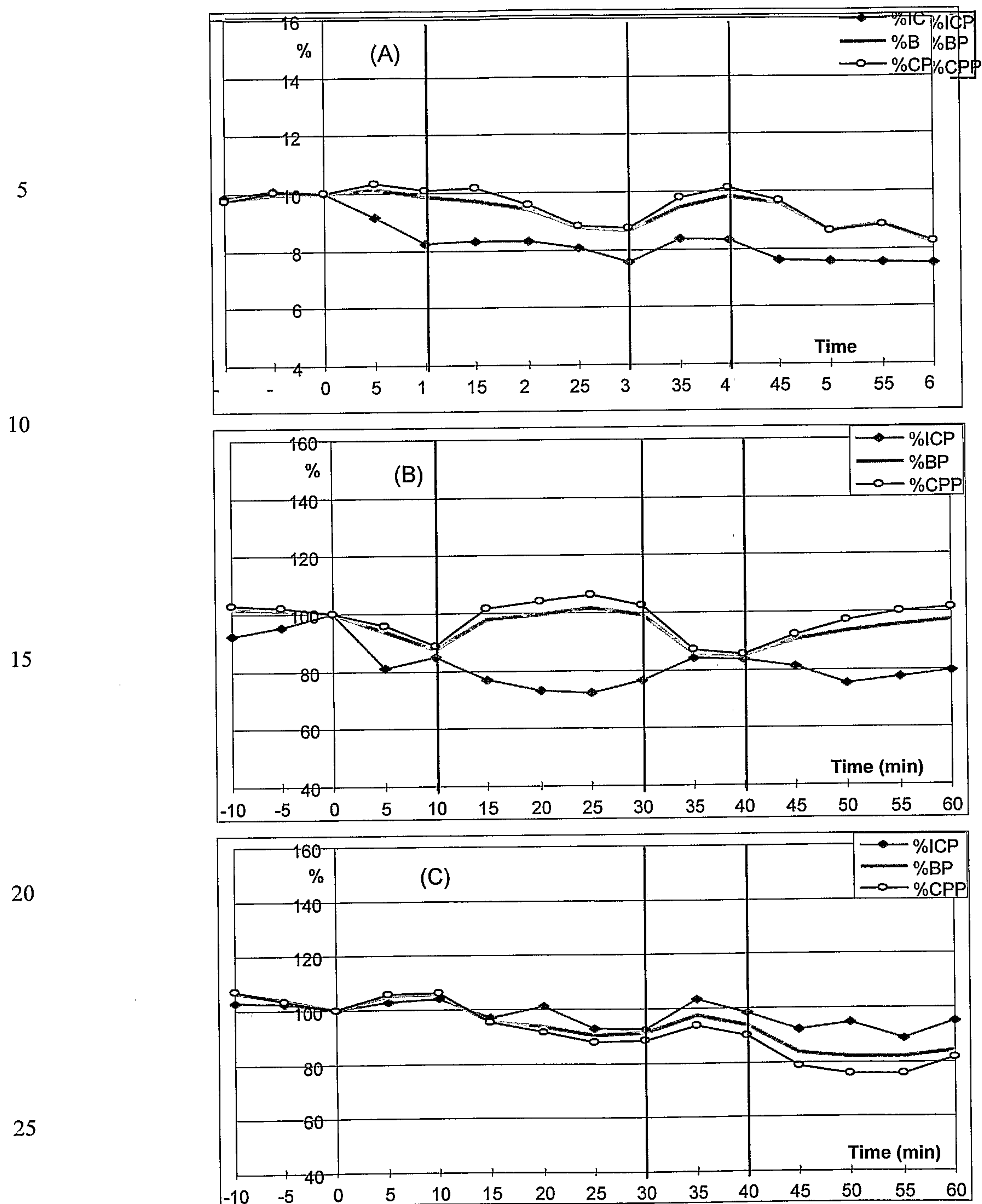
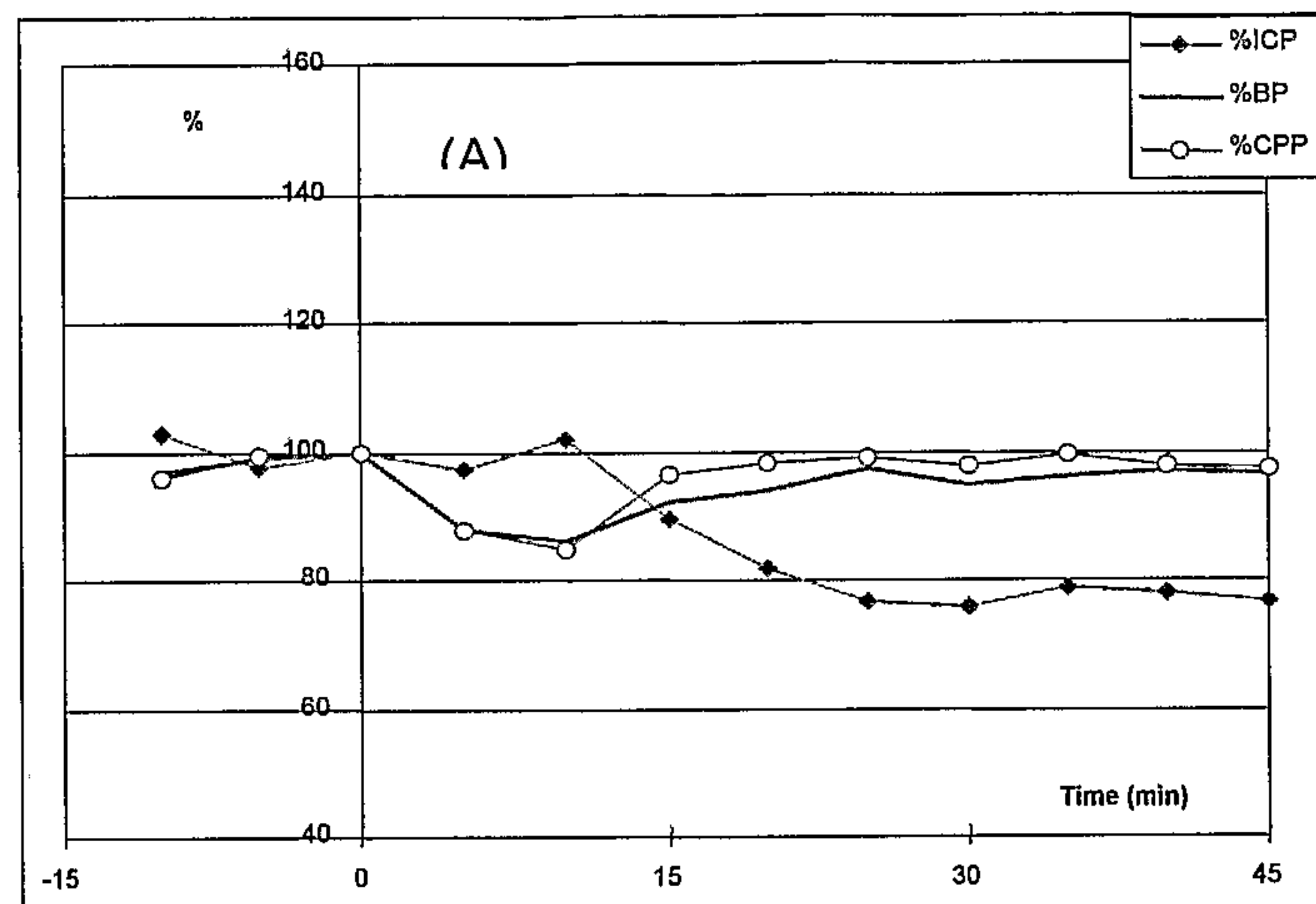


Figure 3: Time course of ICP, MABP and CPP in rats during 3 intermittent treatment periods of 10 min with respectively mannitol (Figure 3-A)(dose: 0.125 g/kg/min), Compound 2 (Figure 3-B) (dose: 1 mg/kg/min) and solvent (Figure 3-C)(10 % HP-beta-CD, pH 4). Treatment was started at 20 min after severe head injury (time = 0) and was repeated at 30 min and 60 min. The curves connect the median value for the subsequent time points. Values are expressed as % of initial value.

-3/4-

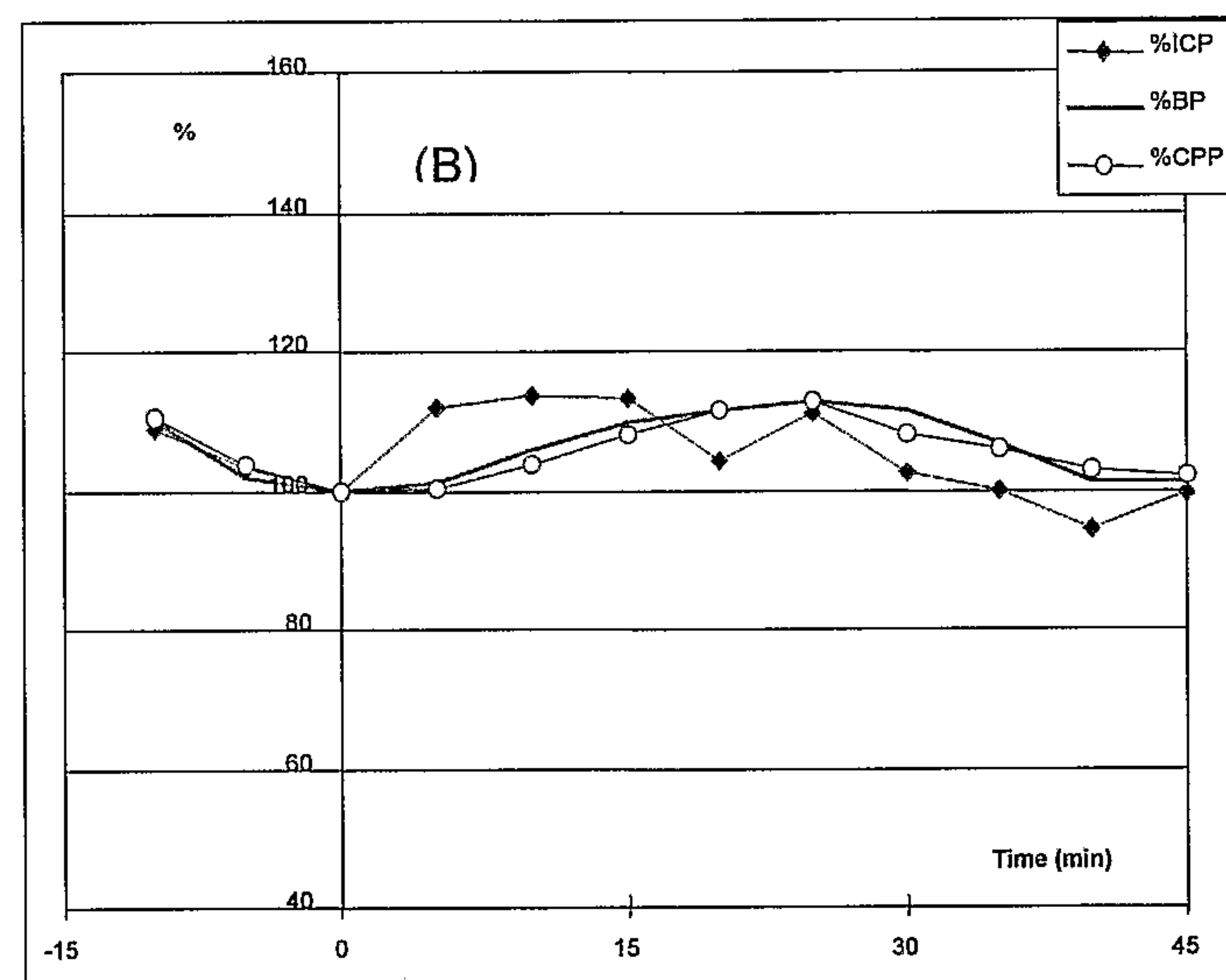
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Figure 4: Time course of ICP, MABP and CPP in rabbits treated respectively with Compound 2 (Figure 4-A)(dose: 2 mg/kg/min during 10 min) or solvent (Figure 4-B)(2ml/min during 10 min). Treatment was started at 24 h after induction of a cortical cold lesion (time = 0). The curves connect the median value for the subsequent time points. Values are expressed as % of initial value.

-4/4-

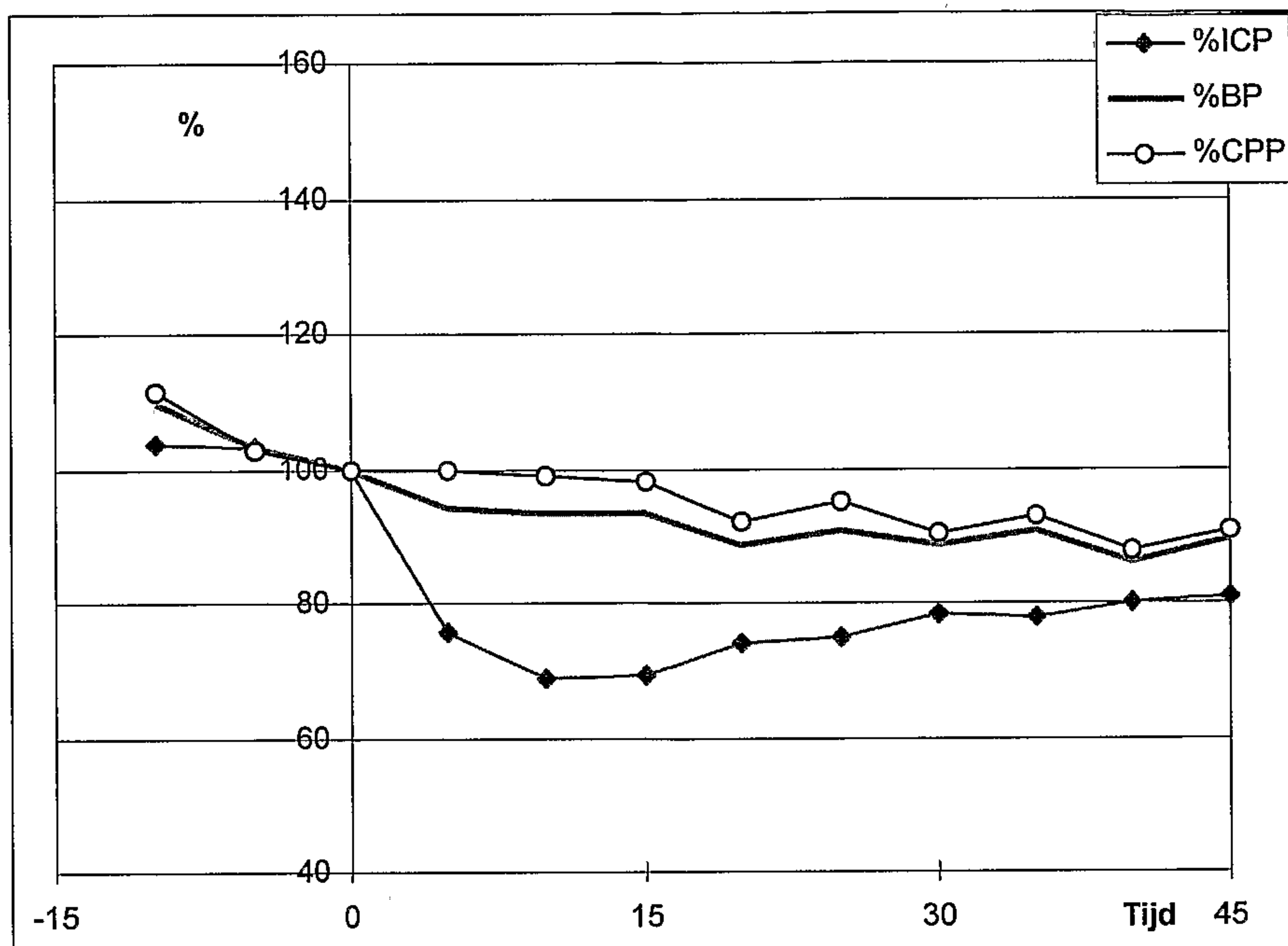


Figure 5 : Time course of ICP, BP and CPP in rabbits (n=6) treated with **Pyrilamine** (dose: 5 mg/kg/min during 10 min) . The curves connect the median value for the subsequent time points. Values are expressed as % of initial value.

