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(54) Title: UNSOLVATED BENZODIAZEPINE COMPOSITIONS AND METHODS

(57) **Abrégé/Abstract:**

The present invention relates to systems and methods for generating new forms of benzodiazepine and benzodiazepine related compounds as well as new compounds and formulations generated by such methods. In particular, the present invention provides high throughput systems and methods for generating and identifying new crystalline benzodiazepine and benzodiazepine related polymorphs and new unsolvated, solvated, and other forms of the compounds that find use as improved drugs and drug formulations.

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(54) Title: UNSOLVATED BENZODIAZEPINE COMPOSITIONS AND METHODS

(57) Abstract: The present invention relates to systems and methods for generating new forms of benzodiazepine and benzodiazepine related compounds as well as new compounds and formulations generated by such methods. In particular, the present invention provides high throughput systems and methods for generating and identifying new crystalline benzodiazepine and benzodiazepine related polymorphs and new unsolvated, solvated, and other forms of the compounds that find use as improved drugs and drug formations.



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UNSOLVATED BENZODIAZEPINE COMPOSITIONS AND METHODS

This invention was supported in part with NIH grant GM046831. The United States government may have rights in this invention.

5 FIELD OF THE INVENTION

The present invention relates to systems and methods for generating new forms of benzodiazepine and benzodiazepine related compounds as well as new compounds and formulations generated by such methods. In particular, the present invention provides high throughput systems and methods for generating and identifying new crystalline
10 benzodiazepine and benzodiazepine related polymorphs and new unsolvated, solvated, and other forms of the compounds that find use as improved drugs and drug formations.

BACKGROUND OF THE INVENTION

Multicellular organisms exert precise control over cell number. A balance between
15 cell proliferation and cell death achieves this homeostasis. Cell death occurs in nearly every type of vertebrate cell via necrosis or through a suicidal form of cell death, known as apoptosis. Apoptosis is triggered by a variety of extracellular and intracellular signals that engage a common, genetically programmed death mechanism.

Multicellular organisms use apoptosis to instruct damaged or unnecessary cells to
20 destroy themselves for the good of the organism. Control of the apoptotic process therefore is very important to normal development, for example, fetal development of fingers and toes requires the controlled removal, by apoptosis, of excess interconnecting tissues, as does the formation of neural synapses within the brain. Similarly, controlled apoptosis is responsible for the sloughing off of the inner lining of the uterus (the endometrium) at the start of
25 menstruation. While apoptosis plays an important role in tissue sculpting and normal cellular maintenance, it is also the primary defense against cells and invaders (*e.g.*, viruses) which threaten the well being of the organism.

Not surprisingly many diseases are associated with dysregulation of the process of cell death. Experimental models have established a cause-effect relationship between
30 aberrant apoptotic regulation and the pathenogenicity of various neoplastic, autoimmune and viral diseases. For instance, in the cell mediated immune response, effector cells (*e.g.*, cytotoxic T lymphocytes "CTLs") destroy virus-infected cells by inducing the infected cells to undergo apoptosis. The organism subsequently relies on the apoptotic process to destroy

the effector cells when they are no longer needed. Autoimmunity is normally prevented by the CTLs inducing apoptosis in each other and even in themselves. Defects in this process are associated with a variety of autoimmune diseases such as lupus erythematosus and rheumatoid arthritis.

5 Multicellular organisms also use apoptosis to instruct cells with damaged nucleic acids (*e.g.*, DNA) to destroy themselves prior to becoming cancerous. Some cancer-causing viruses overcome this safeguard by reprogramming infected (transformed) cells to abort the normal apoptotic process. For example, several human papilloma viruses (HPVs) have been implicated in causing cervical cancer by suppressing the apoptotic removal of
10 transformed cells by producing a protein (E6) which inactivates the p53 apoptosis promoter. Similarly, the Epstein-Barr virus (EBV), the causative agent of mononucleosis and Burkitt's lymphoma, reprograms infected cells to produce proteins that prevent normal apoptotic removal of the aberrant cells thus allowing the cancerous cells to proliferate and to spread throughout the organism.

15 Still other viruses destructively manipulate a cell's apoptotic machinery without directly resulting in the development of a cancer. For example, the destruction of the immune system in individuals infected with the human immunodeficiency virus (HIV) is thought to progress through infected CD4⁺ T cells (about 1 in 100,000) instructing uninfected sister cells to undergo apoptosis.

20 Some cancers that arise by non-viral means have also developed mechanisms to escape destruction by apoptosis. Melanoma cells, for instance, avoid apoptosis by inhibiting the expression of the gene encoding Apaf-1. Other cancer cells, especially lung and colon cancer cells, secrete high levels of soluble decoy molecules that inhibit the initiation of CTL mediated clearance of aberrant cells. Faulty regulation of the apoptotic
25 machinery has also been implicated in various degenerative conditions and vascular diseases.

It is apparent that the controlled regulation of the apoptotic process and its cellular machinery is vital to the survival of multicellular organisms. Typically, the biochemical changes that occur in a cell instructed to undergo apoptosis occur in an orderly procession.
30 However, as shown above, flawed regulation of apoptosis can cause serious deleterious effects in the organism.

There have been various attempts to control and restore regulation of the apoptotic machinery in aberrant cells (*e.g.*, cancer cells). For example, much work has been done to

develop cytotoxic agents to destroy aberrant cells before they proliferate. As such, cytotoxic agents have widespread utility in both human and animal health and represent the first line of treatment for nearly all forms of cancer and hyperproliferative autoimmune disorders like lupus erythematosus and rheumatoid arthritis.

5 Many cytotoxic agents in clinical use exert their effect by damaging DNA (*e.g.*, cis-diaminodichroplatanim(II) cross-links DNA, whereas bleomycin induces strand cleavage). The result of this nuclear damage, if recognized by cellular factors like the p53 system, is to initiate an apoptotic cascade leading to the death of the damaged cell.

10 However, existing cytotoxic chemotherapeutic agents have serious drawbacks. For example, many known cytotoxic agents show little discrimination between healthy and diseased cells. This lack of specificity often results in severe side effects that can limit efficacy and/or result in early mortality. Moreover, prolonged administration of many existing cytotoxic agents results in the expression of resistance genes (*e.g.*, *bcl-2* family or multi-drug resistance (MDR) proteins) that render further dosing either less effective or
15 useless. Some cytotoxic agents induce mutations into p53 and related proteins. Based on these considerations, ideal cytotoxic drugs should only kill diseased cells and not be susceptible to chemo-resistance.

One strategy to selectively kill diseased cells is to develop drugs that selectively recognize molecules expressed in diseased cells. Thus, effective cytotoxic
20 chemotherapeutic agents, would recognize disease indicative molecules and induce (*e.g.*, either directly or indirectly) the death of the diseased cell. Although markers on some types of cancer cells have been identified and targeted with therapeutic antibodies and small molecules, unique traits for diagnostic and therapeutic exploitation are not known for most cancers. Moreover, for diseases like lupus, specific molecular targets for drug development
25 have not been identified.

What are needed are improved compositions and methods for regulating the apoptotic processes in subjects afflicted with diseases and conditions characterized by faulty regulation of these processes (*e.g.*, viral infections, hyperproliferative autoimmune disorders, chronic inflammatory conditions, and cancers).

30 SUMMARY

The present invention relates to systems and methods for generating new forms of benzodiazepine and benzodiazepine related compounds as well as new compounds and

formulations generated by such methods. In particular, the present invention provides high throughput systems and methods for generating and identifying new crystalline benzodiazepine and benzodiazepine related polymorphs and new unsolvated and other forms of the compounds that find use as improved drugs and drug formations.

5 For example, the present invention provides unsolvated forms of benzodiazepine compounds and methods of making such compounds. In some embodiments, the benzodiazepine compounds have orthorhombic crystalline forms. The unsolvated benzodiazepines of the present invention find use in pharmaceutical formulations with enhanced properties (e.g., shelf-life, tabletability, etc.). The present invention is illustrated
10 with the benzodiazepine, Bz-423. However, the present invention is not limited to this particular compound. It will be understood that a variety of related compounds find use in the compositions and methods of the present invention.

In some embodiments, the benzodiazepines of the present invention have orthorhombic crystals (e.g., Bz-423). In some preferred embodiments, the compounds are
15 anhydrous benzodiazepines, an ethanol solvate of benzodiazepines, a succinic acid (2:1) formulation of benzodiazepines, a citric acid (2:1) formulation of benzodiazepines, biphenyl derivate formulations of benzodiazepines, acetic acid formulations of benzodiazepine, CH₃CN formulations of benzodiazepine, methanol formulations of benzodiazepines, ethyl acetate formulations of benzodiazepines, toluene formulations of
20 benzodiazepines, oxalic acid formulations of benzodiazepines, fumaric acid formulations of benzodiazepines, octanol formulations of benzodiazepines, heptanoic acid formulations of benzodiazepines, diphenyl ether formulations of benzodiazepines, and trichlorobenzene formulations of benzodiazepines. Other solvated, unsolvated and salt forms may also be used.

25 In certain embodiments, the present invention provides a composition comprising a solution of dissolved benzodiazepine (e.g., Bz-423) in contact with a crystal form (e.g., orthorhombic) of the same benzodiazepine obtained from the solution, wherein the crystal form, when isolated, is capable of being provided in unsolvated form. In preferred embodiments, the composition further comprises a polymer surface in contact with the
30 crystals. In preferred embodiments, the solution comprises an aqueous solution.

In certain embodiments, the present invention provides a method for producing orthorhombic Bz-423 crystals comprising providing the above described composition and isolating the crystals.

In preferred embodiments, the method further comprises the step of preparing benzodiazepines to generated the above described compositions.

The present invention further provides methods of preparing a pharmaceutical preparation comprising new benzodiazepine compositions (alone, or in combination with other drugs or agents). In preferred embodiments, the pharmaceutical preparation comprises a tablet.

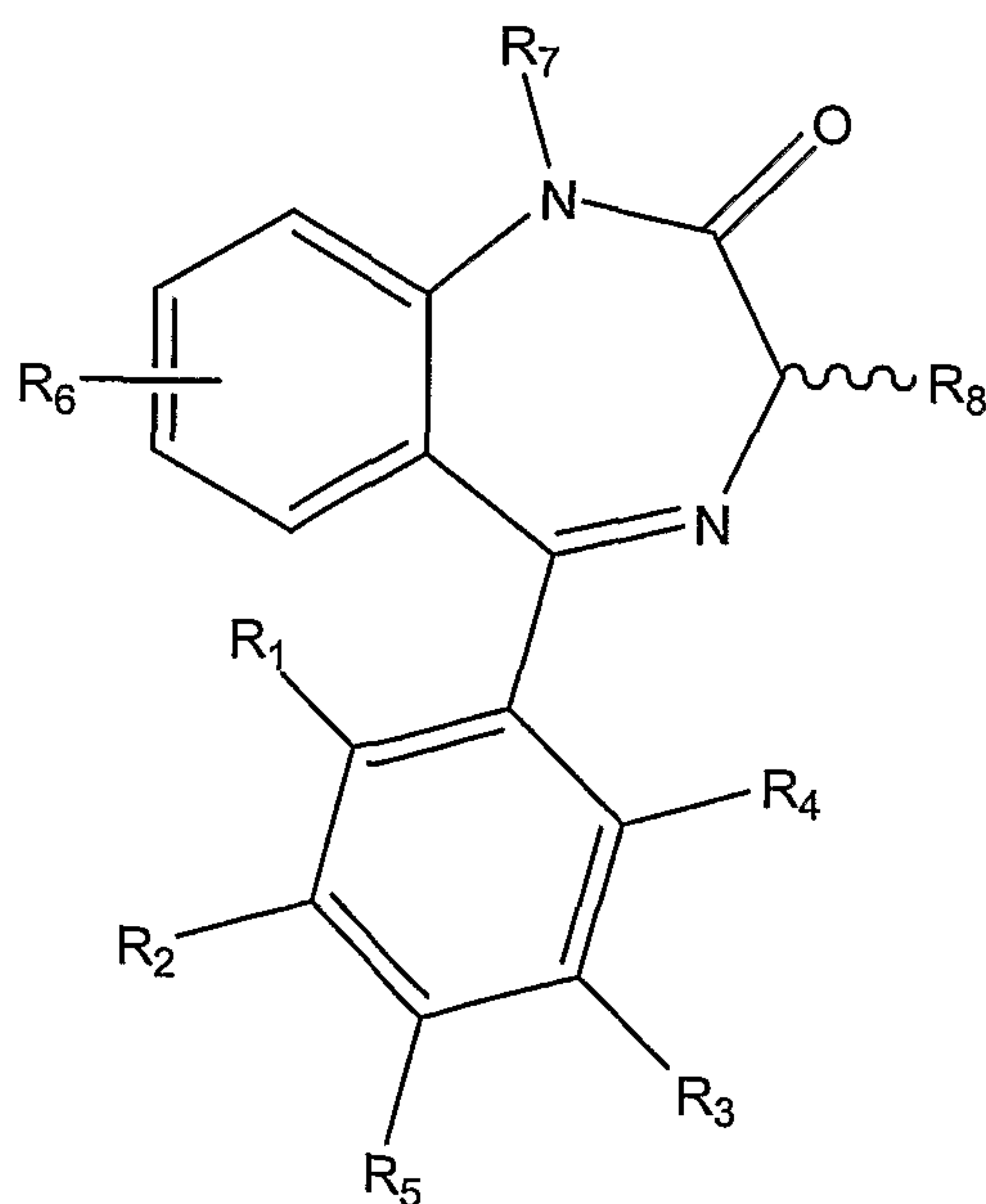
In preferred embodiments, the method further comprises the step of selling the pharmaceutical preparation, prescribing the pharmaceutical preparation to a patient, and/or administering the pharmaceutical preparation to a subject (e.g., human).

In certain embodiments, the present invention provides a method for producing a crystal form of a benzodiazepine described above comprising exposing a solution containing a benzodiazepine to a polymer under conditions that permit crystal formation.

In certain embodiments, the present invention provides a method of treating an autoimmune disorder or hyperproliferative disorder comprising administering to a subject an effective amount of a composition comprising the new benzodiazepine formulations described above.

In preferred embodiments, the composition comprises an oral dose of the new benzodiazepine formulations described above.

In certain embodiments, the present invention provides a composition comprising an unsolvated compound having the structure described by the following formula:

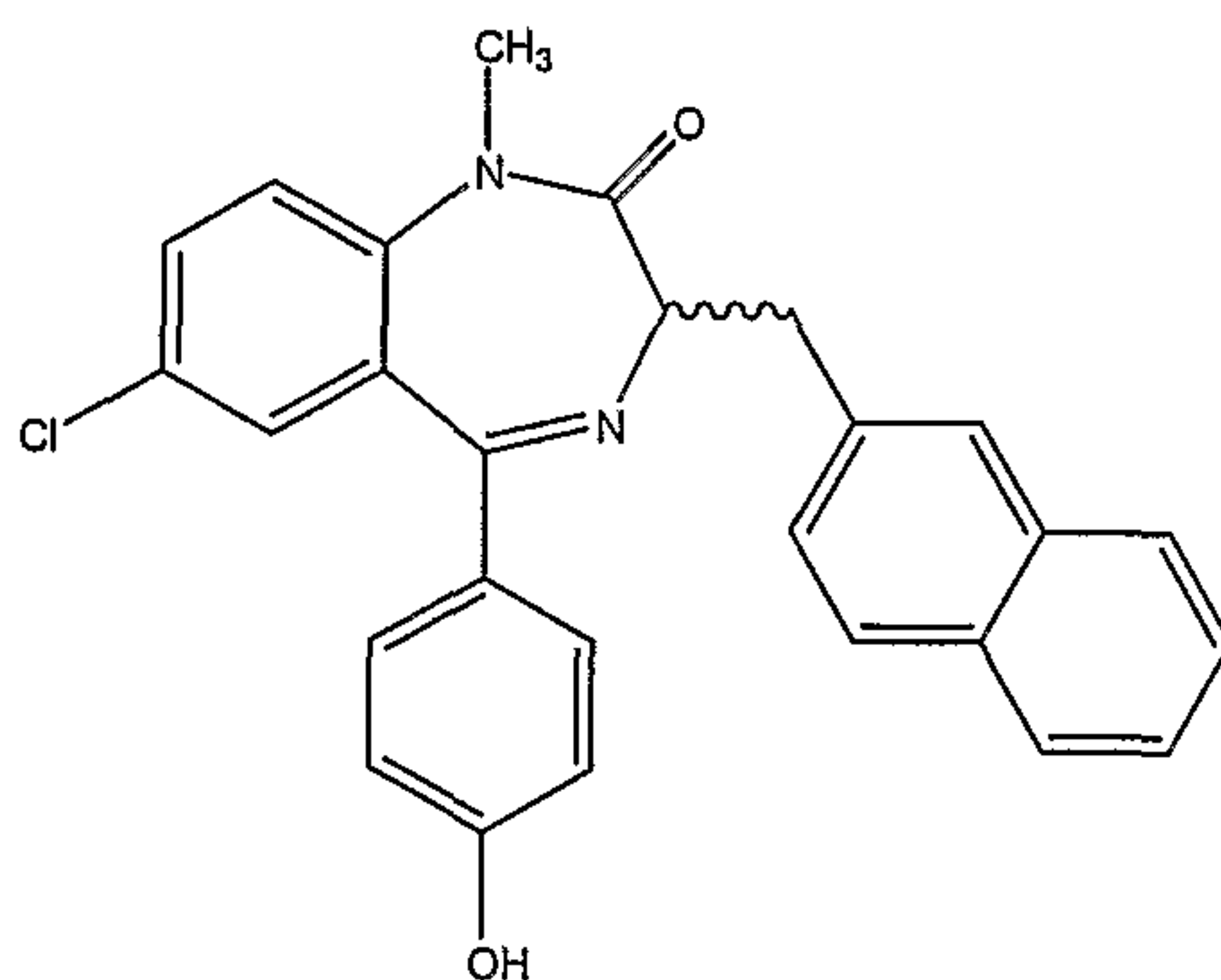


including both R and S enantiomeric forms and racemic mixtures; wherein R1, R2, R3 and R4 are selected from the group consisting of: hydrogen; CH₃; a linear or branched,

saturated or unsaturated aliphatic chain having at least 1 carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxy subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ether subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; wherein R5 is selected from the group consisting of: OH; NO₂; OR'; wherein R' is selected from the group consisting of: a linear or branched, saturated or unsaturated aliphatic chain having at least one carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxyl subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen

containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; wherein R6 is selected from the group consisting of: Hyrdrogen; NO₂; Cl; F; Br; I; SR'; and NR'₂; wherein R' is defined as above in R5; wherein R7 is selected from the group consisting of: Hydrogen; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; and wherein R8 is an aliphatic cyclic group larger than benzene; wherein the larger than benzene comprises any chemical group containing 7 or more non-hydrogen atoms.

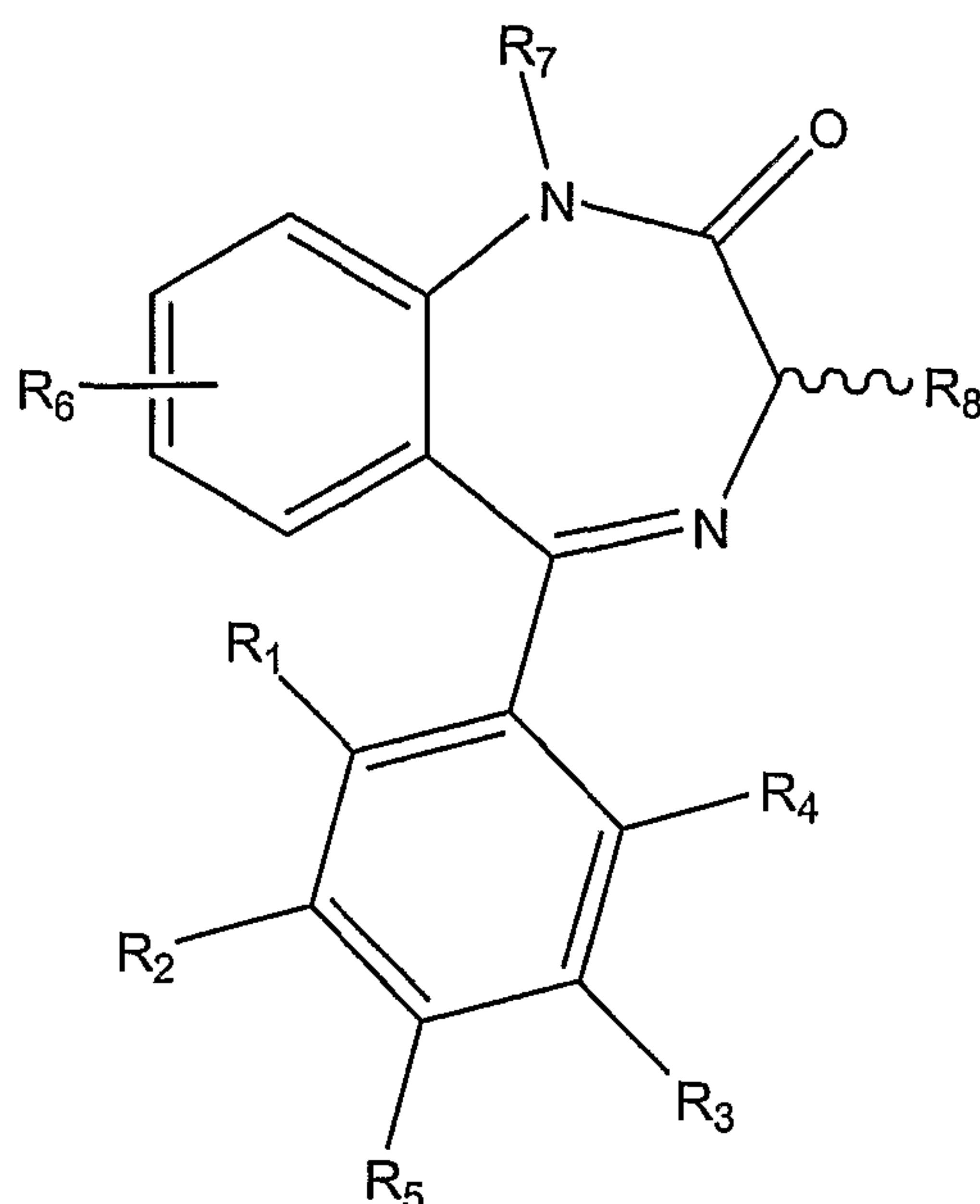
In preferred embodiments, the compound is:



In preferred embodiments, the unsolvated compound is anhydrous. In preferred embodiment, the unsolvated compound has an orthorhombic crystal structure.

In certain embodiments, the present invention provides a composition comprising a compound selected from the group consisting of Bz-423 ethanol solvate, Bz-423 succinic acid, Bz-423 citric acid, Bz-423 biphenyl derivate, BZ-423-acetic acid, BZ-423-CH₃CN, BZ-423-methanol, BZ-423-ethyl acetate, BZ-423-toluene, BZ-423-oxalic acid, BZ-423-fumaric acid, BZ-423-octanol, BZ-423-heptanoic acid, BZ-423-diphenyl ether, Bz-423 1-propanol solvate, Bz-423 2-propanol solvate, Bz-423 1-butanol solvate, Bz-423 2-butanol solvate, Bz-423 1-pentanol solvate, Bz-423 propylene glycol, , Bz-423 1-octanol solvate, Bz-423 acetone glass, and BZ-423-trichlorobenzene.

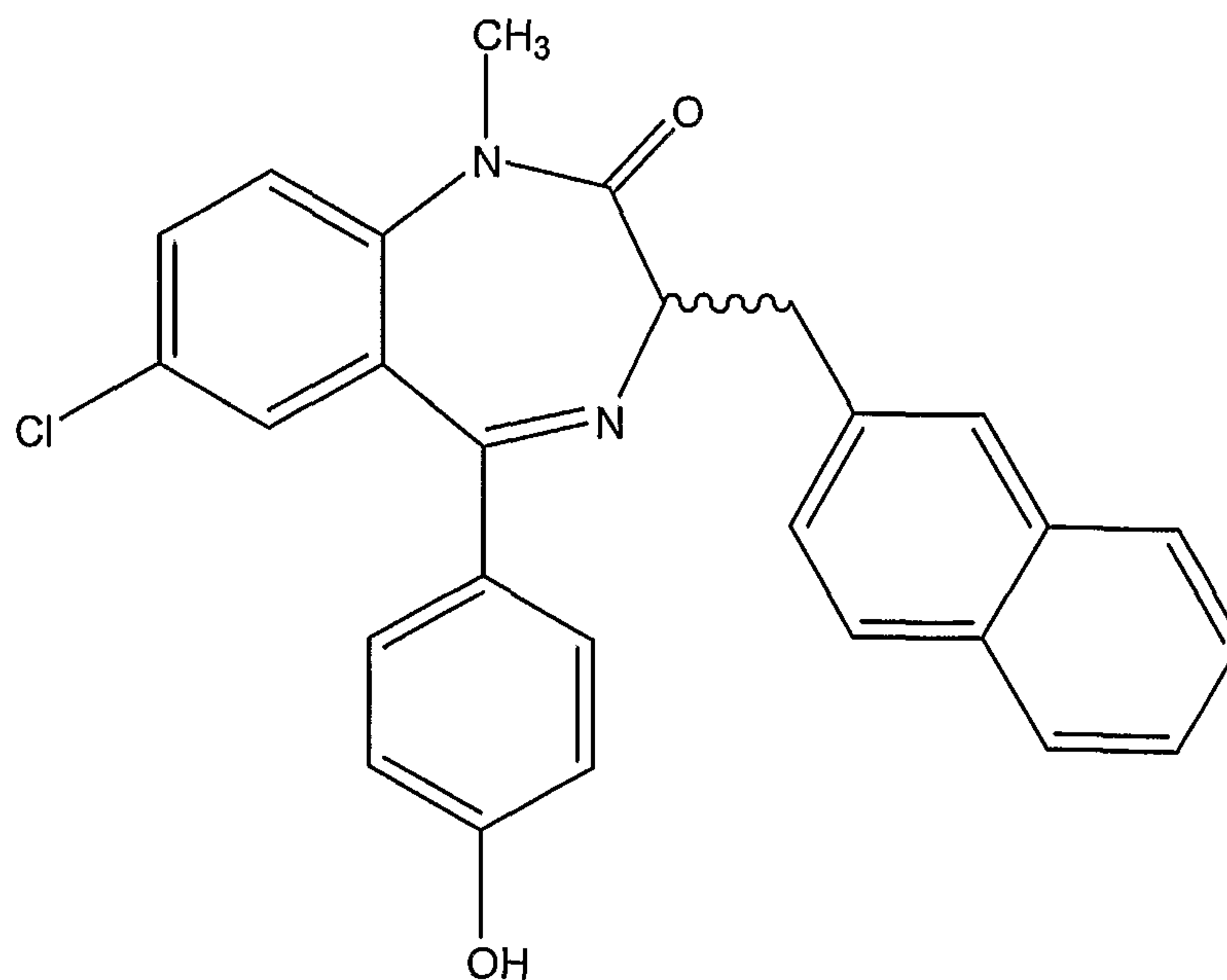
In certain embodiments, the present invention provides a composition comprising an orthorhombic benzodiazepine crystal, the benzodiazepine having the structure:



including both R and S enantiomeric forms and racemic mixtures; wherein R1, R2, R3 and R4 are selected from the group consisting of: hydrogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 1 carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxy subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ether subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; wherein R5 is selected from the group consisting of: OH; NO₂; OR'; wherein R' is selected from the group consisting of: a

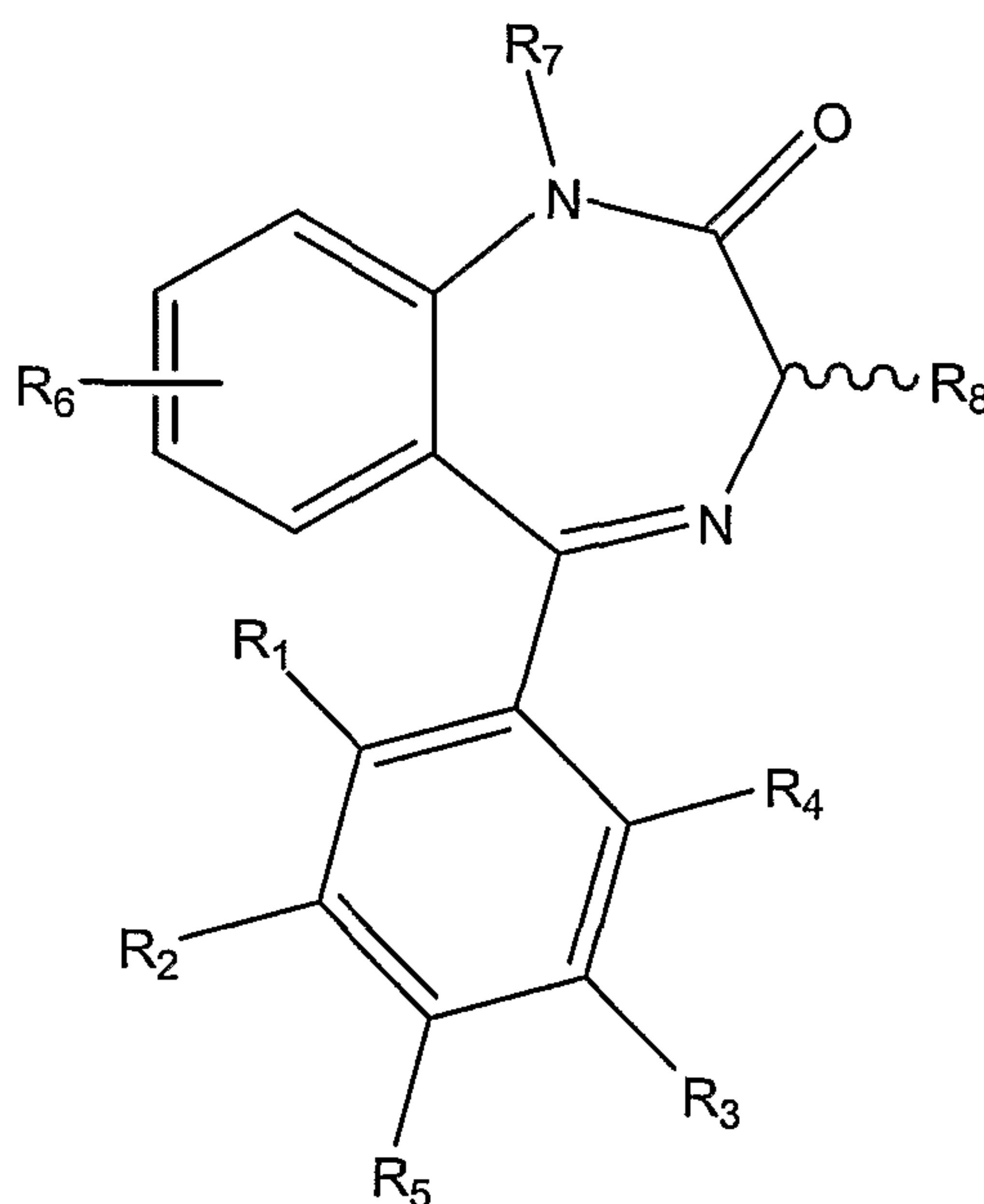
linear or branched, saturated or unsaturated aliphatic chain having at least one carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxyl subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or
5 branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or
10 branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at
15 least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; wherein R6 is selected from the group consisting of: Hyrdrogen; NO₂; Cl; F; Br; I; SR'; and NR'₂; wherein R' is defined as above
20 in R5; wherein R7 is selected from the group consisting of: Hydrogen; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; and wherein R8 is an aliphatic cyclic group larger than benzene; wherein the larger than benzene comprises any chemical group containing 7 or more non-hydrogen atoms.

In preferred embodiments, the compound is:



In preferred embodiments, the orthorhombic benzodiazepine crystal is anhydrous.

In certain embodiments, the present invention provides a composition comprising an
 5 oral dose of a benzodiazepine having the structure:



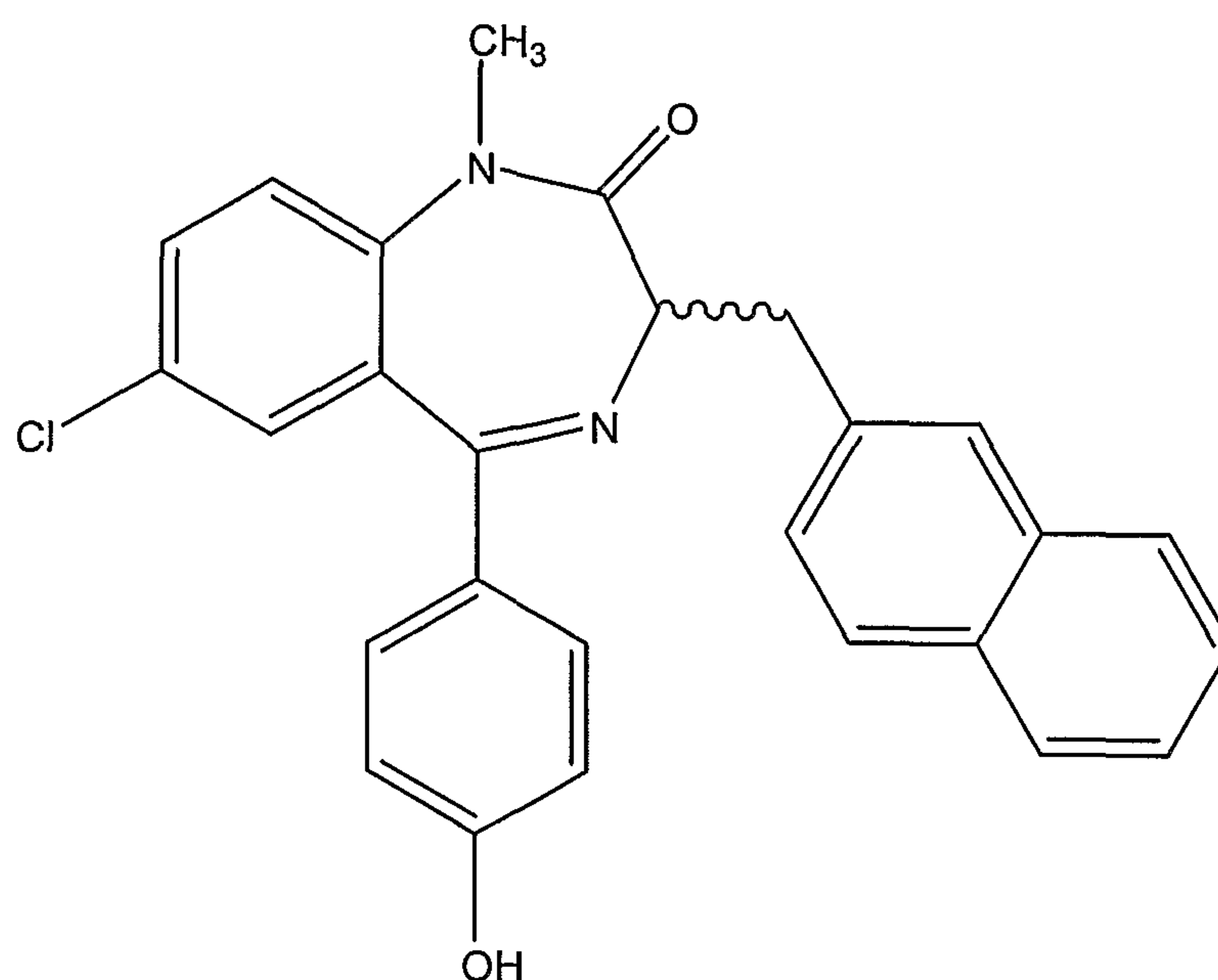
including both R and S enantiomeric forms and racemic mixtures; wherein R1, R2, R3 and
 R4 are selected from the group consisting of: hydrogen; CH₃; a linear or branched, saturated
 or unsaturated aliphatic chain having at least 1 carbon; a linear or branched, saturated or
 10 unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxy
 subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2
 carbons, and having at least one thiol subgroup; a linear or branched, saturated or
 unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates

with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ether subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; wherein R5 is selected from the group consisting of: OH; NO₂; OR'; wherein R' is selected from the group consisting of: a linear or branched, saturated or unsaturated aliphatic chain having at least one carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxyl subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; wherein R6 is selected from the group

consisting of: Hydrogen; NO₂; Cl; F; Br; I; SR'; and NR'₂; wherein R' is defined as above in R5; wherein R7 is selected from the group consisting of: Hydrogen; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; and wherein R8 is an aliphatic cyclic group larger than benzene; wherein the larger than benzene comprises any

5 chemical group containing 7 or more non-hydrogen atoms.

In preferred embodiments, the compound is:



In preferred embodiments, the benzodiazepine compound is anhydrous. In some

10 embodiments, the benzodiazepine compound has an orthorhombic crystal structure.

In certain embodiments, the present invention provides a method of treating an autoimmune disorder or hyperproliferative disorder comprising administering to a subject an effective amount of a composition comprising an unsolvated compound.

15 DESCRIPTION OF THE FIGURES

Figure 1 shows structural data of anhydrous Bz-423.

Figure 2 shows powder x-ray diffraction data for anhydrous Bz-423.

Figure 3 shows Raman spectroscopy data for anhydrous Bz-423.

Figure 4 shows structural data of Bz-423 ethanol solvate.

20 Figure 5 shows powder x-ray diffraction data for Bz-423 ethanol solvate.

Figure 6 shows Raman spectroscopy data for Bz-423 ethanol solvate.

Figure 7 shows Raman spectroscopy data for ball milled Bz-423 succinic acid (2:1).

Figure 8 shows Raman spectroscopy data for ball milled Bz-423 citric acid (2:1).

Figure 9 shows the structural data of Bz-423 biphenyl derivate.

Figure 10 shows solubility (e.g., absorbance) as a function of time for unsolvated Bz-423, anhydrous Bz-423, Bz-423 acetic acid, and Bz-423 citric acid.

Figure 11 shows a comparison of ATP hydrolysis between unsolvated Bz-423 and
5 solvated Bz-423.

Figure 12 shows a comparison of ATP synthesis between unsolvated Bz-423 and solvated Bz-423.

Figure 13 shows a comparison of cell viability between unsolvated Bz-423 and solvated Bz-423.

10 Figure 14 shows a UV-vis spectrum of Bz-423 in simulated gastric fluid.

Figure 15 shows a UV-vis spectrum of Bz-423 in simulated gastric fluid before and after addition of K_2CO_3 .

Figure 16 shows Raman spectroscopy data for Bz-423 ethanol solvate.

Figure 17 shows Raman spectroscopy data for Bz-423 1-propanol solvate.

15 Figure 18 shows Raman spectroscopy data for Bz-423 2-propanol solvate.

Figure 19 shows Raman spectroscopy data for Bz-423 1-butanol solvate.

Figure 20 shows Raman spectroscopy data for Bz-423 2-butanol solvate.

Figure 21 shows Raman spectroscopy data for Bz-423 1-pentanol solvate.

Figure 22 shows Raman spectroscopy data for Bz-423 1-octanol solvate.

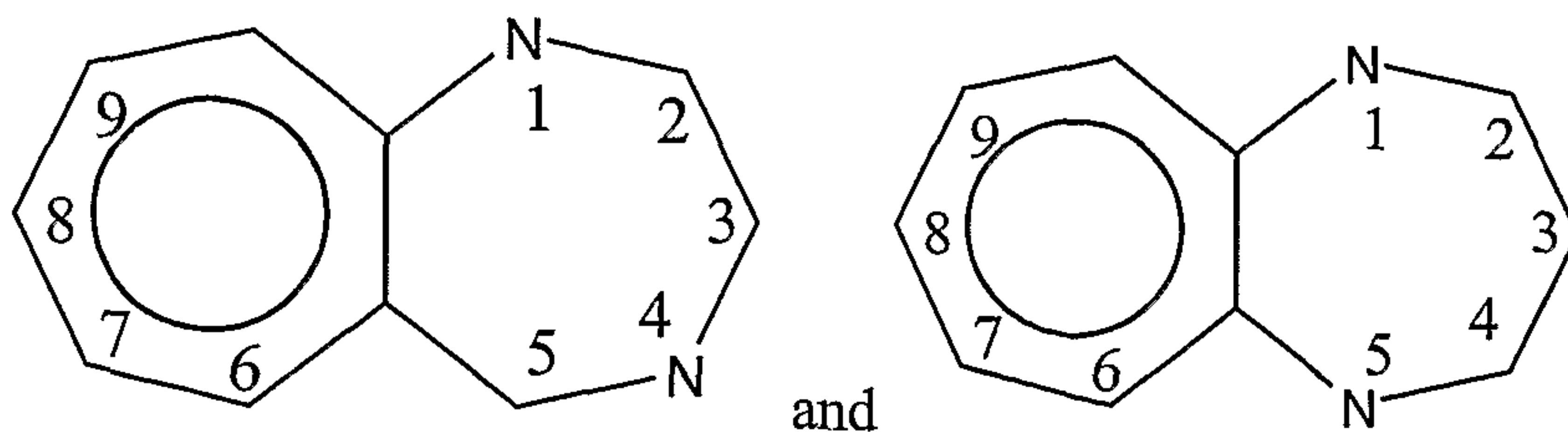
20 Figure 23 shows Raman spectroscopy data for Bz-423 propylene glycol solvate.

Figure 24 shows Raman spectroscopy data for Bz-423 acetone glass.

DEFINITIONS

To facilitate an understanding of the present invention, a number of terms and
25 phrases are defined below.

As used herein, the term "benzodiazepine" refers to a seven membered non-aromatic heterocyclic ring fused to a phenyl ring wherein the seven-membered ring has two nitrogen atoms, as part of the heterocyclic ring. In some aspects, the two nitrogen atoms are in the 1 and 4 positions or the 1 and 5 positions, as shown in the general structures below:



The term "larger than benzene" refers to any chemical group containing 7 or more non-hydrogen atoms.

As used herein, the term "polymorph" refers to a crystalline phase of a substance. Many substances feature polymorphism, which is the ability of a substance to exist as two or more crystalline phases that have different arrangements and/or conformations of the molecules in the crystal lattice. As used herein, the term polymorph includes amorphous phases and solvents/hydrates (i.e., pseudopolymorphs).

As used herein, the term "polymer library" refers to a composition comprising a plurality of different polymers positioned in particular locations so as to allow reactions to occur on the polymers at the particular locations. For example, containers or solid surfaces (e.g., plate, glass, metal, or ceramic surfaces, multi-well plates, dishes, vials, tubes, flasks, etc.) with a plurality of different polymers contained in discrete locations of the surface are polymer libraries. For example, a multi-well plate that contains a first polymer in a first well and a second polymer in a second well, etc. provides a polymer library.

As used herein, the term "tableability" refers to the capacity of a powdered material to be transformed into a tablet of specified strength under the effect of compaction pressure (Joiris et al., Pharm. Res., 15:1122 (1998); herein incorporated by reference in its entirety). Tableability describes the effectiveness of the applied pressure in increasing the tensile strength of the tablet and demonstrates the relationship between the cause, the compaction pressure, and the effect, the strength of the compact.

As used herein, the term "compressibility" refers to the ability of a material to undergo a reduction in volume as a result of an applied pressure (Joiris et al., Pharm. Res., 15:1122 (1998); herein incorporated by reference in its entirety). Compressibility indicates the ease with which a powder undergoes volume reduction under compaction pressure and is often represented by a plot showing the reduction of tablet porosity with increasing compaction pressure.

As used herein, the term "compactibility" refers to the ability of a material to produce tablets with sufficient strength under the effect of densification (Joiris et al., Pharm. Res., 15:1122 (1998); herein incorporated by reference in its entirety). Compactibility shows the tensile strength of tablets normalized by tablet porosity. In many cases, the tensile strength decreases exponentially with increasing porosity (Ryshkewitch, J. Am. Cer. Soc., 36:65 (1953); herein incorporated by reference in its entirety).

As used herein, the term “substituted aliphatic” refers to an alkane possessing less than 10 carbons where at least one of the aliphatic hydrogen atoms has been replaced by a halogen, an amino, a hydroxy, a nitro, a thio, a ketone, an aldehyde, an ester, an amide, a lower aliphatic, a substituted lower aliphatic, or a ring (aryl, substituted aryl, cycloaliphatic, or substituted cycloaliphatic, *etc.*). Examples of such include, but are not limited to, 1-chloroethyl and the like.

As used herein, the term “substituted aryl” refers to an aromatic ring or fused aromatic ring system consisting of no more than three fused rings at least one of which is aromatic, and where at least one of the hydrogen atoms on a ring carbon has been replaced by a halogen, an amino, a hydroxy, a nitro, a thio, a ketone, an aldehyde, an ester, an amide, a lower aliphatic, a substituted lower aliphatic, or a ring (aryl, substituted aryl, cycloaliphatic, or substituted cycloaliphatic). Examples of such include, but are not limited to, hydroxyphenyl and the like.

As used herein, the term “cycloaliphatic” refers to a cycloalkane possessing less than 8 carbons or a fused ring system consisting of no more than three fused cycloaliphatic rings. Examples of such include, but are not limited to, decalin and the like.

As used herein, the term “substituted cycloaliphatic” refers to a cycloalkane possessing less than 10 carbons or a fused ring system consisting of no more than three fused rings, and where at least one of the aliphatic hydrogen atoms has been replaced by a halogen, a nitro, a thio, an amino, a hydroxy, a ketone, an aldehyde, an ester, an amide, a lower aliphatic, a substituted lower aliphatic, or a ring (aryl, substituted aryl, cycloaliphatic, or substituted cycloaliphatic). Examples of such include, but are not limited to, 1-chlorodecalyl, bicyclo-heptanes, octanes, and nonanes (*e.g.*, nonbornyl) and the like.

As used herein, the term “heterocyclic” refers to a cycloalkane and/or an aryl ring system, possessing less than 8 carbons, or a fused ring system consisting of no more than three fused rings, where at least one of the ring carbon atoms is replaced by oxygen, nitrogen or sulfur. Examples of such include, but are not limited to, morpholino and the like.

As used herein, the term “substituted heterocyclic” refers to a cycloalkane and/or an aryl ring system, possessing less than 8 carbons, or a fused ring system consisting of no more than three fused rings, where at least one of the ring carbon atoms is replaced by oxygen, nitrogen or sulfur, and where at least one of the aliphatic hydrogen atoms has been replaced by a halogen, hydroxy, a thio, nitro, an amino, a ketone, an aldehyde, an ester, an

amide, a lower aliphatic, a substituted lower aliphatic, or a ring (aryl, substituted aryl, cycloaliphatic, or substituted cycloaliphatic). Examples of such include, but are not limited to 2-chloropyranyl.

As used herein, the term "linker" refers to a chain containing up to and including eight contiguous atoms connecting two different structural moieties where such atoms are, for example, carbon, nitrogen, oxygen, or sulfur. Ethylene glycol is one non-limiting example.

As used herein, the term "lower-alkyl-substituted-amino" refers to any alkyl unit containing up to and including eight carbon atoms where one of the aliphatic hydrogen atoms is replaced by an amino group. Examples of such include, but are not limited to, ethylamino and the like.

As used herein, the term "lower-alkyl-substituted-halogen" refers to any alkyl chain containing up to and including eight carbon atoms where one of the aliphatic hydrogen atoms is replaced by a halogen. Examples of such include, but are not limited to, chlorethyl and the like.

As used herein, the term "acetylamino" shall mean any primary or secondary amino that is acetylated. Examples of such include, but are not limited to, acetamide and the like.

The term "derivative" of a compound, as used herein, refers to a chemically modified compound wherein the chemical modification takes place either at a functional group of the compound or on the aromatic ring. Non-limiting examples of 1,4-benzodiazepine derivatives of the present invention may include N-acetyl, N-methyl, N-hydroxy groups at any of the available nitrogens in the compound. Additional derivatives may include those having a trifluoromethyl group on the phenyl ring.

The term "epidermal hyperplasia," as used herein, refers to an abnormal multiplication or increase in the number of normal cells in normal arrangement in epidermal tissue. Epidermal hyperplasia is a characteristic of numerous disorders, including but not limited to, psoriasis.

The term "keratinocyte" as used herein, refers to a skin cell of the keratinized layer of the epidermis.

The term "fibroblast" as used herein, refers to mesodermally derived resident cells of connective tissue that secrete fibrillar procollagen, fibronectin and collagenase.

The term “pigment disorder” as used herein, refers to disorders involving skin pigment (e.g., melanin). Examples of pigment disorders include, but are not limited to, all forms of albinism, melasma, pigment loss after skin damage, and vitiligo.

5 The term “stent” or “drug-eluting stent,” as used herein, refers to any device which when placed into contact with a site in the wall of a lumen to be treated, will also place fibrin at the lumen wall and retain it at the lumen wall. This can include especially devices delivered percutaneously to treat coronary artery occlusions and to seal dissections or aneurysms of splenic, carotid, iliac and popliteal vessels. The stent can also have underlying polymeric or metallic structural elements onto which the fibrin is applied or the stent can be
10 a composite of fibrin intermixed with a polymer. For example, a deformable metal wire stent such as that disclosed in U.S. Pat. No.: 4,886,062, herein incorporated by reference, could be coated with fibrin as set forth above in one or more coats (i.e., polymerization of fibrin on the metal framework by application of a fibrinogen solution and a solution of a fibrinogen-coagulating protein) or provided with an attached fibrin preform such as an
15 encircling film of fibrin. The stent and fibrin could then be placed onto the balloon at a distal end of a balloon catheter and delivered by conventional percutaneous means (e.g. as in an angioplasty procedure) to the site of the restriction or closure to be treated where it would then be expanded into contact with the body lumen by inflating the balloon. The catheter can then be withdrawn, leaving the fibrin stent of the present invention in place at
20 the treatment site. The stent may therefore provide both a supporting structure for the lumen at the site of treatment and also a structure supporting the secure placement of fibrin at the lumen wall. Generally, a drug-eluting stent allows for an active release of a particular drug at the stent implementation site.

As used herein, the term “catheter” refers generally to a tube used for gaining access
25 to a body cavity or blood vessel.

As used herein, the term “valve” or “vessel” refers to any lumen within a mammal. Examples include, but are not limited to, arteries, veins, capillaries, and biological lumen.

As used herein, the term “restenosis” refers to any valve which is narrowed. Examples include, but are not limited to, the reclosure of a peripheral or coronary artery
30 following trauma to that artery caused by efforts to open a stenosed portion of the artery, such as, for example, by balloon dilation, ablation, atherectomy or laser treatment of the artery.

As used herein, “angioplasty” or “balloon therapy” or “balloon angioplasty” or “percutaneous transluminal coronary angioplasty” refers to a method of treating blood vessel disorders that involves the use of a balloon catheter to enlarge the blood vessel and thereby improve blood flow.

5 As used herein, “cardiac catheterization” or “coronary angiogram” refers to a test used to diagnose coronary artery disease using a catheterization procedure. Such a procedure may involve, for example, the injection of a contrast dye into the coronary arteries via a catheter, permitting the visualization of a narrowed or blocked artery.

10 As used herein, the term “subject” refers to organisms to be treated by the methods of the present invention. Such organisms preferably include, but are not limited to, mammals (*e.g.*, murines, simians, equines, bovines, porcines, canines, felines, and the like), and most preferably includes humans. In the context of the invention, the term “subject” generally refers to an individual who will receive or who has received treatment (*e.g.*, administration of benzodiazepine compound(s), and optionally one or more other agents)
15 for a condition characterized by the dysregulation of apoptotic processes.

The term “diagnosed,” as used herein, refers to the recognition of a disease by its signs and symptoms (*e.g.*, resistance to conventional therapies), or genetic analysis, pathological analysis, histological analysis, and the like.

20 As used herein, the terms “anticancer agent,” or “conventional anticancer agent” refer to any chemotherapeutic compounds, radiation therapies, or surgical interventions, used in the treatment of cancer.

As used herein the term, “*in vitro*” refers to an artificial environment and to processes or reactions that occur within an artificial environment. *In vitro* environments include, but are not limited to, test tubes and cell cultures. The term “*in vivo*” refers to the
25 natural environment (*e.g.*, an animal or a cell) and to processes or reaction that occur within a natural environment.

As used herein, the term “host cell” refers to any eukaryotic or prokaryotic cell (*e.g.*, mammalian cells, avian cells, amphibian cells, plant cells, fish cells, and insect cells), whether located *in vitro* or *in vivo*.

30 As used herein, the term “cell culture” refers to any *in vitro* culture of cells. Included within this term are continuous cell lines (*e.g.*, with an immortal phenotype), primary cell cultures, finite cell lines (*e.g.*, non-transformed cells), and any other cell population maintained *in vitro*, including oocytes and embryos.

In preferred embodiments, the “target cells” of the compositions and methods of the present invention include, refer to, but are not limited to, lymphoid cells or cancer cells. Lymphoid cells include B cells, T cells, and granulocytes. Granulocytes include eosinophils and macrophages. In some embodiments, target cells are continuously cultured
5 cells or uncultured cells obtained from patient biopsies.

Cancer cells include tumor cells, neoplastic cells, malignant cells, metastatic cells, and hyperplastic cells. Neoplastic cells can be benign or malignant. Neoplastic cells are benign if they do not invade or metastasize. A malignant cell is one that is able to invade and/or metastasize. Hyperplasia is a pathologic accumulation of cells in a tissue or organ, without
10 significant alteration in structure or function.

In one specific embodiment, the target cells exhibit pathological growth or proliferation. As used herein, the term “pathologically proliferating or growing cells” refers to a localized population of proliferating cells in an animal that is not governed by the usual limitations of normal growth.

15 As used herein, the term “un-activated target cell” refers to a cell that is either in the G₀ phase or one in which a stimulus has not been applied.

As used herein, the term “activated target lymphoid cell” refers to a lymphoid cell that has been primed with an appropriate stimulus to cause a signal transduction cascade, or alternatively, a lymphoid cell that is not in G₀ phase. Activated lymphoid cells may
20 proliferate, undergo activation induced cell death, or produce one or more of cytotoxins, cytokines, and other related membrane-associated proteins characteristic of the cell type. They are also capable of recognizing and binding any target cell that displays a particular antigen on its surface, and subsequently releasing its effector molecules.

As used herein, the term “activated cancer cell” refers to a cancer cell that has been
25 primed with an appropriate stimulus to cause a signal transduction. An activated cancer cell may or may not be in the G₀ phase.

An activating agent is a stimulus that upon interaction with a target cell results in a signal transduction cascade. Examples of activating stimuli include, but are not limited to, small molecules, radiant energy, and molecules that bind to cell activation cell surface
30 receptors. Responses induced by activation stimuli can be characterized by changes in, among others, intracellular Ca²⁺, superoxide, or hydroxyl radical levels; the activity of enzymes like kinases or phosphatases; or the energy state of the cell. For cancer cells, activating agents also include transforming oncogenes.

In one aspect, the activating agent is any agent that binds to a cell surface activation receptor. These can be selected from the group consisting of, but not limited to, a T cell receptor ligand, a B cell activating factor, a TNF, a Fas ligand, a proliferation inducing ligand, a cytokine, a chemokine, a hormone, an amino acid, a steroid, a B cell receptor ligand, gamma irradiation, UV irradiation, an agent or condition that enhances cell stress, or an antibody that specifically recognizes and binds a cell surface activation receptor. Antibodies include monoclonal or polyclonal or a mixture thereof.

Examples of a T cell ligand include, but are not limited to, a peptide that binds to an MHC molecule, a peptide MHC complex, or an antibody that recognizes components of the T cell receptor.

Examples of a B cell ligand include, but are not limited to, a molecule or antibody that binds to or recognizes components of the B cell receptor.

Examples of agents or conditions that enhance cell stress include heat, radiation, oxidative stress, or growth factor withdrawal and the like. Examples of growth factors include, but are not limited to serum, IL-2, platelet derived growth factor ("PDGF"), and the like.

As used herein, the term "effective amount" refers to the amount of a compound (*e.g.*, benzodiazepine) sufficient to effect beneficial or desired results. An effective amount can be administered in one or more administrations, applications or dosages and is not limited intended to be limited to a particular formulation or administration route.

As used herein, the term "dysregulation of the process of cell death" refers to any aberration in the ability of (*e.g.*, predisposition) a cell to undergo cell death via either necrosis or apoptosis. Dysregulation of cell death is associated with or induced by a variety of conditions, including for example, autoimmune disorders (*e.g.*, systemic lupus erythematosus, rheumatoid arthritis, graft-versus-host disease, myasthenia gravis, Sjögren's syndrome, *etc.*), chronic inflammatory conditions (*e.g.*, psoriasis, asthma and Crohn's disease), hyperproliferative disorders (*e.g.*, tumors, B cell lymphomas, T cell lymphomas, *etc.*), viral infections (*e.g.*, herpes, papilloma, HIV), and other conditions such as osteoarthritis and atherosclerosis.

It should be noted that when the dysregulation is induced by or associated with a viral infection, the viral infection may or may not be detectable at the time dysregulation occurs or is observed. That is, viral-induced dysregulation can occur even after the disappearance of symptoms of viral infection.

A "hyperproliferative disorder," as used herein refers to any condition in which a localized population of proliferating cells in an animal is not governed by the usual limitations of normal growth. Examples of hyperproliferative disorders include tumors, neoplasms, lymphomas and the like. A neoplasm is said to be benign if it does not undergo, invasion or metastasis and malignant if it does either of these. A metastatic cell or tissue means that the cell can invade and destroy neighboring body structures. Hyperplasia is a form of cell proliferation involving an increase in cell number in a tissue or organ, without significant alteration in structure or function. Metaplasia is a form of controlled cell growth in which one type of fully differentiated cell substitutes for another type of differentiated cell. Metaplasia can occur in epithelial or connective tissue cells. A typical metaplasia involves a somewhat disorderly metaplastic epithelium.

The pathological growth of activated lymphoid cells often results in an autoimmune disorder or a chronic inflammatory condition. As used herein, the term "autoimmune disorder" refers to any condition in which an organism produces antibodies or immune cells which recognize the organism's own molecules, cells or tissues. Non-limiting examples of autoimmune disorders include autoimmune hemolytic anemia, autoimmune hepatitis, Berger's disease or IgA nephropathy, Celiac Sprue, chronic fatigue syndrome, Crohn's disease, dermatomyositis, fibromyalgia, graft versus host disease, Grave's disease, Hashimoto's thyroiditis, idiopathic thrombocytopenia purpura, lichen planus, multiple sclerosis, myasthenia gravis, psoriasis, rheumatic fever, rheumatic arthritis, scleroderma, Sjorgren syndrome, systemic lupus erythematosus, type 1 diabetes, ulcerative colitis, vitiligo, and the like.

As used herein, the term "chronic inflammatory condition" refers to a condition wherein the organism's immune cells are activated. Such a condition is characterized by a persistent inflammatory response with pathologic sequelae. This state is characterized by infiltration of mononuclear cells, proliferation of fibroblasts and small blood vessels, increased connective tissue, and tissue destruction. Examples of chronic inflammatory diseases include, but are not limited to, Crohn's disease, psoriasis, chronic obstructive pulmonary disease, inflammatory bowel disease, multiple sclerosis, and asthma. Autoimmune diseases such as rheumatoid arthritis and systemic lupus erythematosus can also result in a chronic inflammatory state.

As used herein, the term "co-administration" refers to the administration of at least two agent(s) (*e.g.*, benzodiazepines) or therapies to a subject. In some embodiments, the co-

administration of two or more agents/therapies is concurrent. In other embodiments, a first agent/therapy is administered prior to a second agent/therapy. Those of skill in the art understand that the formulations and/or routes of administration of the various agents/therapies used may vary. The appropriate dosage for co-administration can be readily determined by one skilled in the art. In some embodiments, when agents/therapies are co-administered, the respective agents/therapies are administered at lower dosages than appropriate for their administration alone. Thus, co-administration is especially desirable in embodiments where the co-administration of the agents/therapies lowers the requisite dosage of a known potentially harmful (*e.g.*, toxic) agent(s).

As used herein, the term “toxic” refers to any detrimental or harmful effects on a cell or tissue as compared to the same cell or tissue prior to the administration of the toxicant.

As used herein, the term “pharmaceutical composition” refers to the combination of an active agent with a carrier, inert or active, making the composition especially suitable for diagnostic or therapeutic use *in vivo*, *in vivo* or *ex vivo*.

As used herein, the term “pharmaceutically acceptable carrier” refers to any of the standard pharmaceutical carriers, such as a phosphate buffered saline solution, water, emulsions (*e.g.*, such as an oil/water or water/oil emulsions), and various types of wetting agents. The compositions also can include stabilizers and preservatives. For examples of carriers, stabilizers and adjuvants. (*See e.g.*, Martin, Remington's Pharmaceutical Sciences, 15th Ed., Mack Publ. Co., Easton, PA [1975]).

As used herein, the term “pharmaceutically acceptable salt” refers to any pharmaceutically acceptable salt (*e.g.*, acid or base) of a compound of the present invention which, upon administration to a subject, is capable of providing a compound of this invention or an active metabolite or residue thereof. As is known to those of skill in the art, “salts” of the compounds of the present invention may be derived from inorganic or organic acids and bases. Examples of acids include, but are not limited to, hydrochloric, hydrobromic, sulfuric, nitric, perchloric, fumaric, maleic, phosphoric, glycolic, lactic, salicylic, succinic, toluene-p-sulfonic, tartaric, acetic, citric, methanesulfonic, ethanesulfonic, formic, benzoic, malonic, naphthalene-2-sulfonic, benzenesulfonic acid, and the like. Other acids, such as oxalic, while not in themselves pharmaceutically acceptable, may be employed in the preparation of salts useful as intermediates in obtaining the compounds of the invention and their pharmaceutically acceptable acid addition salts.

Examples of bases include, but are not limited to, alkali metals (*e.g.*, sodium) hydroxides, alkaline earth metals (*e.g.*, magnesium), hydroxides, ammonia, and compounds of formula NW_4^+ , wherein W is C_{1-4} alkyl, and the like.

Examples of salts include, but are not limited to: acetate, adipate, alginate, aspartate,
5 benzoate, benzenesulfonate, bisulfate, butyrate, citrate, camphorate, camphorsulfonate, cyclopentanepropionate, digluconate, dodecylsulfate, ethanesulfonate, fumarate, flucoheptanoate, glycerophosphate, hemisulfate, heptanoate, hexanoate, hydrochloride, hydrobromide, hydroiodide, 2-hydroxyethanesulfonate, lactate, maleate, methanesulfonate, 2-naphthalenesulfonate, nicotinate, oxalate, palmoate, pectinate, persulfate,
10 phenylpropionate, picrate, pivalate, propionate, succinate, tartrate, thiocyanate, tosylate, undecanoate, and the like. Other examples of salts include anions of the compounds of the present invention compounded with a suitable cation such as Na^+ , NH_4^+ , and NW_4^+ (wherein W is a C_{1-4} alkyl group), and the like.

For therapeutic use, salts of the compounds of the present invention are
15 contemplated as being pharmaceutically acceptable. However, salts of acids and bases that are non-pharmaceutically acceptable may also find use, for example, in the preparation or purification of a pharmaceutically acceptable compound.

As used herein, the terms "solid phase supports" or "solid supports," are used in their broadest sense to refer to a number of supports that are available and known to those of
20 ordinary skill in the art. Solid phase supports include, but are not limited to, silica gels, resins, derivatized plastic films, glass beads, cotton, plastic beads, alumina gels, and the like. As used herein, "solid supports" also include synthetic antigen-presenting matrices, cells, liposomes, and the like. A suitable solid phase support may be selected on the basis of desired end use and suitability for various protocols. For example, for peptide synthesis,
25 solid phase supports may refer to resins such as polystyrene (*e.g.*, PAM-resin obtained from Bachem, Inc., Peninsula Laboratories, *etc.*), POLYHIPE) resin (obtained from Aminotech, Canada), polyamide resin (obtained from Peninsula Laboratories), polystyrene resin grafted with polyethylene glycol (TENTAGEL, Rapp Polymere, Tubingen, Germany) or polydimethylacrylamide resin (obtained from Milligen/Biosearch, California).

30 As used herein, the term "pathogen" refers a biological agent that causes a disease state (*e.g.*, infection, cancer, *etc.*) in a host. "Pathogens" include, but are not limited to, viruses, bacteria, archaea, fungi, protozoans, mycoplasma, prions, and parasitic organisms.

The terms "bacteria" and "bacterium" refer to all prokaryotic organisms, including those within all of the phyla in the Kingdom Procaryotae. It is intended that the term encompass all microorganisms considered to be bacteria including Mycoplasma, Chlamydia, Actinomyces, Streptomyces, and Rickettsia. All forms of bacteria are included within this definition including cocci, bacilli, spirochetes, spheroplasts, protoplasts, *etc.* Also included within this term are prokaryotic organisms which are gram negative or gram positive. "Gram negative" and "gram positive" refer to staining patterns with the Gram-staining process which is well known in the art. (*See e.g.*, Finegold and Martin, Diagnostic Microbiology, 6th Ed., CV Mosby St. Louis, *pp.* 13-15 [1982]). "Gram positive bacteria" are bacteria which retain the primary dye used in the Gram stain, causing the stained cells to appear dark blue to purple under the microscope. "Gram negative bacteria" do not retain the primary dye used in the Gram stain, but are stained by the counterstain. Thus, gram negative bacteria appear red.

As used herein, the term "microorganism" refers to any species or type of microorganism, including but not limited to, bacteria, archaea, fungi, protozoans, mycoplasma, and parasitic organisms. The present invention contemplates that a number of microorganisms encompassed therein will also be pathogenic to a subject.

As used herein, the term "fungi" is used in reference to eukaryotic organisms such as the molds and yeasts, including dimorphic fungi.

As used herein, the term "virus" refers to minute infectious agents, which with certain exceptions, are not observable by light microscopy, lack independent metabolism, and are able to replicate only within a living host cell. The individual particles (*i.e.*, virions) typically consist of nucleic acid and a protein shell or coat; some virions also have a lipid containing membrane. The term "virus" encompasses all types of viruses, including animal, plant, phage, and other viruses.

The term "sample" as used herein is used in its broadest sense. A sample suspected of indicating a condition characterized by the dysregulation of apoptotic function may comprise a cell, tissue, or fluids, chromosomes isolated from a cell (*e.g.*, a spread of metaphase chromosomes), genomic DNA (in solution or bound to a solid support such as for Southern blot analysis), RNA (in solution or bound to a solid support such as for Northern blot analysis), cDNA (in solution or bound to a solid support) and the like. A sample suspected of containing a protein may comprise a cell, a portion of a tissue, an extract containing one or more proteins and the like.

As used herein, the terms "purified" or "to purify" refer, to the removal of undesired components from a sample. As used herein, the term "substantially purified" refers to molecules that are at least 60% free, preferably 75% free, and most preferably 90%, or more, free from other components with which they usually associated.

5 As used herein, the term "antigen binding protein" refers to proteins which bind to a specific antigen. "Antigen binding proteins" include, but are not limited to, immunoglobulins, including polyclonal, monoclonal, chimeric, single chain, and humanized antibodies, Fab fragments, F(ab')₂ fragments, and Fab expression libraries. Various procedures known in the art are used for the production of polyclonal antibodies. For the
10 production of antibody, various host animals can be immunized by injection with the peptide corresponding to the desired epitope including but not limited to rabbits, mice, rats, sheep, goats, *etc.* In a preferred embodiment, the peptide is conjugated to an immunogenic carrier (*e.g.*, diphtheria toxoid, bovine serum albumin (BSA), or keyhole limpet hemocyanin [KLH]). Various adjuvants are used to increase the immunological response, depending on
15 the host species, including but not limited to Freund's (complete and incomplete), mineral gels such as aluminum hydroxide, surface active substances such as lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, keyhole limpet hemocyanins, dinitrophenol, and potentially useful human adjuvants such as BCG (Bacille Calmette-Guerin) and *Corynebacterium parvum*.

20 For preparation of monoclonal antibodies, any technique that provides for the production of antibody molecules by continuous cell lines in culture may be used (*See e.g.*, Harlow and Lane, *Antibodies: A Laboratory Manual*, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY). These include, but are not limited to, the hybridoma technique originally developed by Köhler and Milstein (Köhler and Milstein, *Nature*, 256:495-497
25 [1975]), as well as the trioma technique, the human B-cell hybridoma technique (*See e.g.*, Kozbor *et al.*, *Immunol. Today*, 4:72 [1983]), and the EBV-hybridoma technique to produce human monoclonal antibodies (Cole *et al.*, in *Monoclonal Antibodies and Cancer Therapy*, Alan R. Liss, Inc., *pp.* 77-96 [1985]).

According to the invention, techniques described for the production of single chain
30 antibodies (U.S. 4,946,778; herein incorporated by reference) can be adapted to produce specific single chain antibodies as desired. An additional embodiment of the invention utilizes the techniques known in the art for the construction of Fab expression libraries

(Huse *et al.*, Science, 246:1275-1281 [1989]) to allow rapid and easy identification of monoclonal Fab fragments with the desired specificity.

Antibody fragments that contain the idiotype (antigen binding region) of the antibody molecule can be generated by known techniques. For example, such fragments
5 include but are not limited to: the F(ab')₂ fragment that can be produced by pepsin digestion of an antibody molecule; the Fab' fragments that can be generated by reducing the disulfide bridges of an F(ab')₂ fragment, and the Fab fragments that can be generated by treating an antibody molecule with papain and a reducing agent.

Genes encoding antigen binding proteins can be isolated by methods known in the
10 art. In the production of antibodies, screening for the desired antibody can be accomplished by techniques known in the art (*e.g.*, radioimmunoassay, ELISA (enzyme-linked immunosorbant assay), "sandwich" immunoassays, immunoradiometric assays, gel diffusion precipitin reactions, immunodiffusion assays, *in situ* immunoassays (using colloidal gold, enzyme or radioisotope labels, for example), Western Blots, precipitation
15 reactions, agglutination assays (*e.g.*, gel agglutination assays, hemagglutination assays, *etc.*), complement fixation assays, immunofluorescence assays, protein A assays, and immunoelectrophoresis assays, *etc.*) *etc.*

As used herein, the term "immunoglobulin" or "antibody" refer to proteins that bind a specific antigen. Immunoglobulins include, but are not limited to, polyclonal,
20 monoclonal, chimeric, and humanized antibodies, Fab fragments, F(ab')₂ fragments, and includes immunoglobulins of the following classes: IgG, IgA, IgM, IgD, IgE, and secreted immunoglobulins (sIg). Immunoglobulins generally comprise two identical heavy chains and two light chains. However, the terms "antibody" and "immunoglobulin" also encompass single chain antibodies and two chain antibodies.

25 The term "epitope" as used herein refers to that portion of an antigen that makes contact with a particular immunoglobulin. When a protein or fragment of a protein is used to immunize a host animal, numerous regions of the protein may induce the production of antibodies which bind specifically to a given region or three-dimensional structure on the protein; these regions or structures are referred to as "antigenic determinants". An antigenic
30 determinant may compete with the intact antigen (*i.e.*, the "immunogen" used to elicit the immune response) for binding to an antibody.

The terms "specific binding" or "specifically binding" when used in reference to the interaction of an antibody and a protein or peptide means that the interaction is dependent

upon the presence of a particular structure (*i.e.*, the antigenic determinant or epitope) on the protein; in other words the antibody is recognizing and binding to a specific protein structure rather than to proteins in general. For example, if an antibody is specific for epitope "A," the presence of a protein containing epitope A (or free, unlabelled A) in a reaction containing labeled "A" and the antibody will reduce the amount of labeled A bound to the antibody.

As used herein, the terms "non-specific binding" and "background binding" when used in reference to the interaction of an antibody and a protein or peptide refer to an interaction that is not dependent on the presence of a particular structure (*i.e.*, the antibody is binding to proteins in general rather than a particular structure such as an epitope).

As used herein, the term "modulate" refers to the activity of a compound (*e.g.*, benzodiazepine compound) to affect (*e.g.*, to promote or retard) an aspect of cellular function, including, but not limited to, cell growth, proliferation, apoptosis, and the like.

As used herein, the term "competes for binding" is used in reference to a first molecule (*e.g.*, a first benzodiazepine derivative) with an activity that binds to the same substrate (*e.g.*, the oligomycin sensitivity conferring protein in mitochondrial ATP synthase) as does a second molecule (*e.g.*, a second benzodiazepine derivative or other molecule that binds to the oligomycin sensitivity conferring protein in mitochondrial ATP synthase, *etc.*). The efficiency (*e.g.*, kinetics or thermodynamics) of binding by the first molecule may be the same as, or greater than, or less than, the efficiency of the substrate binding to the second molecule. For example, the equilibrium binding constant (K_D) for binding to the substrate may be different for the two molecules.

As used herein, the term "instructions for administering said compound to a subject," and grammatical equivalents thereof, includes instructions for using the compositions contained in a kit for the treatment of conditions characterized by the dysregulation of apoptotic processes in a cell or tissue (*e.g.*, providing dosing, route of administration, decision trees for treating physicians for correlating patient-specific characteristics with therapeutic courses of action). The term also specifically refers to instructions for using the compositions contained in the kit to treat autoimmune disorders (*e.g.*, systemic lupus erythematosus, rheumatoid arthritis, graft-versus-host disease, myasthenia gravis, Sjögren's syndrome, *etc.*), chronic inflammatory conditions (*e.g.*, psoriasis, asthma and Crohn's disease), hyperproliferative disorders (*e.g.*, tumors, B cell

lymphomas, T cell lymphomas, *etc.*), viral infections (*e.g.*, herpes virus, papilloma virus, HIV), and other conditions such as osteoarthritis and atherosclerosis, and the like.

The term “test compound” refers to any chemical entity, pharmaceutical, drug, and the like, that can be used to treat or prevent a disease, illness, sickness, or disorder of bodily function, or otherwise alter the physiological or cellular status of a sample (*e.g.*, the level of dysregulation of apoptosis in a cell or tissue). Test compounds comprise both known and potential therapeutic compounds. A test compound can be determined to be therapeutic by using the screening methods of the present invention. A “known therapeutic compound” refers to a therapeutic compound that has been shown (*e.g.*, through animal trials or prior experience with administration to humans) to be effective in such treatment or prevention. In preferred embodiments, “test compounds” are agents that modulate apoptosis in cells.

As used herein, the term “third party” refers to any entity engaged in selling, warehousing, distributing, or offering for sale a test compound contemplated for administered with a compound for treating conditions characterized by the dysregulation of apoptotic processes.

GENERAL DESCRIPTION OF THE INVENTION

Pharmaceutical companies expend much of their resources in attempts to find new blockbuster drugs (greater than \$1 billion/year sales) (D. Eric Walters, BC 5220, Techniques in Biomedical Research, “The Rational Basis of Drug Design”). In order to be successful, a new drug should satisfy several criteria: safe to use; effective for the intended use; stable (chemically and metabolically); good solubility profile; synthetically feasible; and novel (*i.e.*, patentable). An important aspect of drug development is the identification of leads. A lead is any chemical compound that shows the biological activity sought. A lead is not the same as a drug however—as it should meet the criteria listed above prior to use as a drug. There are two broad tasks in drug discovery. The first is lead-finding. Here the task is to find a chemical compound that has a desired bioactivity. The second is lead-optimization, modifying the lead structure to build in the other desirable properties (safety, solubility, stability, *etc.*).

There are many ways to find lead compounds. In the beginning, plants and other natural products were the source of most medicinal substances. As the science of medicinal chemistry evolved, it was discovered that the plants and natural products contained specific

compounds that are responsible for the therapeutic effect. It became possible to isolate the active components, so that dosage could be more accurately regulated.

Other medicines came about because of accidental observations and discoveries (e.g., penicillin). The discovery of penicillin led to a large-scale screening effort, in which
5 thousands of soil microorganisms were grown and tested to see whether they could produce other substances that kill bacteria. Antibiotics such as streptomycin, neomycin, gentamicin, erythromycin, and the tetracyclines resulted from these efforts.

Chemical modification of known drugs can often lead to improved drugs. For example, naturally occurring penicillin G is broken down by bacterial beta-lactamases.
10 Addition of two $-OCH_3$ groups produces methicillin, which is resistant to lactamase. Another example of chemical modification is found in the opiate analgesics. The parent compound is morphine, which occurs in opium poppies. Morphine is a powerful analgesic, but it has serious side effects: respiratory depression, constipation, and dependence liability. Thousands of analogs (related chemical structures) have been synthesized in an
15 effort to find analgesics with lower incidence of side effects. In some cases, small changes in chemical structure may have a big influence on the activity. For example, nalorphine is a partial agonist (shows some morphine-like activity, and at higher concentration, antagonizes morphine effects), and naloxone is an antagonist. Considerable simplification of the molecule is possible. For example, meperidine has only two rings instead of four, but it
20 maintains strong analgesic activity. It has better oral absorption than morphine, and shows less GI side effects. Methadone is an analgesic in which the original piperidine ring (6-membered ring containing a nitrogen atom) is completely absent. It retains analgesic activity, has good oral activity, and has a long half-life in the body. Dextromethorphan is constructed on a mirror image of the morphine ring system. It has no opiate analgesic
25 effects or side effects, but is a useful anti-tussive agent.

Some drugs are discovered by observing side effects of existing drugs. For example, minoxidil was found to grow hair on bald men as a side effect in a study of its antihypertensive effects. Viagra's effect on penile dysfunction was discovered in clinical trials for treatment of angina; it had originally been designed as an antihypertensive drug.

30 In the modern era, most leads are discovered using various screening processes. For example, over a couple of decades, the National Cancer Institute has put hundreds of thousands of different chemical compounds through a battery of anti-cancer assays. Current screening assays often employ miniaturization and automation with robots for high

throughput screening, allowing hundreds of thousands of compounds to be screened in a short period of time.

Structure-based molecular design is yet another method to identify lead molecules for drug design. This method is based on the premise that desired drug candidates possess significant structural and chemical complementarity with their target molecules. This design method can create molecules with specific properties that make them conducive for binding to the target site. The molecular structures that are designed by the structure-based design process are meant to interact with biochemical targets, for example, whose three-dimensional structures are known.

Even with the extensive resources expended in drug discovery and design, there are no systematic methods for generating drugs with desired properties. Thus, the art is in need of additional systems and methods to facilitate the discovery and optimization of therapeutic and other useful compounds.

As a class of drugs, benzodiazepine compounds have been widely studied and reported to be effective medicaments for treating a number of disease. For example, U.S. 4,076,823, 4,110,337, 4,495,101, 4,751,223 and 5,776,946, each incorporated herein by reference in its entirety, report that certain benzodiazepine compounds are effective as analgesic and anti-inflammatory agents. Similarly, U.S. 5,324,726 and U.S. 5,597,915, each incorporated by reference in its entirety, report that certain benzodiazepine compounds are antagonists of cholecystokinin and gastrin and thus might be useful to treat certain gastrointestinal disorders.

Other benzodiazepine compounds have been studied as inhibitors of human neutrophil elastase in the treating of human neutrophil elastase-mediated conditions such as myocardial ischemia, septic shock syndrome, among others (*See e.g.*, U.S. 5,861,380 incorporated herein by reference in its entirety). U.S. 5,041,438, incorporated herein by reference in its entirety, reports that certain benzodiazepine compounds are useful as anti-retroviral agents.

Despite the attention benzodiazepine compounds have drawn, it will become apparent from the description below, that the present invention provides novel uses for benzodiazepine compounds and related and other compounds and methods of using the compounds, as well as known compounds, for treating a variety of diseases.

Benzodiazepine compounds are known to bind to benzodiazepine receptors in the central nervous system (CNS) and thus have been used to treat various CNS disorders

including anxiety and epilepsy. Peripheral benzodiazepine receptors have also been identified, which receptors may incidentally also be present in the CNS. The present invention demonstrates that benzodiazepines and related compounds have pro-apoptotic and cytotoxic properties. The route of action of these compounds is not through the previously
5 identified benzodiazepine receptors.

Thus, in some embodiments, the present invention provides a number of compounds. In other embodiments, the present invention provides methods for using such compounds to regulate biological processes. The present invention also provides drug-screening methods to identify and optimize compounds. In preferred embodiments, the
10 present invention provides unsolvated benzodiazepine structures and benzodiazepine related structures with long term storage capability, and stability under high pressures (e.g., storage pressures necessary in generating drug tablets). These and other research and therapeutic utilities are described below.

15 DETAILED DESCRIPTION OF THE INVENTION

Exemplary compositions and methods of the present invention are described in more detail in the following sections: I. Modulators of Cell Death; II. Modulators of Cell Growth and Proliferation; III. Benzodiazepine and Benzodiazepine Related Crystal Forms; IV. Pharmaceutical compositions, formulations, and exemplary administration routes and
20 dosing considerations; V. Drug screens; VI. Therapeutic Applications; and VII. ATPase Inhibitors And Methods For Identifying Therapeutic Inhibitors.

The present invention herein incorporates by reference U.S. Provisional Patent Nos. 60/131,761, 60/165,511, 60/191,855, 60/312,560, 60/313,689, 60/396,670, 60/565,788, 60/607,599, 60/641,040, and U.S. Patent Application Nos. 11/324,419, 11/176,719,
25 11/110,228, 10/935,333, 10/886,450, 10/795,535, 10/634,114, 10/427,211, 10/427,212, 10/217,878, 09/767,283, 09/700,101, and related applications. All compounds and uses described in the above mentioned cases are contemplated to be part of the present invention. Additionally, all other known uses of benzodiazepines may be used with the new formulations of the invention. Additional references include, but are not limited to, Otto,
30 M.W., et al., (2005) J. Clin. Psychiatry 66 Suppl. 2:34-38; Yoshii, M., et al., (2005) Nippon Yakurigaku Zasshi 125(1):33-36; Yasuda, K. (2004) Nippon Rinsho. 62 Suppl. 12:360-363; Decaudin, D. (2004) 15(8):737-745; Bonnot, O., et al. (2003) Encephale. 29(6):553-559; Sugiyama, T. (2003) Ryoikibetsu Shokogun Shirizu. 40:489-492; Lacapere, J.J., et al., (2003) Steroids. 68(7-8):569-585; Galiegue, S., et al., (2003) Curr. Med. Chem.

10(16):1563-1572; Papadopoulo, V. (2003) *Ann. Pharm. Fr.* 61(1):30-50; Goethals, I., et al., (2002) *Eur. J. Nucl. Med. Mol. Imaging* 30(2):325-328; Castedo, M., et al., (2002) *J. Exp. Med.* 196(9):1121-1125; Buffett-Jerrott, S.E., et al., (2002) *Curr. Pharm. Des.* 8(1):45-58; Beurdeley-Thomas, A., et al., (2000) *J. Neurooncol.* 46(1):45-56; Smyth, W.F., et al.,
 5 (1998) *Electrophoresis* 19(16-17):2870-2882; Yoshii, M., et al., (1998) *Nihon Shinkei Seishin Yakurigaku Zasshi.* 18(2):49-54; Trimble, M. and Hindmarch, I. (2000) *Benzodiazepines*, published by Wrighton Biomedical Publishing; and Salamone, S.J. (2001) *Benzodiazepines and GHB – Detection and Pharmacology*, published by Humana Press; each herein incorporated by reference in their entireties.

10 The practice of the present invention employs, unless otherwise indicated, conventional techniques of organic chemistry, pharmacology, molecular biology (including recombinant techniques), cell biology, biochemistry, and immunology, which are within the skill of the art. Such techniques are explained fully in the literature, such as, “Molecular cloning: a laboratory manual” Second Edition (Sambrook *et al.*, 1989); “Oligonucleotide
 15 synthesis” (M.J. Gait, ed., 1984); “Animal cell culture” (R.I. Freshney, ed., 1987); the series “Methods in enzymology” (Academic Press, Inc.); “Handbook of experimental immunology” (D.M. Weir & C.C. Blackwell, eds.); “Gene transfer vectors for mammalian cells” (J.M. Miller & M.P. Calos, eds., 1987); “Current protocols in molecular biology” (F.M. Ausubel *et al.*, eds., 1987, and periodic updates); “PCR: the polymerase chain
 20 reaction” (Mullis *et al.*, eds., 1994); and “Current protocols in immunology” (J.E. Coligan *et al.*, eds., 1991), each of which is herein incorporated by reference in its entirety.

I. Modulators of Cell Death

In preferred embodiments, the compounds of the present invention regulate
 25 apoptosis through the exposure of cells to compounds. The effect of compounds can be measured by detecting any number of cellular changes. Cell death may be assayed as described herein and in the art. In preferred embodiments, cell lines are maintained under appropriate cell culturing conditions (*e.g.*, gas (CO₂), temperature and media) for an appropriate period of time to attain exponential proliferation without density dependent
 30 constraints. Cell number and or viability are measured using standard techniques, such as trypan blue exclusion/hemo-cytometry, or MTT dye conversion assay. Alternatively, the cell may be analyzed for the expression of genes or gene products associated with aberrations in apoptosis or necrosis.

In preferred embodiments, exposing the compounds of the present invention to a cell induces apoptosis. In some embodiments, the present invention causes an initial increase in cellular ROS levels (*e.g.*, O_2^-). In further embodiments, exposure of the compounds of the present invention to a cell causes an increase in cellular O_2^- levels. In still further
5 embodiments, the increase in cellular O_2^- levels resulting from the compounds of the present invention is detectable with a redox-sensitive agent that reacts specifically with O_2^- (*e.g.*, dihydroethidium (DHE)).

In other embodiments, increased cellular O_2^- levels resulting from compounds of the present invention diminish after a period of time (*e.g.*, 10 minutes). In other embodiments,
10 increased cellular O_2^- levels resulting from the compounds of the present invention diminish after a period of time and increase again at a later time (*e.g.*, 10 hours). In further embodiments, increased cellular O_2^- levels resulting from the compounds of the present invention diminish at 1 hour and increase again after 4 hours. In preferred embodiments, an early increase in cellular O_2^- levels, followed by a diminishing in cellular O_2^- levels,
15 followed by another increase in cellular O_2^- levels resulting from the compounds of the present invention is due to different cellular processes (*e.g.*, bimodal cellular mechanisms).

In some embodiments, the compounds of the present invention cause a collapse of a cell's mitochondrial $\Delta\Psi_m$. In preferred embodiments, a collapse of a cell's mitochondrial $\Delta\Psi_m$ resulting from the present invention is detectable with a mitochondria-selective
20 potentiometric probe (*e.g.*, DiOC₆). In further embodiments, a collapse of a cell's mitochondrial $\Delta\Psi_m$ resulting from the present invention occurs after an initial increase in cellular O_2^- levels.

In some embodiments, the compounds of the present invention enable caspase activation. In other embodiments, the compounds of the present invention cause the release
25 of cytochrome *c* from mitochondria. In further embodiments, the compounds of the present invention alter cytosolic cytochrome *c* levels. In still other embodiments, altered cytosolic cytochrome *c* levels resulting from the compounds of the present invention are detectable with immunoblotting cytosolic fractions. In preferred embodiments, diminished cytosolic cytochrome *c* levels resulting from the compounds of the present invention are detectable
30 after a period of time (*e.g.*, 10 hours). In further preferred embodiments, diminished cytosolic cytochrome *c* levels resulting from the compounds of the present invention are detectable after 5 hours.

In other embodiments, the compounds of the present invention cause the opening of the mitochondrial PT pore. In preferred embodiments, the cellular release of cytochrome *c* resulting from the compounds of the present invention are consistent with a collapse of mitochondrial $\Delta\Psi_m$. In still further preferred embodiments, the compounds of the present invention cause an increase in cellular O_2^- levels after a mitochondrial $\Delta\Psi_m$ collapse and a release of cytochrome *c*. In further preferred embodiments, a rise in cellular O_2^- levels is caused by a mitochondrial $\Delta\Psi_m$ collapse and release of cytochrome *c* resulting from the compounds of the present invention.

In other embodiments, the compounds of the present invention cause cellular caspase activation. In preferred embodiments, caspase activation resulting from the compounds of the present invention is measurable with a pan-caspase sensitive fluorescent substrate (*e.g.*, FAM-VAD-fmk). In still further embodiments, caspase activation resulting from the compounds of the present invention tracks with a collapse of mitochondrial $\Delta\Psi_m$. In other embodiments, the compounds of the present invention cause an appearance of hypodiploid DNA. In preferred embodiments, an appearance of hypodiploid DNA resulting from the compounds of the present invention is slightly delayed with respect to caspase activation.

In some embodiments, the molecular target for the compounds of the present invention is found within mitochondria. In further embodiments, the molecular target of the compounds of the present invention involves the mitochondrial ATPase. The primary sources of cellular ROS include redox enzymes and the mitochondrial respiratory chain (hereinafter MRC). In preferred embodiments, cytochrome *c* oxidase (complex IV of the MRC) inhibitors (*e.g.*, NaN_3) preclude a dependent increase in cellular ROS levels for the compounds of the present invention. In other preferred embodiments, the ubiquinol-cytochrome *c* reductase component of MRC complex III inhibitors (*e.g.*, FK506) preclude a dependent increase in ROS levels for the compounds of the present invention.

In some embodiments, an increase in cellular ROS levels due to the compounds of the present invention result from the binding of the compounds of the present invention to a target within mitochondria. In preferred embodiments, the compounds of the present invention oxidizes 2',7'-dichlorodihydrofluorescein (hereinafter DCF) diacetate to DCF. DCF is a redox-active species capable of generating ROS. In further embodiments, the rate of DCF production resulting from the present invention increases after a lag period.

Antimycin A generates O_2^- by inhibiting ubiquinol-cytochrome *c* reductase. In preferred embodiments, the compounds of the present invention increase the rate of ROS production in an equivalent manner to antimycin A. In further embodiments, the compounds of the present invention increase the rate of ROS production in an equivalent manner to antimycin A under aerobic conditions supporting state 3 respiration. In further
5 embodiments, the compounds of the present invention do not directly target the MPT pore. In additional embodiments, the compounds of the present invention do not generate substantial ROS in the subcellular S15 fraction (*e.g.*, cytosol; microsomes). In even further embodiments, the compounds of the present invention do not stimulate ROS if
10 mitochondria are in state 4 respiration.

MRC complexes I – III are the primary sources of ROS within mitochondria. In preferred embodiments, the primary source of an increase in cellular ROS levels resulting from the dependent invention emanates from these complexes as a result of inhibiting the mitochondrial F_1F_0 -ATPase. Indeed, in still further embodiments, the present invention
15 inhibits mitochondrial ATPase activity of bovine sub-mitochondrial particles (hereinafter SMPs). In particularly preferred embodiments, the compounds of the present invention bind to the OSCP component of the mitochondrial F_1F_0 -ATPase.

Oligomycin is a macrolide natural product that binds to the mitochondrial F_1F_0 -ATPase, induces a state 3 to 4 transition, and as a result, generates ROS (*e.g.*, O_2^-). In
20 preferred embodiments, the compounds of the present invention bind the OSCP component of the mitochondrial F_1F_0 -ATPase. In preferred embodiments, the compounds of the present invention bind the junction between the OSCP and the F_1 subunit of the mitochondrial F_1F_0 -ATPase. In some embodiments, the compounds of the present invention bind the F_1 subunit. In certain embodiments, screening assays of the present
25 invention permit detection of binding partners of the OSCP, F_1 , or OSCP/ F_1 junction. OSCP is an intrinsically fluorescent protein. In certain embodiments, titrating a solution of test compounds of the present invention into an *E. Coli* sample overexpressed with OSCP results in quenching of the intrinsic OSCP fluorescence. In other embodiments, fluorescent or radioactive test compounds can be used in direct binding assays. In other embodiments,
30 competition binding experiments can be conducted. In this type of assay, test compounds are assessed for their ability to compete with Bz-423 for binding to the OSCP. In some embodiments, the compounds of the present invention cause a reduced increase in cellular ROS levels and reduced apoptosis in cells through regulation of the OSCP gene (*e.g.*,

altering expression of the OSCP gene). In further embodiments, the present invention functions by altering the molecular motions of the ATPase motor.

II. Modulators of Cellular Proliferation and Cell Growth

5 In some embodiments, the compounds and methods of the present invention cause decreased cellular proliferation. In other embodiments, the compounds and methods of the present invention cause decreased cellular proliferation and apoptosis. For example, cell culture cytotoxicity assays conducted during the development of the present invention demonstrated that the compounds and methods of the present invention prevents cell growth
10 after an extended period in culture (*e.g.*, 3 days).

III. Benzodiazepine and Benzodiazepine Related Crystal Forms

The present invention relates to systems and methods for generating new formulations of benzodiazepine compounds and benzodiazepine related compounds. The
15 present invention also provides high throughput systems and methods for generating and identifying new crystalline benzodiazepine and benzodiazepine related compounds.

For example, the present invention provides libraries of polymers from which crystals are nucleated by exposing solutions (*e.g.*, supersaturated solutions), the melt or vapor of the compound to the polymers. Growth of crystals on a plurality of polymers
20 provides new methods for obtaining desired polymorphs of compounds and for generating previously unidentified polymorphs of compounds. For example, the systems and methods of the present invention have been used to identify novel polymorphs of benzodiazepine compounds. For example, the systems and methods have also been used to generate efficient methods for producing orthorhombic benzodiazepine compounds from solution.
25 The novel polymorphs identified by the systems and methods of the present invention find use in identifying drugs with enhanced properties, compared to previously available polymorphs of the compound. Thus, the systems and methods of the present invention provide means for finding drug leads and/or optimizing existing drugs (see, *e.g.*, U.S. Patent Application No. 10/269,190; herein incorporated by reference in its entirety).

30 Many pharmaceutical solids exhibit polymorphism (*e.g.*, the ability of a substance to exist as two or more crystalline phases that have different arrangements and/or conformations of the molecules in the crystal lattice). Because of their structural differences, polymorphs have different solid-state properties. Consequently, polymorphism

can exert profound effects on pharmaceutical processing, including, but not limited to, milling, granulation, and tableting (Conte et al., *Il Farmaco* (Ed. Pr.) 30:194 (1974); Otsuka et al., *Chem. Pharm. Bull.*, 45:894 (1997); Otsuka et al., *J. Pharm. Sci.*, 84:614 (1995); Tuladhar et al., *J. Pharm. Pharmacol.*, 35:269 (1982); and Wong and Mitchell, *Int. J. Pharm.*, 88:261 (1992)).

Despite the fact that upon dissolution, two polymorphs will yield identical solutions, the crystalline form affects the rate of dissolution, equilibrium solubility, shelf life and ultimately bioavailability. This has implications for isolation, clinical trials, and mass production and is therefore an important aspect of creating a viable therapeutic. With a greater number of polymorphs to choose between for a solid dosage, it is more likely that an optimal mixture of properties can be achieved leading to more efficacious drugs.

In its most simple form, the process of crystallization can be considered to start from a supersaturated solution, produced by evaporation, cooling, or addition of a nonsolvent, by formation of nuclei. These species must achieve a sufficient size in order to proceed on to bulk crystals and it is the arrangement of the molecules in these nanometer-sized structures that leads to the macroscopic crystal. Thus the formation of unstable polymorphs can be attributed to their success in forming viable nuclei, a kinetic effect. Additives designed by consideration of functional groups and lattice parameters (derived from diffraction methods) can also interact with these nuclei to stabilize or destabilize them, and this approach of using designed additives has met with success in some cases (Weissbuch et al., *Acta Crystallogr. Sect. B-Struct. Sci.* 51:115 (1995); Chen et al., *J. Cryst. Growth* 144:297 (1994); and Davey et al., *J. Am. Chem. Soc.*, 119:1767 (1997)). However, this method is best suited for modifying the crystallization behavior of known polymorphs and is not readily adapted to the generation of new forms with unknown lattice parameters.

Even in crystallizations where no additives are used, it is recognized that spontaneous (homogenous) nucleation is not very common, and in most cases impurities on vessel walls function as heteronuclei to induce crystal formation. The reluctance of saturated solutions to undergo homogeneous nucleation can be explained by the energetic barrier to building a species with a high surface area to volume ratio where many of the molecules do not experience the full stabilization of the bulk. A heteronucleus reduces this barrier by providing stabilization of a growing face of the crystal.

The present invention provides systems and methods for utilizing a combinatorial library of functionalized polymers for crystal formation. Both the types of functional

groups and the spacing of these groups are altered to produce surfaces that facilitate polymorph generation. By varying these parameters (e.g., systematically) throughout the library, these polymers produce crystal forms without prior knowledge of the polymorph's structure and allow the discovery of new forms of compounds (e.g., pharmaceutical compounds, etc.) with improved properties over previously available structures. Properties that differ among polymorphs include, but are not limited to: packing properties (e.g., molar volume and density, refractive index, electrical conductivity, thermal conductivity, hygroscopicity); thermodynamic properties (e.g., melting and sublimation temperatures, internal [e.g., structural] energy, enthalpy, heat capacity, entropy, free energy and chemical potential, thermodynamic activity, vapor pressure, solubility); spectroscopic properties (e.g., electronic transitions such as ultraviolet to visible absorption spectra, vibrational transitions such as infrared absorption and Raman spectra, rotational transitions such as far infrared and microwave absorption spectra, nuclear spin transitions such as nuclear magnetic resonance spectra); kinetic properties (e.g., dissolution rate, rates of solid state reactions, and stability); surface properties (e.g., surface free energy, interfacial tensions, habit); and mechanical properties (e.g., hardness, tensile strength, compactibility, tableting, handling, flow, and blending) (See e.g., "Polymorphism in Pharmaceutical Solids," ed. Harry G. Brittain, Marcel Dekker, Inc., New York (1999)).

In some embodiments of the present invention, a plurality of polymers are provided with (e.g., placed onto or into) a solid surface or vessel to facilitate high throughput crystal growth and analysis. In some embodiments, the solid surface or vessel is a multi-chamber plate (e.g., a 96-well or 384-well plate). However, the present invention is not limited by the solid surface or vessel employed. As used herein, the terms "solid support" or "support" refer to any material that provides a solid or semi-solid structure with which another material (e.g., a polymer) can be associated. Such materials include smooth supports (e.g., metal, glass, plastic, silicon, and ceramic surfaces) as well as textured and porous materials. Such materials also include, but are not limited to, gels, rubbers, polymers, and other non-rigid materials. Solid supports need not be flat. Supports include any type of shape including spherical shapes (e.g., beads). Materials associated with the solid support may be associated with any portion of the solid support (e.g., may be attached, enclosed, or in contact with an interior portion of a porous solid support material).

The present invention is not limited by the nature of the polymer used to promote crystal growth. In preferred embodiments, the plurality of polymers used in screening

methods of the present invention comprise two or more polymers (e.g., three or more, four or more, five or more, . . . , ten or more, . . . , twenty or more, . . . , fifty or more different polymers). The maximum number of polymers employed in the systems and methods of the present invention is constrained only by the availability of polymer materials (i.e., any
5 of the thousands of known polymers may be employed, as well as new polymers that are identified in the future) and by physical and space limitations of the testing area. However, the systems and methods of the present invention may be employed at very large scales. For example, in some embodiments, 384-well plates are used wherein the bottom surface of each well contains a different polymer material. Dozens of such plates may be arranged on
10 shelves and dozens of shelves may be placed in racks. A single laboratory space can hold hundreds of racks. Thus, a single room can house tens of millions of different polymers, wherein a solution with a candidate compound is applied to each of the polymers and crystals are grown and analyzed to identify the properties of the crystals. Further miniaturization allows even more reactions to be run simultaneously in a single run.

15 While the present invention is not limited by the nature of the polymer, commercially available polymers that find use with the present invention include, but are not limited to, acrylonitrile/butadiene/styrene resin, alginic acid (sodium salt), butyl/isobutyl methacrylate copolymer, cellulose acetate, cellulose acetate butyrate, cellulose propionate, cellulose triacetate, ethyl cellulose, ethylene/acrylic acid copolymer, ethylene/ethyl acrylate
20 copolymer, ethylene/propylene copolymer, ethylene/vinyl acetate (14, 18, 25, 28, 33% and 40% VA) copolymer, hydroxybutyl methyl cellulose, hydroxypropyl cellulose, hydroxypropyl methyl cellulose, methyl cellulose, methyl vinyl ether/maleic acid copolymer, methyl vinyl ether/maleic anhydride copolymer, nylon 6, nylon 6/6, nylon 6/9, nylon 6/10, nylon 6/12, nylon 6(3)T, nylon 11, nylon 12, phenoxy resin, polyacetal,
25 polyacrylamide, polyacrylamide carboxyl modified (low), polyacrylamide carboxyl modified (high), poly(acrylic acid), polyamide resin, 1,2-polybutadiene, poly(1-butene) isotactic, poly(n-butyl methacrylate), polycarbonate resin, poly(diallyl isophthalate), poly(diallyl phthalate), poly(2,6-dimethyl-p-phenylene oxide), poly(4,4-dipropoxy-2,2-diphenyl propane fumarate), poly(ethyl methacrylate), polyethylene high density,
30 polyethylene low density, polyethylene chlorinated (25, 36, 42, and 48% chlorine), polyethylene chlorosulfonated, poly(ethylene oxide), polyethylene oxidized, poly(ethylene terephthalate), poly(2-hydroxyethyl methacrylate), poly(isobutyl methacrylate), polyisoprene chlorinated, poly(methyl methacrylate), poly(4-methyl-1-pentene), poly(alpha-

methylstyrene), poly(p-phenylene ether-sulphone), poly(phenylene sulfide), polypropylene
 isotactic chlorinated, polypropylene isotactic, polystyrene, polysulfone resin,
 poly(tetrafluoroethylene), poly(2,4,6-tribromostyrene), poly(vinyl acetate), poly(vinyl
 alcohol) 100% hydrolyzed, poly(vinyl alcohol) 98% hydrolyzed, poly(vinyl butyral),
 5 poly(vinyl chloride), poly(vinyl chloride) 1.8% carboxylated, poly(vinyl formal),
 polyvinylpyrrolidone, poly(vinyl stearate), poly(vinylidene fluoride), styrene/acrylonitrile
 copolymer (75/25), styrene/acrylonitrile copolymer (70/30), styrene/allyl alcohol
 copolymer, styrene/butadiene ABA block copolymer, styrene/butyl methacrylate
 copolymer, styrene/ethylene-butylene ABA block copolymer, styrene/maleic anhydride
 10 copolymer, vinyl alcohol/vinyl butyral copolymer, vinyl chloride/vinyl acetate (10, 12, and
 19% VA) copolymer, vinyl chloride/vinyl acetate copolymer 1% carboxylated, vinyl
 chloride/vinyl acetate/hydroxypropyl acrylate terpolymer, and vinyl chloride/vinyl
 acetate/vinyl alcohol terpolymer, as well as, functionalized polybutadienes, poly(ethylene-
 co-propylene-co-5-methylene-2-norbornene), poly(perfluoropropylene oxide)-co-
 15 poly(perfluoroformaldehyde), metaldehyde, pectic acid, polyethylenimine, poly(ethylene-
 co-carbon monoxide), poly(3-hydroxybutyric acid) and copolymers with valeric acid,
 polylactide, polyaminoacids, polyacenaphthylene, poly(dimethylsiloxane), poly[(dibenzo-
 18-crown-6)-co-formaldehyde] and other polymers containing metal chelating groups,
 poly[(phenyl isocyanate)-co-formaldehyde], poly(vinylsulfonic acid), poly(melamine-co-
 20 formaldehyde), polyphosphates, polyphosphazenes, tributyltin fluoride,
 polysaccharides, and other organic and inorganic polymers.

Polymers at each location in the library can comprise a mixture of two or more
 different polymers in one or more different locations. The combination of polymers in
 different ratios dramatically expands the diversity of conditions available in the libraries.

25 Solutions containing the compound to be screened are applied to the polymers and
 incubated under conditions that facilitate crystal growth. The present invention is not
 limited by the manner in which the compounds are applied to the polymers. In some
 embodiments, a solution is used to supply each region of a polymer library. Solution may
 be delivered by pouring, transfer through tubing, injection, or any other means. Where
 30 thousands to millions of individual polymers are used, in preferred embodiments, an
 automated delivery system is used. The present invention is not limited to the use of
 solutions. Melts of materials and vapors onto the polymers also find use in the system and
 methods of the present invention.

The solvent used to solubilize any particular compound may be varied. In some embodiments, a variety of solvents are used for each compound, wherein each different solvent type is exposed to each type of polymer to increase the range of crystallization conditions used in the library. In such embodiments, multiple regions (e.g., zones) of each polymer are created in the library to allow each solvent type to be combined with each polymer type. In addition to different solvent types, a variety of different ingredients (e.g., salts) may be placed in the solution, yet further expanding the array of choices for library analysis. Such applications find use in the generation and isolation of new pseudopolymorphs (solvates and salts) that may be used as drugs.

Crystals formed on each polymer are analyzed using any suitable method. In some embodiments, analysis is conducted directly on the polymer surface, without removing the crystals. In other embodiments, crystals are removed and analyzed. Analysis includes, but is not limited to, crystal structure analysis, analysis of spectroscopic, packing, density, thermodynamic, properties, kinetic, surface, and mechanical properties. In some embodiments, analysis includes functional analysis such as testing bioavailability or biological activity after administration to a test organism (e.g., an animal or plant). For example, in some embodiments, rapid screening is conducted using the D8 Discover with GADDS X-ray diffraction system (Bruker AXS, Madison, WI) or similar systems.

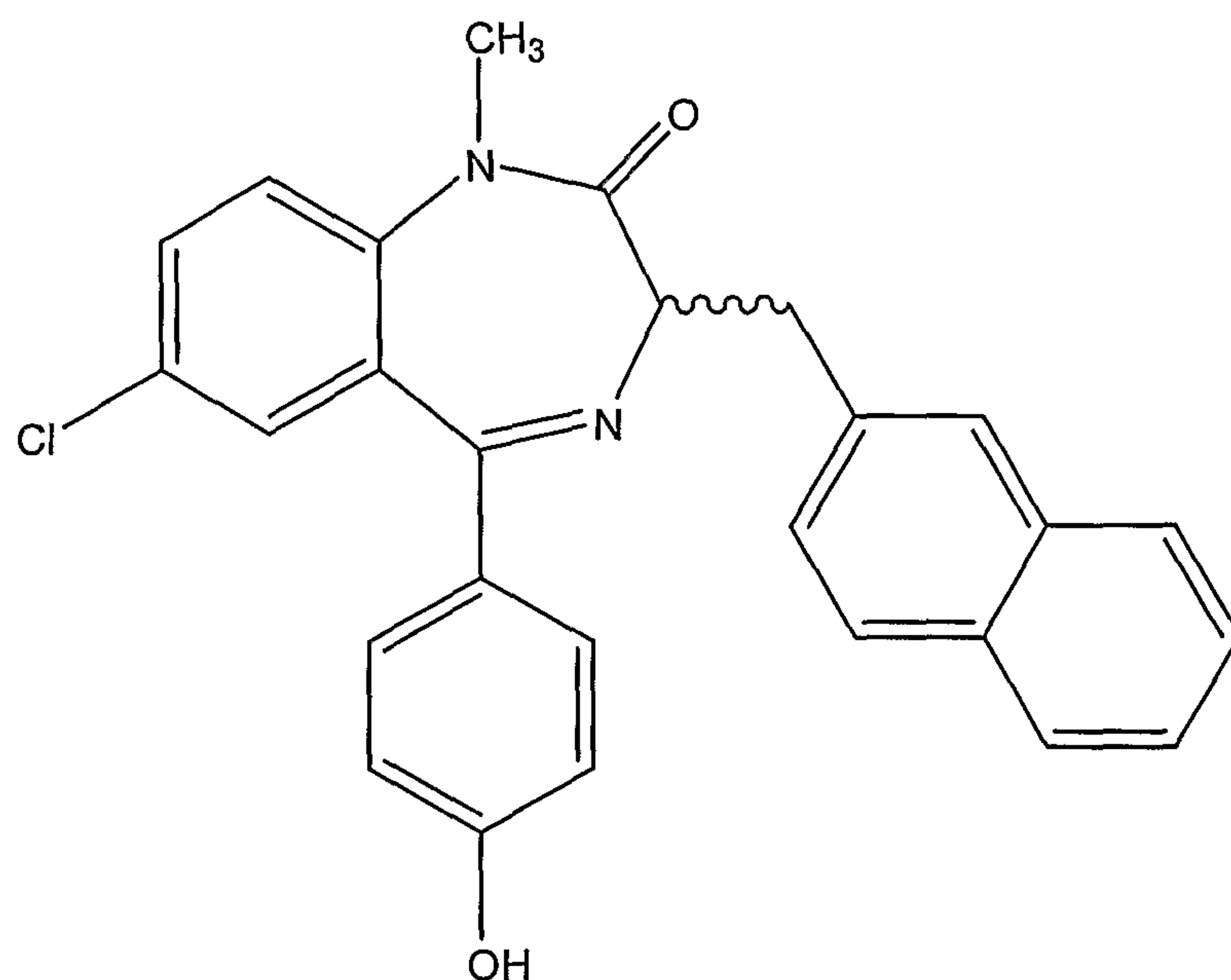
Polymorphs identified in the screening method are compared to existing polymorphs. Where a new polymorph is identified, the polymorph is characterized to identify properties that differ from previously known polymorphs (e.g., to identify improved drugs). Known polymorphs generated using the systems and methods of the present invention also are compared to existing production methods to identify whether the polymer-based method of the present invention provides advantages over existing production methods (e.g., less expensive or easier to produce, greater purity, superior crystals, ability to produce from aqueous solution, etc.).

Polymorphs identified by the present invention can be produced in large quantities. In some embodiments, crystals are collected and used to seed further solutions of the compound. However, in some cases the presence of the polymer surface may be required to generate crystals. In such embodiments, large or multiple surfaces or vessels are provided with the polymer known to generate the crystal to allow large-scale production.

Polymorphs produced by the methods of the present invention may be used in the generation of pharmaceutical formulations. The novel polymorphs identified increase the

available choices for designing drugs with desired properties, both in biological activity and in handling. For example, in order for many drugs to take action, they must dissolve in the gut and be absorbed in the blood stream. In many cases the rate at which the drug dissolves can limit its effectiveness. The polymorphs of the present invention, either alone, or in
5 combination with other forms of the drug find use in optimizing effectiveness, generally, or for particular patients or patient groups (e.g., age groups, genders, species, etc.). In some cases, the novel polymorphs provide advantages in shelf-life or the ability of the compound to be included in tablets (See e.g., Sun and Grant, Pharm. Res., 18:274 (2001)).

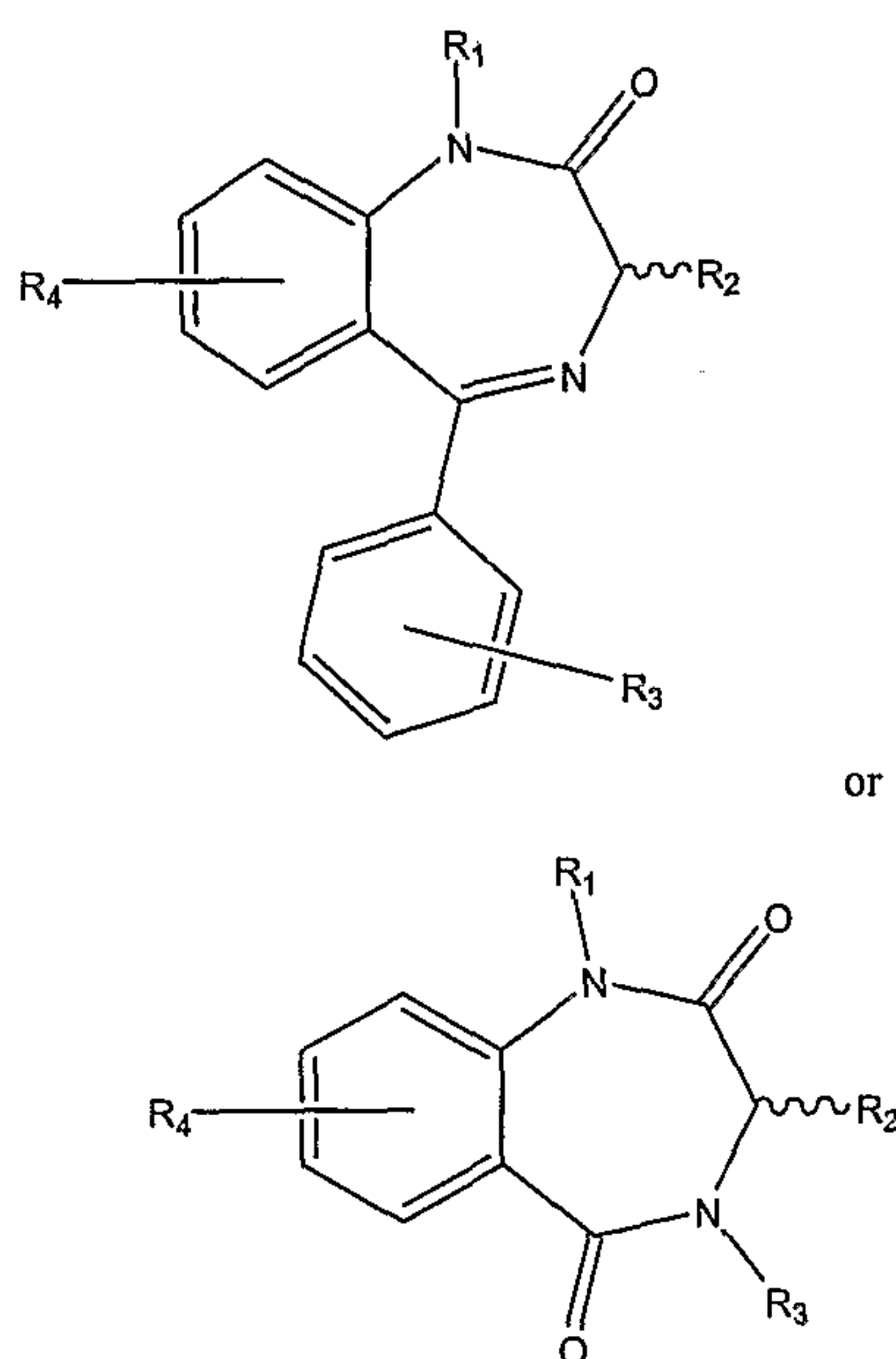
Exemplary benzodiazepine compounds provided by the present invention include
10 crystal forms and formulations of Bz-423:



Bz-423 differs from benzodiazepines in clinical use by the presence of a hydrophobic substituent at C-3. This substitution renders binding to the peripheral benzodiazepine receptor ("PBR") weak (K_d ca. 1 μ M) and prevents binding to the central
15 benzodiazepine receptor so that Bz-423 is not a sedative. Additionally, compositions of the present invention comprising benzodiazepine compounds (e.g., Bz-423) have been shown to bind to the oligomycin sensitivity conferring protein (OSCP) portion of the mitochondrial F_0F_1 ATPase synthase complex, to the OSCP junction, or to the F1 subunit (see, e.g., U.S. Provisional Patent Nos. 60/131,761, 60/165,511, 60/191,855, 60/312,560, 60/313,689,
20 60/396,670, 60/565,788, 60/607,599, 60/641,040, and U.S. Patent Application Nos. 11/324,419, 11/176,719, 11/110,228, 10/935,333, 10/886,450, 10/795,535, 10/634,114, 10/427,211, 10/427,212, 10/217,878, 09/767,283, 09/700,101, and related applications; each herein incorporated by reference in their entireties).

Exemplary crystal forms of Bz-423 and related compounds provided by the present invention include, but are not limited to, anhydrous Bz-423, Bz-423 ethanol solvate, Bz-423 succinic acid (2:1), Bz-423 citric acid (2:1), Bz-423 biphenyl derivate, BZ-423-acetic acid, BZ-423-CH₃CN, BZ-423-methanol, BZ-423-ethyl acetate, BZ-423-toluene, BZ-423-oxalic acid, BZ-423-fumaric acid, BZ-423-octanol, BZ-423-heptanoic acid, BZ-423-diphenyl ether, Bz-423-1-propanol solvate, Bz-423 2-propanol solvate, Bz-423 1-butanol solvate, Bz-423 2-butanol solvate, Bz-423 1-pentanol solvate, Bz-423 propylene glycol, Bz-423 acetone glass, and BZ-423-trichlorobenzene.

Additional exemplary compounds of the present invention also include, but are not limited to, crystal forms and formulations of:



or its enantiomer, wherein, R₁ is aliphatic or aryl; R₂ is aliphatic, aryl, -NH₂, -NHC(=O)-R₅; or a moiety that participates in hydrogen bonding, wherein R₅ is aryl, heterocyclic, -R₆-NH-C(=O)-R₇ or -R₆-C(=O)-NH-R₇, wherein R₆ is an aliphatic linker of 1-6 carbons and R₇ is aliphatic, aryl, or heterocyclic, each of R₃ and R₄ is independently a hydroxy, alkoxy, halo, amino, lower-alkyl-substituted-amino, acetylamino, hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, aryl, or heterocyclic; or a pharmaceutically acceptable salt, prodrug or derivative thereof.

In the above structures, R₁ is a hydrocarbyl group of 1-20 carbons and 1-20 hydrogens. Preferably, R₁ has 1-15 carbons, and more preferably, has 1-12 carbons. Preferably, R₁ has 1-12 hydrogens, and more preferably, 1-10 hydrogens. Thus R₁ can be an aliphatic group or an aryl group.

The term "aliphatic" represents the groups commonly known as alkyl, alkenyl, alkynyl, alicyclic. The term "aryl" as used herein represents a single aromatic ring such as a phenyl ring, or two or more aromatic rings that are connected to each other (*e.g.*, bisphenyl) or fused together (*e.g.*, naphthalene or anthracene). The aryl group can be optionally substituted with a lower aliphatic group (*e.g.*, C₁-C₄ alkyl, alkenyl, alkynyl, or C₃-C₆ alicyclic). Additionally, the aliphatic and aryl groups can be further substituted by one or more functional groups such as -NH₂, -NHCOCH₃, -OH, lower alkoxy (C₁-C₄), halo (-F, -Cl, -Br, or -I). It is preferable that R₁ is primarily a nonpolar moiety.

In the above structures, R₂ can be aliphatic, aryl, -NH₂, -NHC(=O)-R₅, or a moiety that participates in hydrogen bonding, wherein R₅ is aryl, heterocyclic, R₆-NH-C(=O)-R₇ or -R₆-C(=O)-NH-R₇, wherein R₆ is an aliphatic linker of 1-6 carbons and R₇ is an aliphatic, aryl, or heterocyclic. The terms "aliphatic" and "aryl" are as defined above.

The term "a moiety that participates in hydrogen bonding" as used herein represents a group that can accept or donate a proton to form a hydrogen bond thereby.

Some specific non-limiting examples of moieties that participate in hydrogen bonding include a fluoro, oxygen-containing and nitrogen-containing groups that are well-known in the art. Some examples of oxygen-containing groups that participate in hydrogen bonding include: hydroxy, lower alkoxy, lower carbonyl, lower carboxyl, lower ethers and phenolic groups. The qualifier "lower" as used herein refers to lower aliphatic groups (C₁-C₄) to which the respective oxygen-containing functional group is attached.

Thus, for example, the term "lower carbonyl" refers to *inter alia*, formaldehyde, acetaldehyde.

Some nonlimiting examples of nitrogen-containing groups that participate in hydrogen bond formation include amino and amido groups. Additionally, groups containing both an oxygen and a nitrogen atom can also participate in hydrogen bond formation. Examples of such groups include nitro, N-hydroxy and nitrous groups.

It is also possible that the hydrogen-bond acceptor in the present invention can be the π electrons of an aromatic ring. However, the hydrogen bond participants of this invention do not include those groups containing metal atoms such as boron. Further the hydrogen bonds formed within the scope of practicing this invention do not include those formed between two hydrogens, known as "dihydrogen bonds." (*See*, R.H. Crabtree, *Science*, 282:2000-2001 [1998], for further description of such dihydrogen bonds).

The term "heterocyclic" represents, for example, a 3-6 membered aromatic or nonaromatic ring containing one or more heteroatoms. The heteroatoms can be the same or different from each other. Preferably, at least one of the heteroatom's is nitrogen. Other heteroatoms that can be present on the heterocyclic ring include oxygen and sulfur.

5 Aromatic and nonaromatic heterocyclic rings are well-known in the art. Some nonlimiting examples of aromatic heterocyclic rings include pyridine, pyrimidine, indole, purine, quinoline and isoquinoline. Nonlimiting examples of nonaromatic heterocyclic compounds include piperidine, piperazine, morpholine, pyrrolidine and pyrazolidine. Examples of oxygen containing heterocyclic rings include, but not limited to furan, oxirane,
10 2H-pyran, 4H-pyran, 2H-chromene, and benzofuran. Examples of sulfur-containing heterocyclic rings include, but are not limited to, thiophene, benzothiophene, and parathiazine.

 Examples of nitrogen containing rings include, but not limited to, pyrrole, pyrrolidine, pyrazole, pyrazolidine, imidazole, imidazoline, imidazolidine, pyridine,
15 piperidine, pyrazine, piperazine, pyrimidine, indole, purine, benzimidazole, quinoline, isoquinoline, triazole, and triazine.

 Examples of heterocyclic rings containing two different heteroatoms include, but are not limited to, phenothiazine, morpholine, parathiazine, oxazine, oxazole, thiazine, and thiazole.

20 The heterocyclic ring is optionally further substituted with one or more groups selected from aliphatic, nitro, acetyl (*i.e.*, -C(=O)-CH₃), or aryl groups.

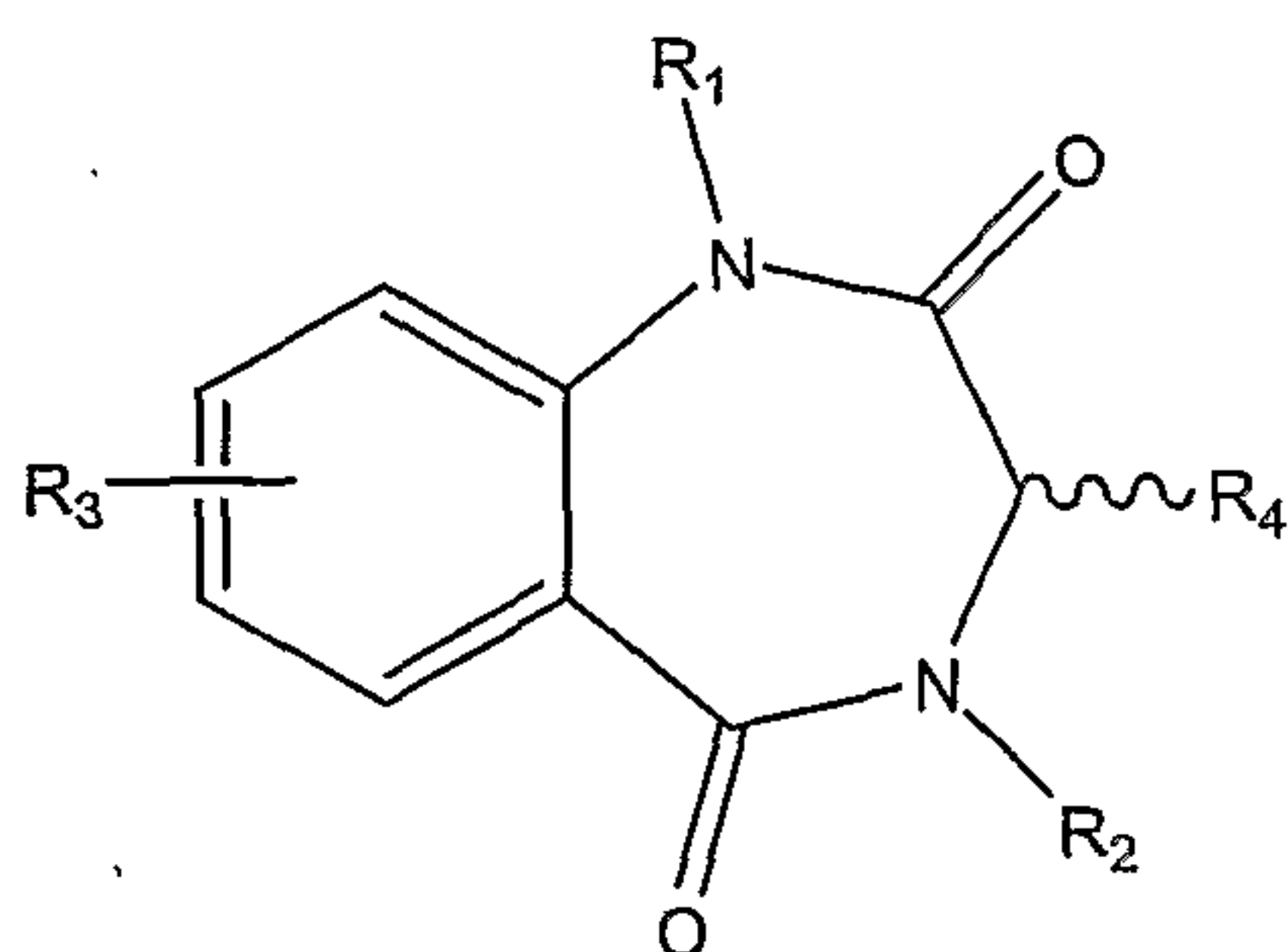
 Each of R₃ and R₄ can be independently a hydroxy, alkoxy, halo, amino, or substituted amino (such as lower-alkyl-substituted-amino, or acetylamino or hydroxyamino), or an aliphatic group having 1-8 carbons and 1-20 hydrogens. When each
25 of R₃ and R₄ is an aliphatic group, it can be further substituted with one or more functional groups such as a hydroxy, alkoxy, halo, amino or substituted amino groups as described above. The terms "aliphatic" is defined above. Alternatively, each of R₃ and R₄ can be hydrogen.

 It is well-known that many 1,4-benzodiazepines exist as optical isomers due to the
30 chirality introduced into the heterocyclic ring at the C₃ position. The optical isomers are sometimes described as L- or D-isomers in the literature. Alternatively, the isomers are also referred to as R- and S- enantiomorphs. For the sake of simplicity, these isomers are referred to as enantiomorphs or enantiomers. The 1,4-benzodiazepine compounds described

herein include their enantiomeric forms as well as racemic mixtures. Thus, the usage "benzodiazepine or its enantiomers" herein refers to the benzodiazepine as described or depicted, including all its enantiomorphs as well as their racemic mixture.

From the above description, it is apparent that many specific examples are represented by the generic formulas presented above. Thus, in one example, R₁ is aliphatic, R₂ is aliphatic, whereas in another example, R₁ is aryl and R₂ is a moiety that participates in hydrogen bond formation. Alternatively, R₁ can be aliphatic, and R₂ can be an -NHC(=O)-R₅, or a moiety that participates in hydrogen bonding, wherein R₅ is aryl, heterocyclic, -R₆-NH-C(=O)-R₇ or -R₆-C(=O)-NH-R₇, wherein R₆ is an aliphatic linker of 1-6 carbons and R₇ is an aliphatic, aryl, or heterocyclic. A wide variety of sub combinations arising from selecting a particular group at each substituent position are possible and all such combinations are within the scope of this invention.

Additional exemplary compounds of the present invention also include, but are not limited to, crystal forms and formulations of:



enantiomers and pharmaceutically acceptable salts thereof:

wherein R₁ is an aliphatic group having 1 to 20 carbon atoms and 1 to

20 hydrogen atoms or an aryl group having up to 20 carbon atoms and up to 20 hydrogen atoms;

wherein each of R₂ and R₃ is independently selected from the group consisting of hydrogen, hydroxy, C₁₋₄alkoxy, halo, amino, C₁₋₄alkyl-substituted-amino, acetylamino, hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, an aryl group having from 6 to 14 carbon atoms, and a heterocyclic group having a 3-6 membered aromatic or nonaromatic ring containing one or more heteroatoms selected from nitrogen, oxygen, and sulfur;

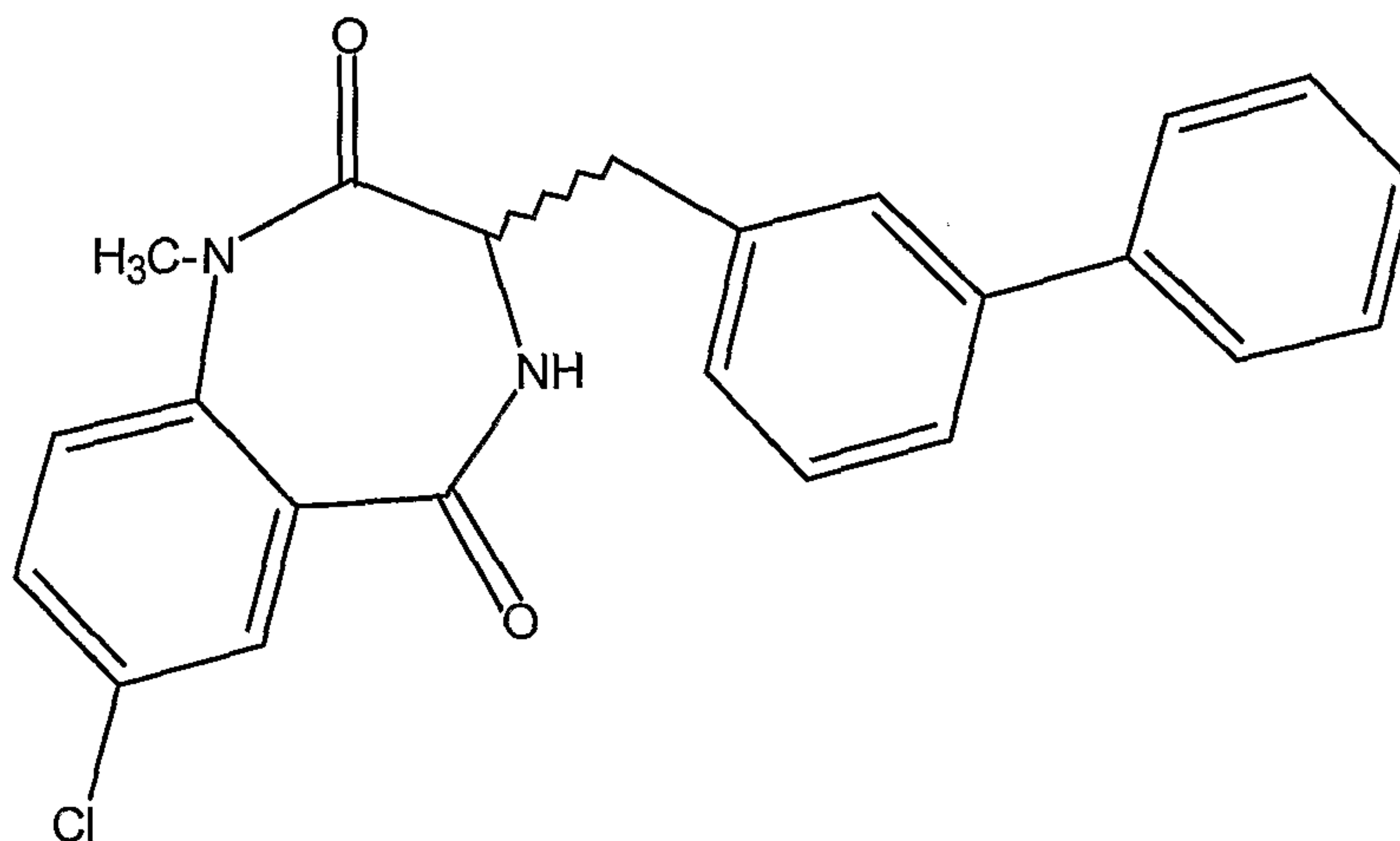
wherein R₄ is aliphatic or aryl;

wherein R_5 is selected from the group consisting of an aryl group having from 6 to 14 carbon atoms, a heterocyclic group having a 3-6 membered aromatic or nonaromatic ring containing one or more heteroatoms selected from nitrogen, oxygen, and sulfur, $-R_6-NH-C(=O)-R_7$, and $-R_6-C(=O)-NH-R_7$;

5 wherein R_6 is an aliphatic linker group of 1-6 carbons; and,

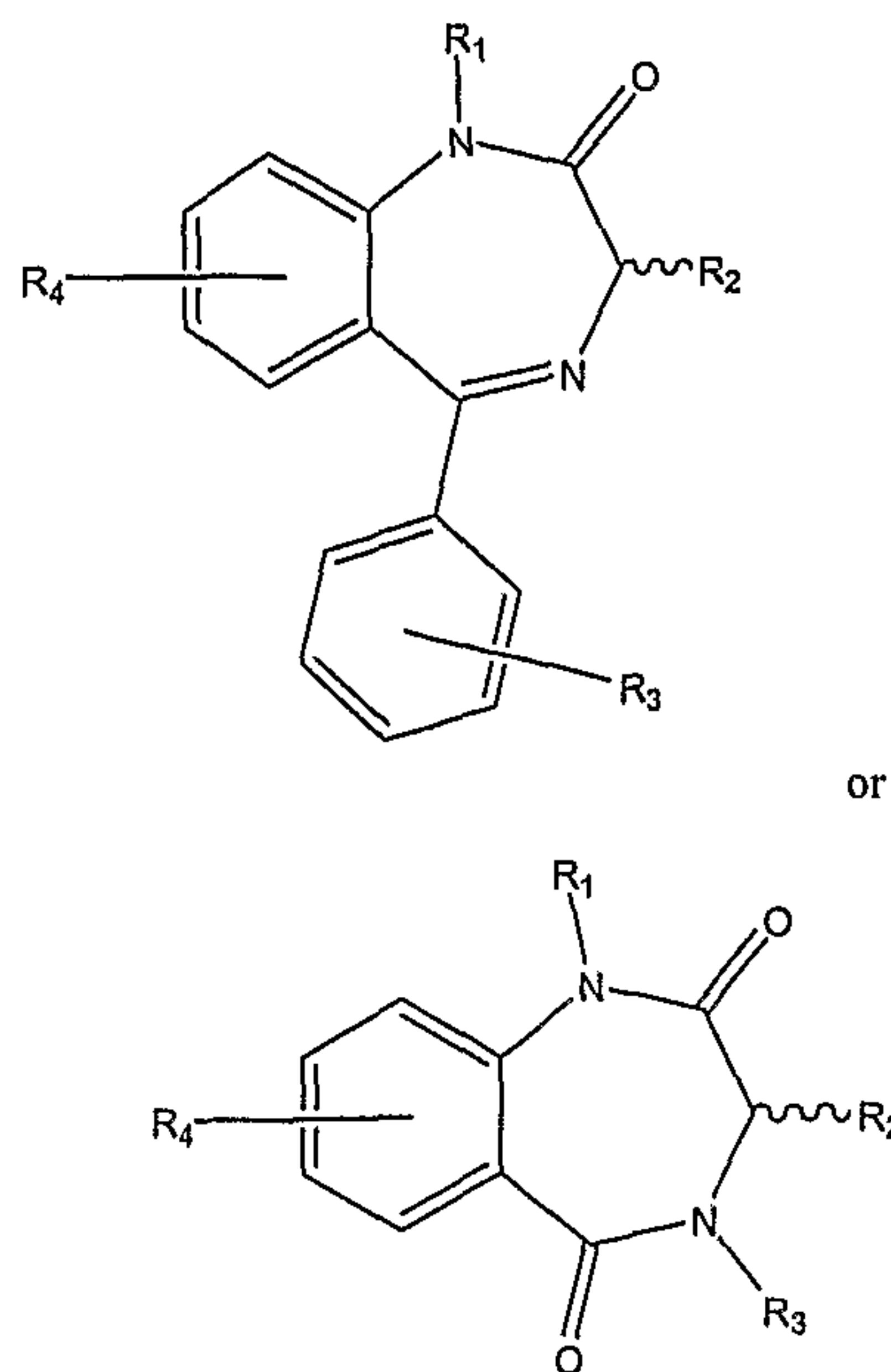
wherein R_7 is selected from the group consisting of an aliphatic group having 1-8 carbons and 1-20 hydrogens, an aryl group having from 6 to 14 carbon atoms and a heterocyclic group having a 3-6 membered aromatic or nonaromatic ring containing one or more heteroatoms selected from nitrogen, oxygen, and sulfur.

10 In some preferred embodiments, crystal forms and formulations of the following exemplary compound are provided:



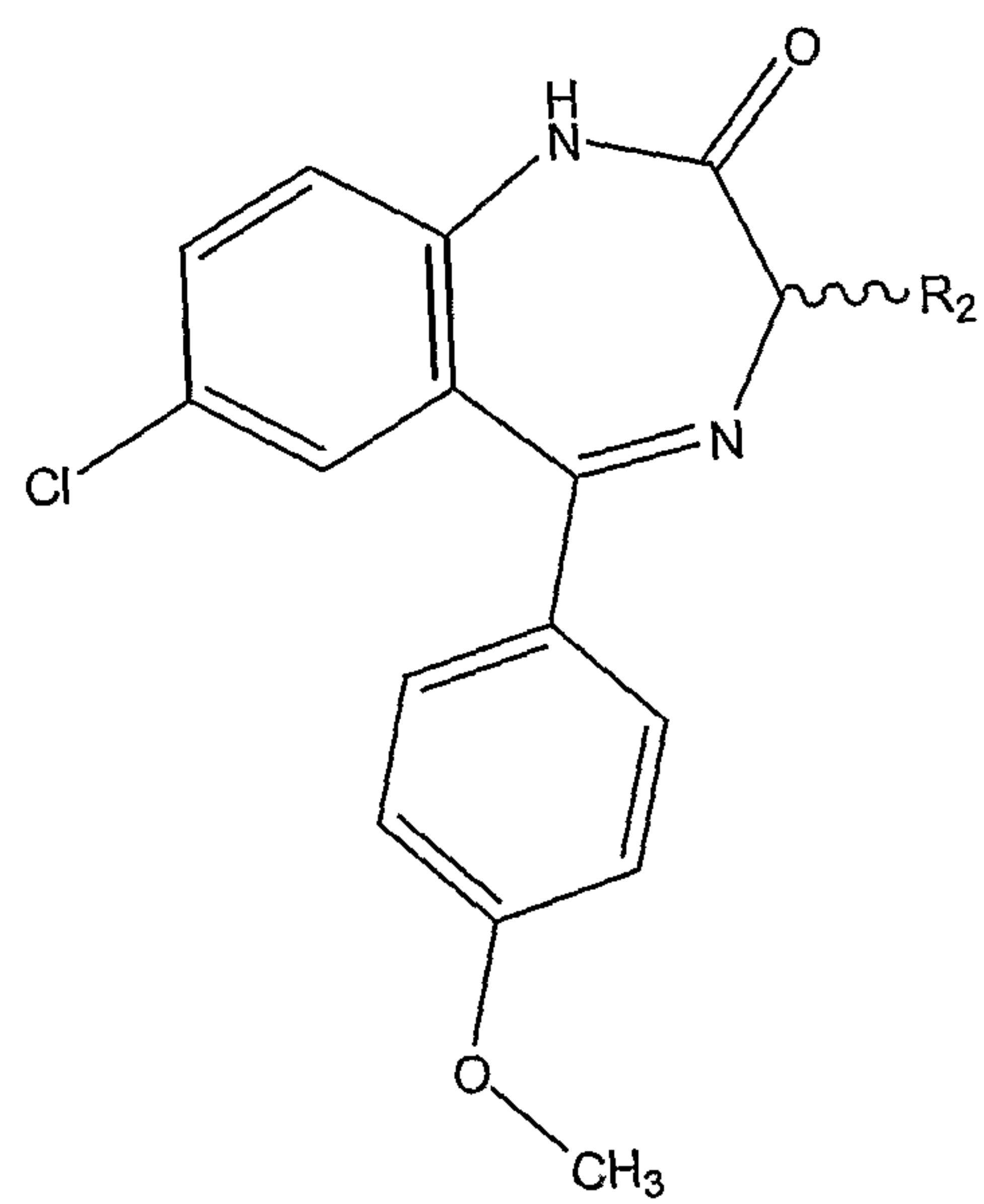
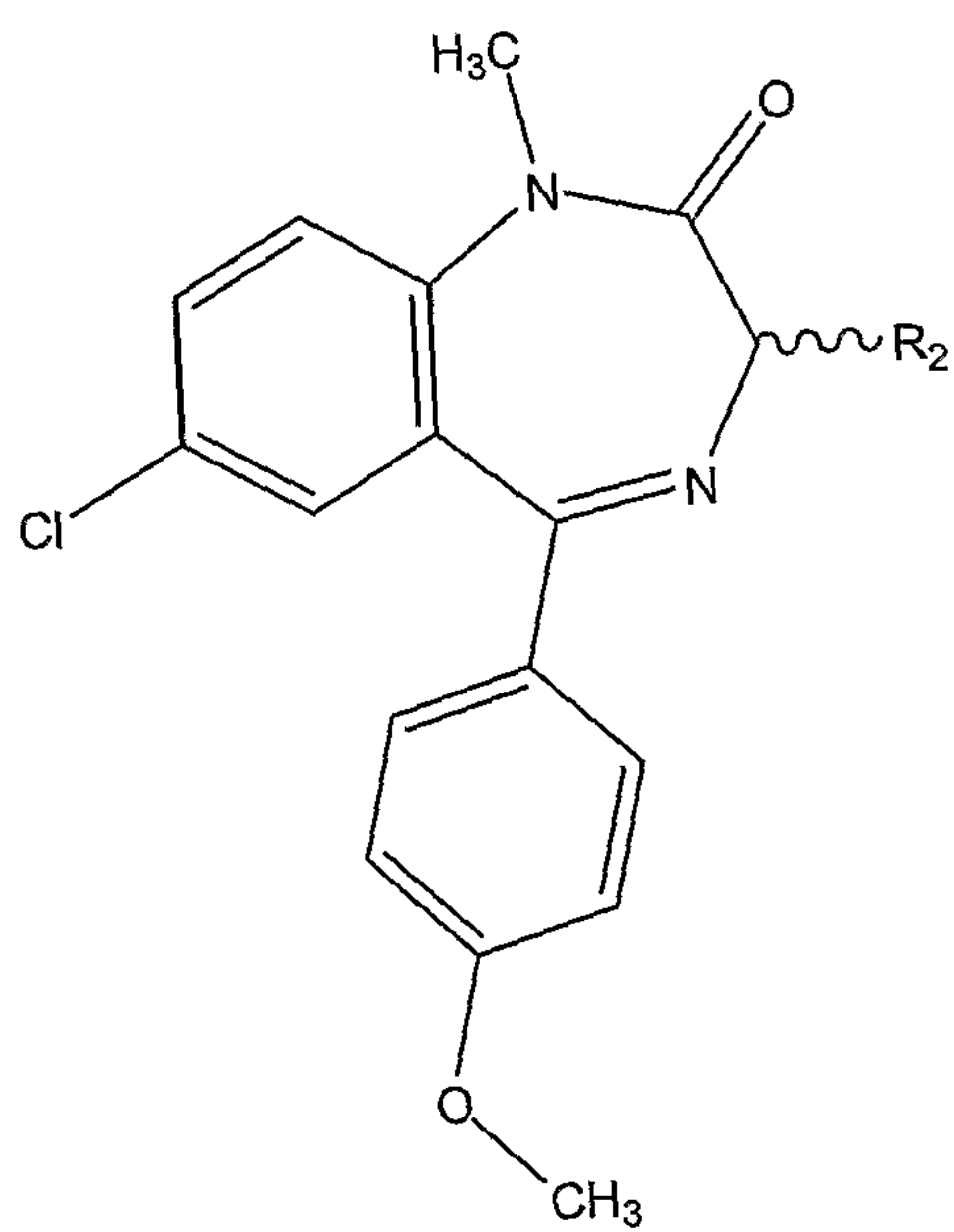
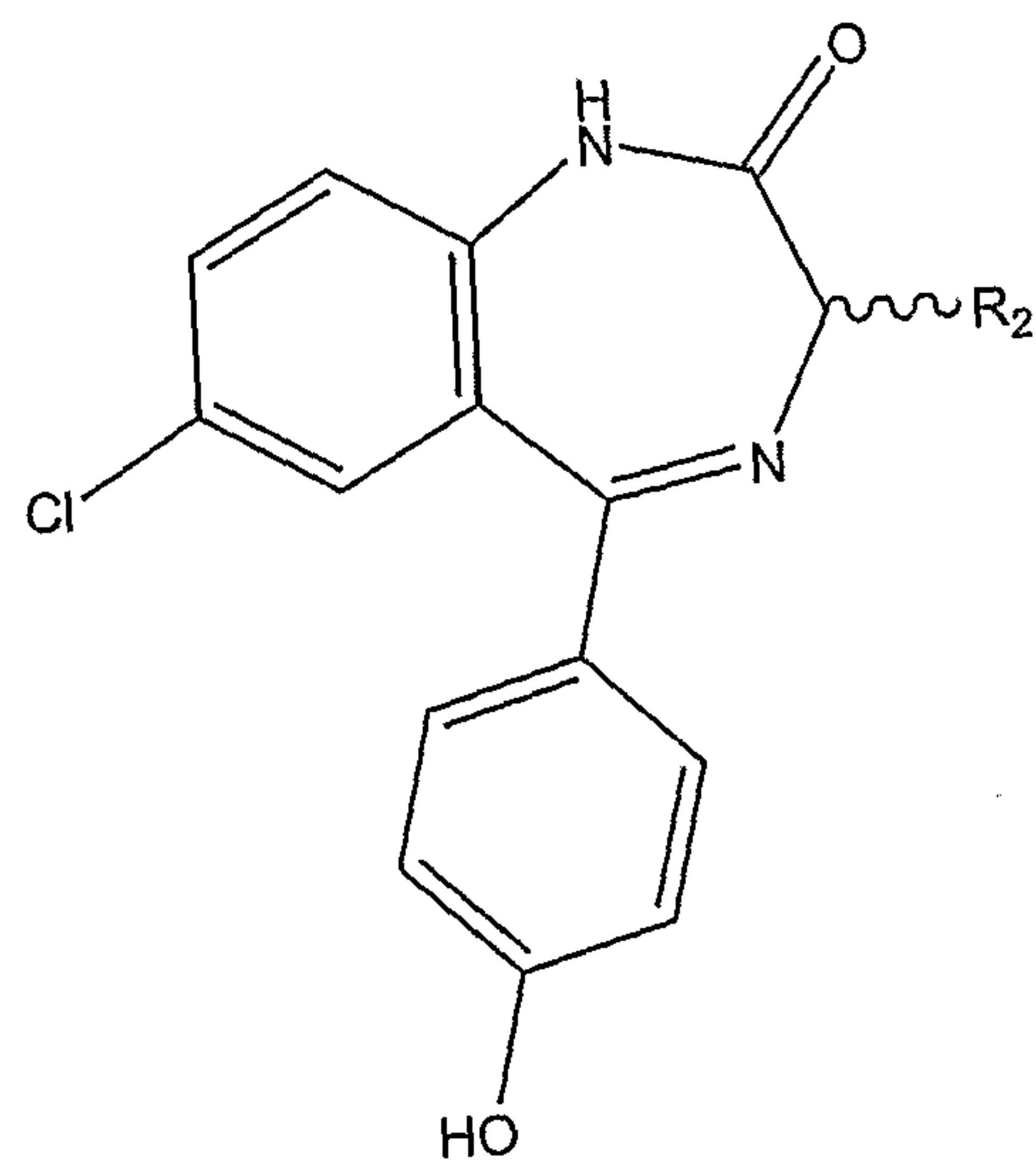
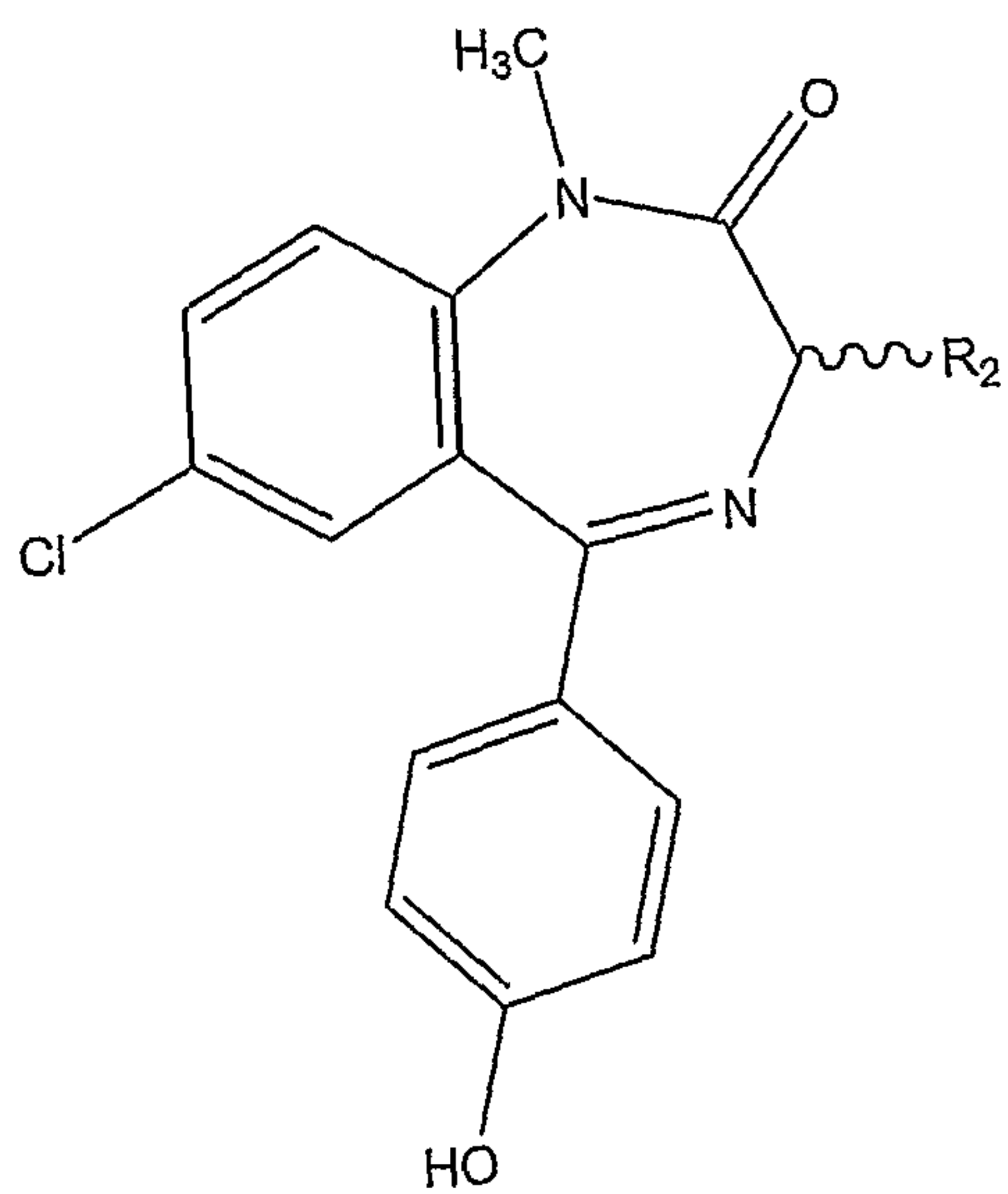
In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:

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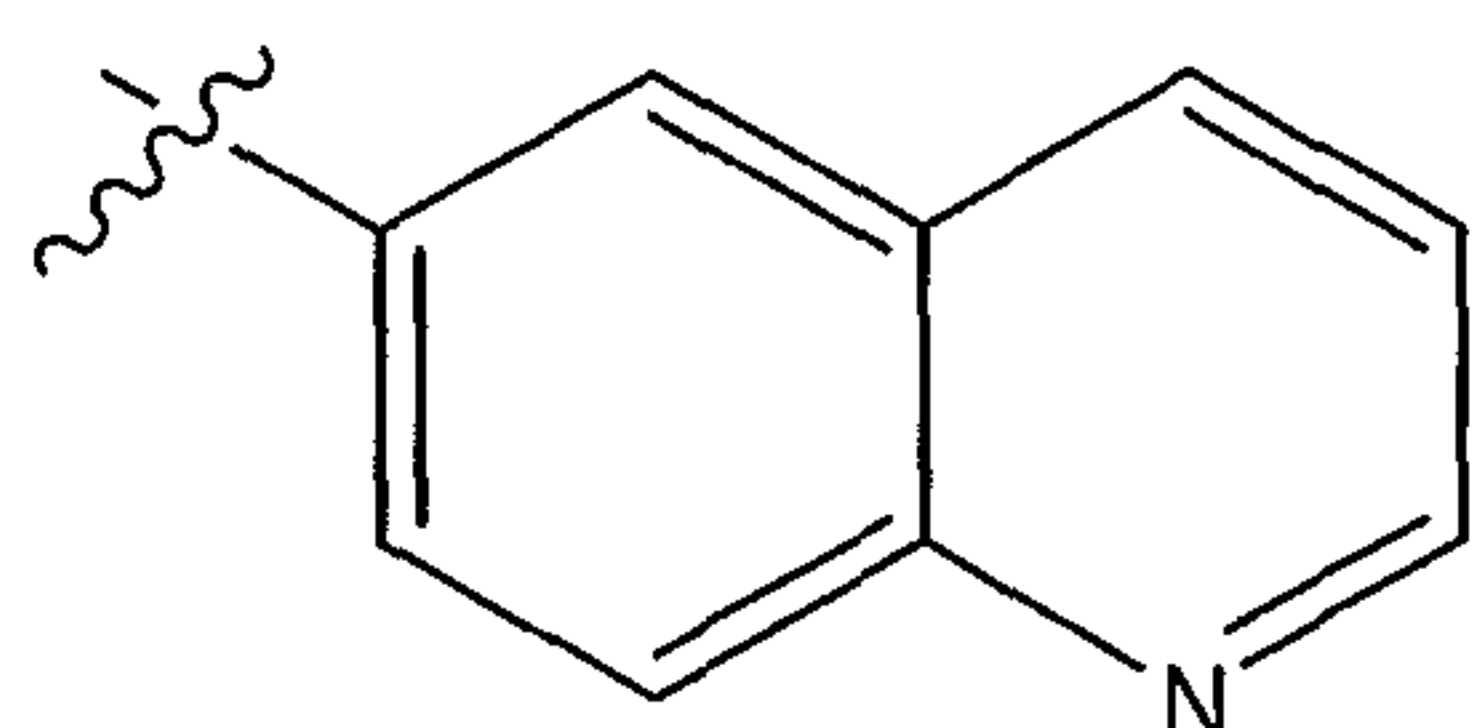
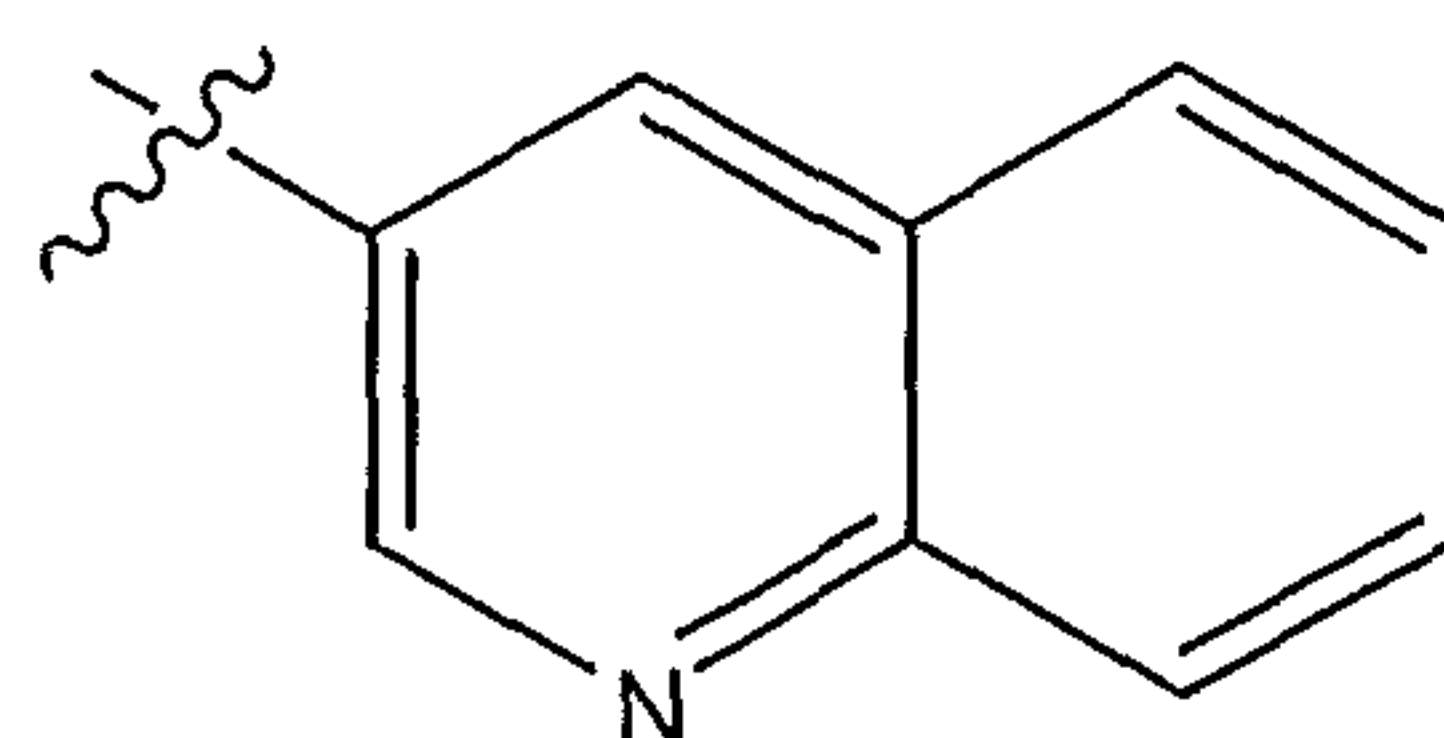
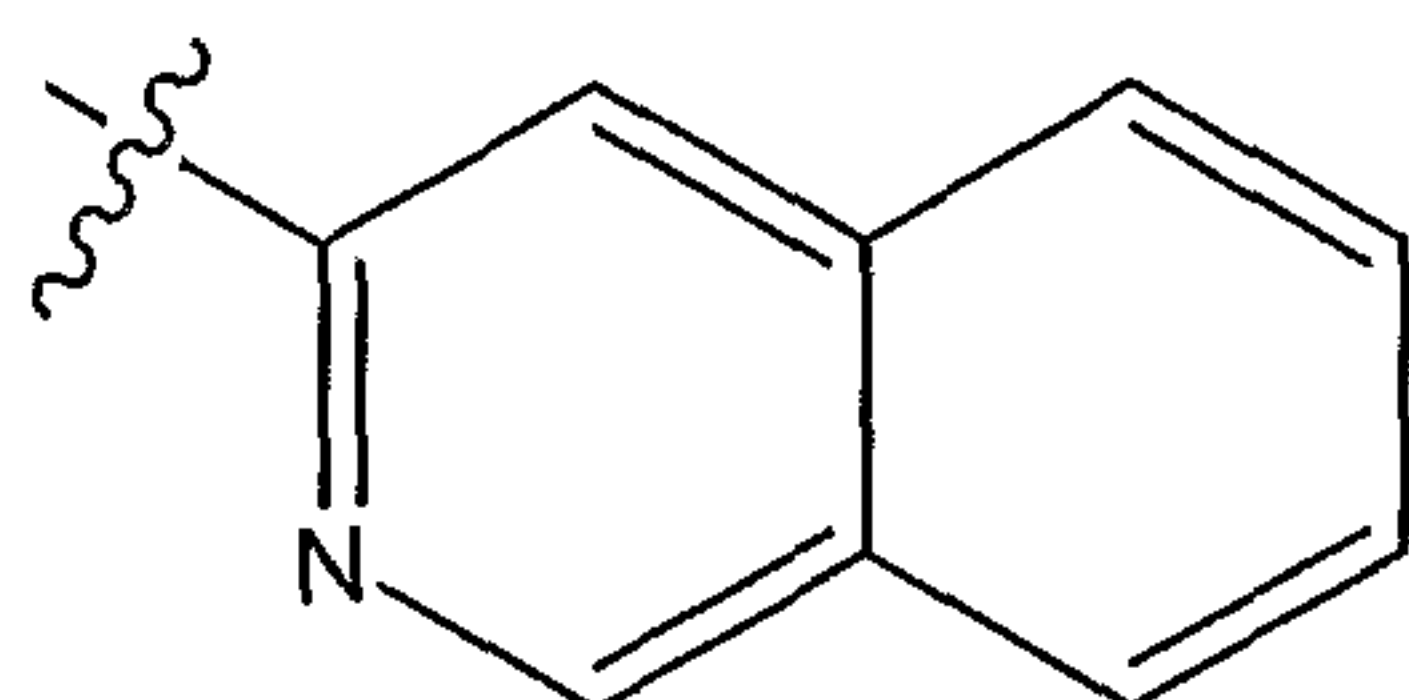
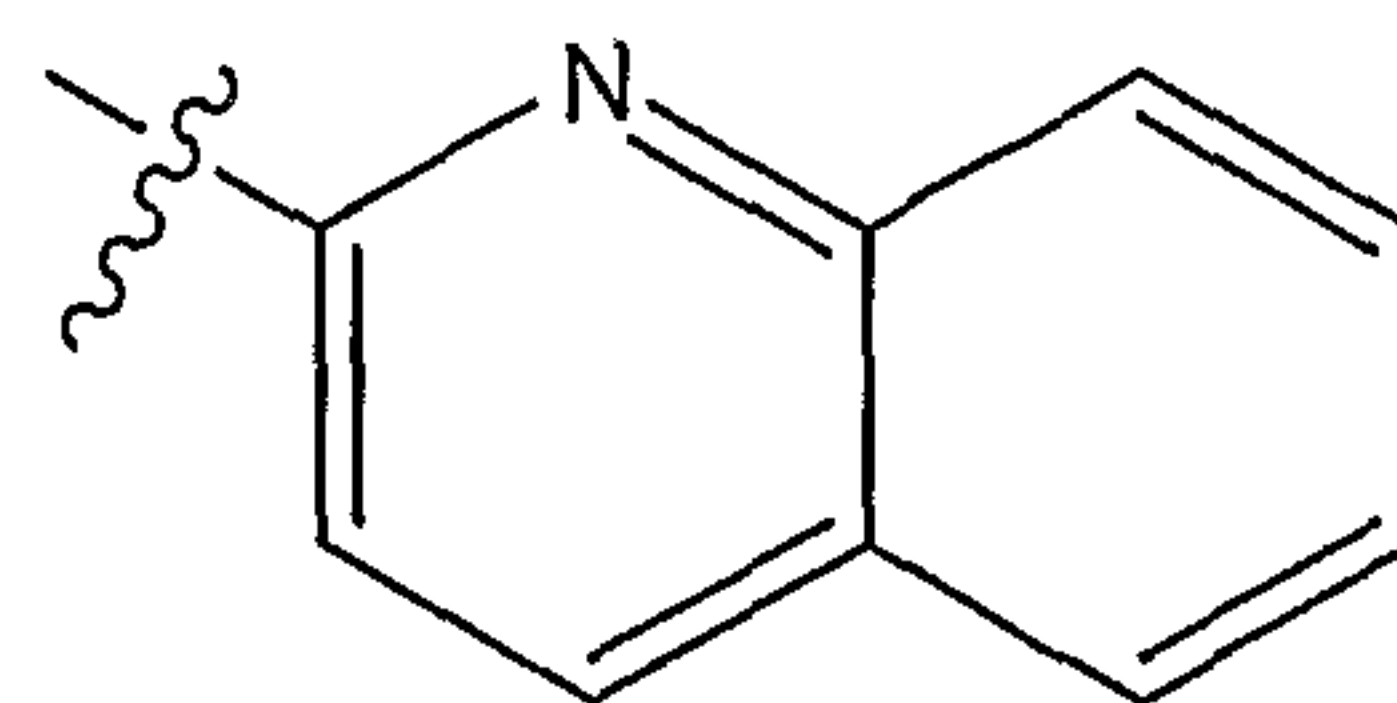
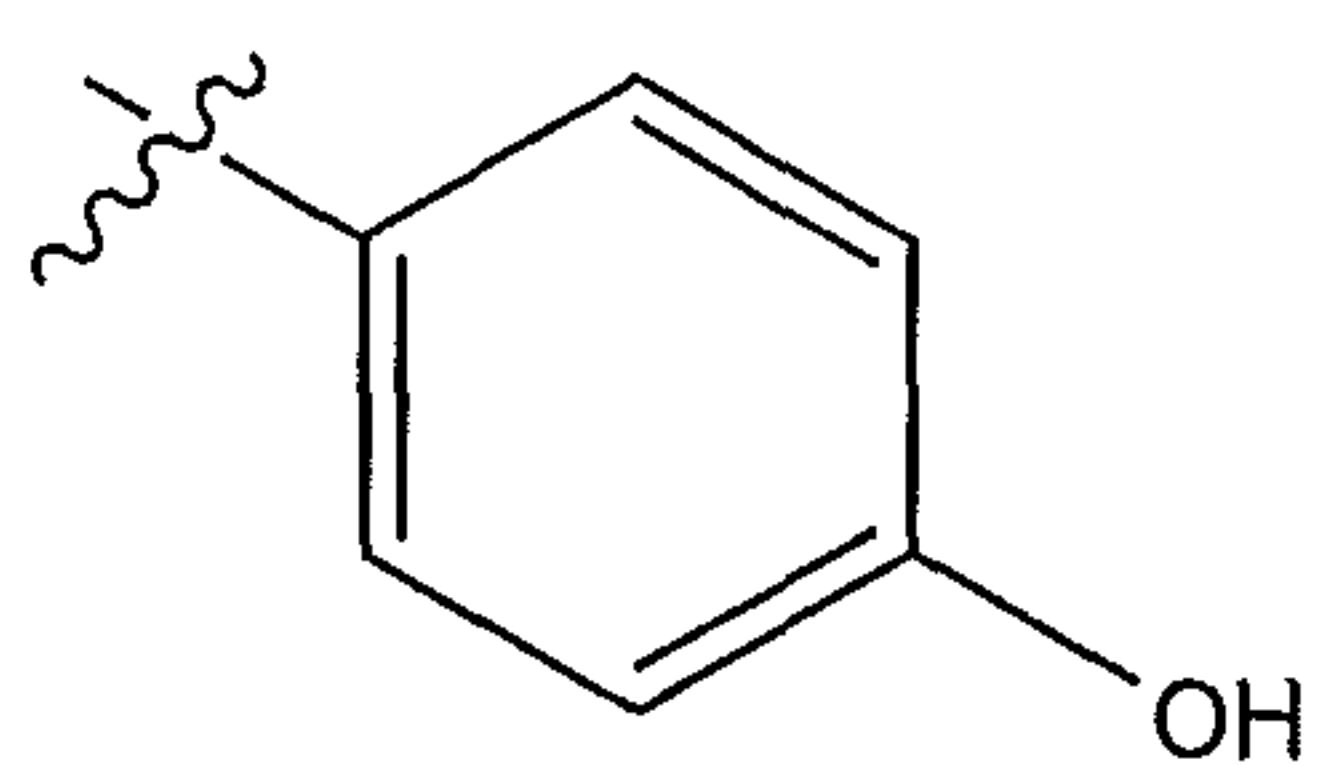
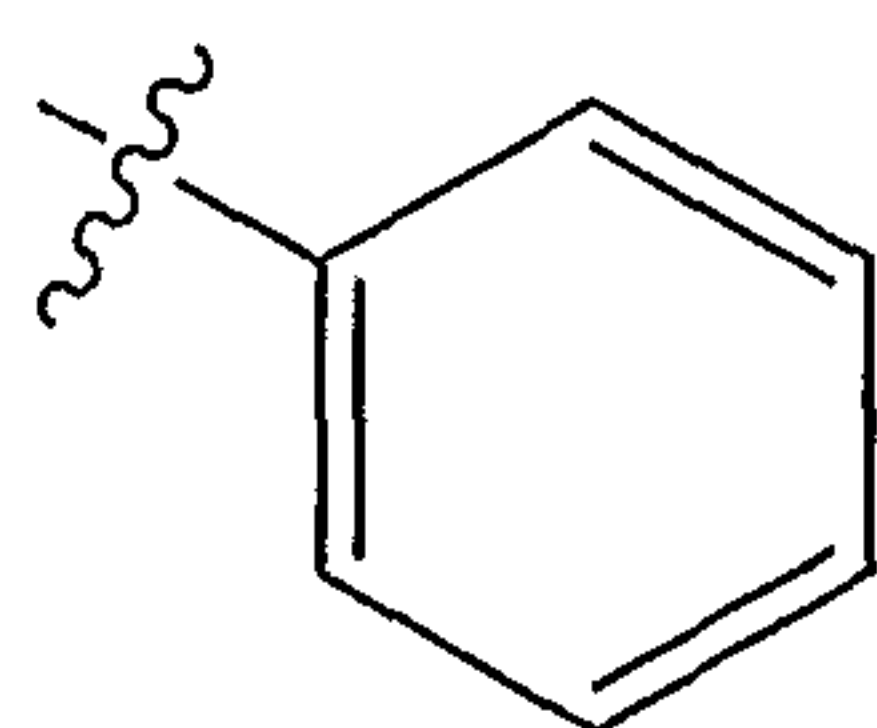


or its enantiomer, wherein, R_1 is aliphatic or aryl; R_2 is aliphatic, aryl, $-NH_2$, $-NHC(=O)-R_5$; or a moiety that participates in hydrogen bonding, wherein R_5 is aryl, heterocyclic, $-R_6-NH-C(=O)-R_7$ or $-R_6-C(=O)-NH-R_7$, wherein R_6 is an aliphatic linker of 1-6 carbons and R_7 is
 5 aliphatic, aryl, or heterocyclic, each of R_3 and R_4 is independently a hydroxy, alkoxy, halo, amino, lower-alkyl-substituted-amino, acetylamino, hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, aryl, or heterocyclic; or a pharmaceutically acceptable salt, prodrug or derivative thereof.

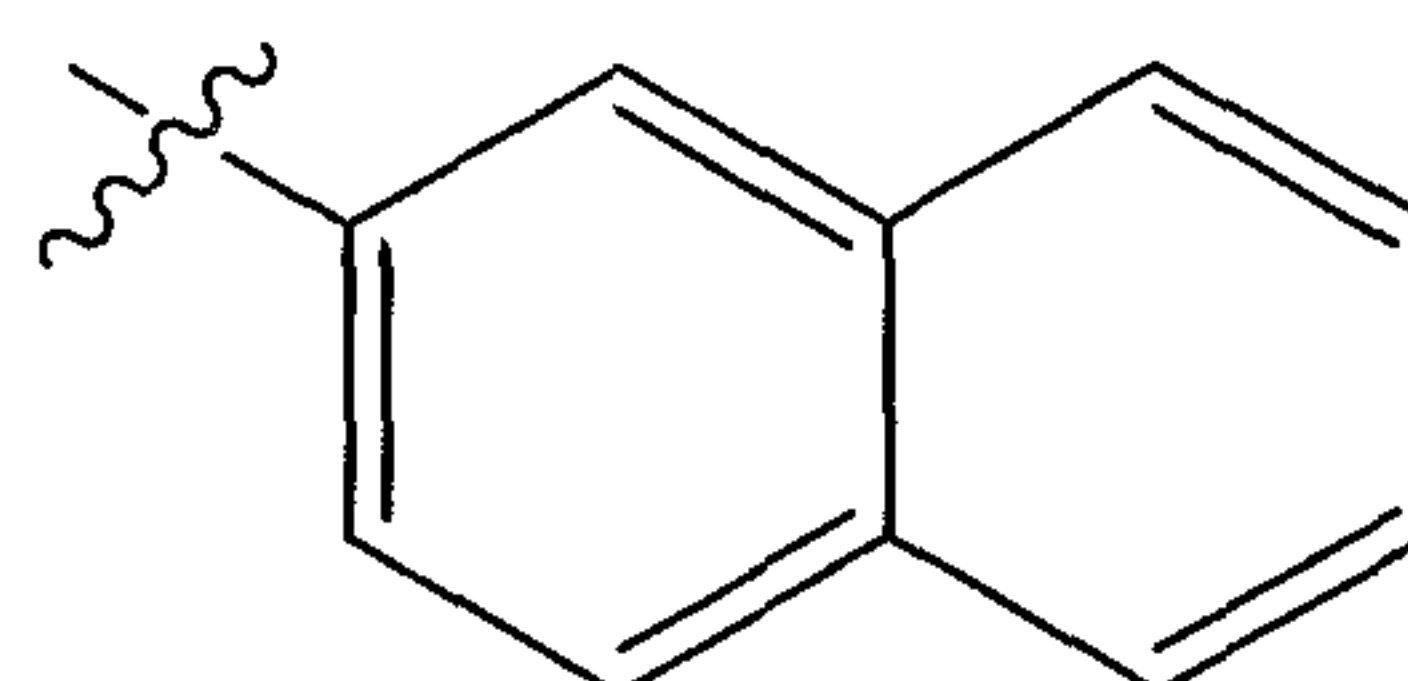
In some preferred embodiments, crystal forms and formulations of the following
 10 exemplary compounds are provided:



wherein R₂ is

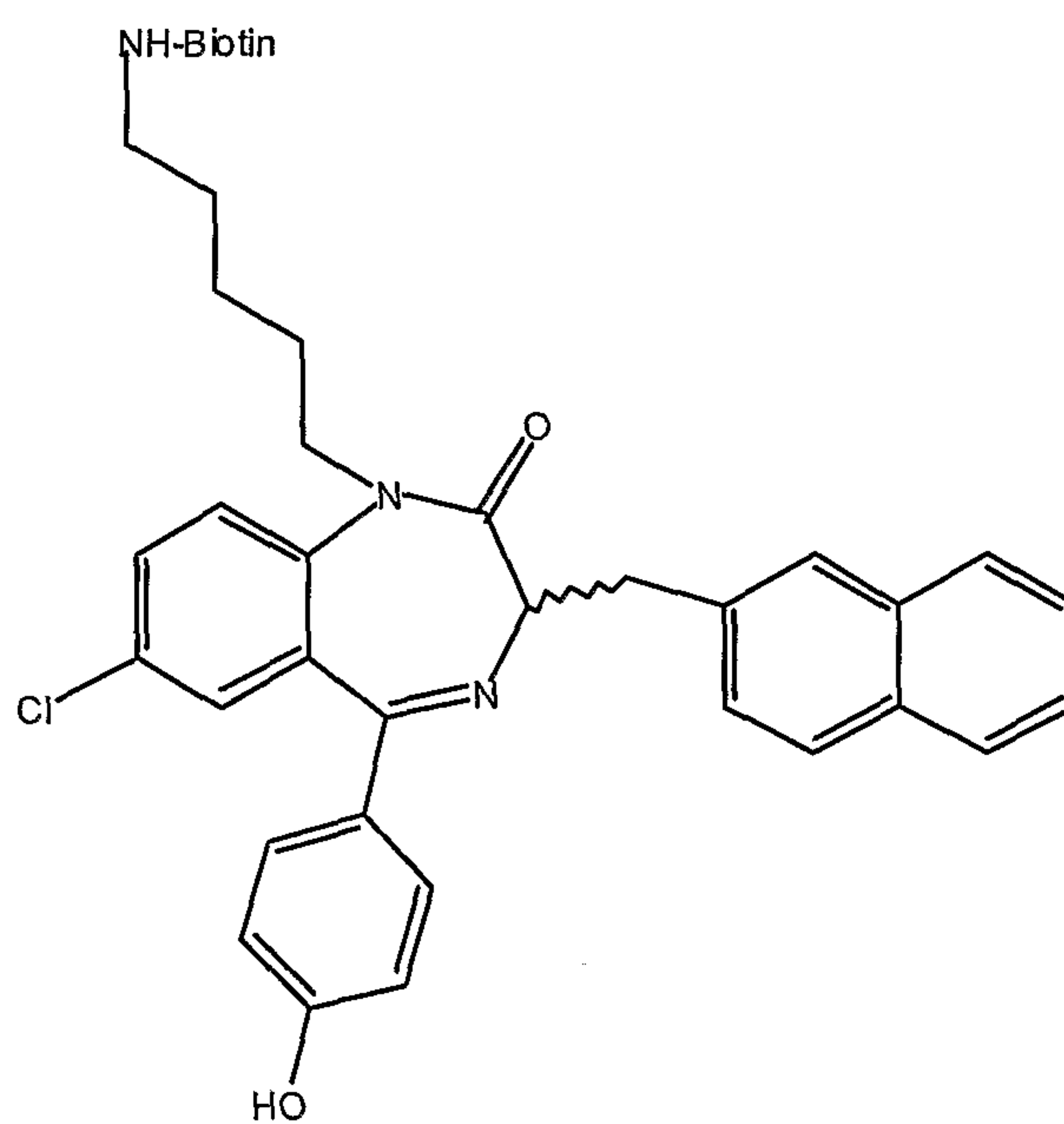
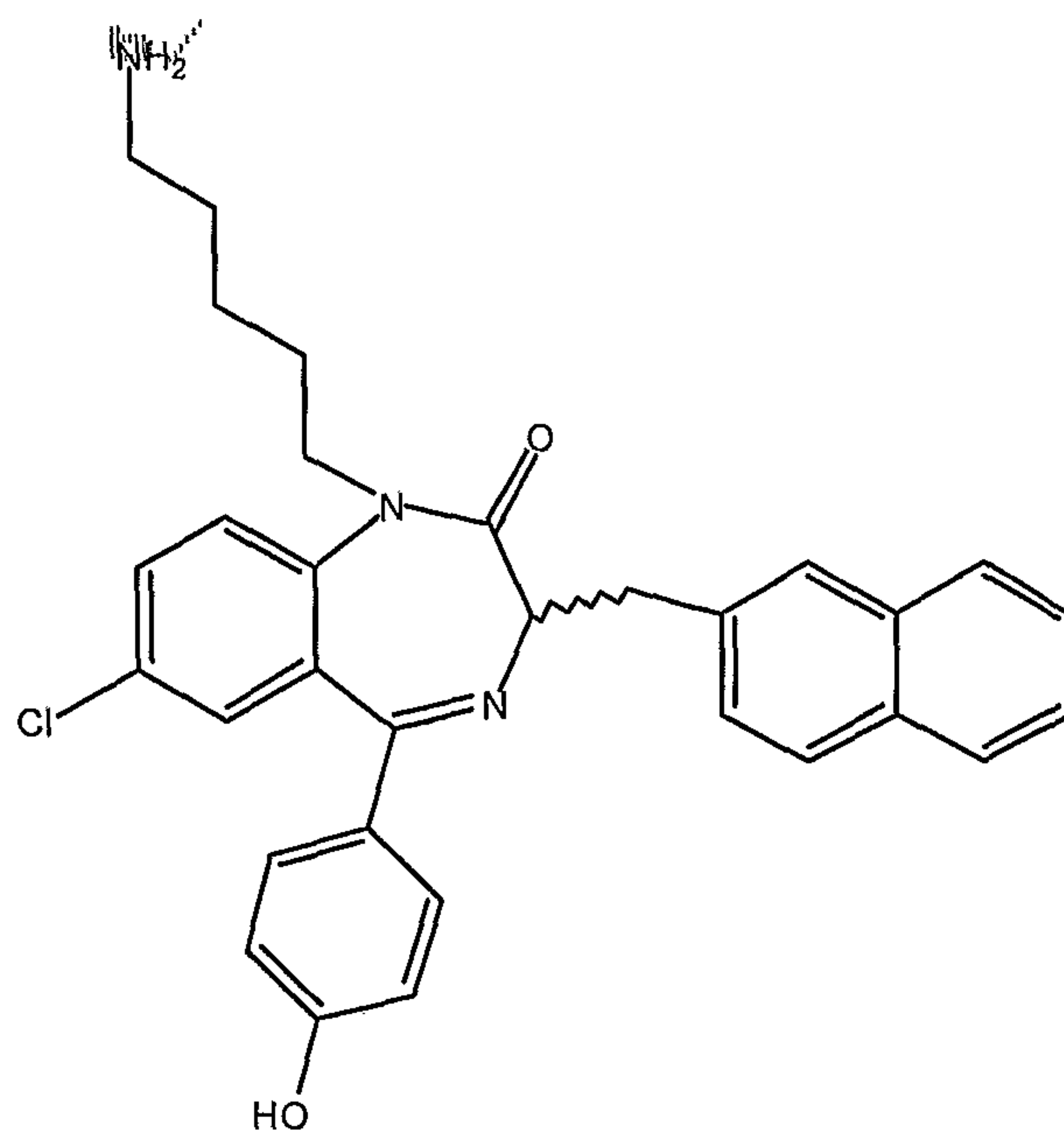


and

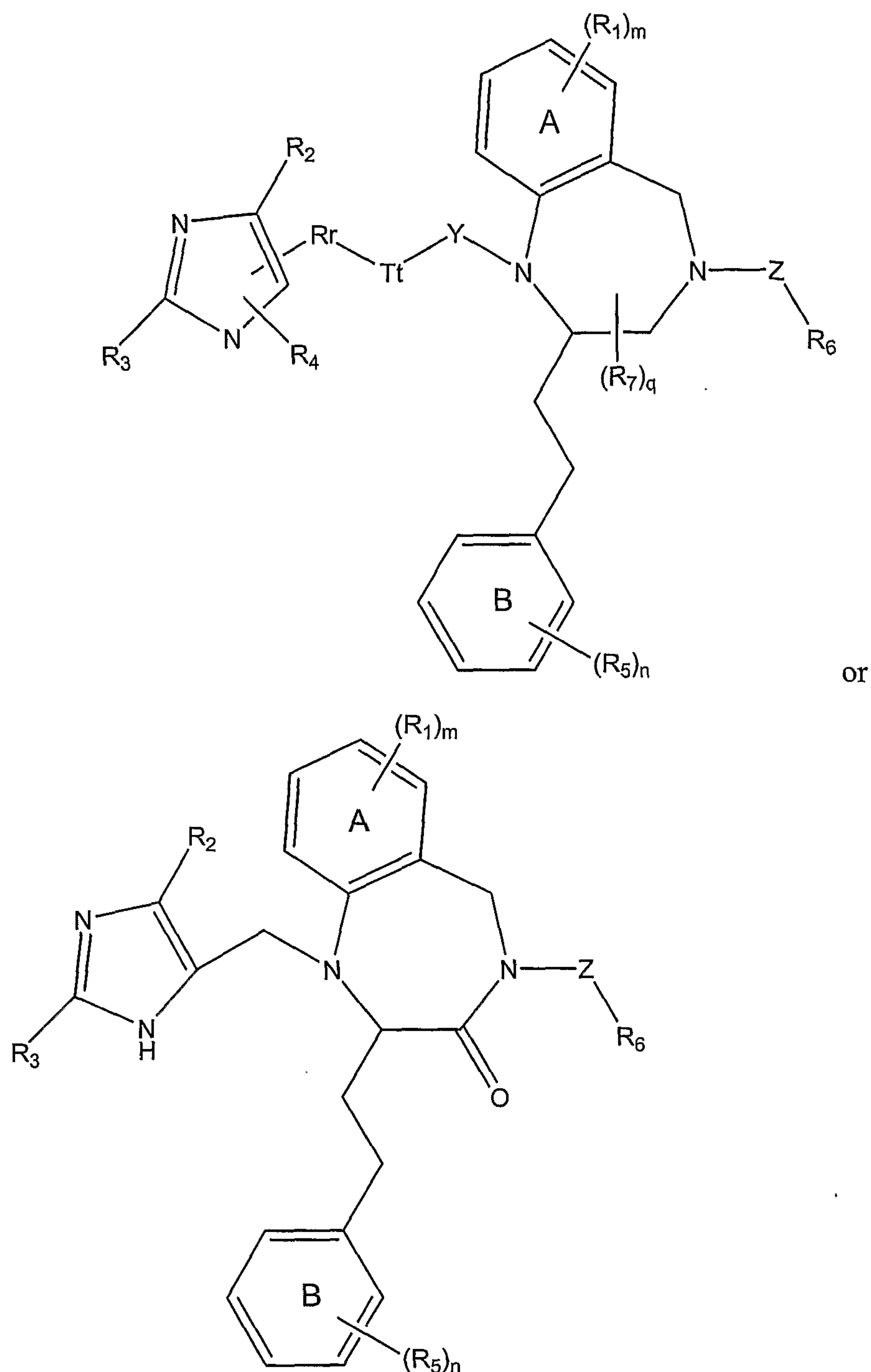


and dimethylphenyl (all isomers) and ditrifluoromethyl (all isomers).

- 5 In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:

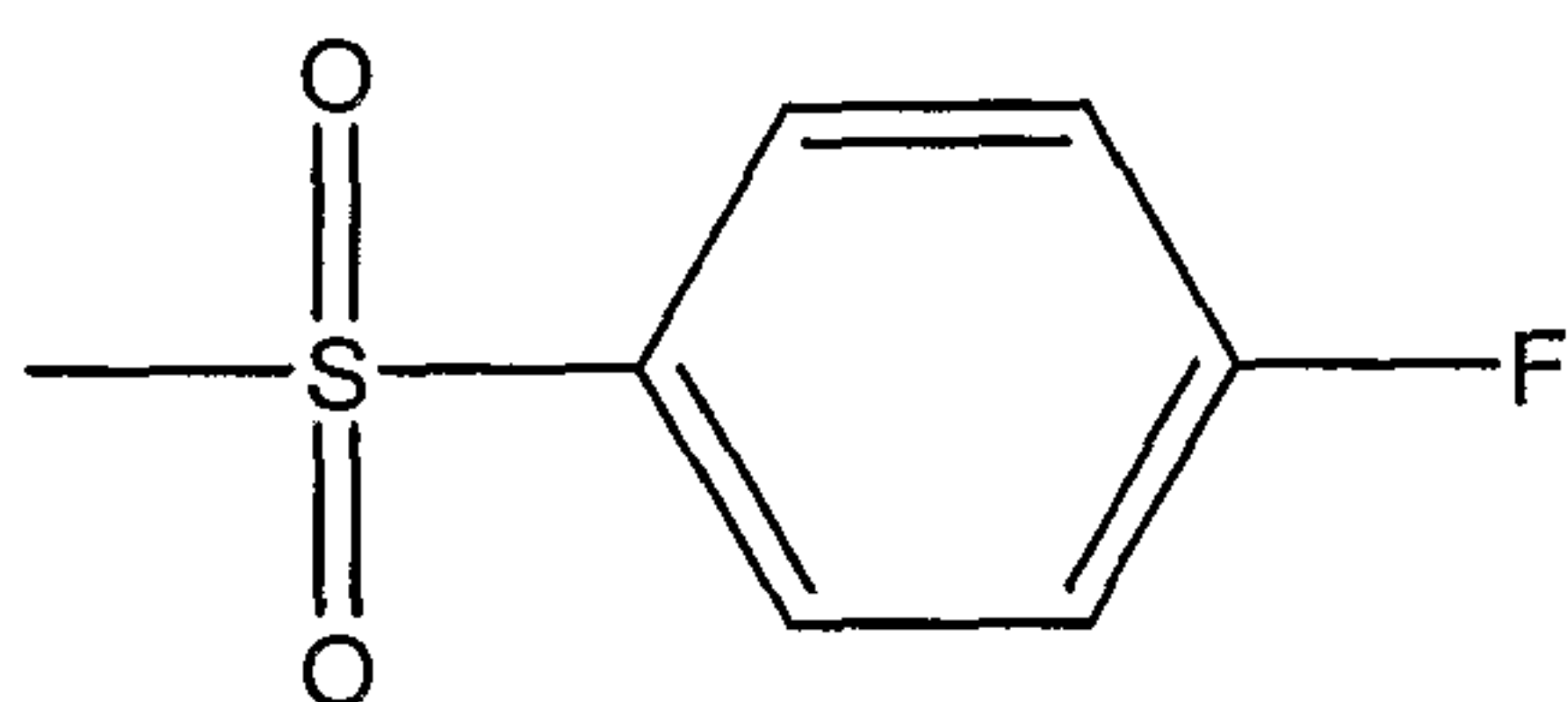


In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



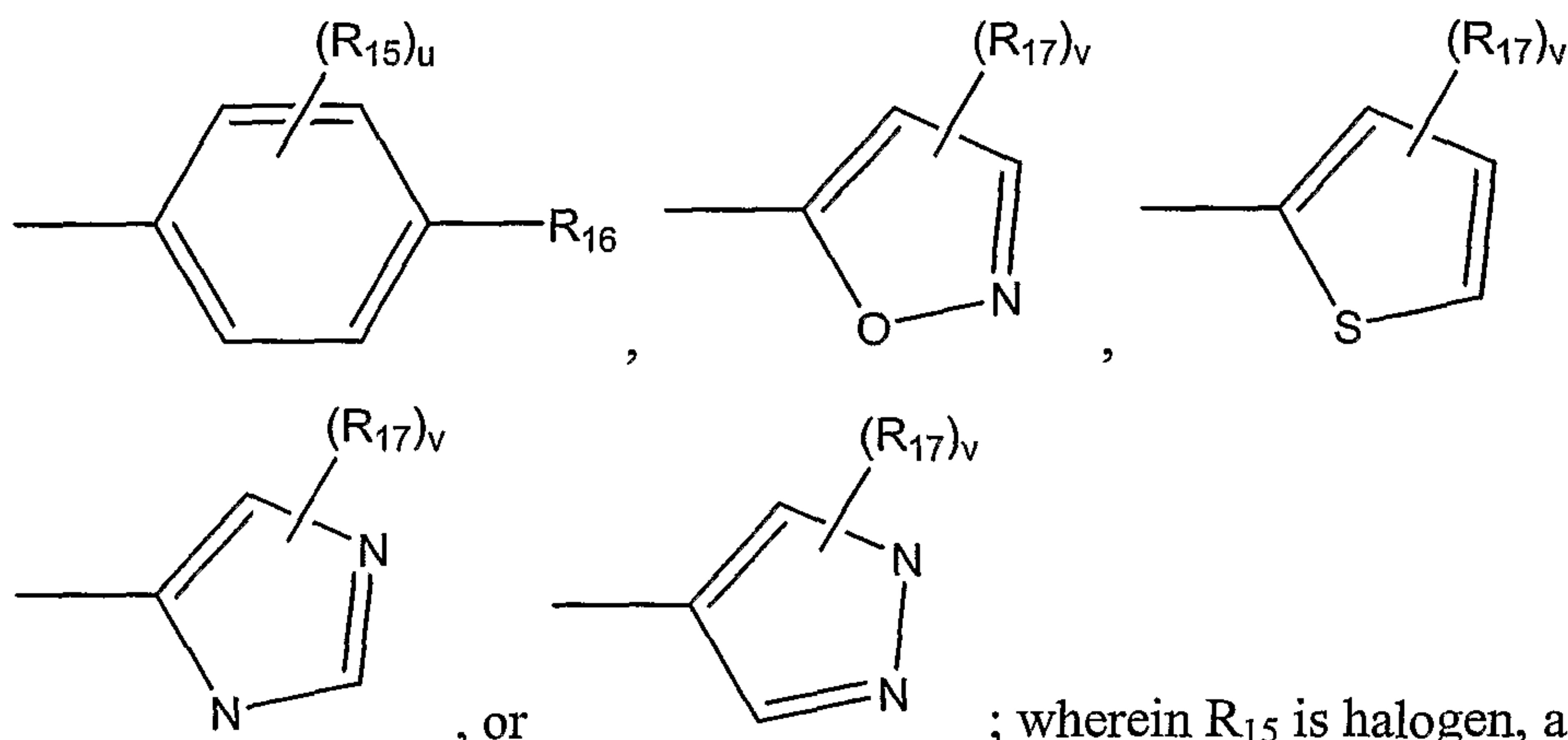
- or a stereoisomer, a pharmaceutically-acceptable salt, hydrate, or prodrug thereof, wherein:
- 5 R_1 and R_5 are attached to any available carbon atom of phenyl rings A and B, respectively, and at each occurrence are independently selected from alkyl, substituted alkyl, halogen, cyano, nitro, OR_8 , NR_8R_9 , $C(=O)R_8$, CO_2R_8 , $C(=O)NR_8R_9$, $NR_8C(=O)R_9$, $NR_8C(=O)OR_9$, $S(O)_0R_9$, $NR_8SO_2R_9$, $SO_2NR_8R_9$, cycloalkyl, heterocycle, aryl, and heteroaryl, and/or two of R_1 and/or two of R_5 join together to form a fused benzo ring; R_2 , R_3 and R_4 are

independently selected from hydrogen, alkyl, and substituted alkyl, or one of R_2 , R_3 and R_4 is a bond to R, T or Y and the other of R_2 , R_3 and R_4 is selected from hydrogen, alkyl, and substituted alkyl; Z and Y are independently selected from $C(=O)$, $-CO_2-$, $-SO_2-$, $-CH_2-$, $-CH_2C(=O)-$, and $-C(=O)C(=O)-$, or Z may be absent; R and T are selected from $-CH_2-$, $-C(=O)-$, and $-CH[(CH_2)_p(Q)]-$, wherein Q is $NR_{10}R_{11}$, OR_{10} or CN; R_6 is selected from alkyl, alkenyl, substituted alkyl, substituted alkenyl, aryl, cycloalkyl, heterocyclo, and heteroaryl; provided that where R_2 is hydrogen, Z- R_6 together are not $-SO_2-Me$ or



; R_7 is selected from hydrogen, alkyl, substituted alkyl, alkenyl, substituted alkenyl, aminoalkyl, halogen, cyano, nitro, keto ($=O$), hydroxy, alkoxy, alkylthio, $C(=O)H$, acyl, CO_2H , alkoxycarbonyl, carbamyl, sulfonyl, sulfonamidyl, cycloalkyl, heterocycle, aryl, and heteroaryl; R_8 and R_9 are independently selected from hydrogen, alkyl, substituted alkyl, cycloalkyl, heterocycle, aryl, and heteroaryl, or R_8 and R_9 taken together to form a heterocycle or heteroaryl, except R_9 is not hydrogen when attached to a sulfonyl group as in SO_2R_9 ; R_{10} and R_{11} are independently selected from hydrogen, alkyl, and substituted alkyl; m and n are independently selected from 0, 1, 2 and 3; o, p and q are independently 0, 1 or 2; and r and t are 0 or 1.

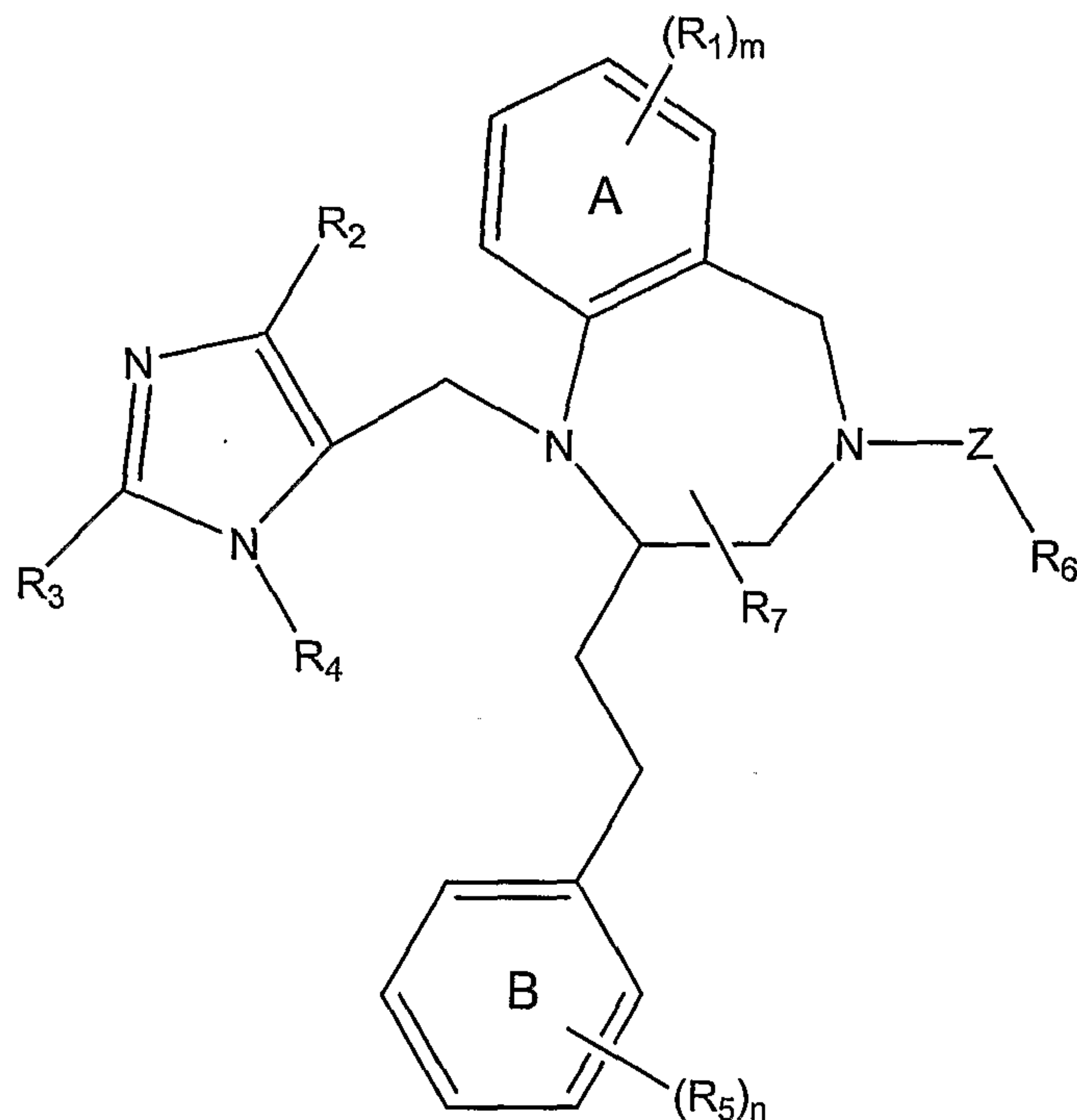
In further exemplary compounds, Z- R_6 taken together are selected from: i. thiophenyl optionally substituted with R_{14} ; ii. imidazolyl optionally substituted with R_{14} ; iii. $-CH(aryl)(CO_2C_{1-6}alkyl)$; iv. $-CO_2-alkyl$; v. $-SO_2-alkyl$ optionally substituted with up to three of halogen and/or phenyl; vi. $-SO_2-alkenyl$ optionally substituted with phenyl; and vii.



; wherein R_{15} is halogen, alkyl, nitro, cyano, hydroxy, alkoxy, $NHC(=O)alkyl$, and/or two R_{15} groups are taken together to form a fused

benzo ring or a five to six membered heteroaryl; R_{16} is selected from hydrogen, halogen, alkyl, nitro, cyano, hydroxy, alkoxy, $\text{NHC}(=\text{O})\text{alkyl}$, and phenyloxy or benzyloxy in turn optionally substituted with 1 to 3 of halogen, cyano, and $\text{C}_{1-4}\text{alkoxy}$; R_{17} is selected from alkyl, alkoxy, $\text{CO}_2\text{C}_{1-6}\text{alkyl}$, and SO_2phenyl ; and u and v are independently 0, 1 or 2.

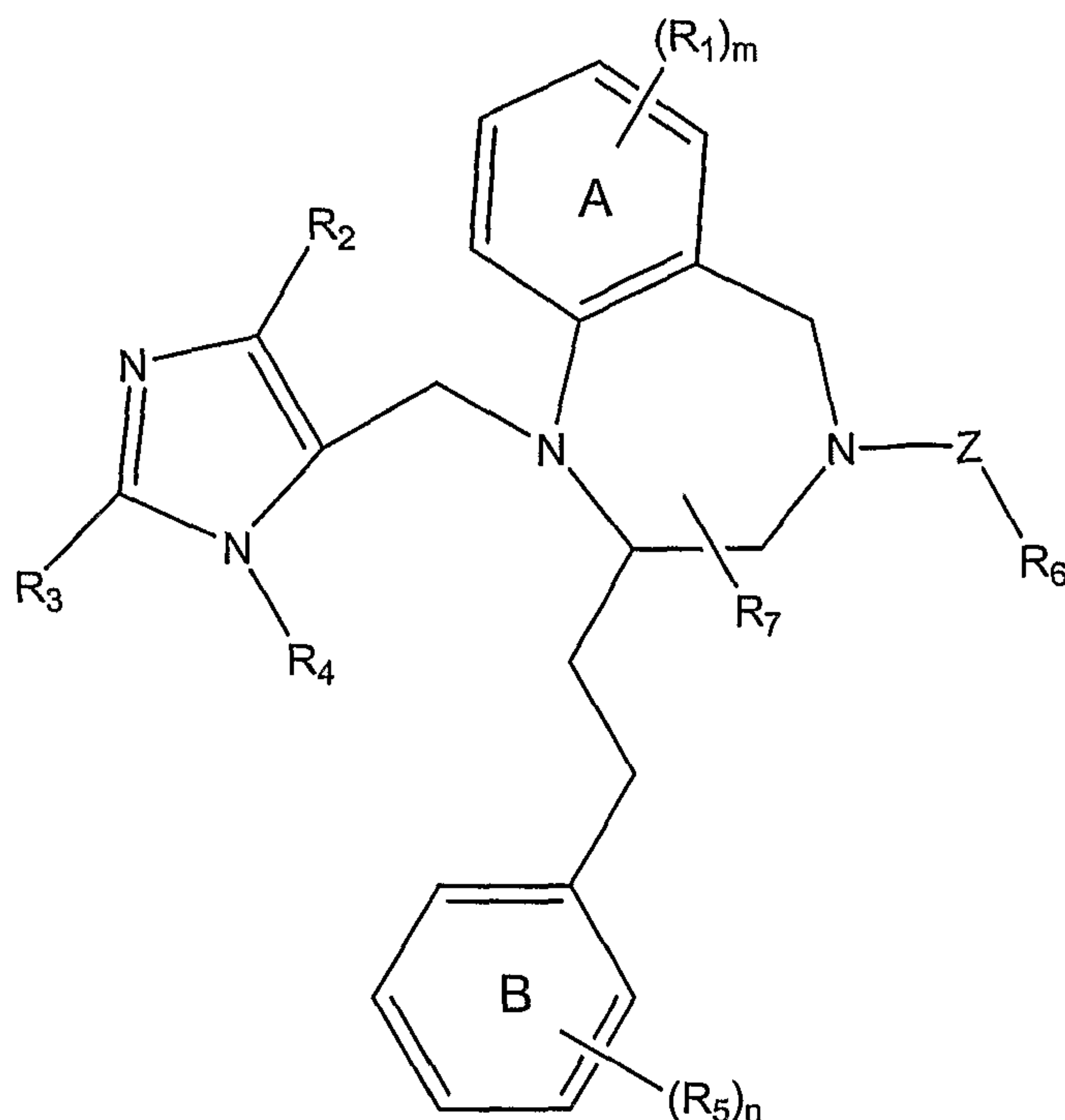
- 5 In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



or a stereoisomer, a pharmaceutically-

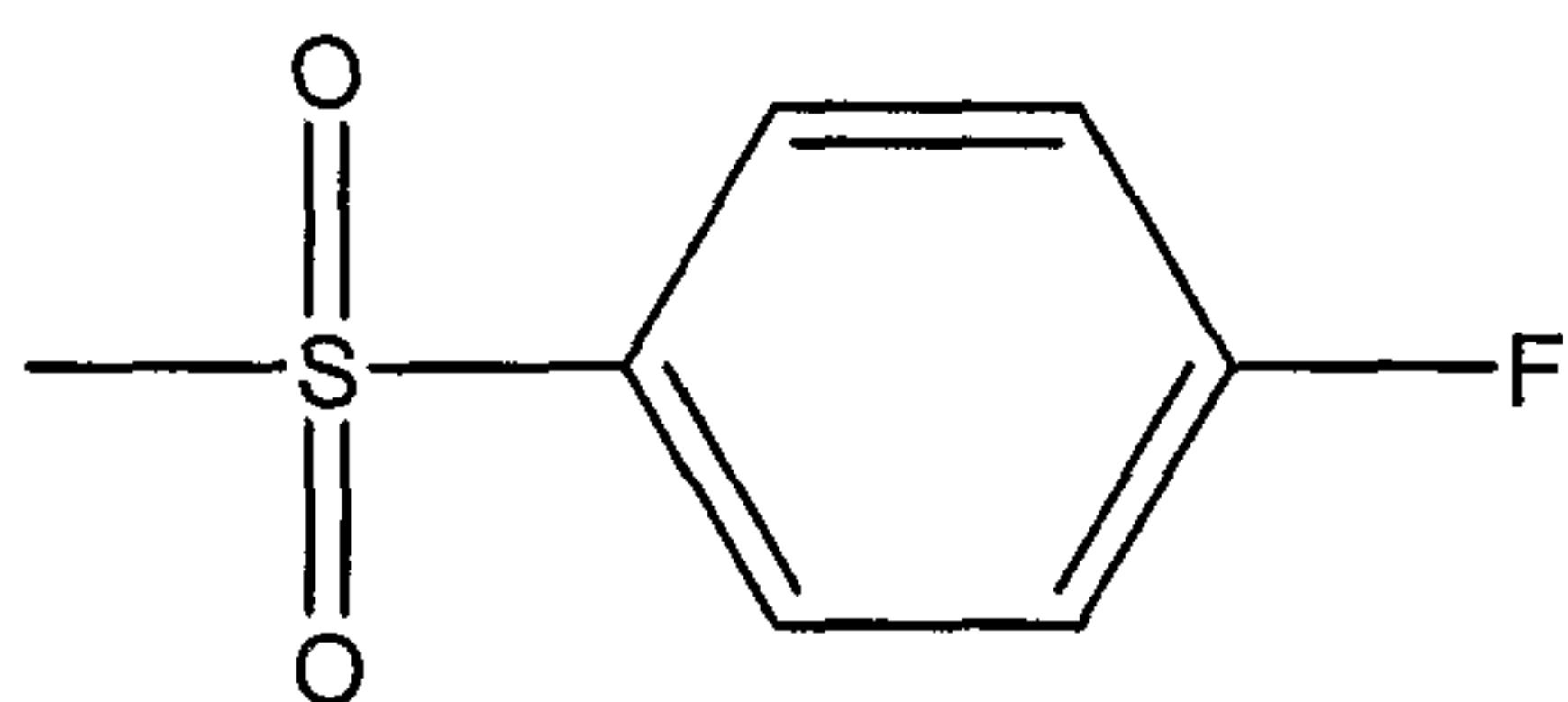
- acceptable salt, hydrate, or prodrug thereof, in which: R_1 and R_5 are attached to any available carbon atom of phenyl ring A and phenyl ring B, respectively, and at each
- 10 occurrence are independently selected from $\text{C}_{1-6}\text{alkyl}$, substituted $\text{C}_{1-6}\text{alkyl}$, halogen, cyano, $\text{O}(\text{C}_{1-6}\text{alkyl})$, $\text{O}(\text{phenyl})$, $\text{O}(\text{benzyl})$, NH_2 , $\text{NH}(\text{C}_{1-6}\text{alkyl})$, $\text{N}(\text{C}_{1-6}\text{alkyl})_2$, $\text{C}(=\text{O})\text{H}$, $\text{C}(=\text{O})(\text{C}_{1-6}\text{alkyl})$, CO_2H , $\text{CO}_2(\text{C}_{1-6}\text{alkyl})$, $\text{C}(=\text{O})\text{NH}_2$, $\text{C}(=\text{O})\text{NH}(\text{C}_{1-6}\text{alkyl})$, $\text{C}(=\text{O})\text{N}(\text{C}_{1-6}\text{alkyl})_2$, $\text{NHC}(=\text{O})(\text{C}_{1-6}\text{alkyl})$, $\text{S}(\text{O})_2(\text{C}_{1-6}\text{alkyl})$, $\text{NHSO}_2(\text{C}_{1-6}\text{alkyl})$, SO_2NH_2 , $\text{SO}_2\text{NH}(\text{C}_{1-6}\text{alkyl})$, $\text{SO}_2\text{N}(\text{C}_{1-6}\text{alkyl})_2$, $\text{C}_{3-7}\text{cycloalkyl}$, phenyl, five or six membered heteroaryl, or four to seven
- 15 membered heterocyclo, and/or two of R_1 and/or two of R_5 join together to form a fused benzo ring; R_2 and R_3 are independently selected from hydrogen and $\text{C}_{1-4}\text{alkyl}$; Z is $-\text{CO}_2-$, $-\text{SO}_2-$, or is absent; R_6 is selected from optionally-substituted alkyl, alkenyl, aryl, and heteroaryl; m and n are independently selected from 0, 1, and 2; and q is 0 or 1.

- 20 In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



, or a stereoisomer, a pharmaceutically-

acceptable salt, hydrate, or prodrug thereof, wherein: R_1 and R_5 are attached to any available carbon atom of phenyl ring A and phenyl ring B, respectively, and at each occurrence are independently selected from alkyl, substituted alkyl, halogen, cyano, nitro, hydroxy, alkoxy, alkylthio, alkylamino, $C(=O)H$, acyl, CO_2H , alkoxycarbonyl, carbamyl, sulfonyl, sulfonamidyl, cycloalkyl, heterocycle, aryl, and heteroaryl, and/or two of R_1 and/or two of R_5 join together to form a fused benzo ring; R_2 , R_3 and R_4 are independently selected from hydrogen and alkyl; Z is $-CO_2-$, $-SO_2-$, or is absent; R_6 is selected from: a) C_{1-4} alkyl or C_{1-4} alkenyl optionally substituted with up to three of halogen, aryl and CO_2C_{1-6} alkyl; b) phenyl optionally substituted with up to three R_{12} and/or having fused thereto a benzo-ring or a five to six membered heteroaryl; c) heteroaryl selected from thiophenyl, imidazolyl, pyrazolyl, and isoxazolyl, wherein said heteroaryl is optionally substituted with up to two R_{12} , provided that where R_2 is hydrogen, Z- R_6 together are not $-SO_2-Me$ or



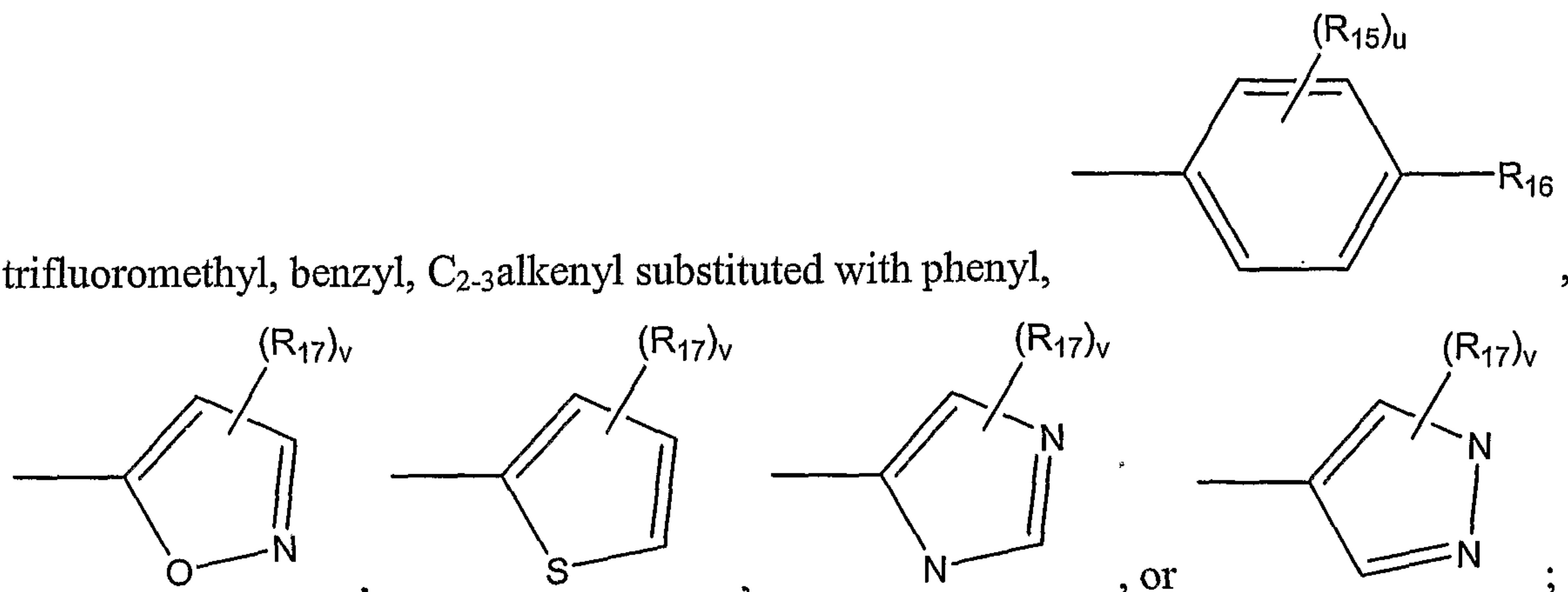
; R_7 is selected from hydrogen, keto ($=O$), C_{1-6} alkyl,

substituted C_{1-6} alkyl, halogen, cyano, $O(C_{1-6}$ alkyl), $O(phenyl)$, $O(benzyl)$, NH_2 , $NH(C_{1-6}$ alkyl), $N(C_{1-6}alkyl)_2$, $C(=O)H$, $C(=O)(C_{1-6}alkyl)$, CO_2H , $CO_2(C_{1-6}alkyl-)$; R_{12} at each occurrence is independently selected from each other R_{12} from the group consisting of C_{1-6} alkyl, halogen, nitro, cyano, hydroxy, alkoxy, $NHC(=O)alkyl$, $-CO_2alkyl$, $-SO_2phenyl$,

five to six membered monocyclic heteroaryl, and phenyloxy or benzyloxy in turn optionally substituted with halogen, C₁₋₄alkyl, and/or O(C₁₋₄alkyl); and m and n are independently selected from 0, 1, or 2.

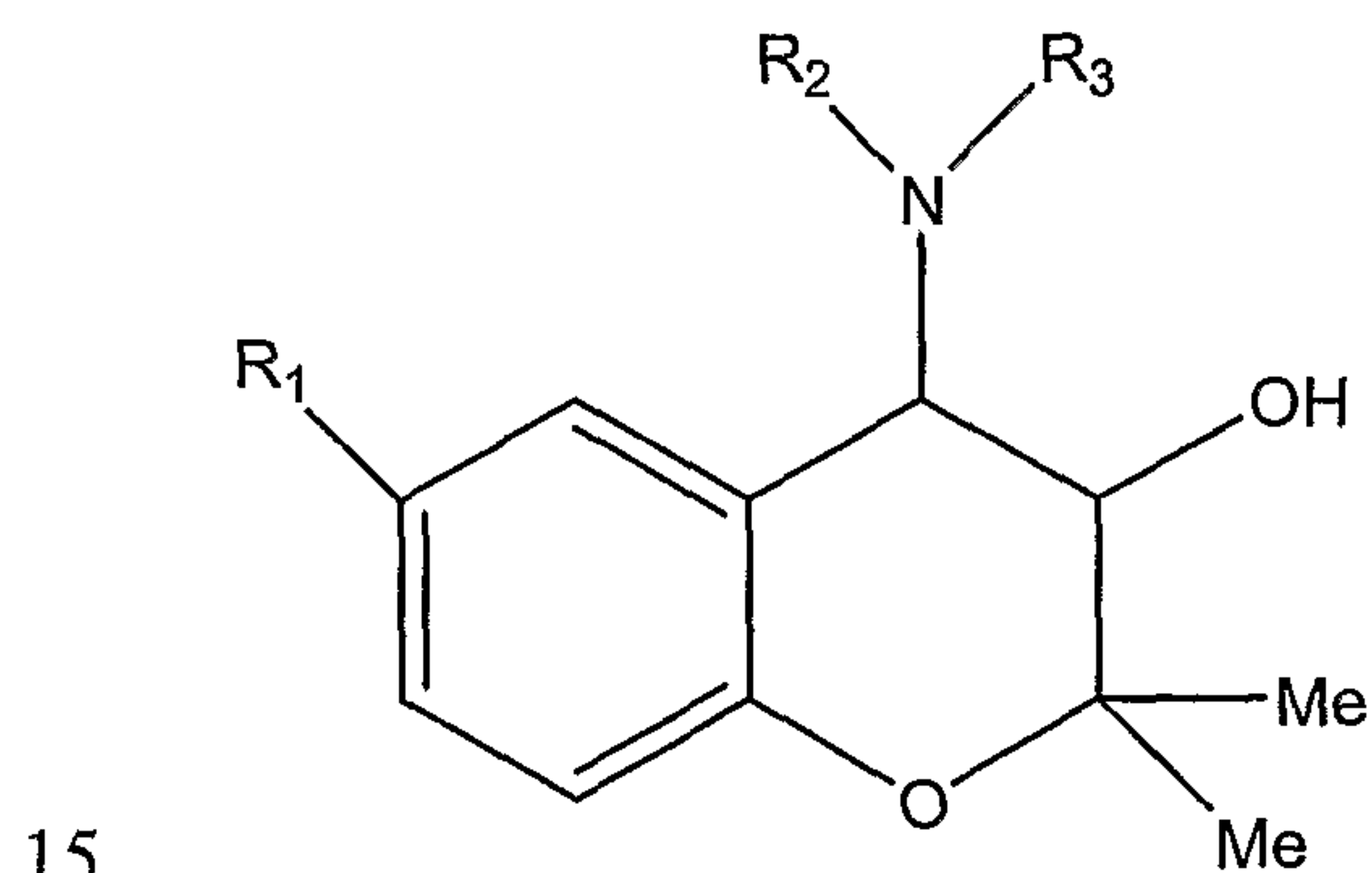
In further exemplary compounds, Z is -SO₂--; R₆ is selected from C₁₋₄alkyl,

5 trifluoromethyl, benzyl, C₂₋₃alkenyl substituted with phenyl,



R₁₅ is halogen, alkyl, nitro, cyano, hydroxy, alkoxy, NHC(=O)alkyl, and/or two R₁₅ groups are taken together to form a fused benzo ring or a five to six membered heteroaryl; R₁₆ is selected from hydrogen, halogen, alkyl, nitro, cyano, hydroxy, alkoxy, NHC(=O)alkyl, and
 10 phenyloxy or benzyloxy in turn optionally substituted with 1 to 3 of halogen, cyano, and C₁₋₄alkoxy; R₁₇ is selected from alkyl, alkoxy, CO₂C₁₋₆alkyl, and SO₂phenyl; and u and v are independently 0, 1 or 2.

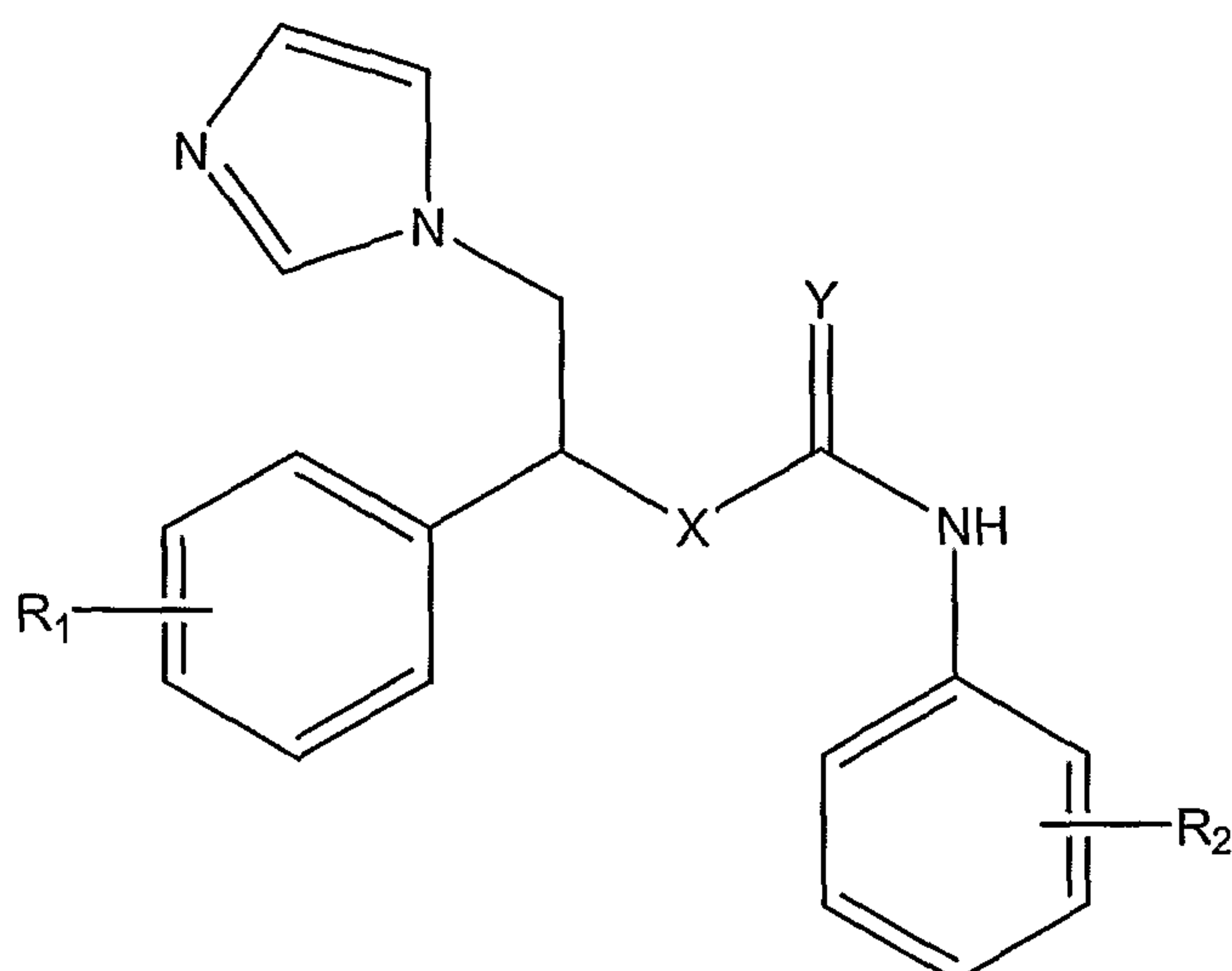
In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



15

or a stereoisomer, a pharmaceutically-acceptable salt, hydrate, or prodrug thereof, wherein: R₁ is selected from the group consisting of H, CN and SO₂-piperidine; R₂ is selected from the group consisting of H, 4-Cl-Ph, Ph, and 2-Me-imidazole; R₃ is selected from the group
 20 consisting of H, CH₂-2-imidazole, and CH₂-2-oxazole.

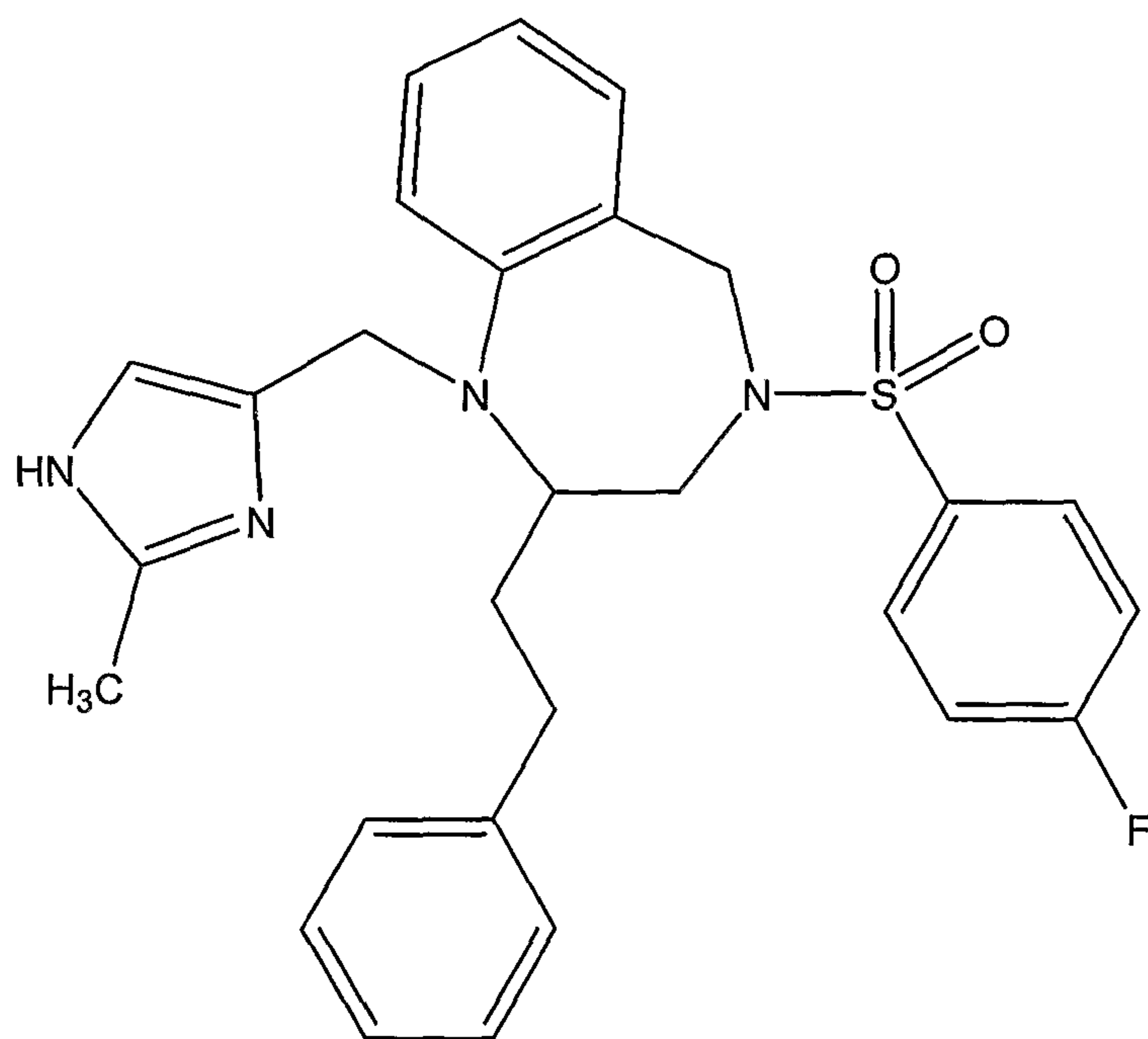
In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



or a stereoisomer, a pharmaceutically-acceptable salt, hydrate, or prodrug thereof, wherein:

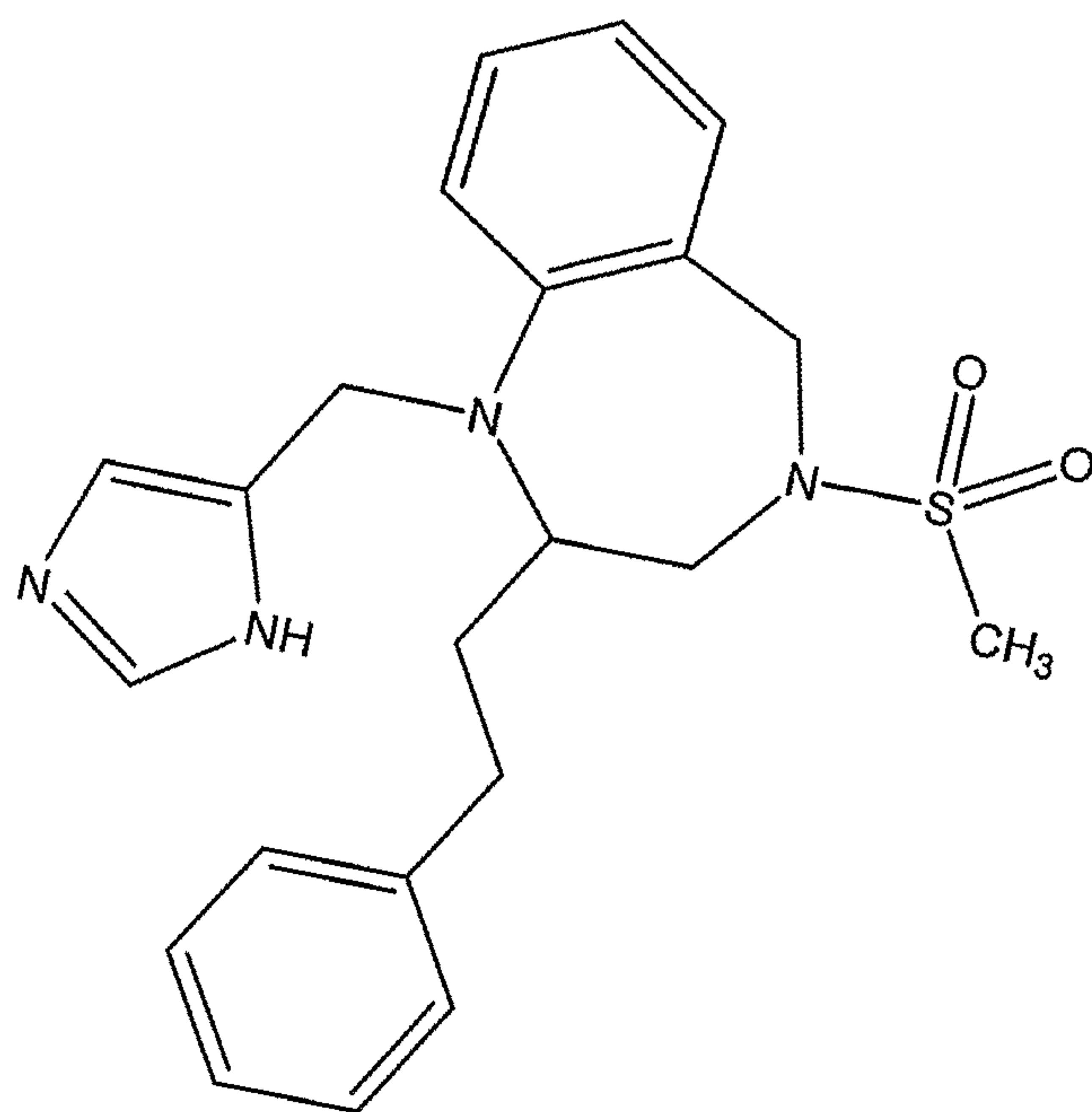
R₁ is selected from the group consisting of H, 2,4-Cl₂, 2,4-Me₂, and 2,5-(CF₃)₂; R₂ is
 5 selected from the group consisting of H, 4-Cl, 4-Me, 2,4-Cl₂, 2,4-Me₂, 3-Cl; X is selected from the group consisting of O and NH; Y is selected from the group consisting of S, O, NCN, CO(3-CN-Ph), CO(4-CN-Ph), CO(4-Cl-Ph), and COEt.

In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:

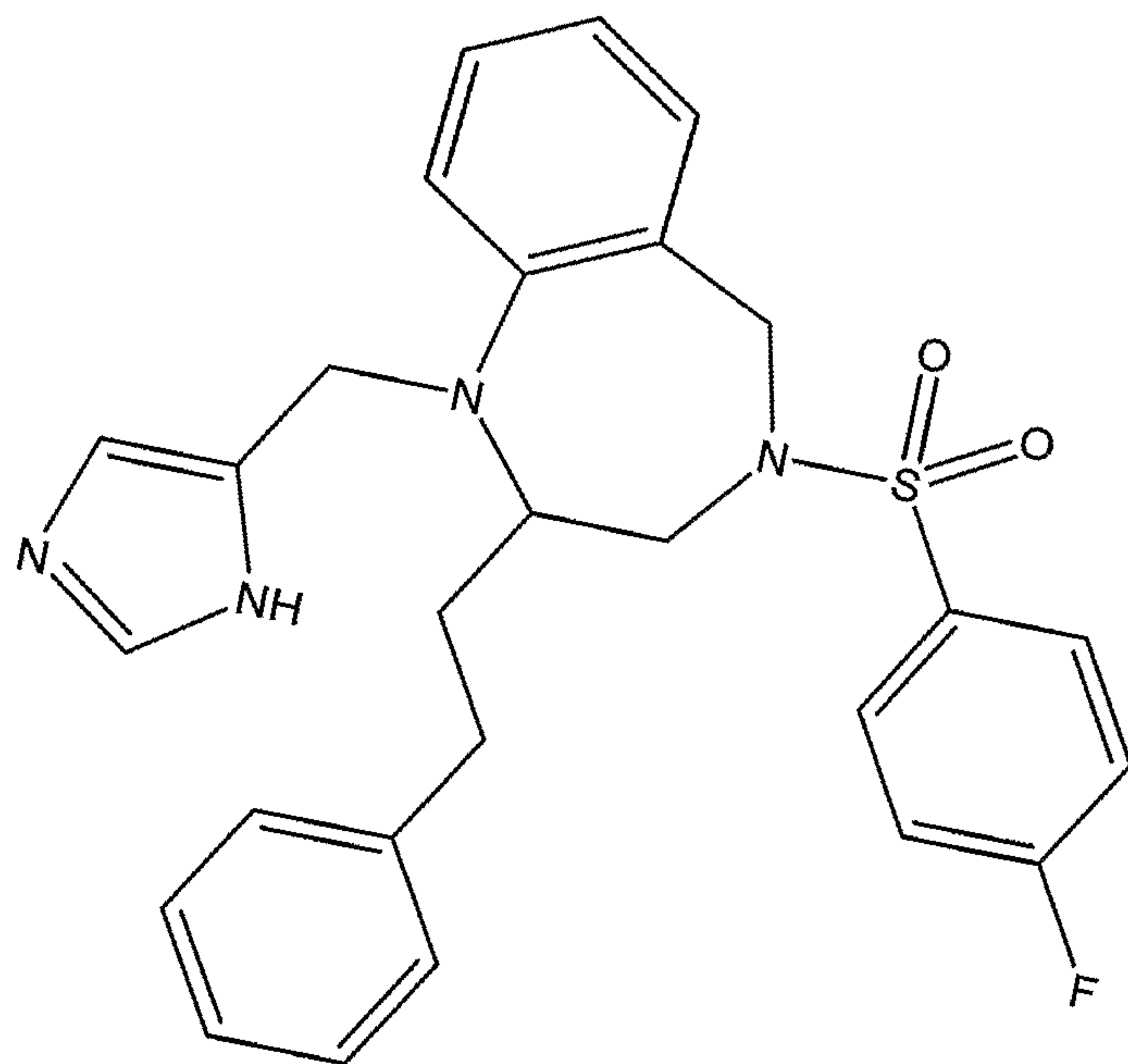


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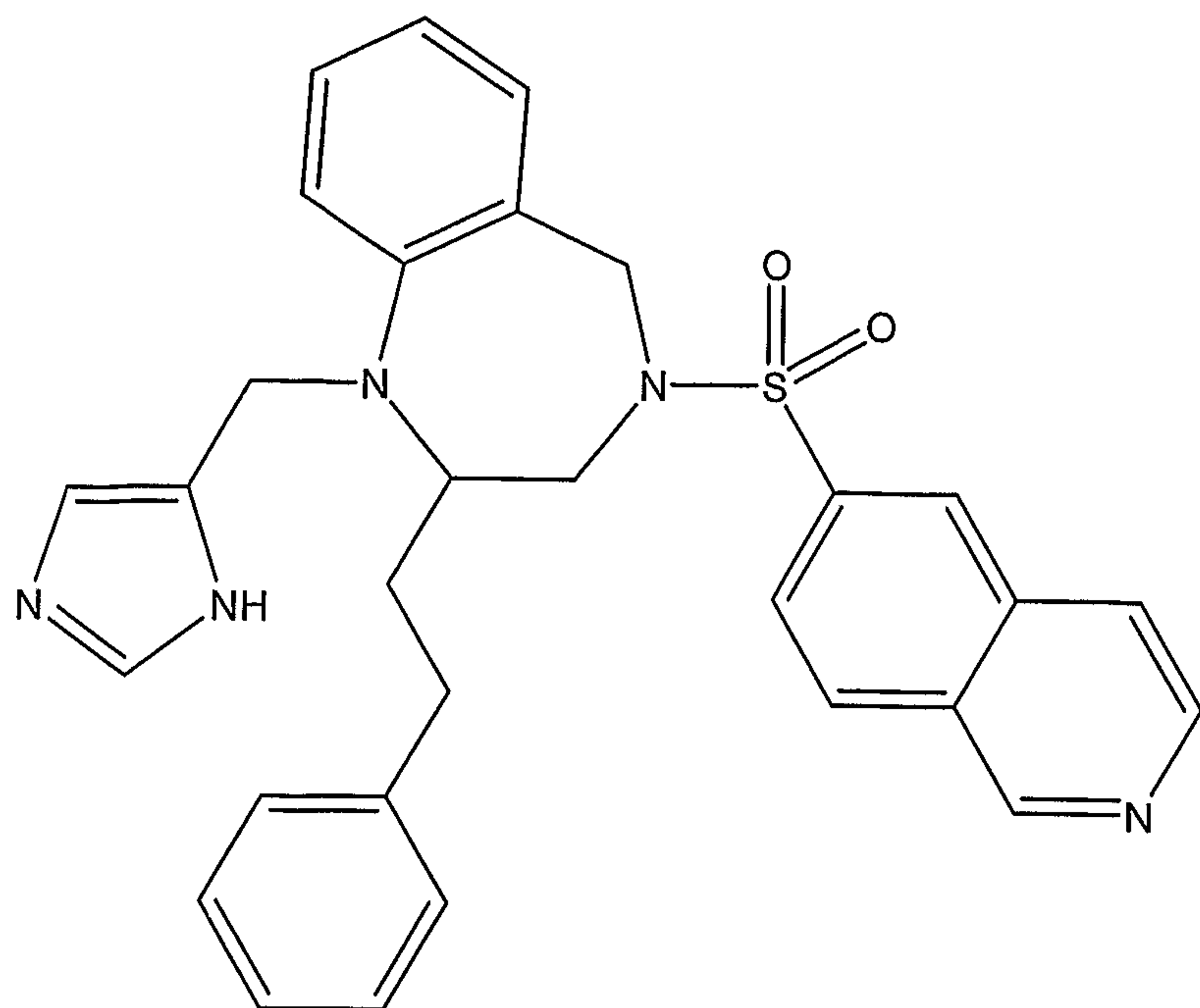
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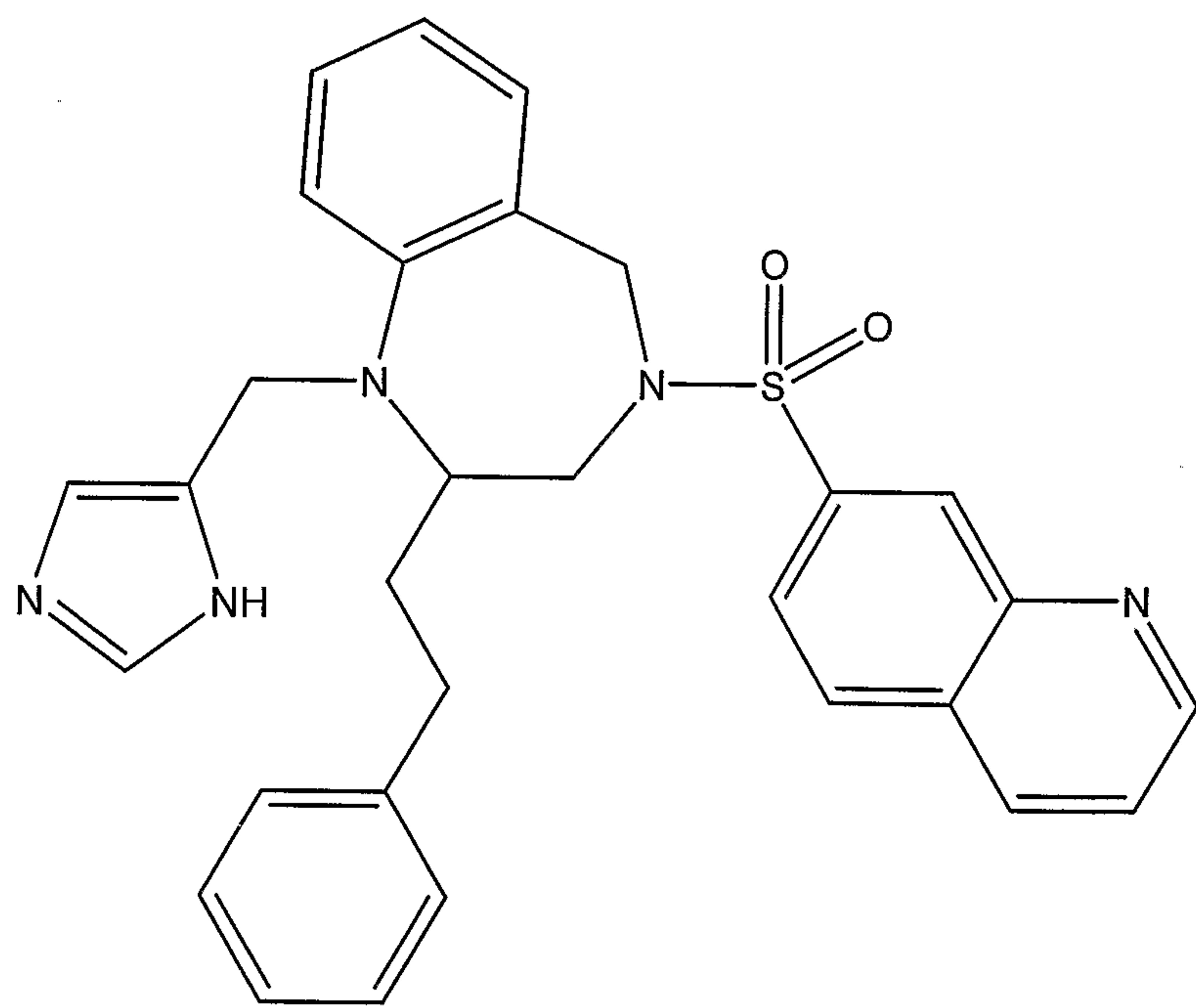
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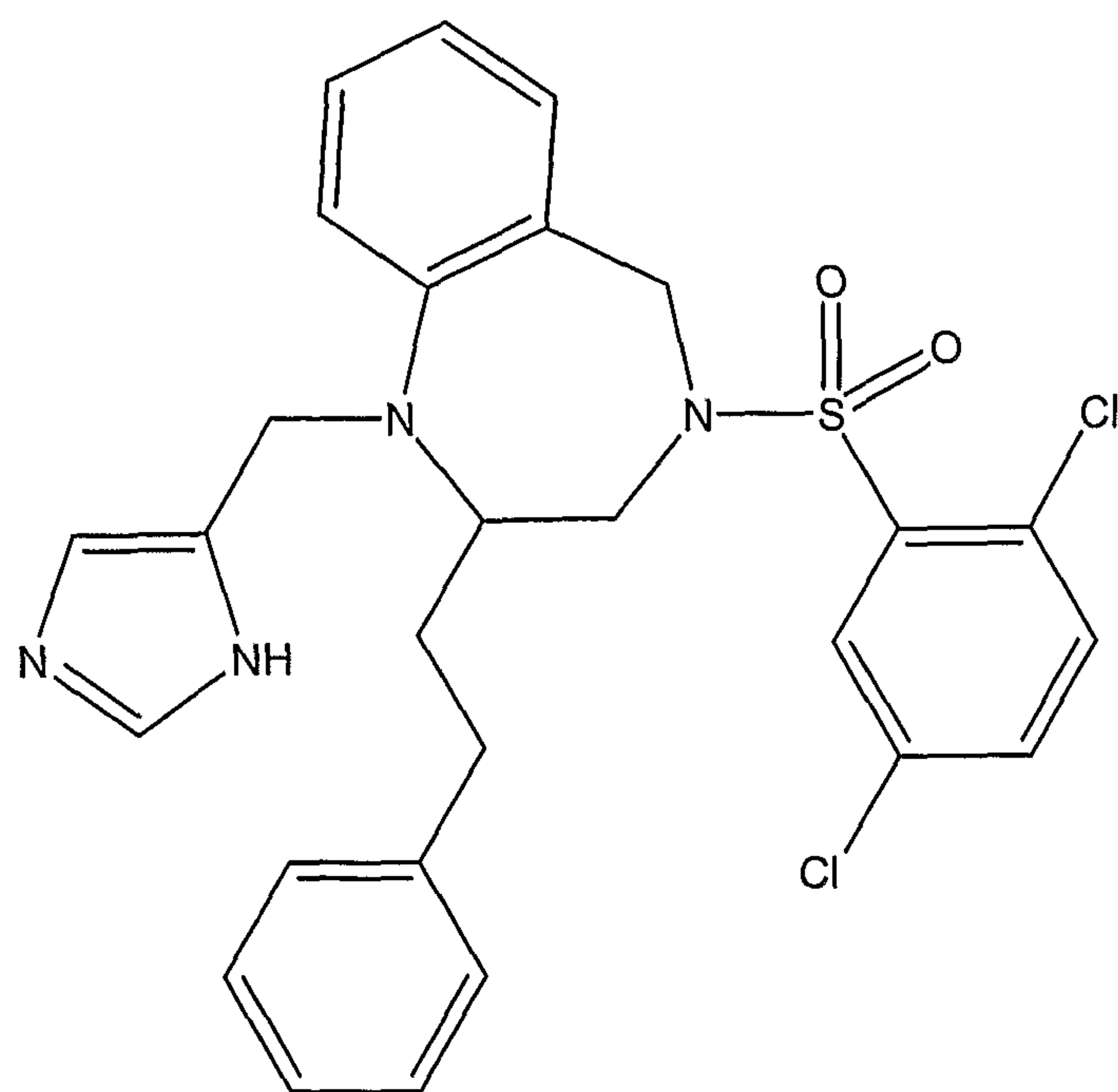
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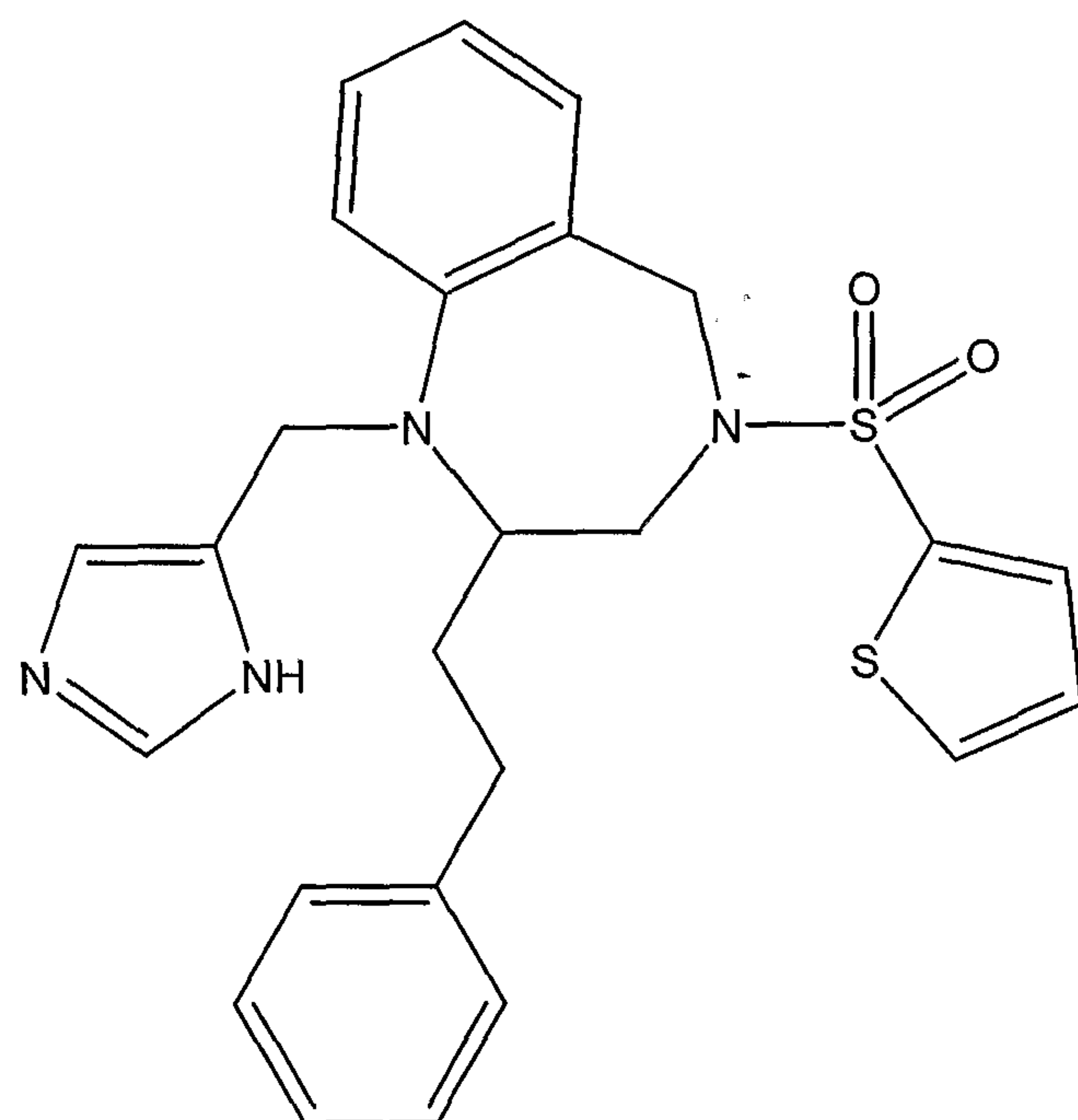
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;



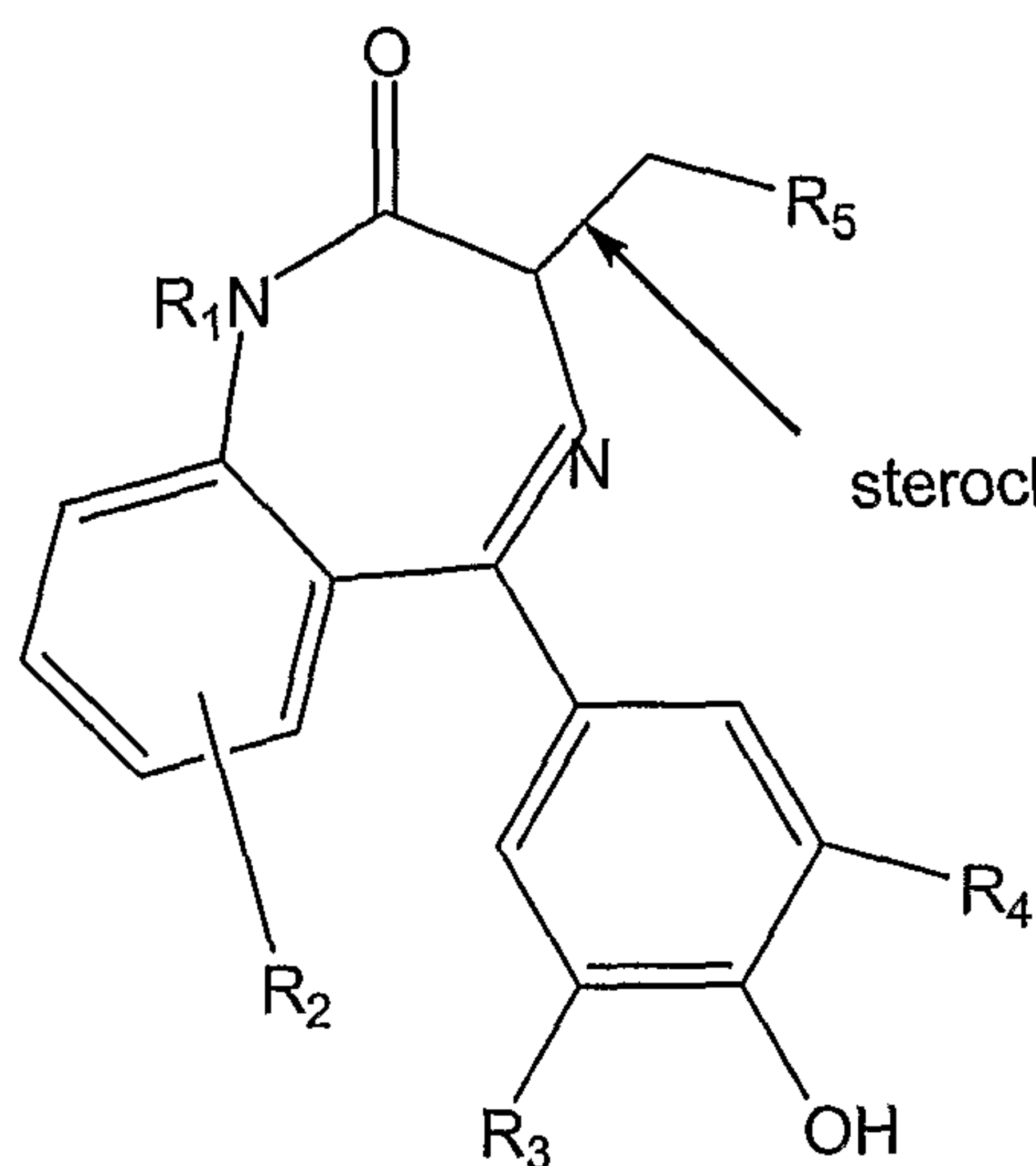
;



Additional exemplary compounds of the present invention are described in U.S.

5 Provisional Patent Nos. U.S. Provisional Patent Nos. 60/131,761, 60/165,511, 60/191,855, 60/312,560, 60/313,689, 60/396,670, 60/565,788, 60/607,599, 60/641,040, and U.S. Patent Application Nos. 11/324,419, 11/176,719, 11/110,228, 10/935,333, 10/886,450, 10/795,535, 10/634,114, 10/427,211, 10/427,212, 10/217,878, 09/767,283, 09/700,101, and related applications; each herein incorporated by reference in their entireties.

10 In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



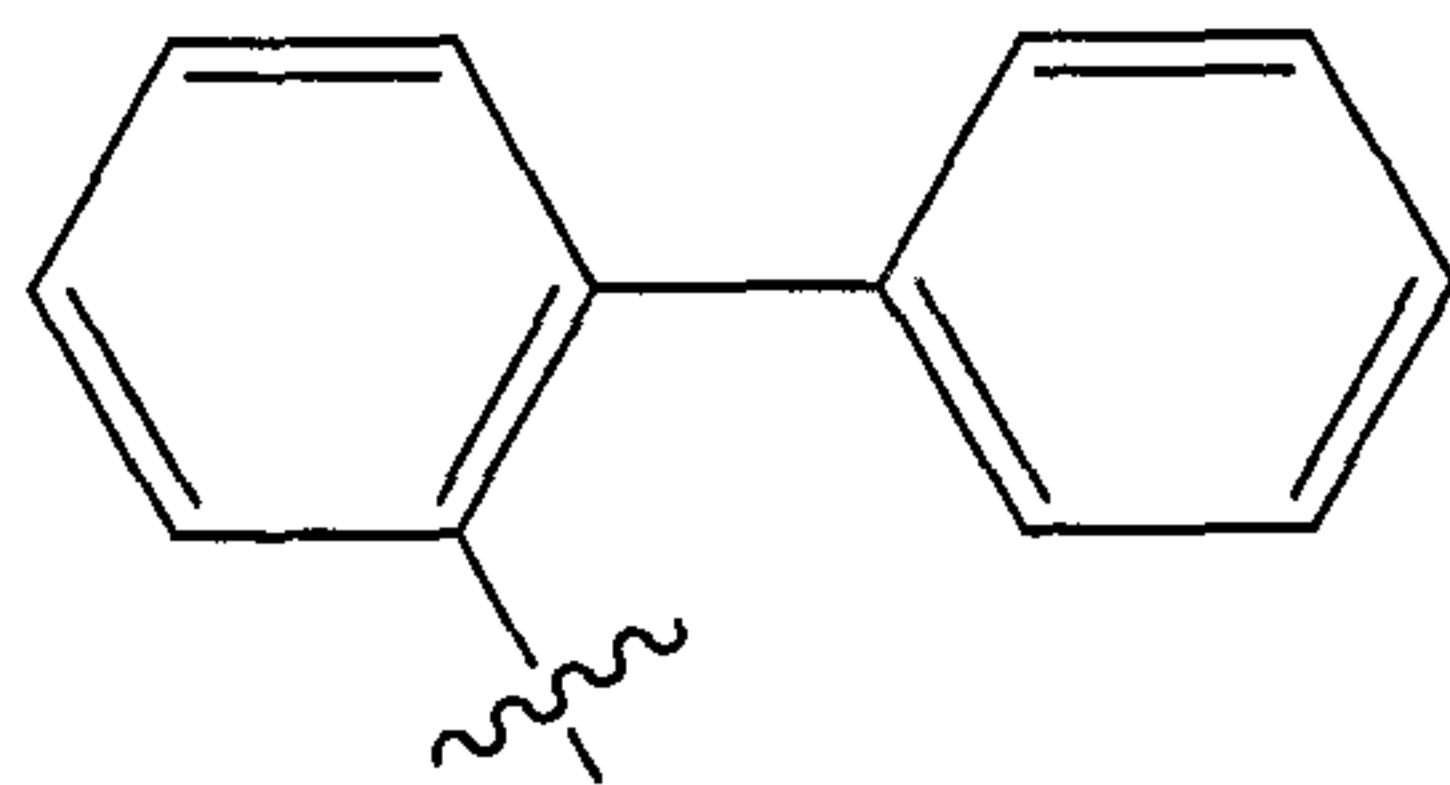
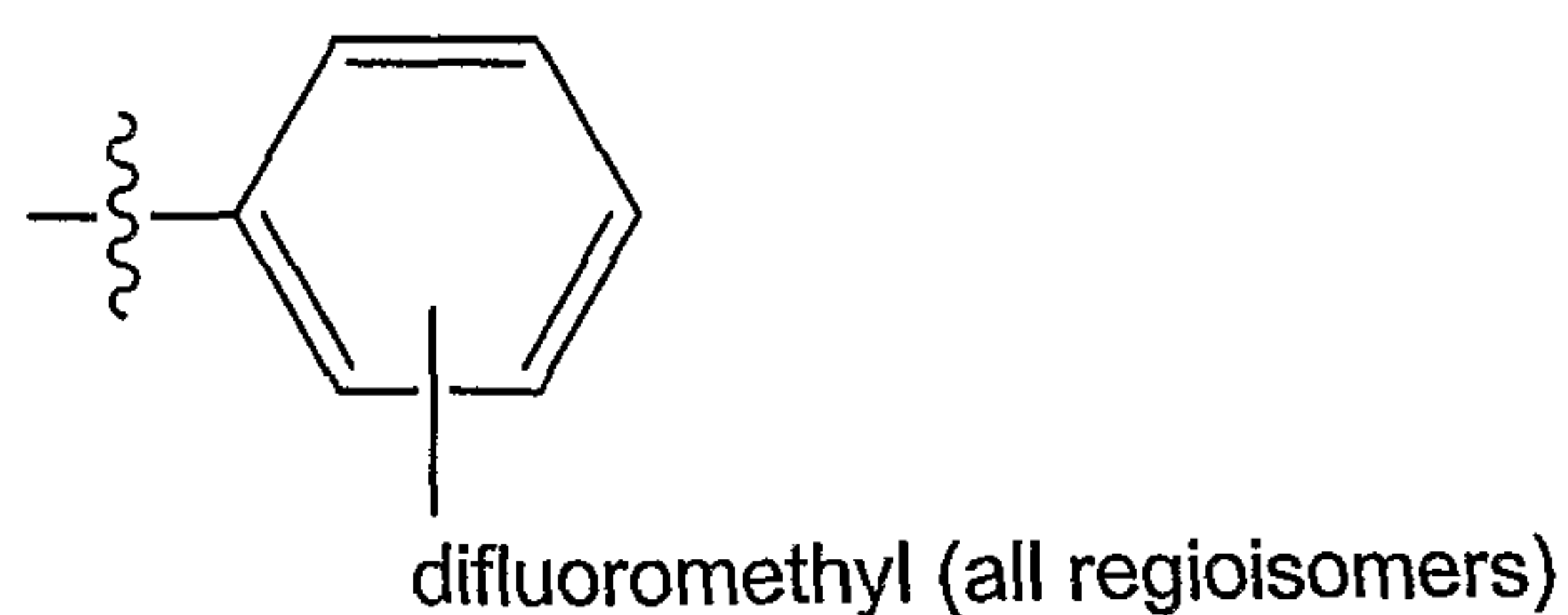
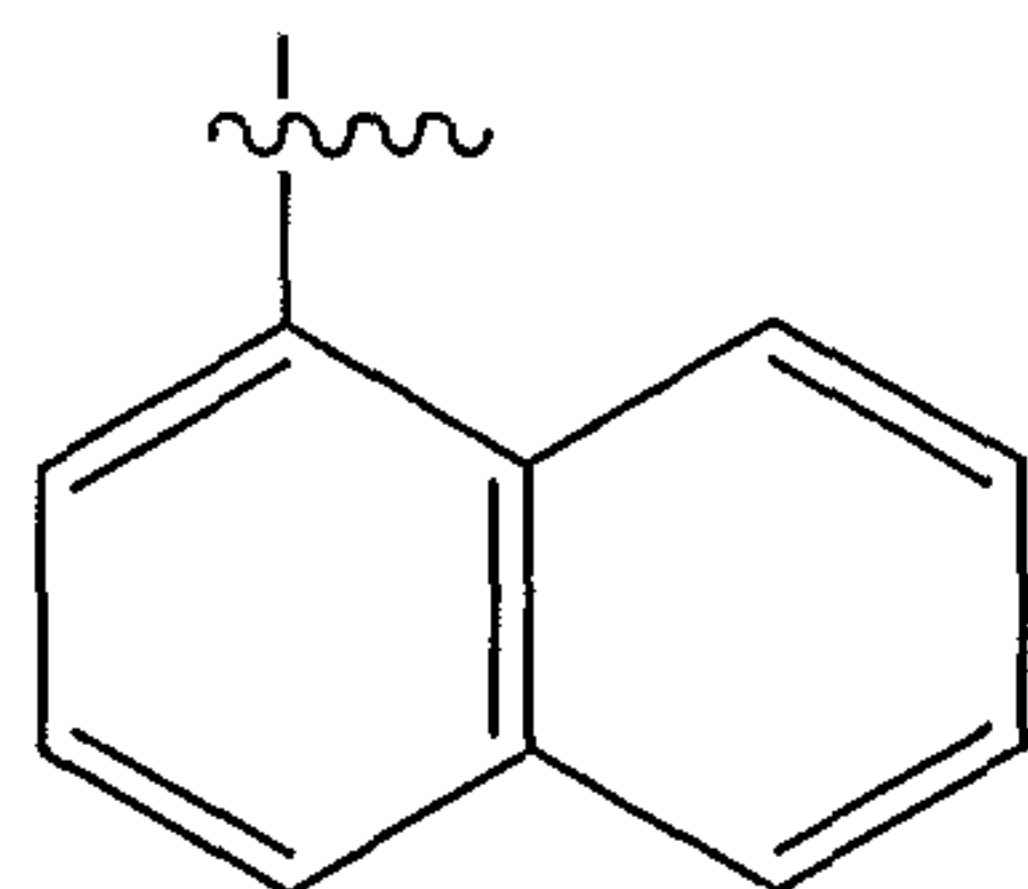
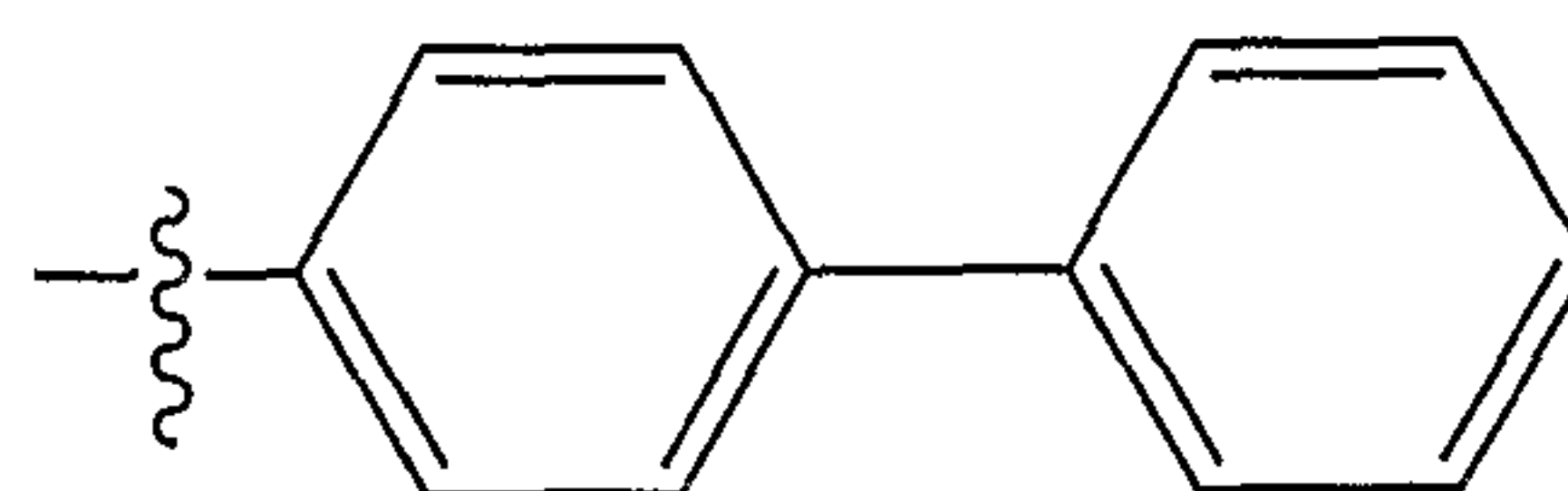
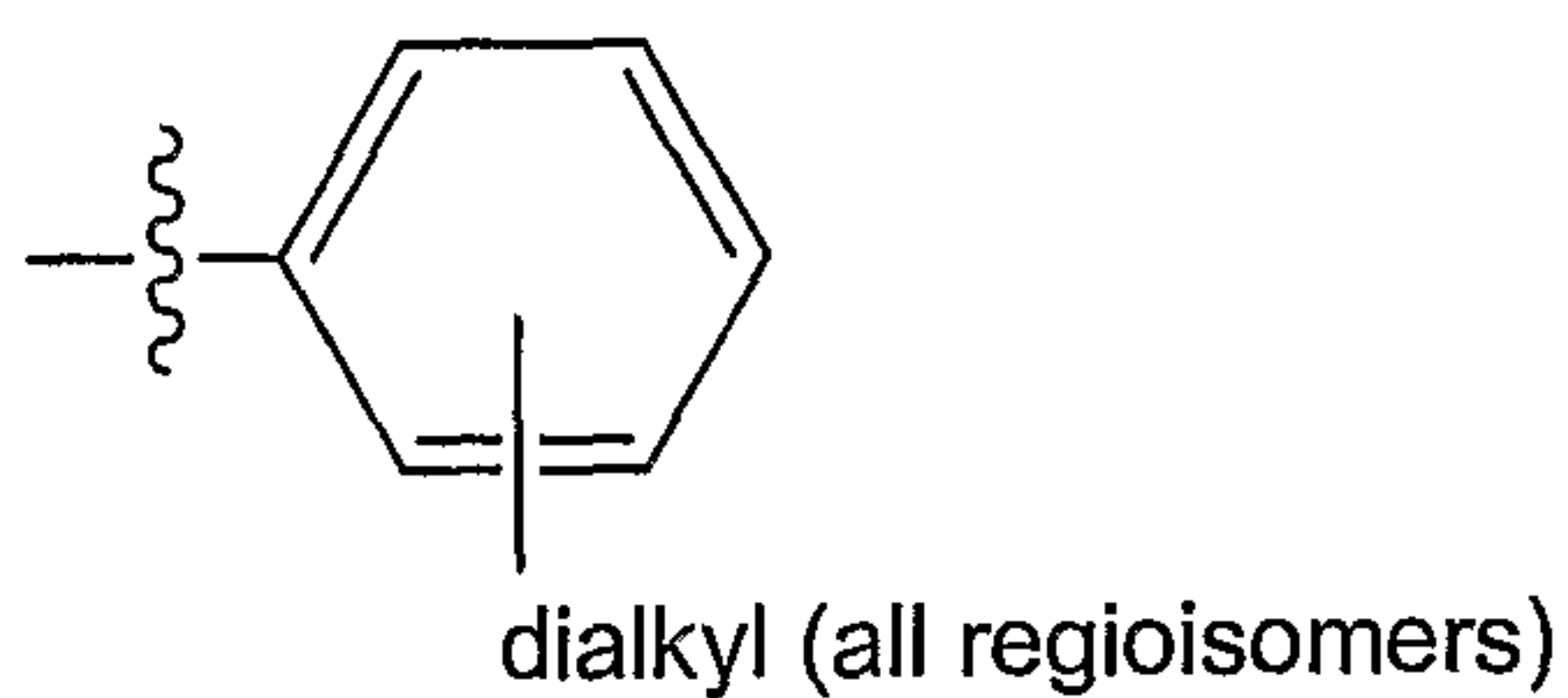
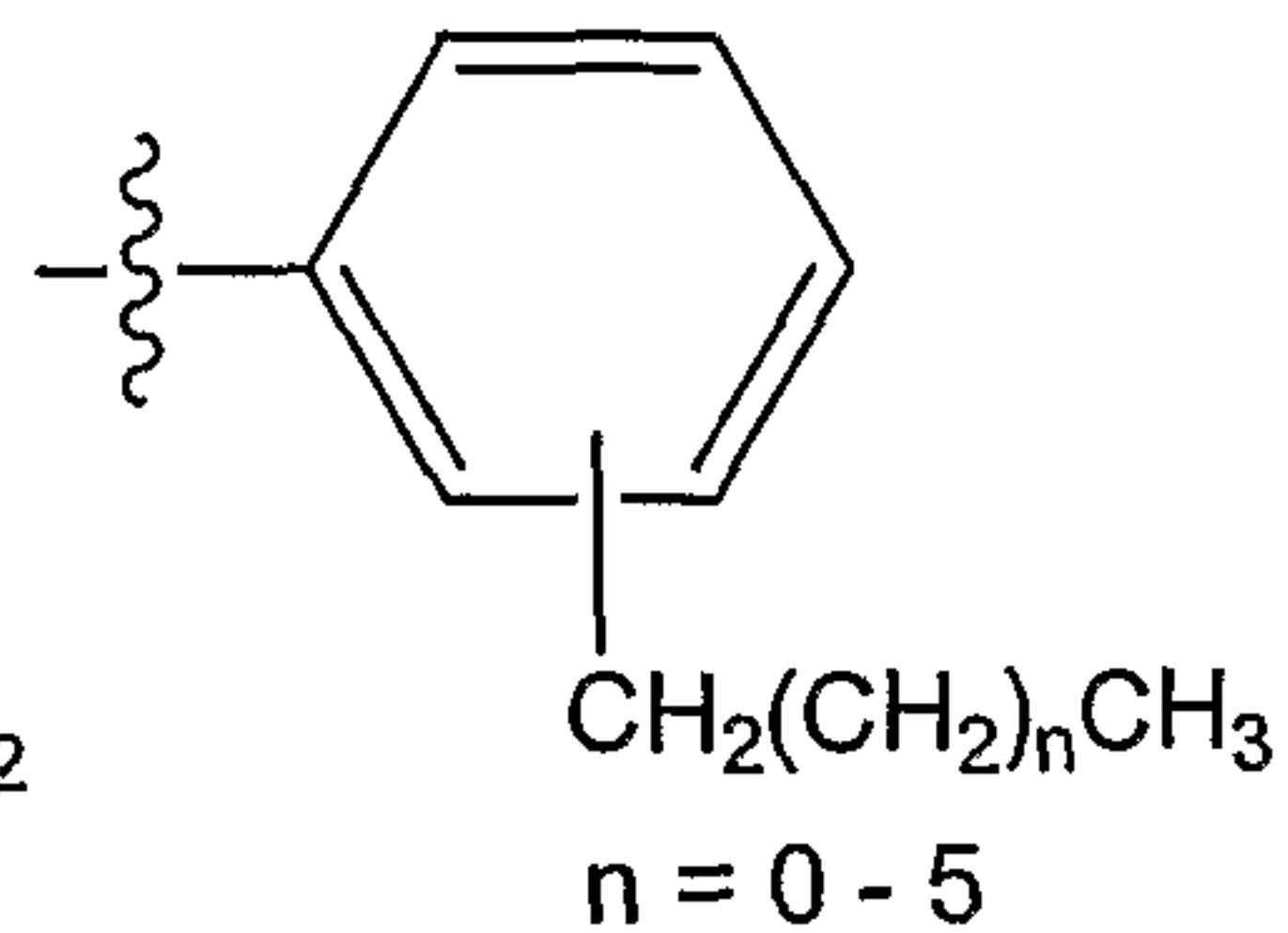
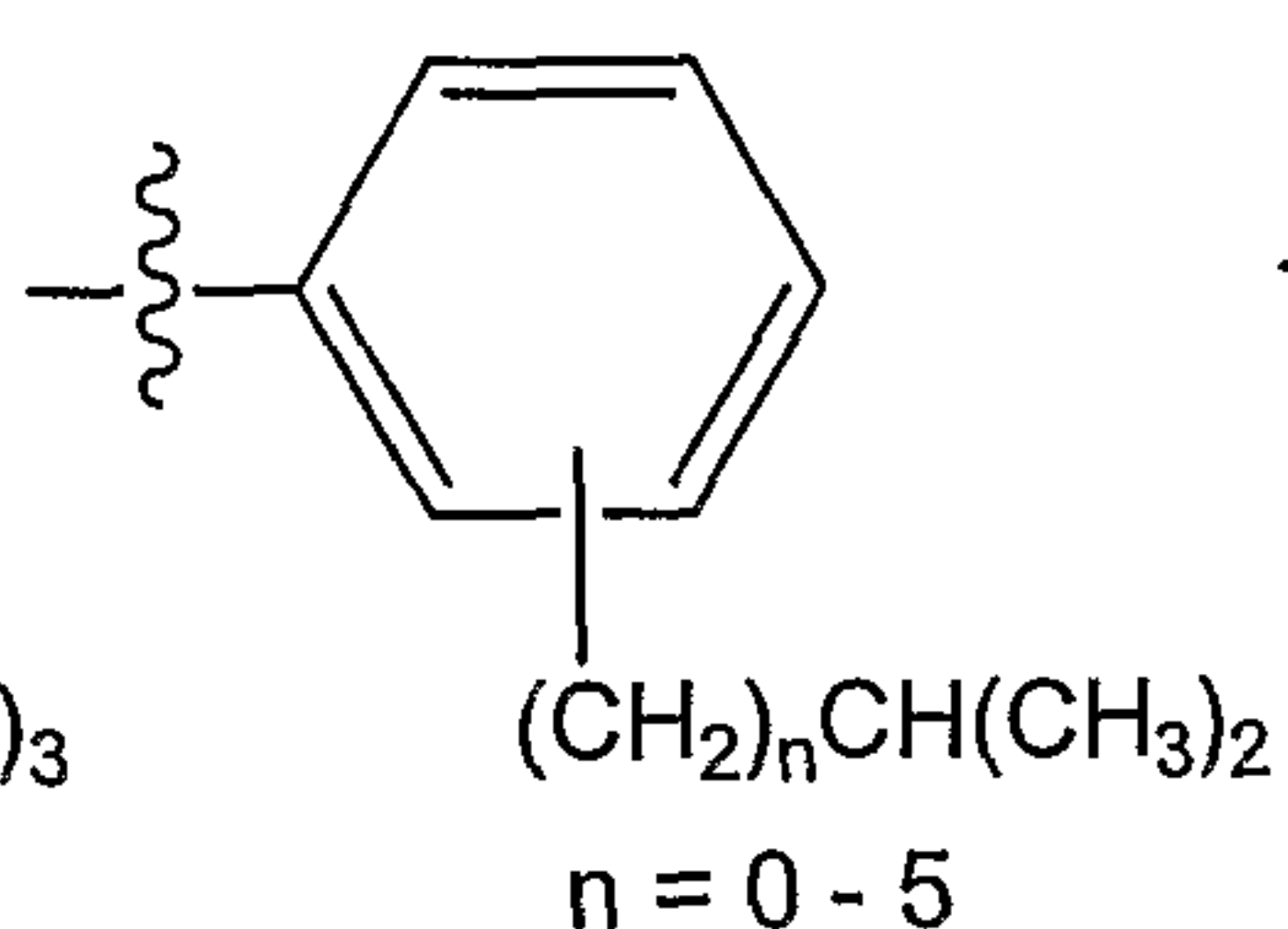
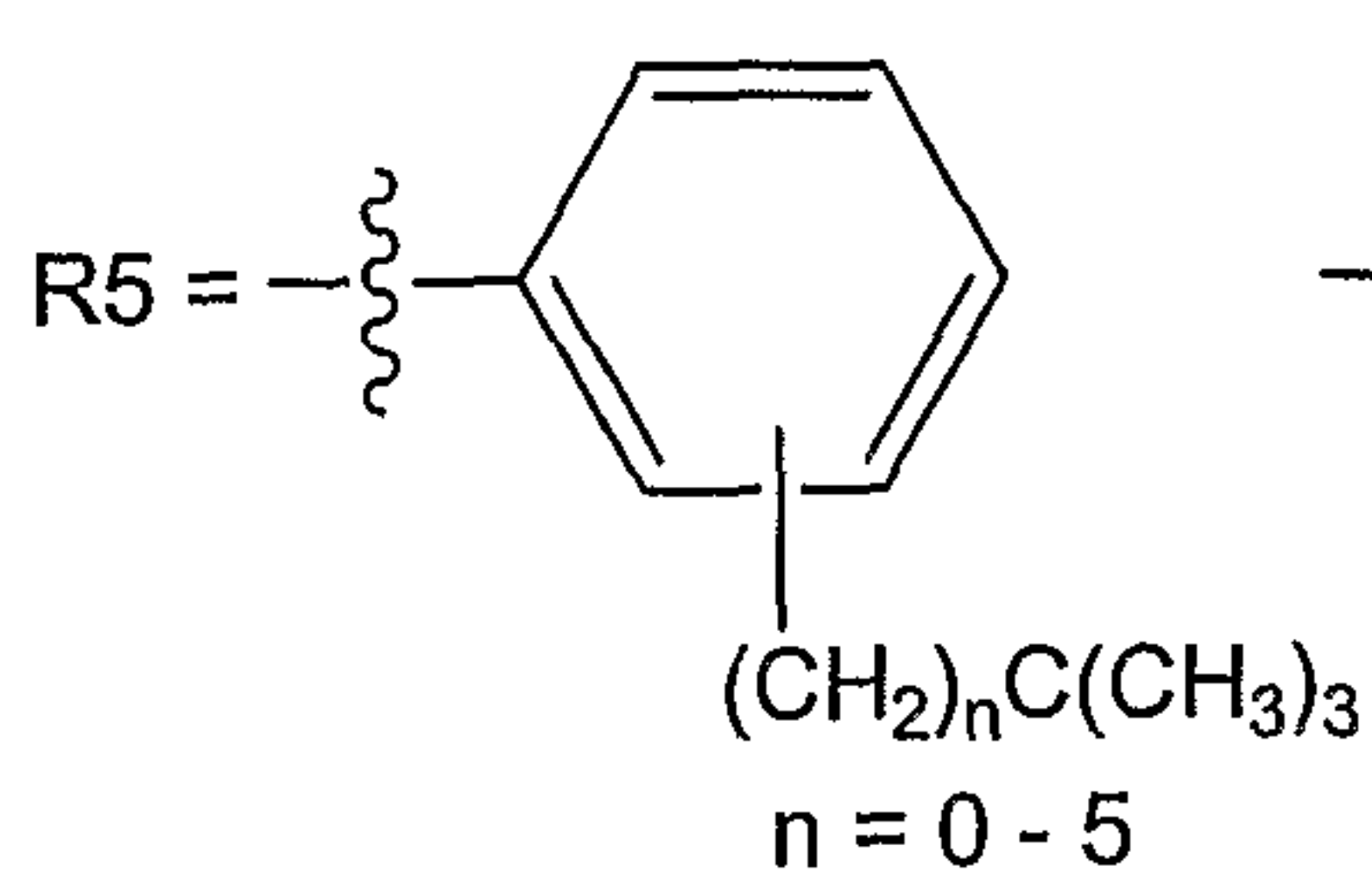
R1 = H, alkyl, or substituted alkyl

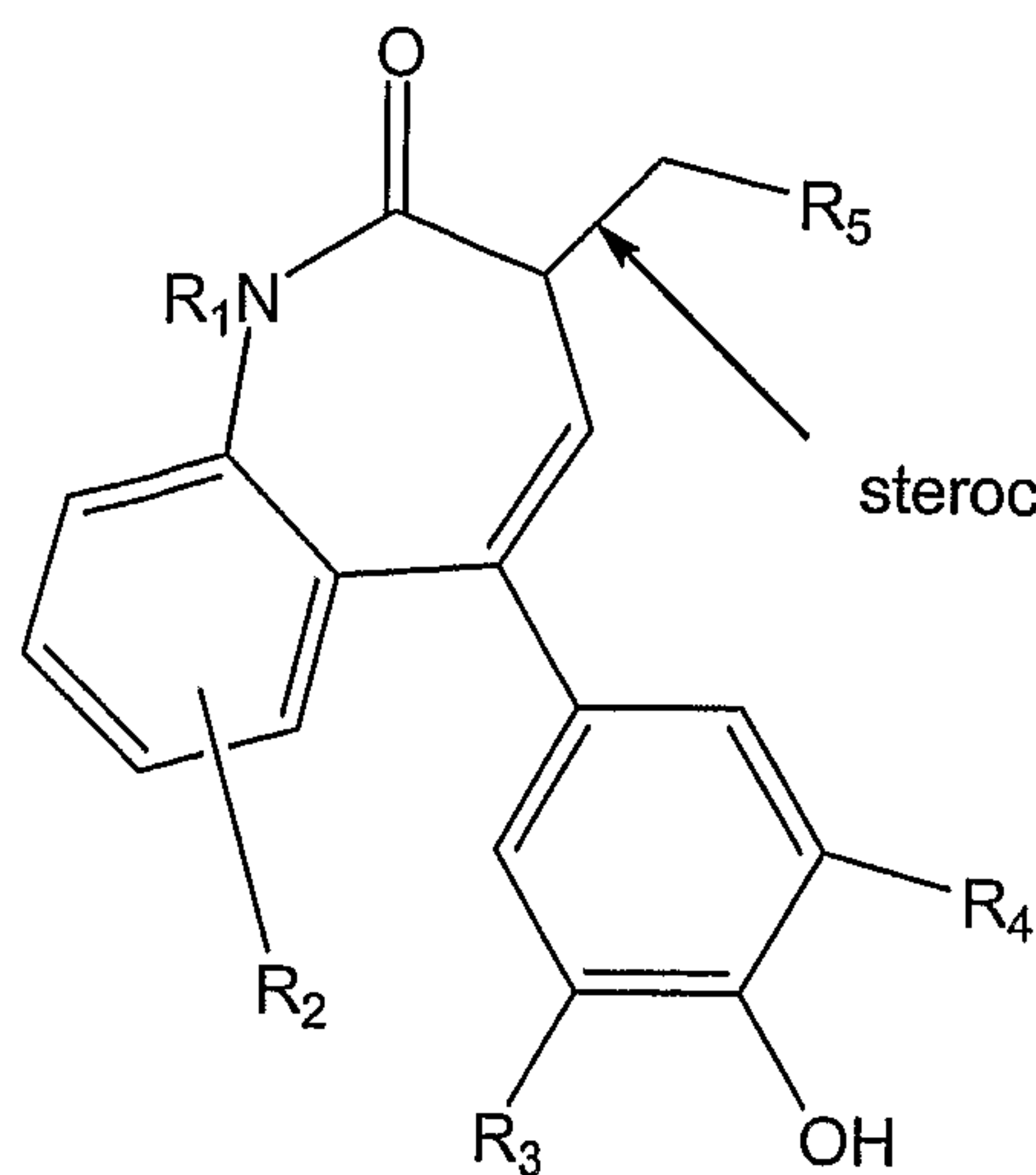
R3 = H, alkyl, or substituted alkyl

R4 = H, alkyl, or substituted alkyl

stereochemistry is R, S, or racemic

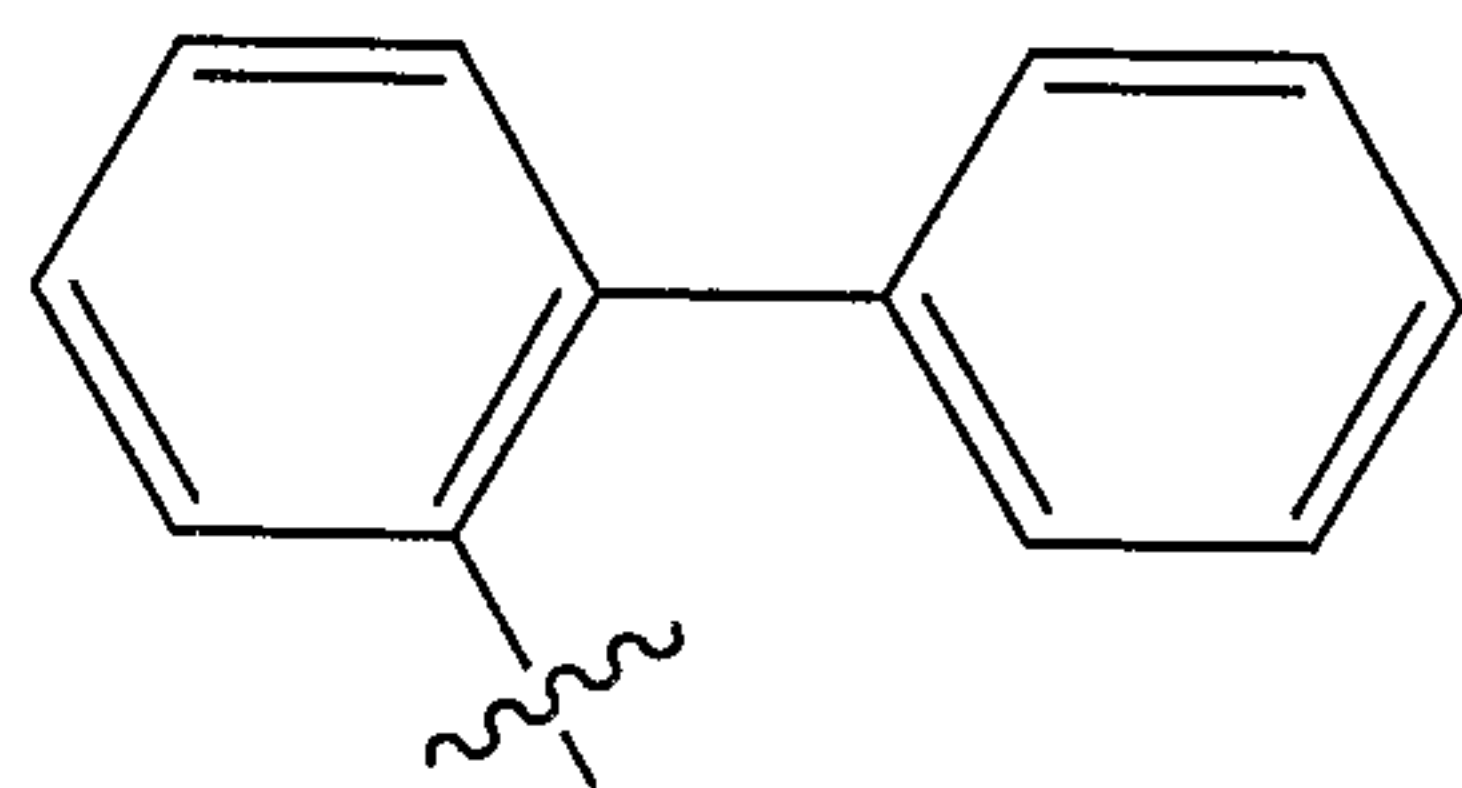
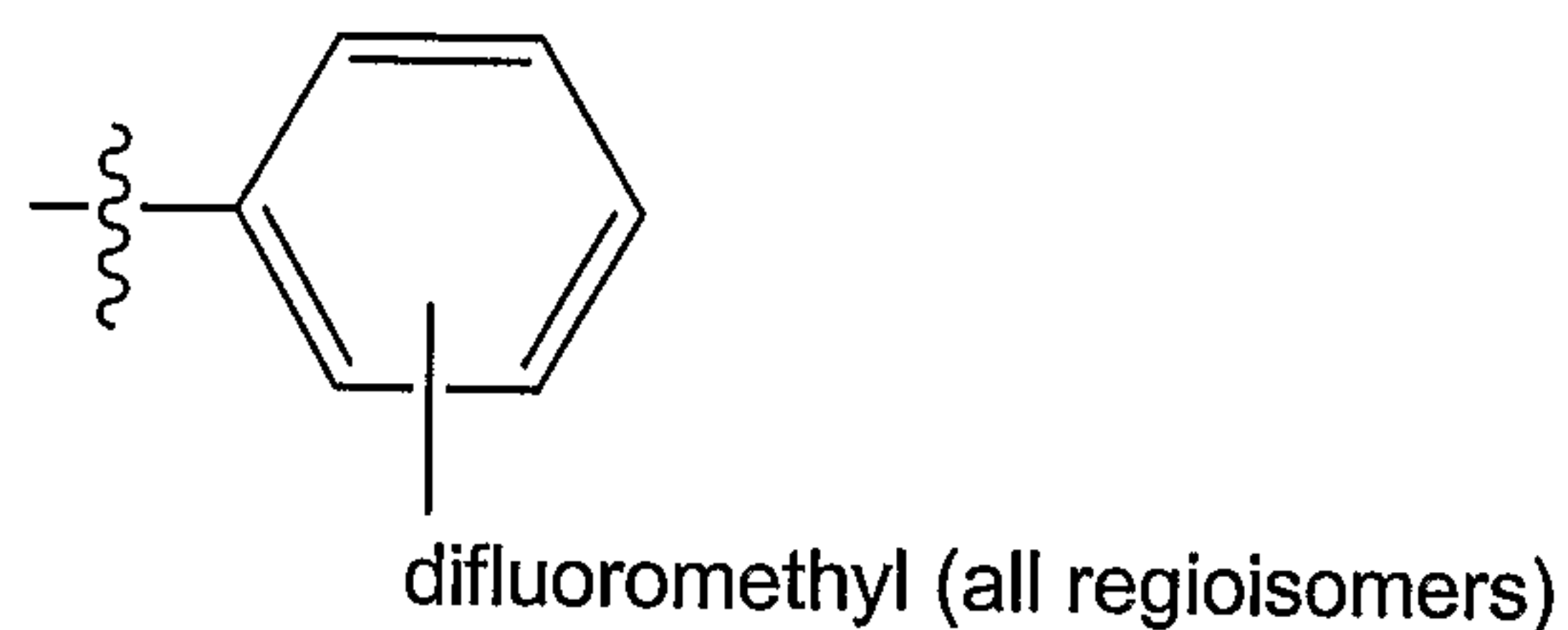
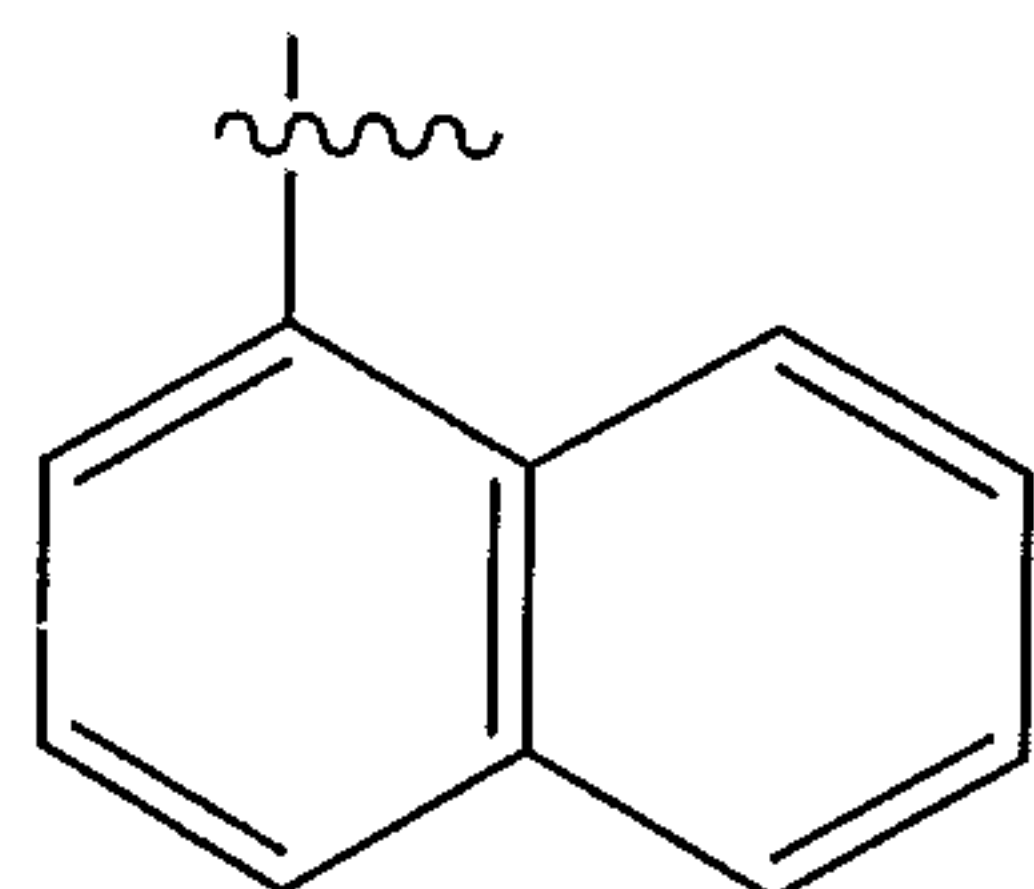
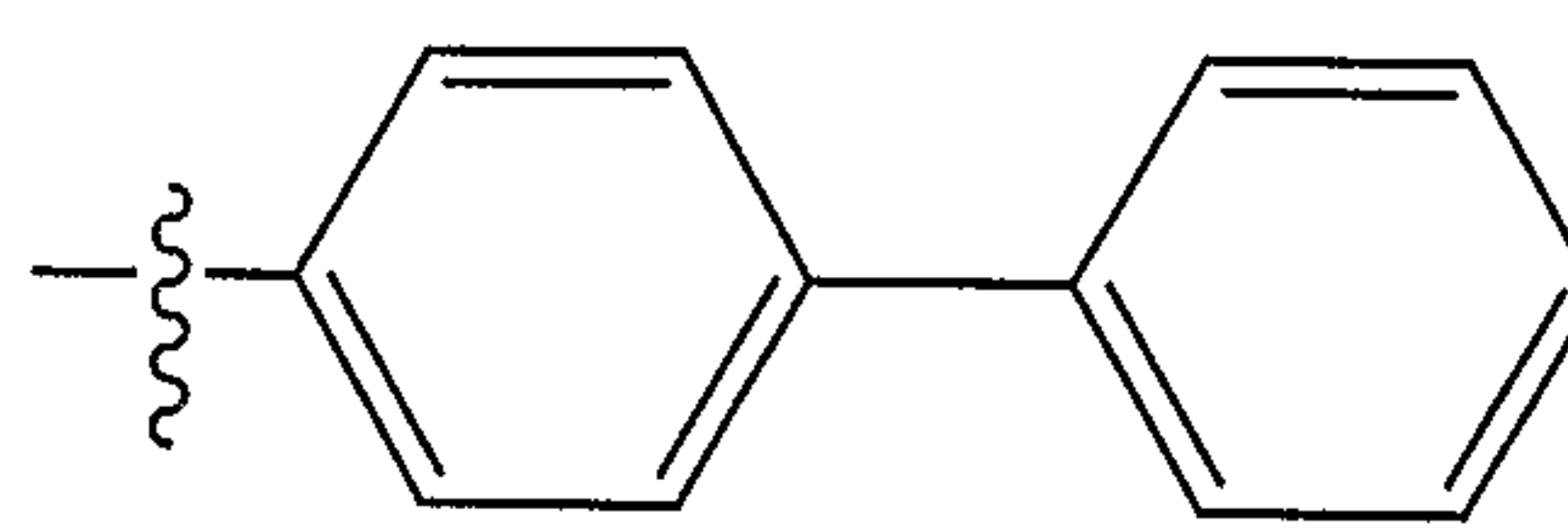
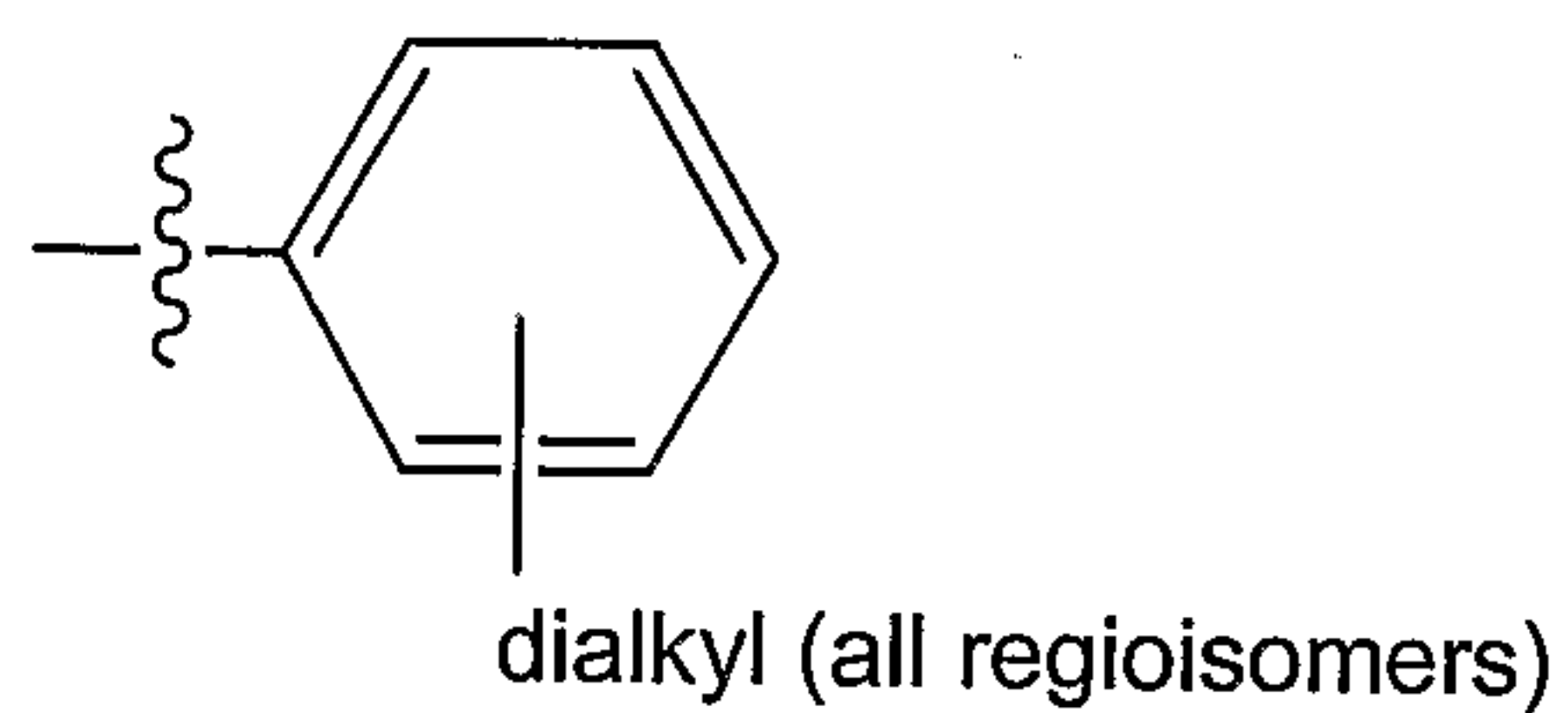
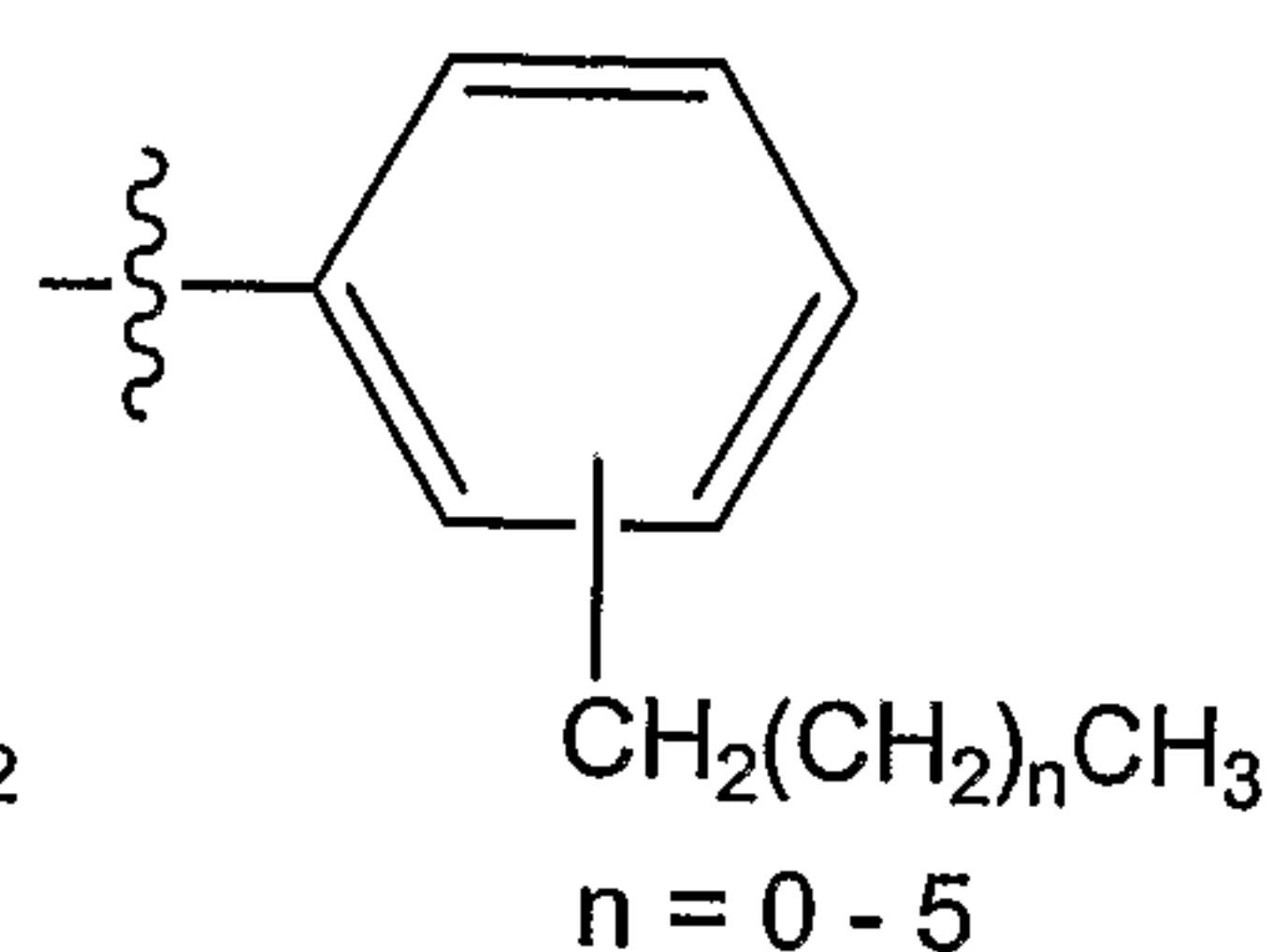
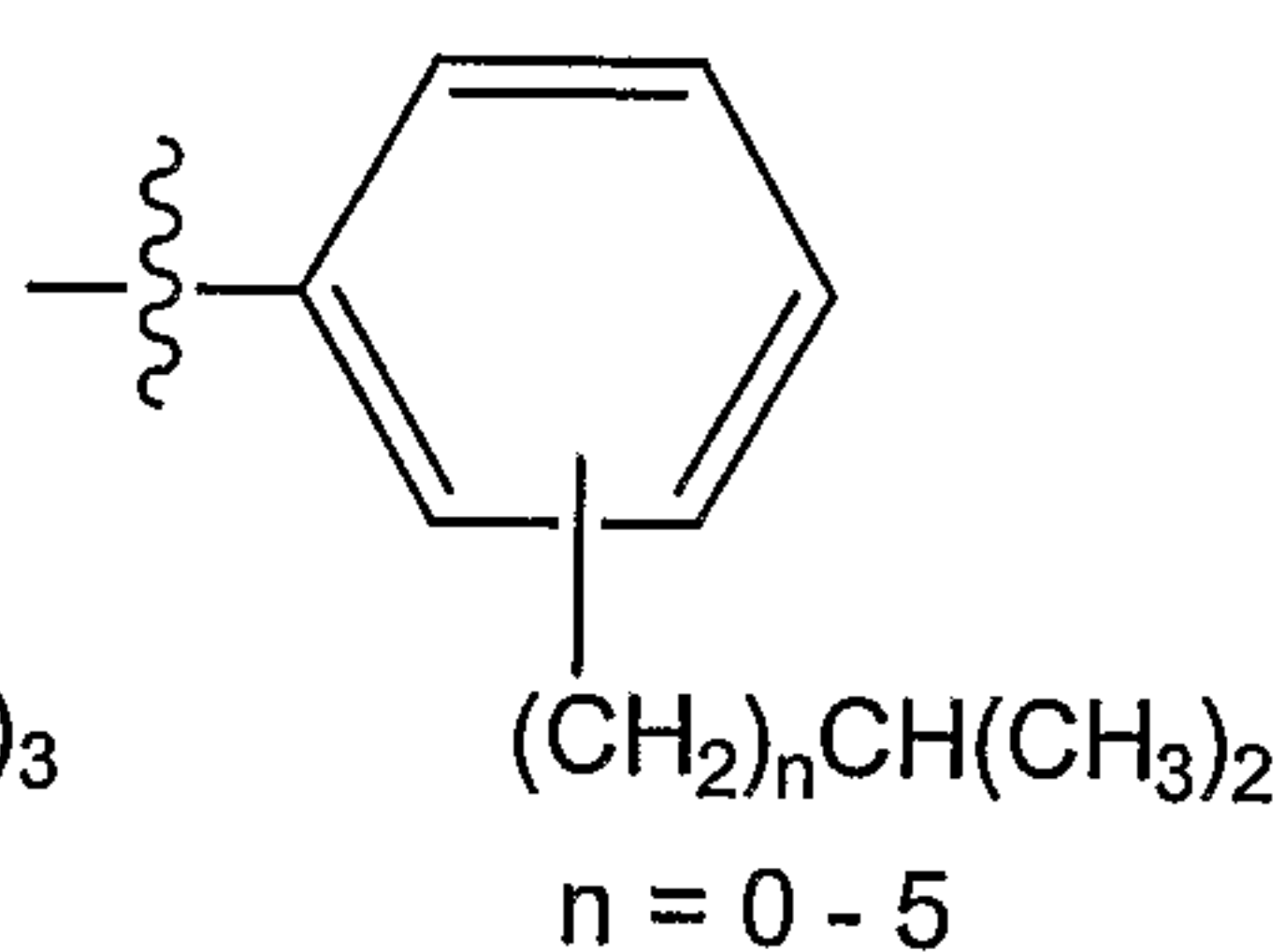
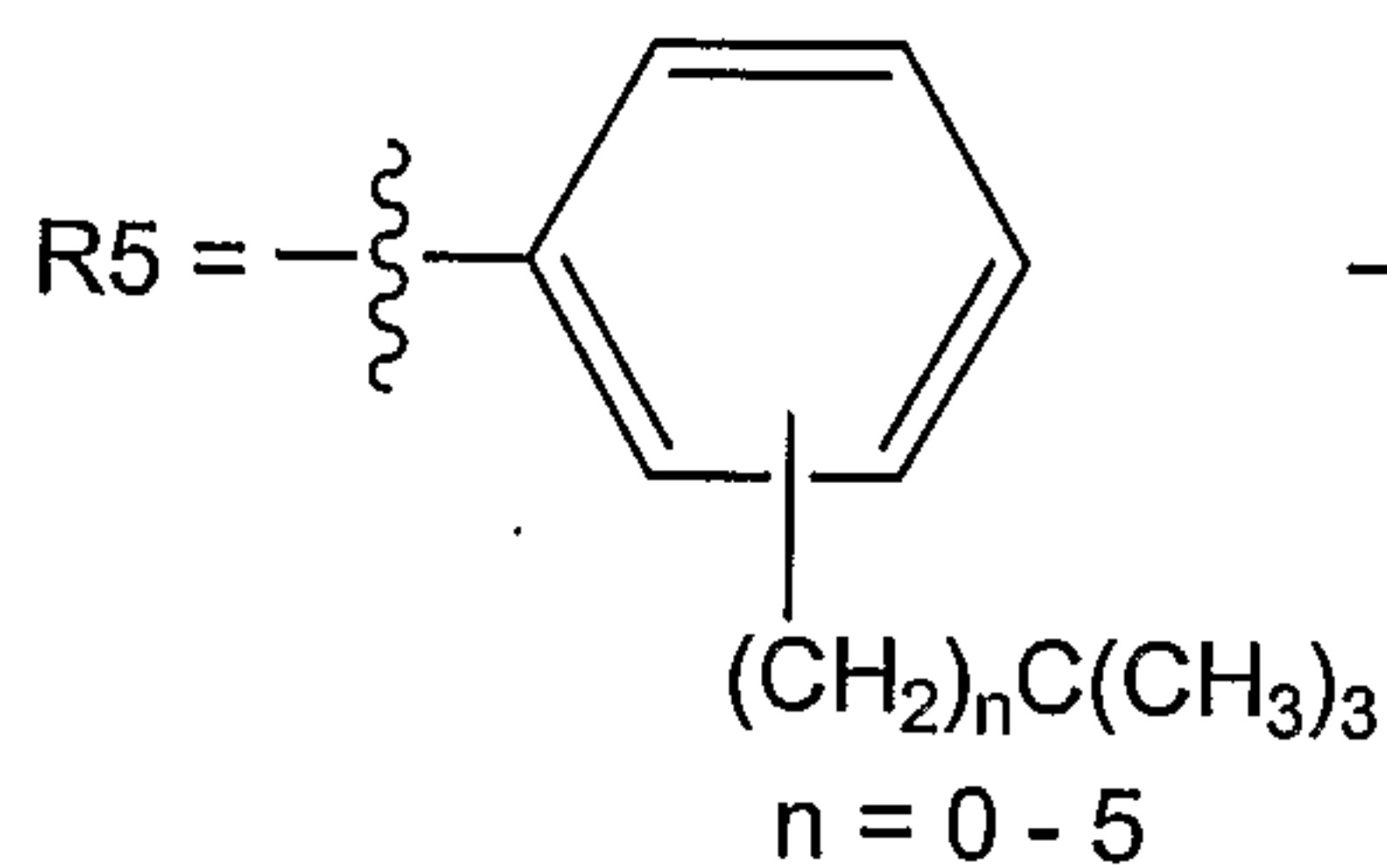
R2 is selected from hydrogen, a hydroxy, an alkoxy, a halo, an amino, a lower-alkyl-a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic

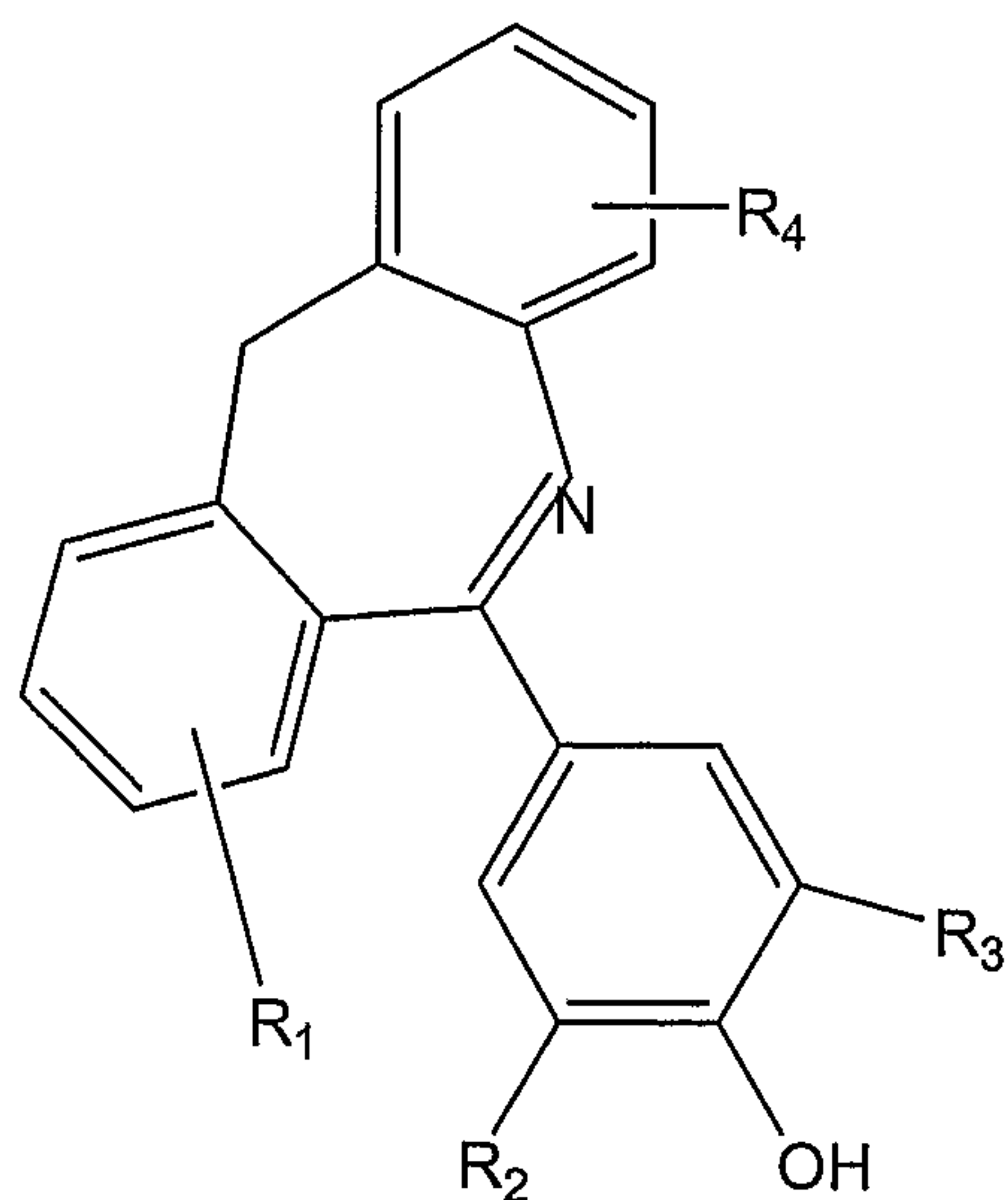


R₁ = H, alkyl, or substituted alkylR₃ = H, alkyl, or substituted alkylR₄ = H, alkyl, or substituted alkyl

stereochemistry is R, S, or racemic

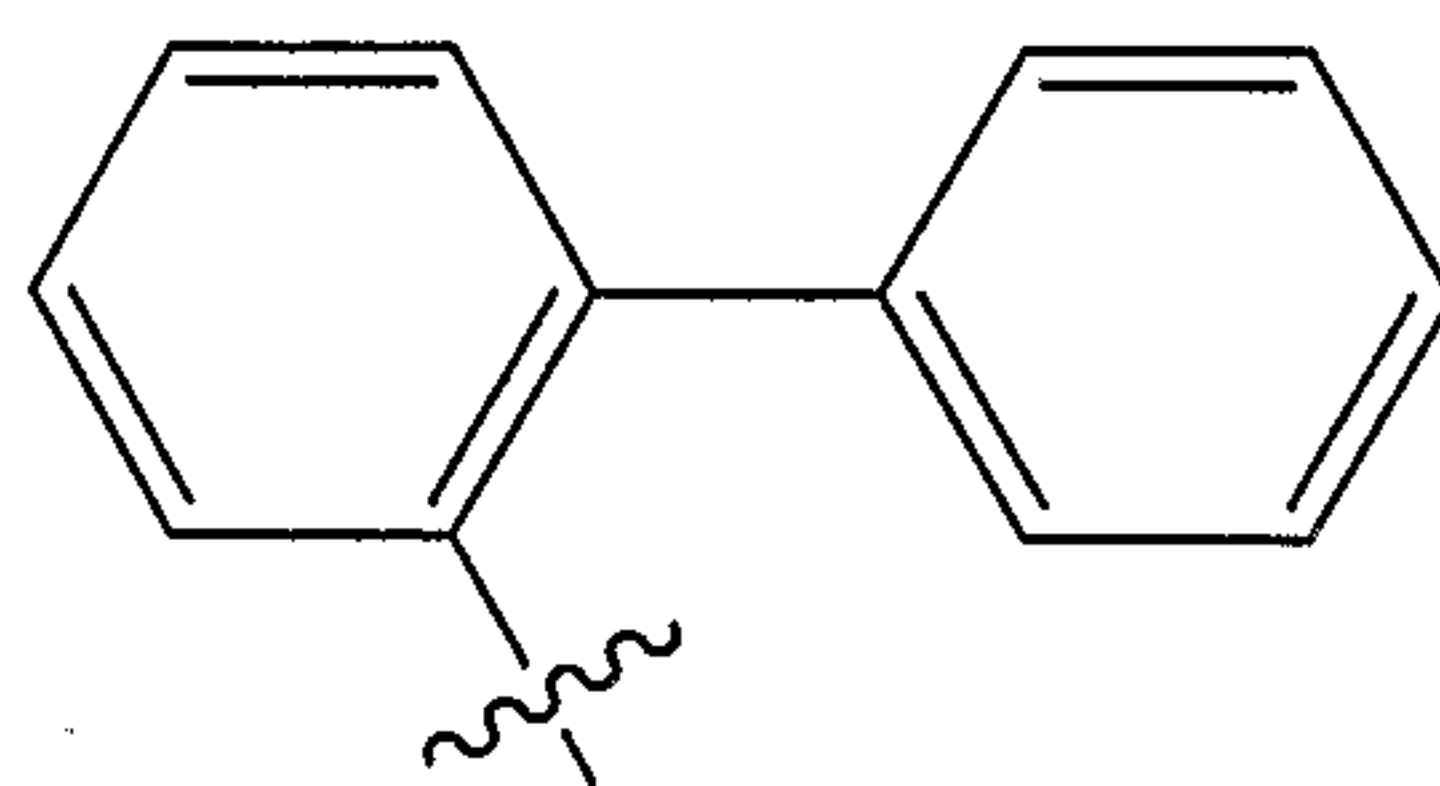
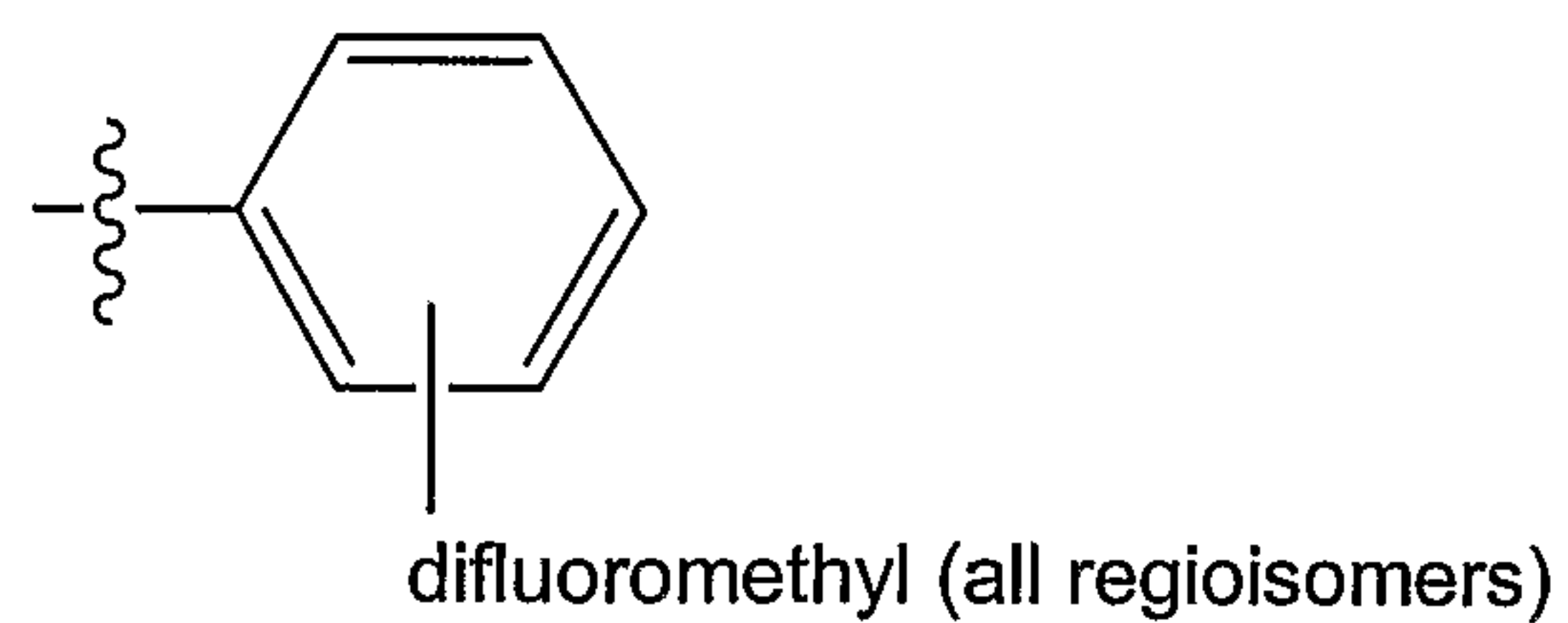
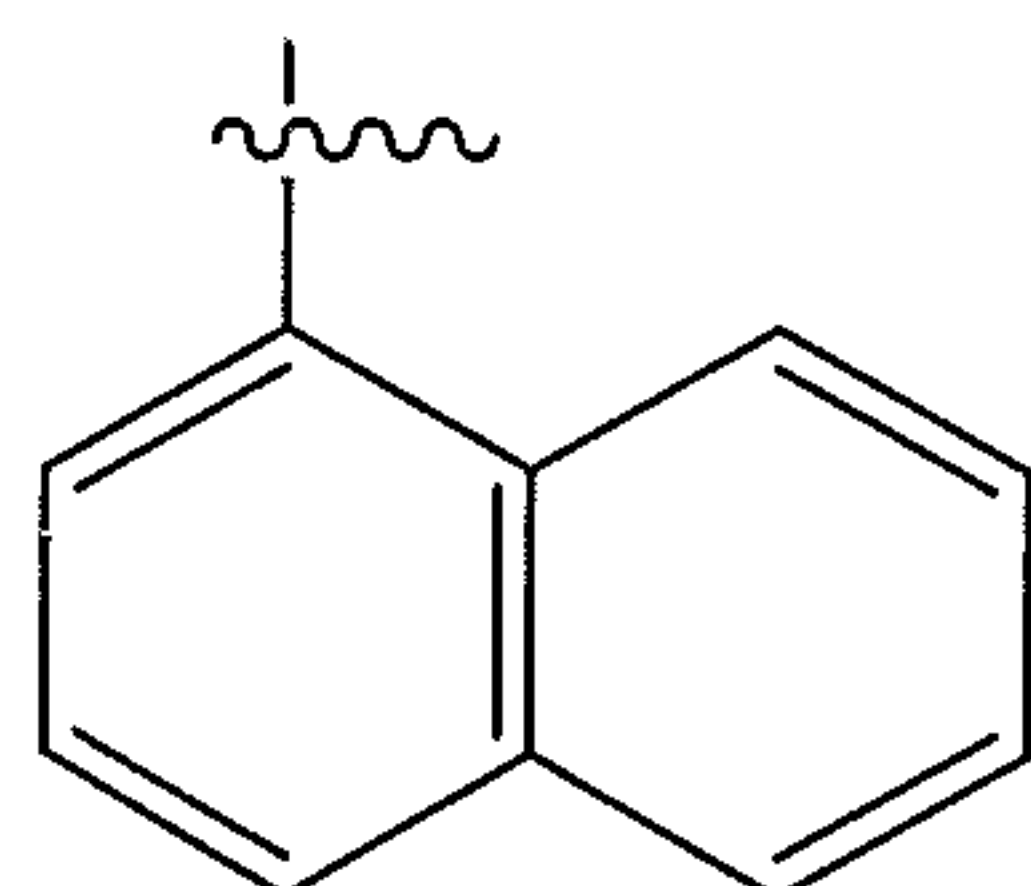
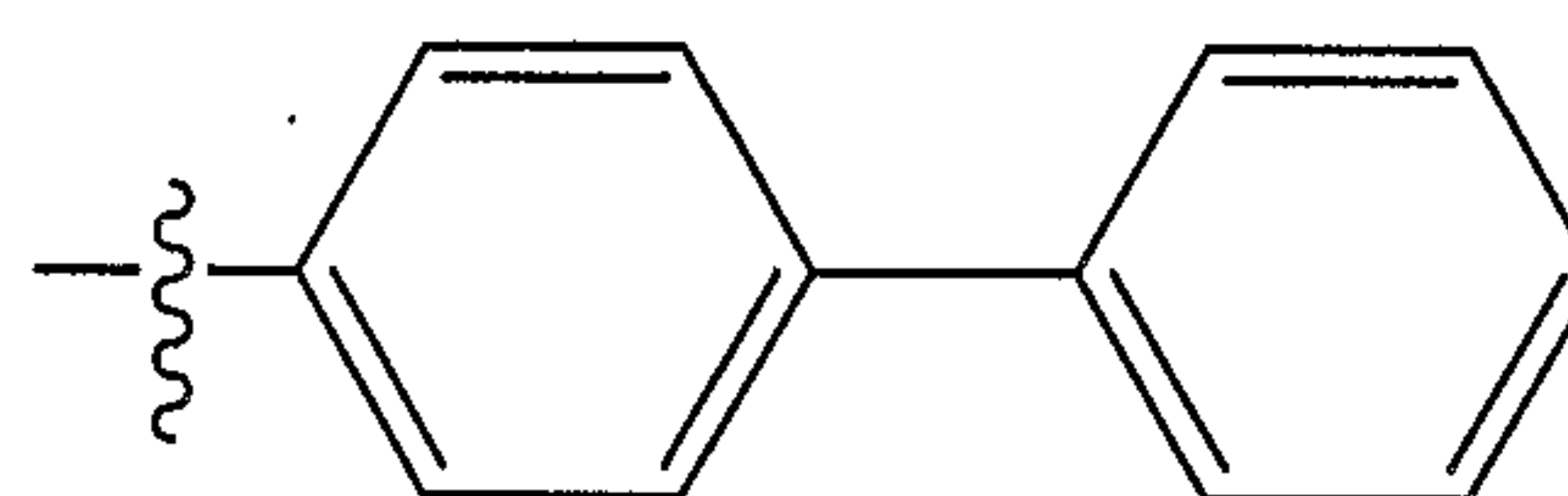
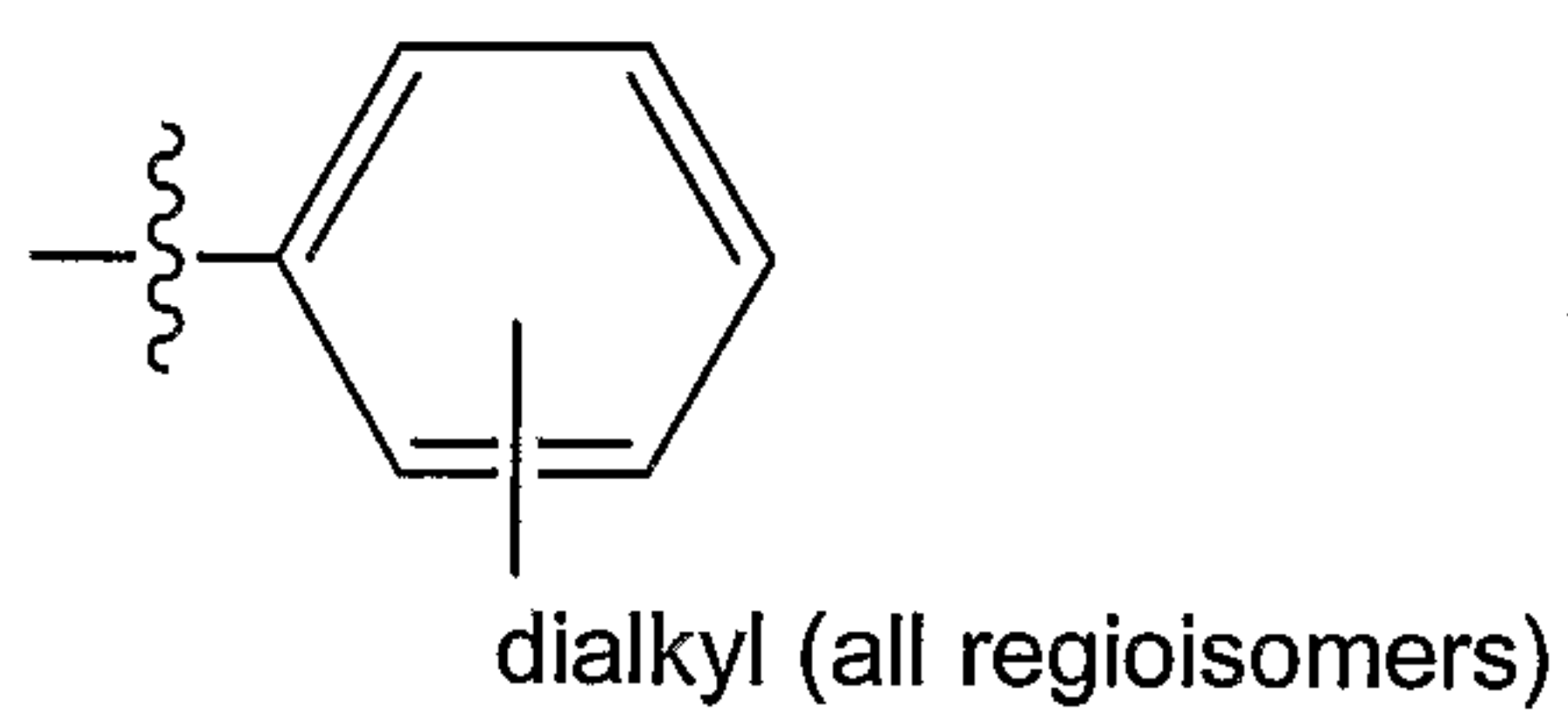
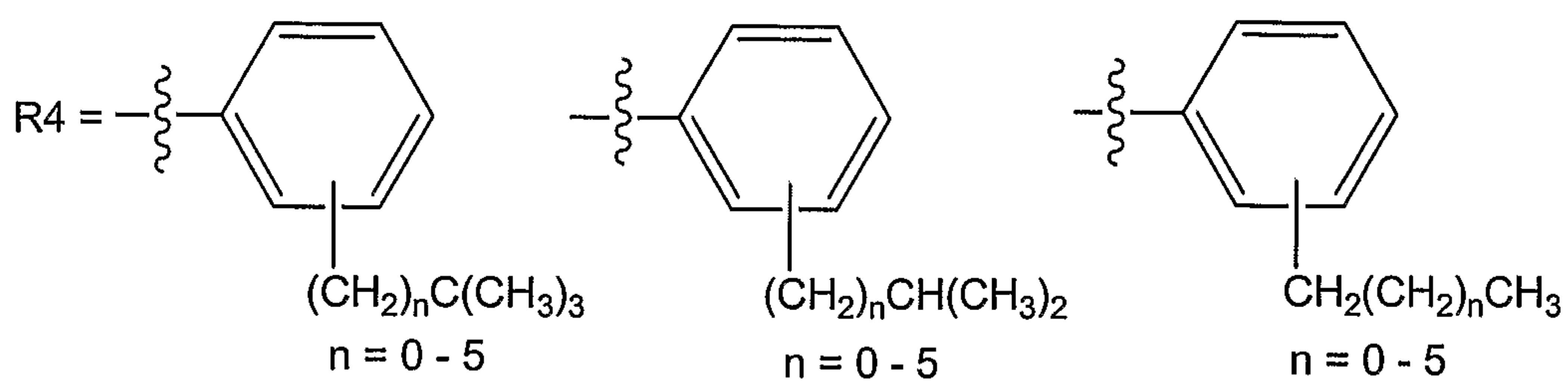
R₂ is selected from hydrogen, a hydroxy, an alkoxy, a halo, an amino, a lower-alkyl-a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic

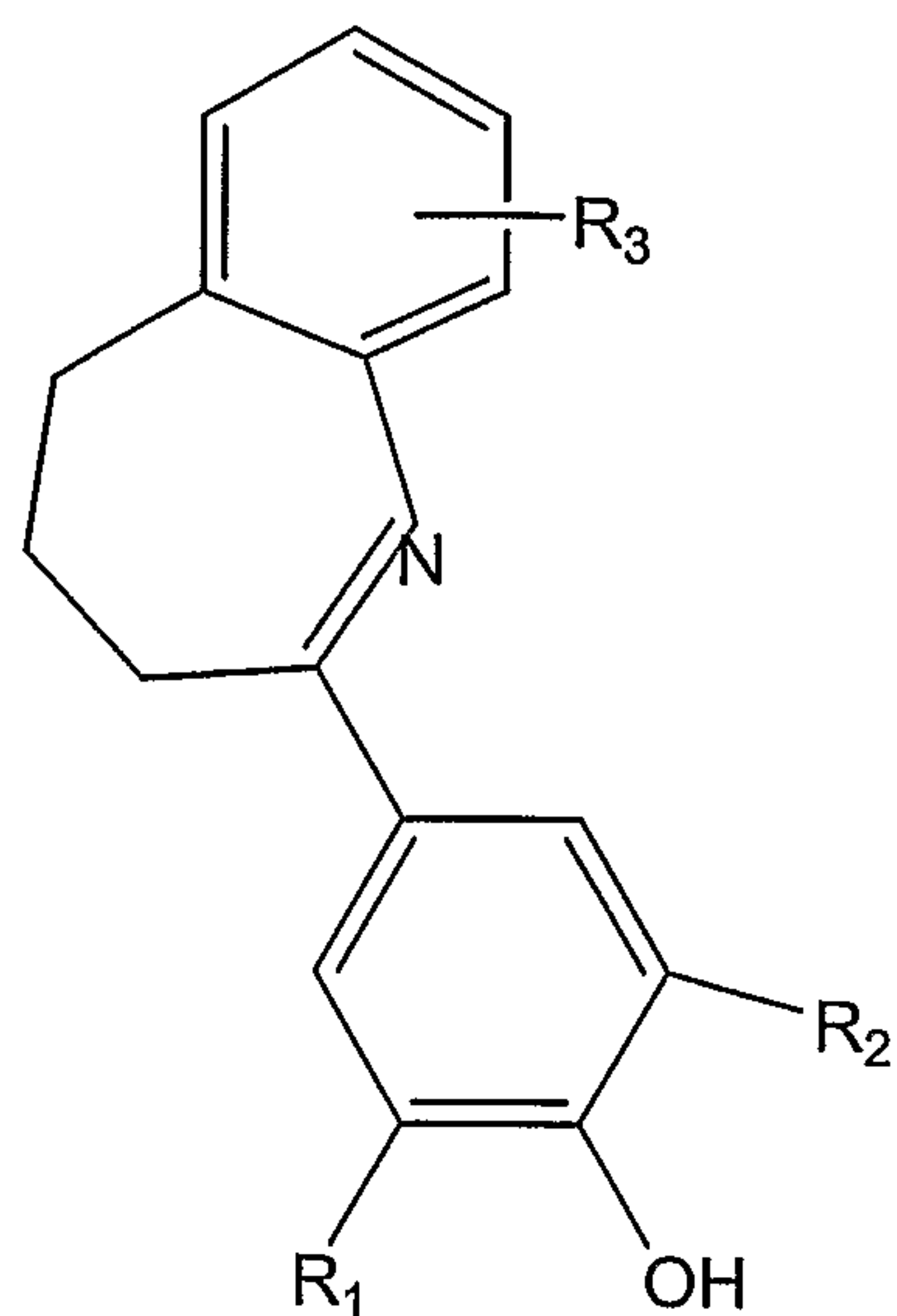




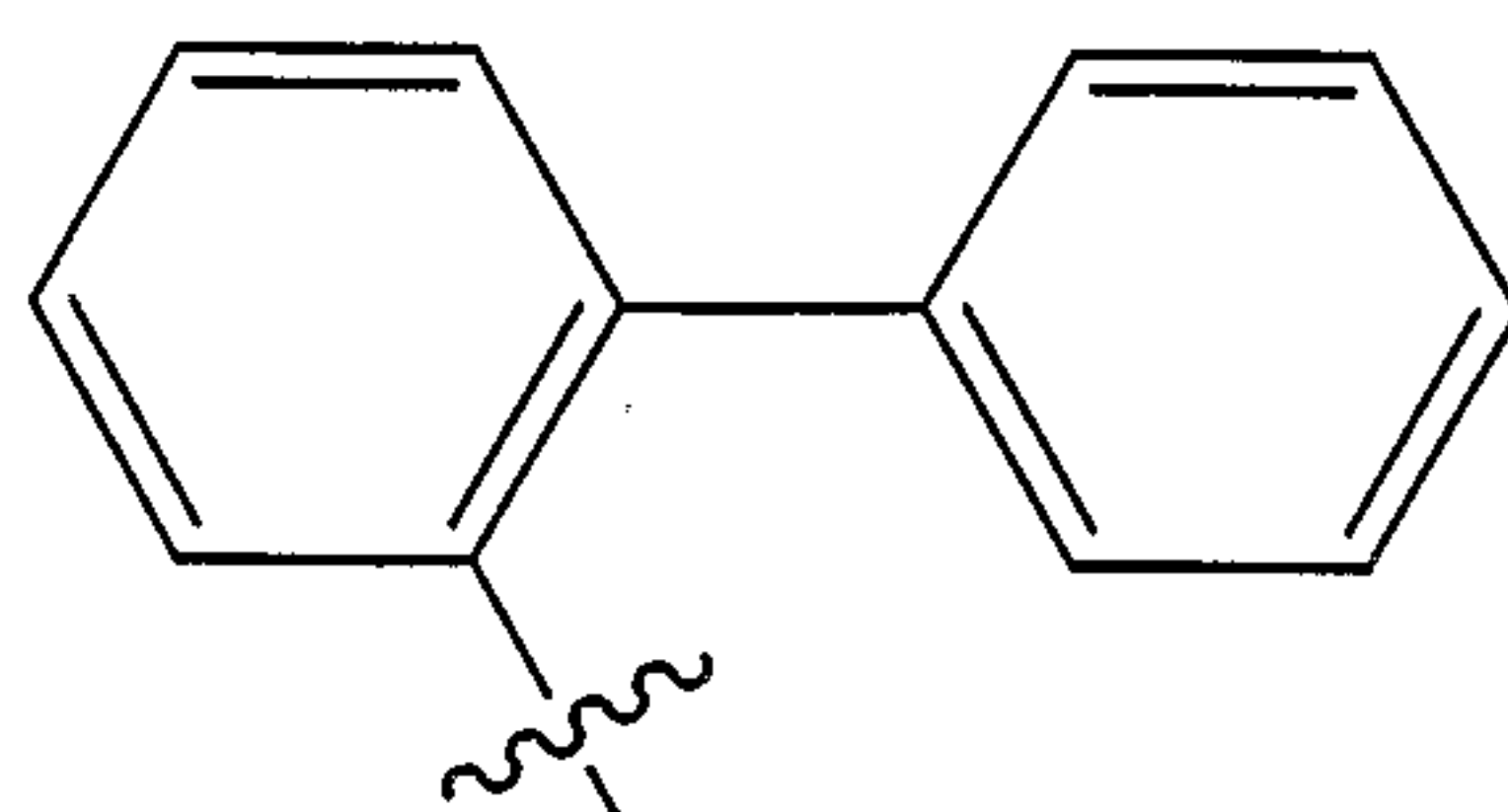
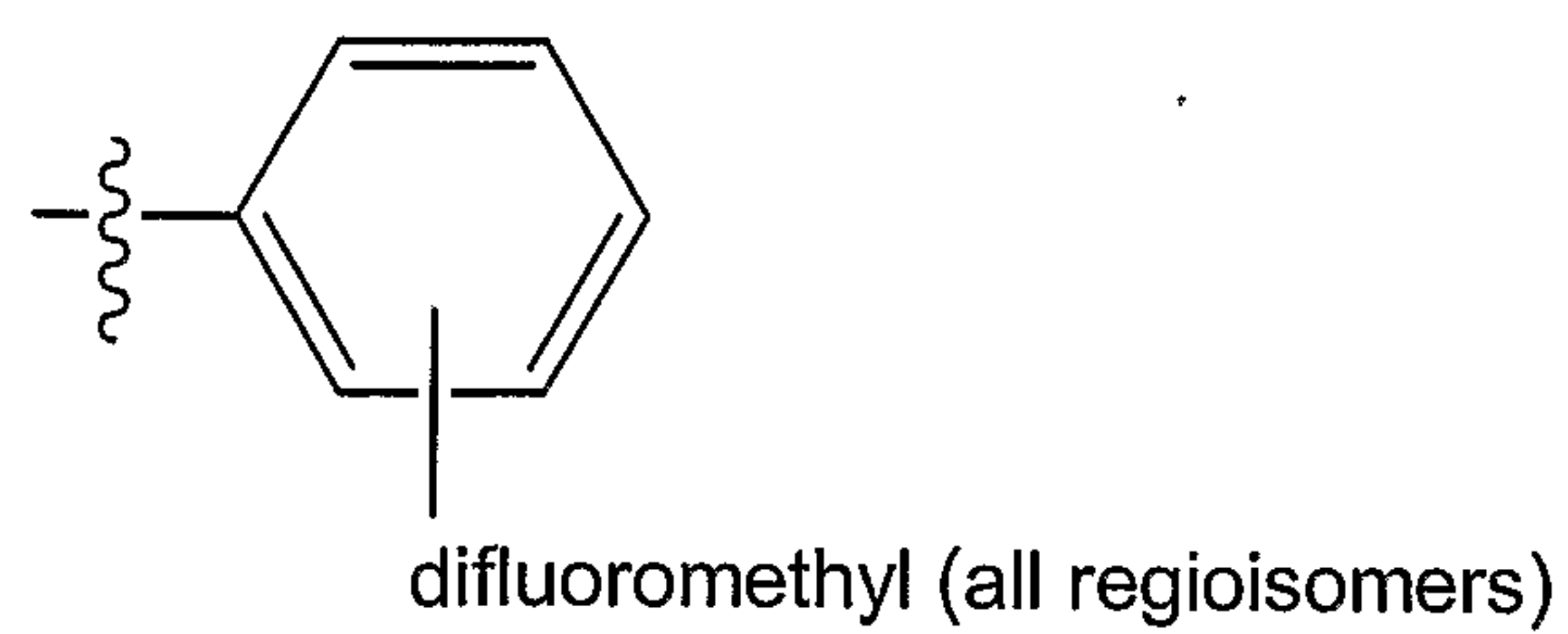
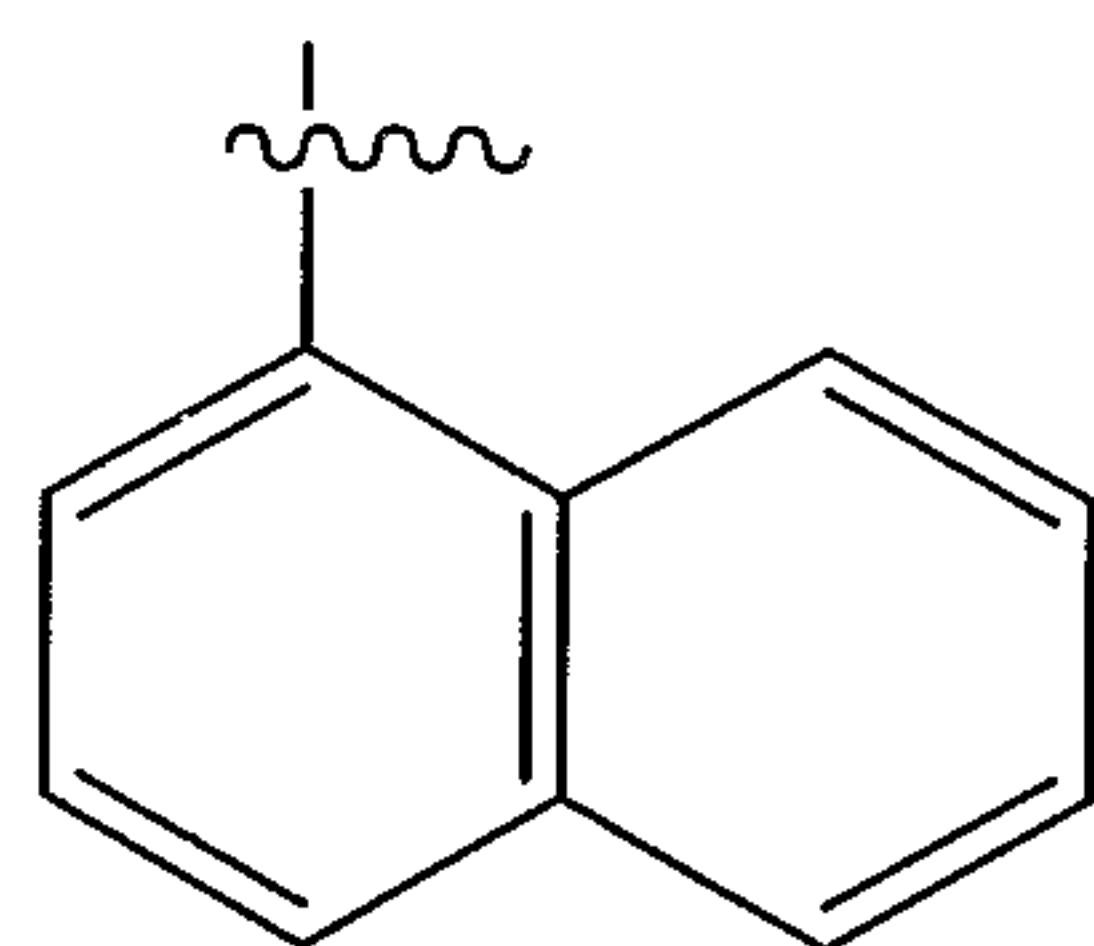
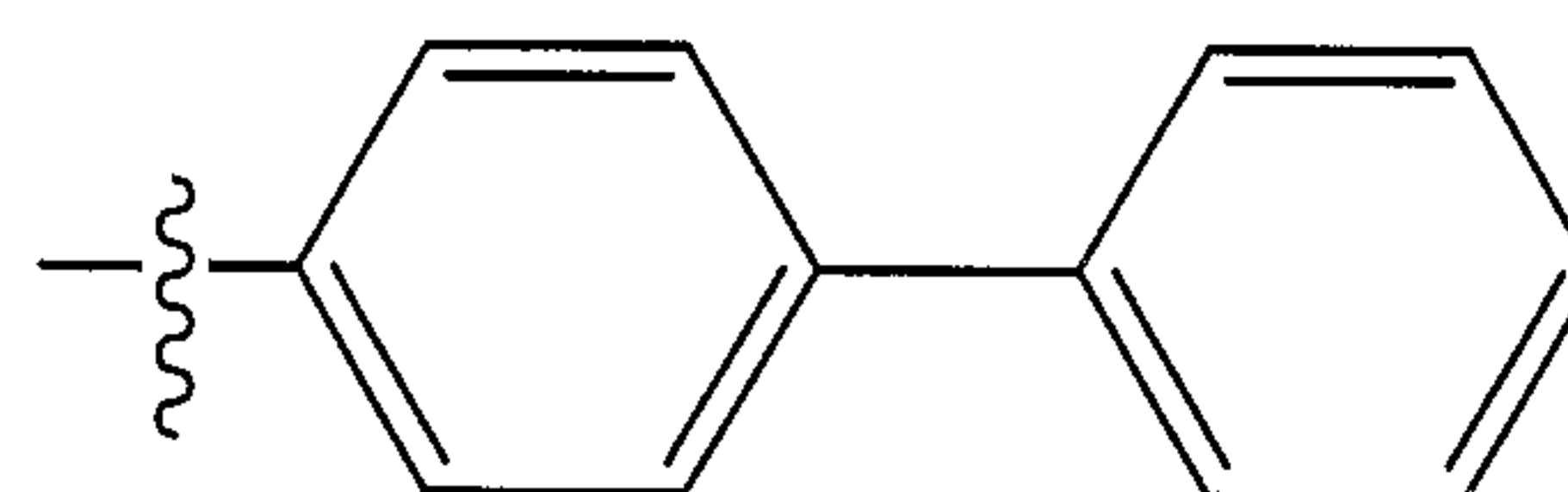
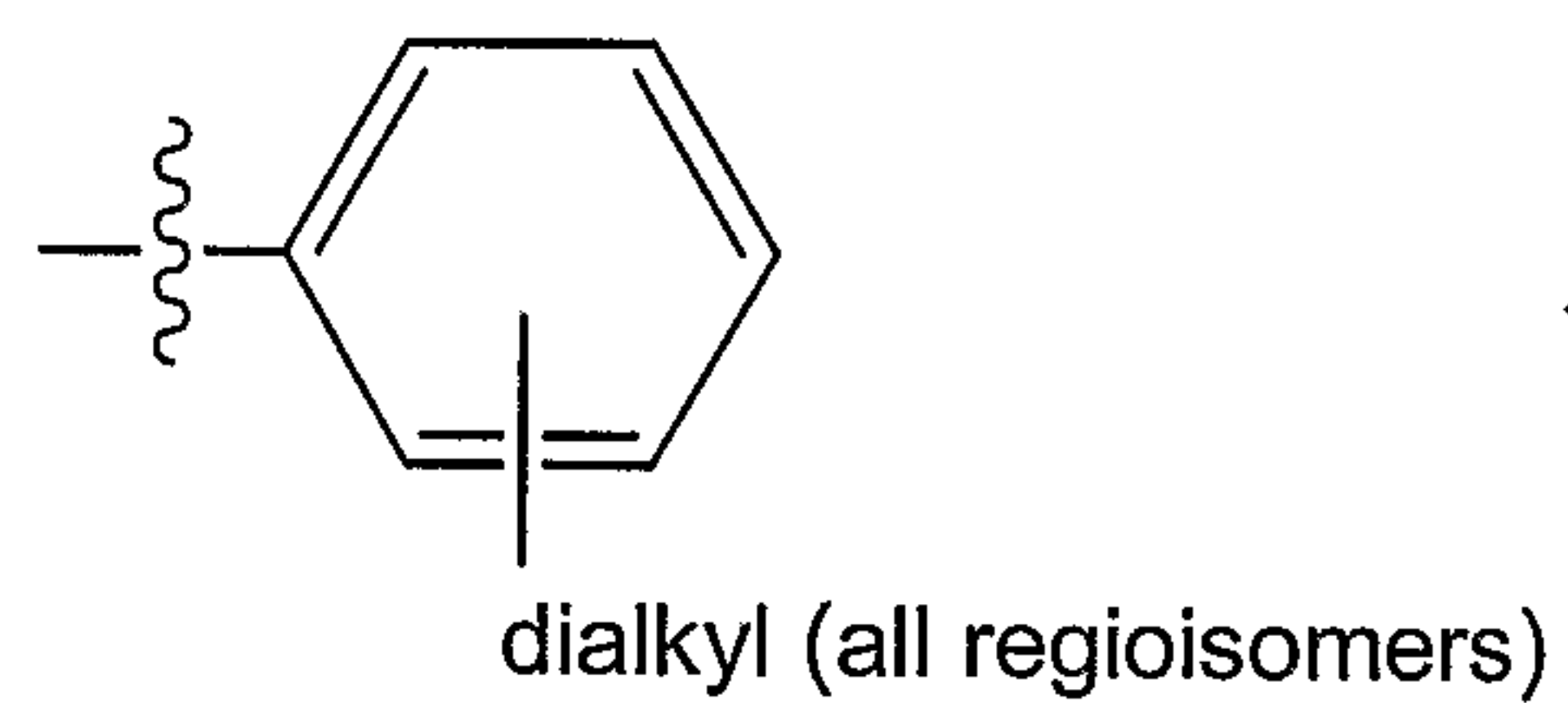
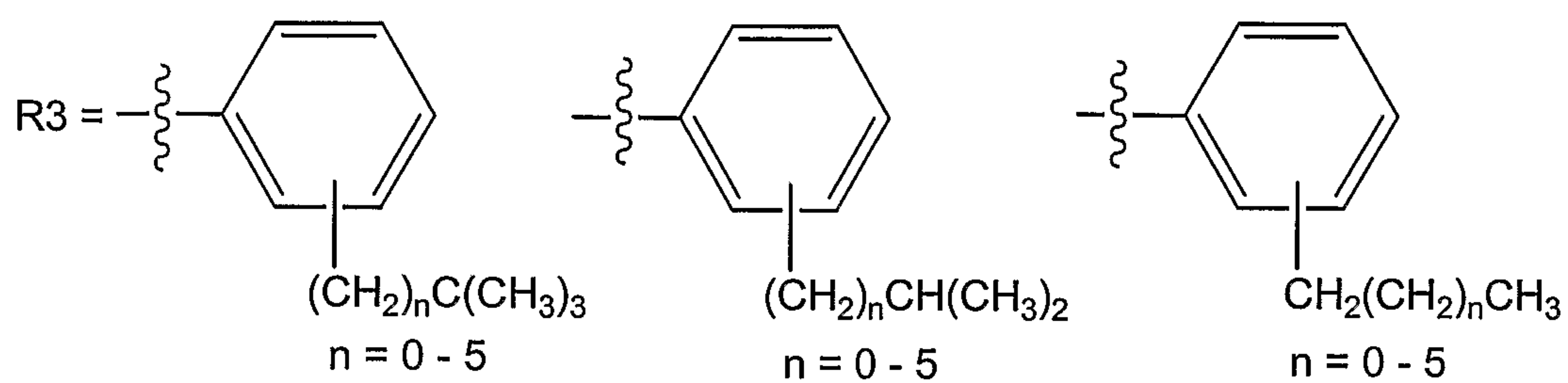
R₂ = H, alkyl, or substituted alkyl
 R₃ = H, alkyl, or substituted alkyl

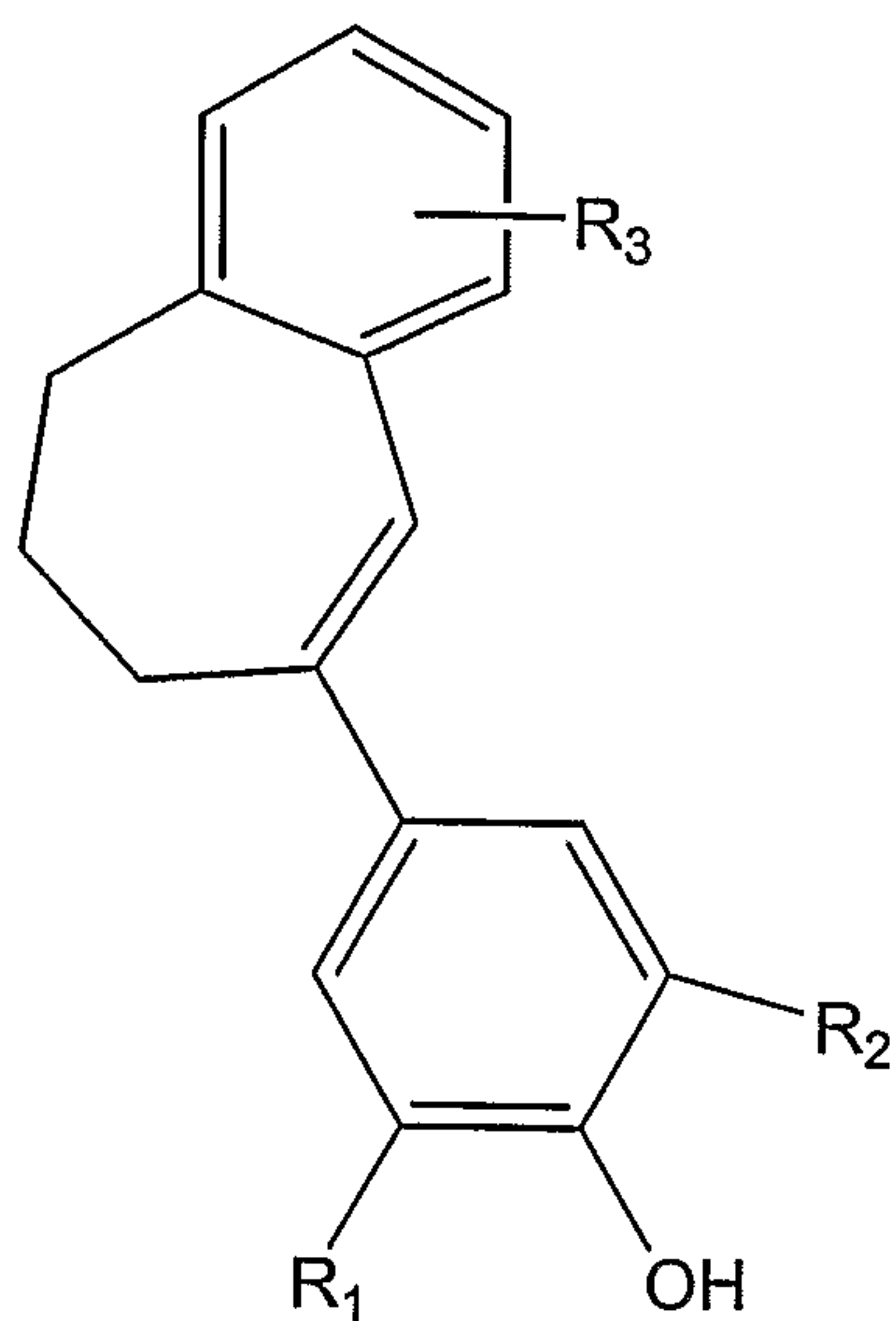
R₁ is selected hydrogen, a hydroxy, an alkoxy, a halo, an amino, a lower-alkyl-a substituted-amino, an acetamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic



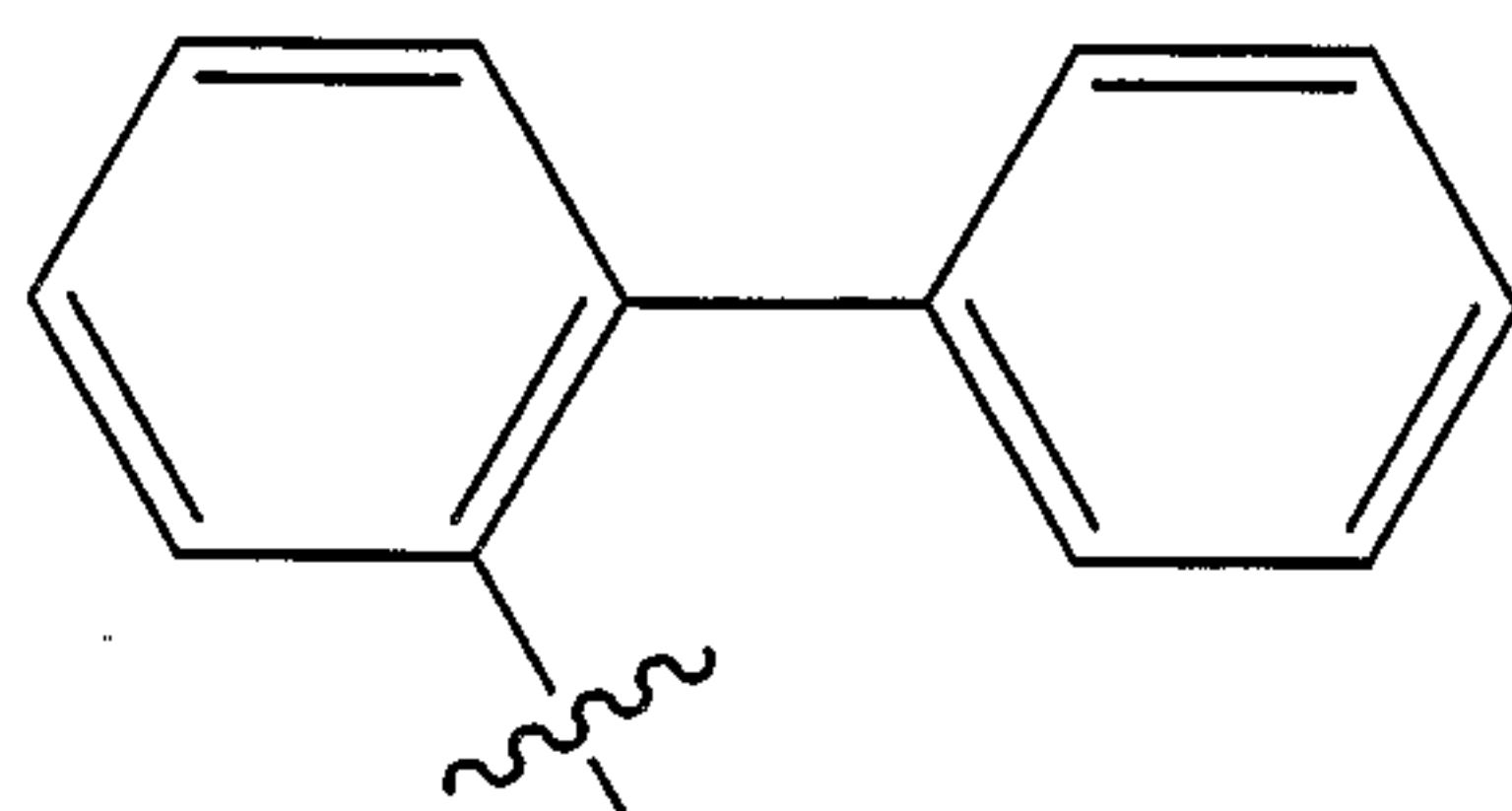
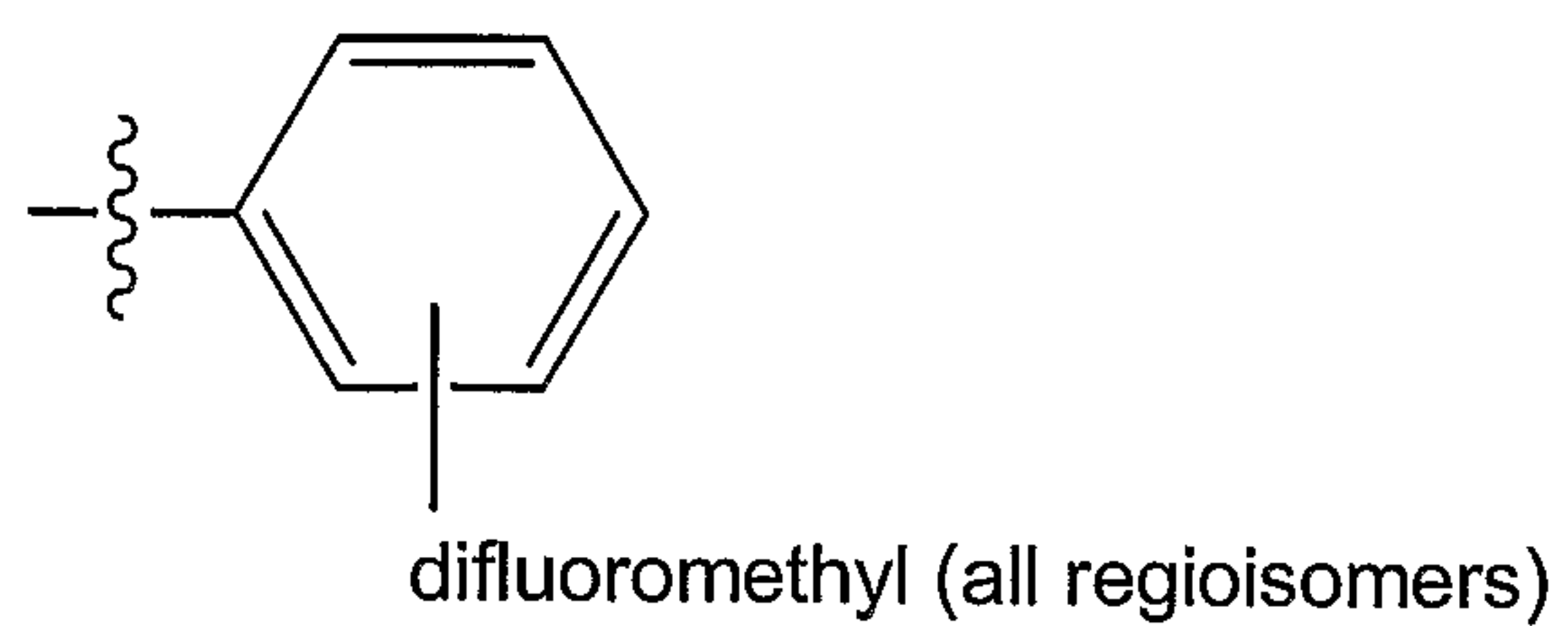
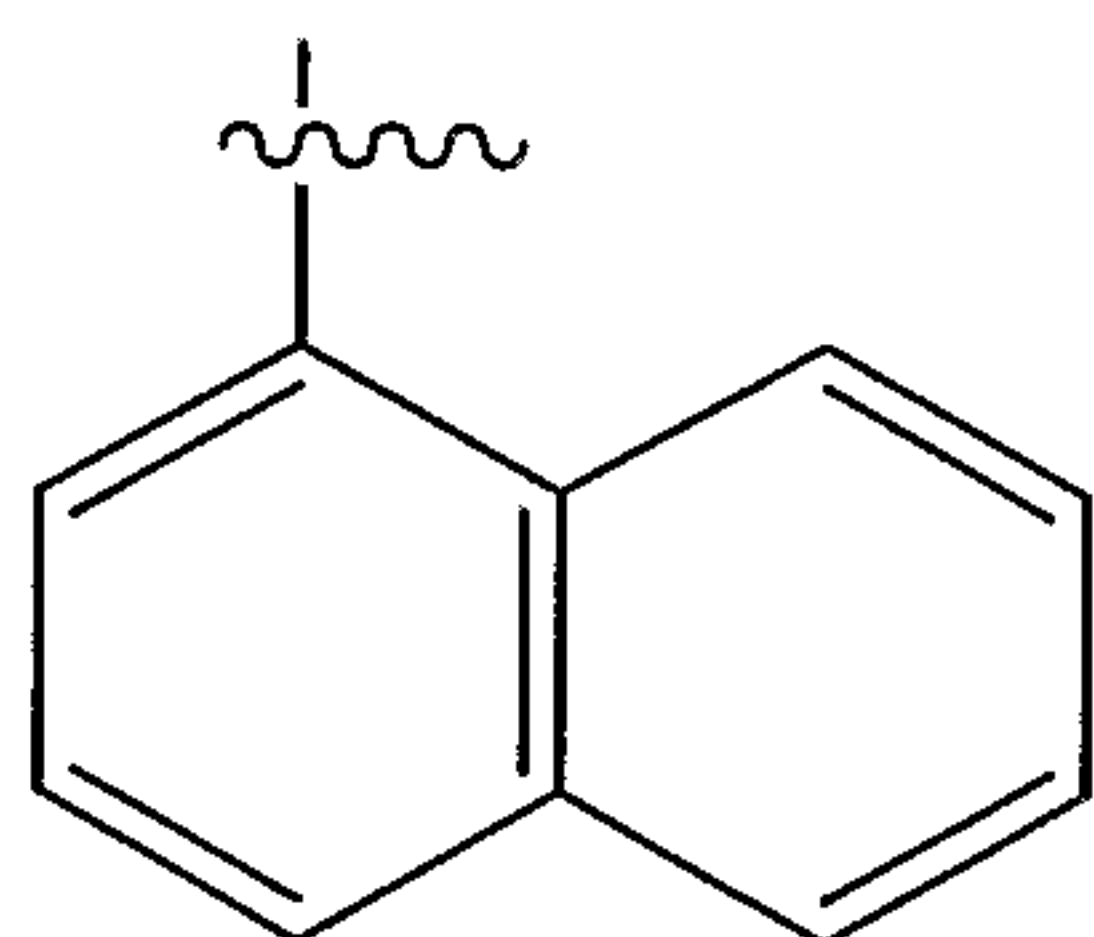
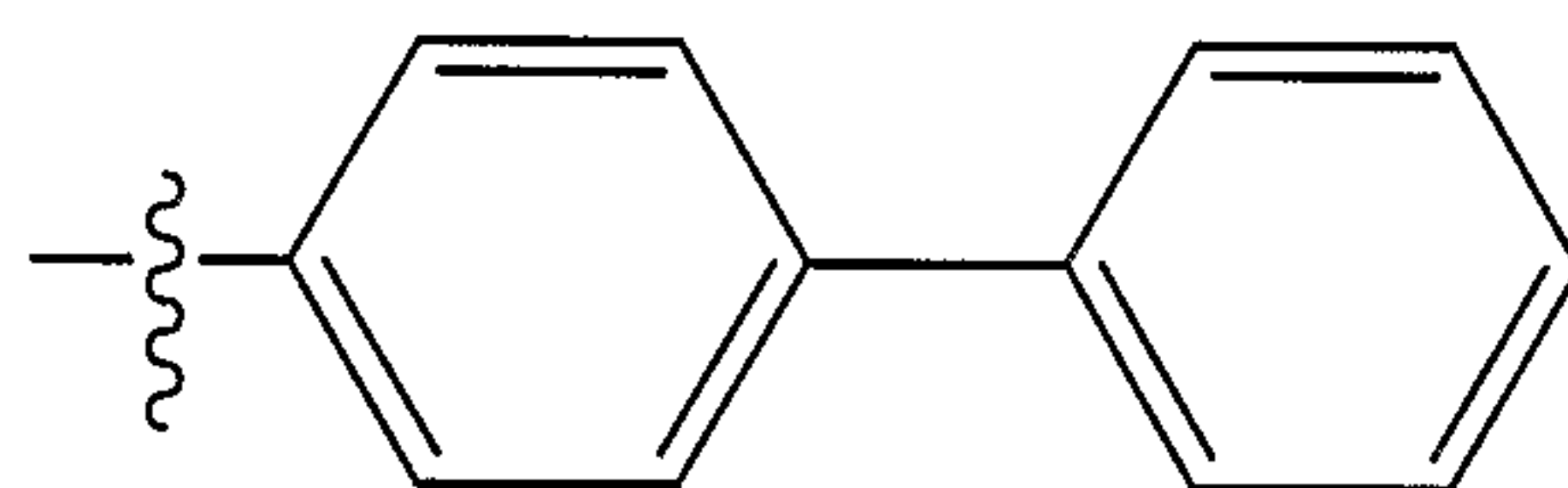
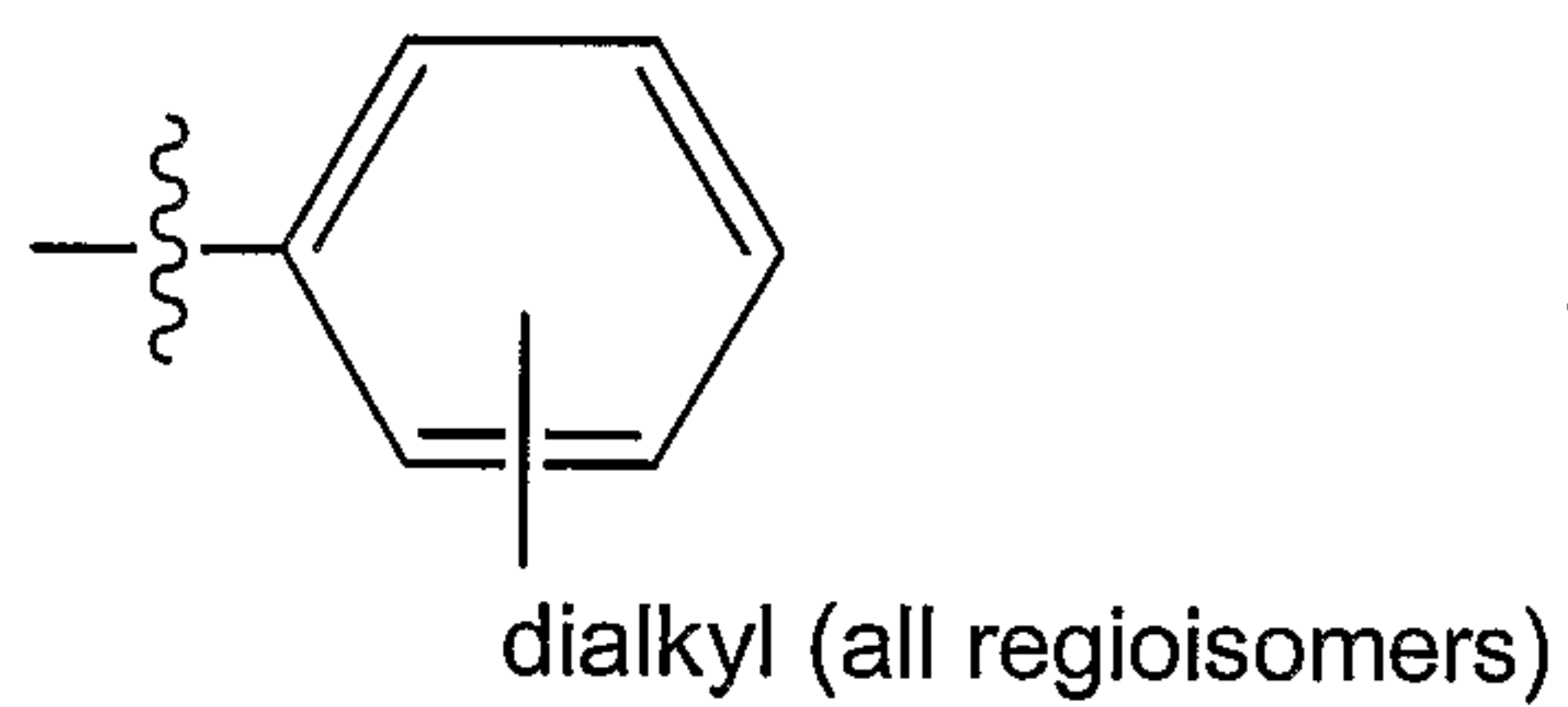
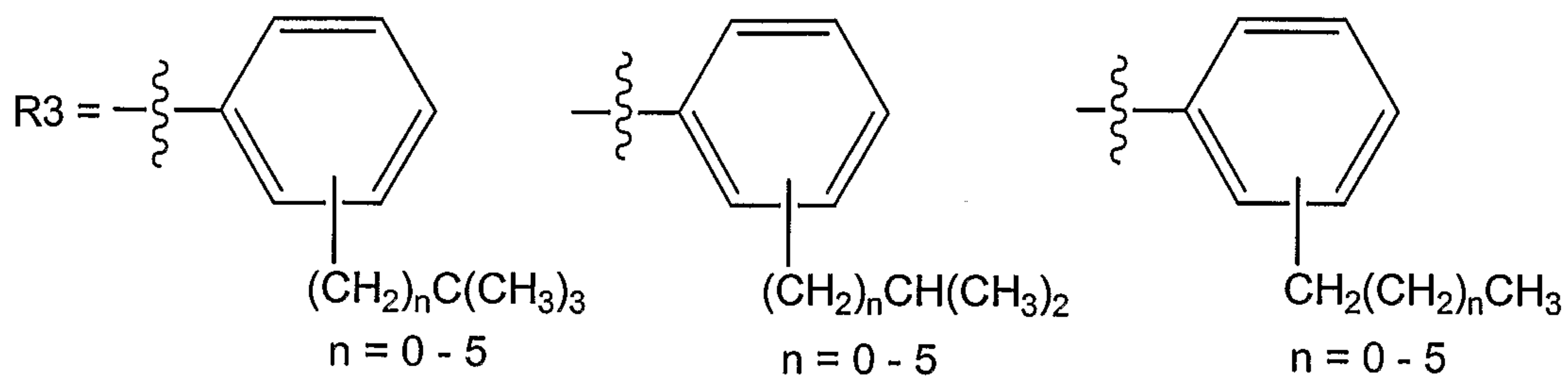


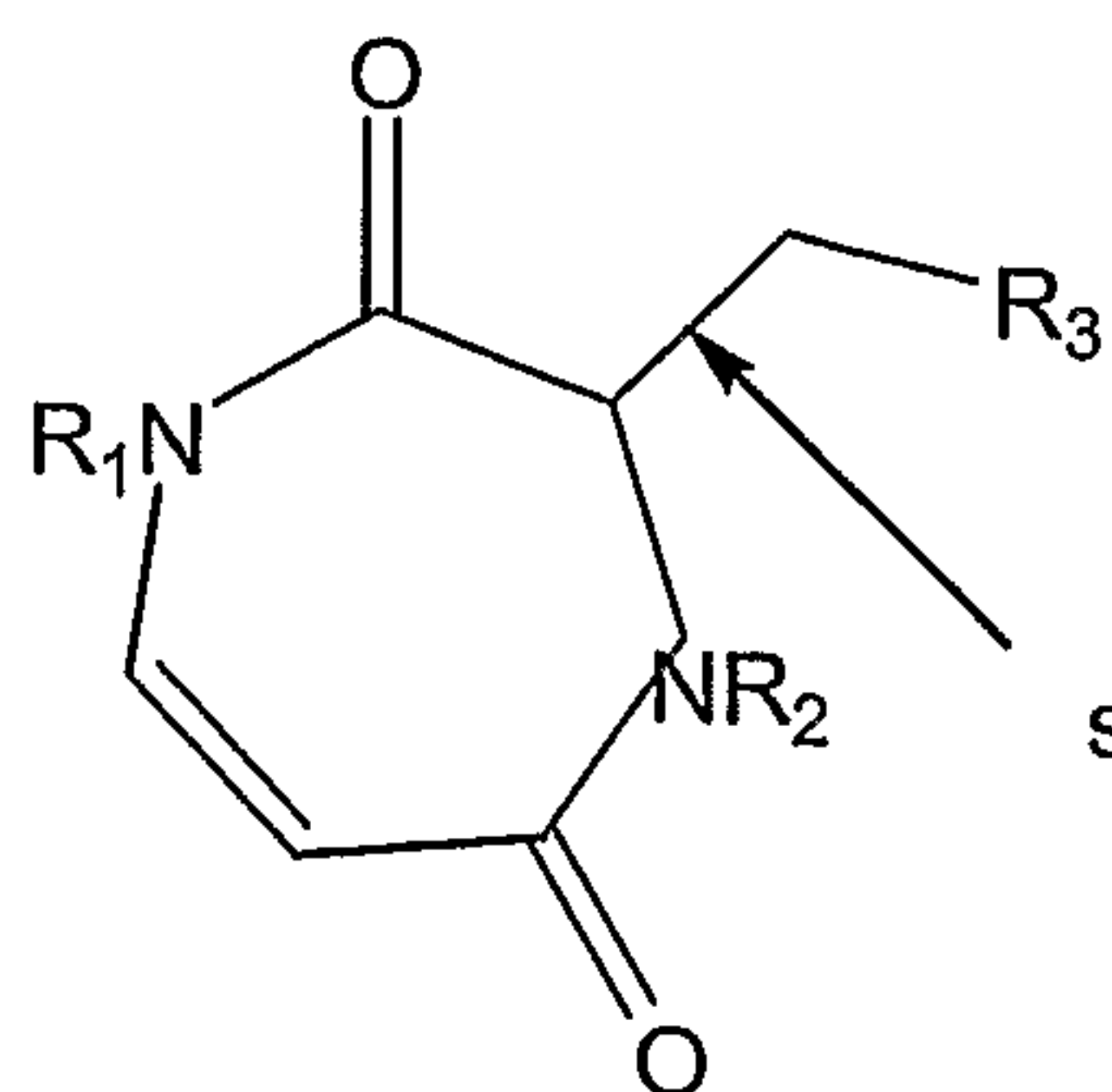
R_1 = H, alkyl, or substituted alkyl
 R_2 = H, alkyl, or substituted alkyl





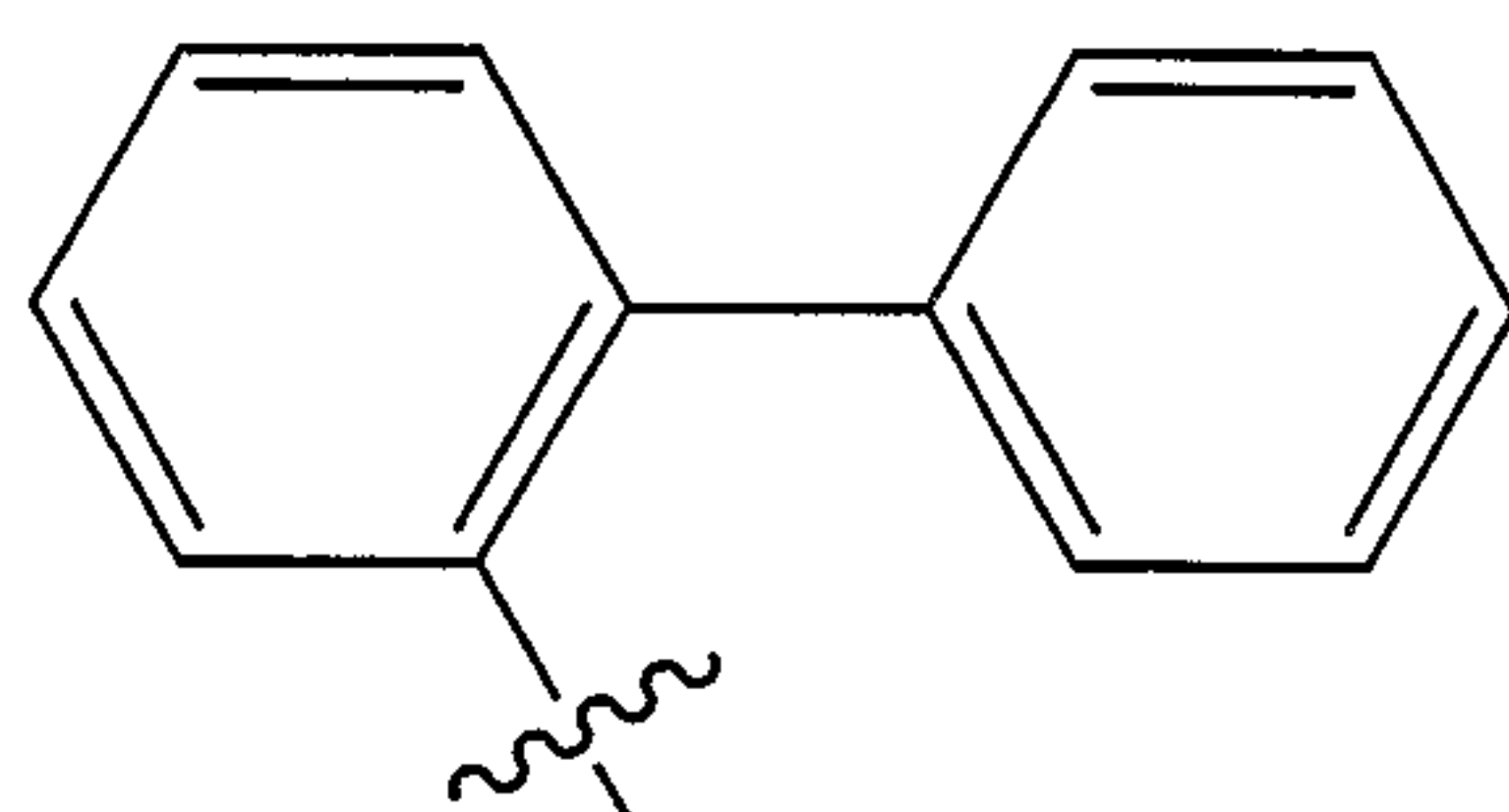
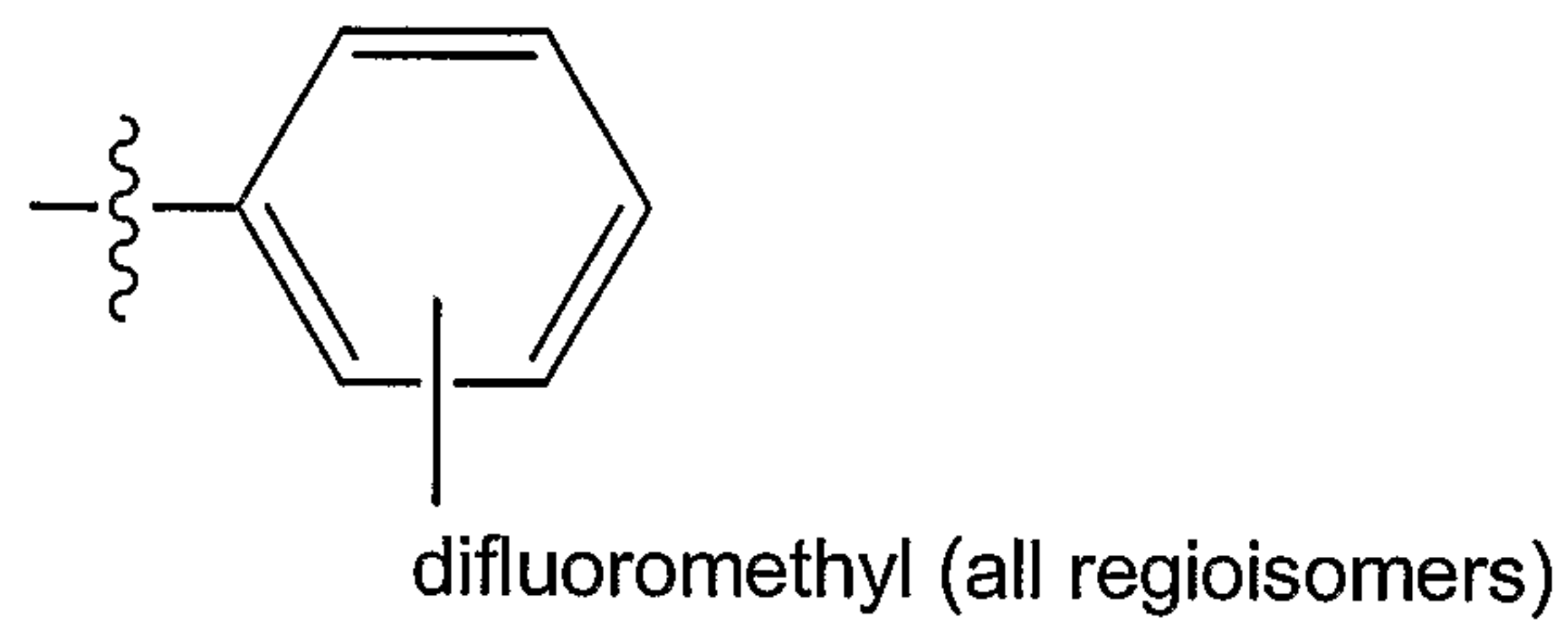
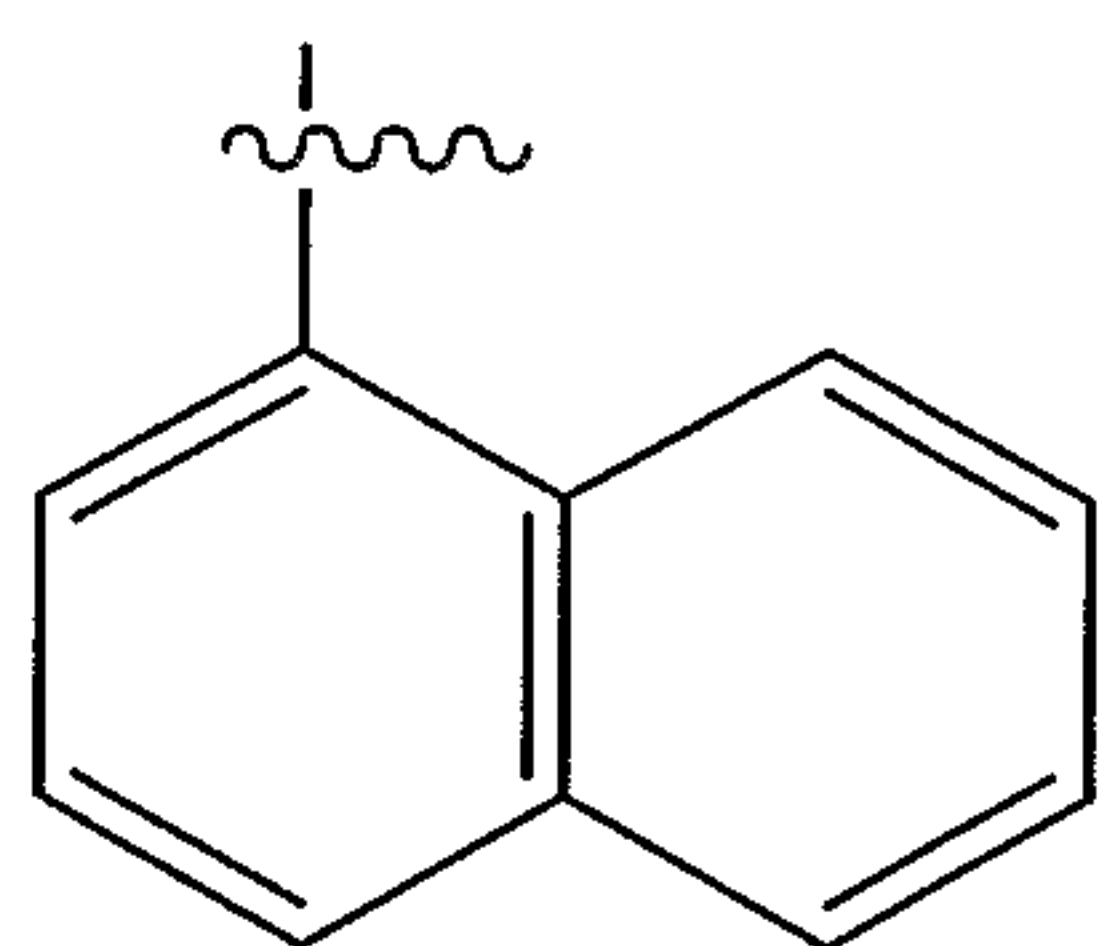
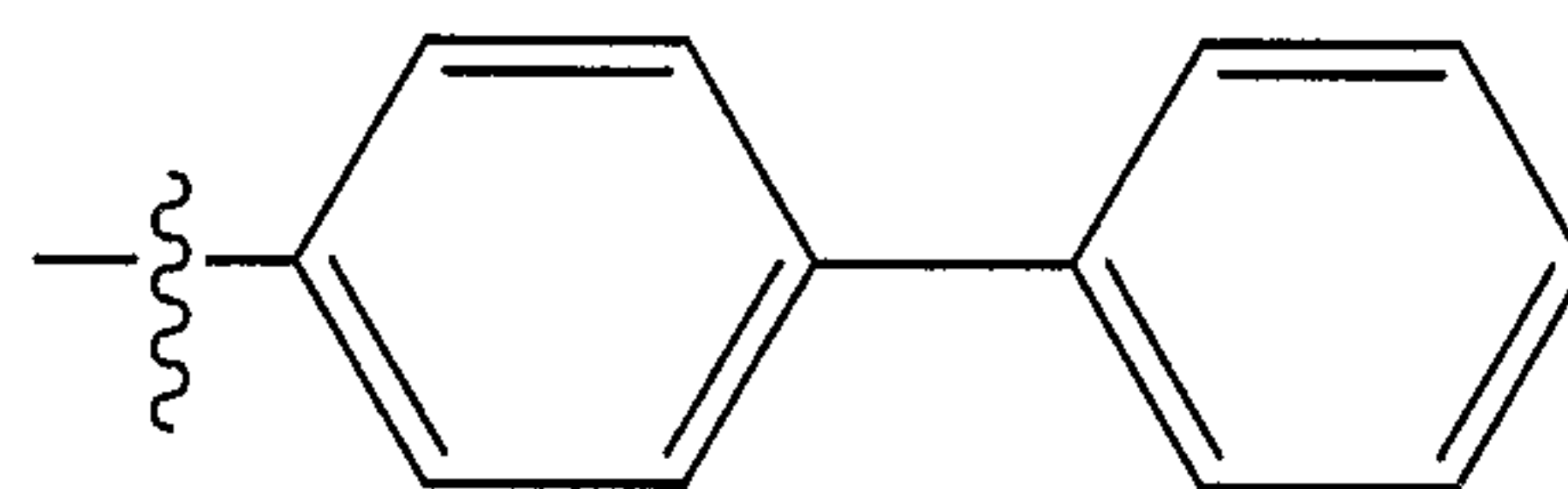
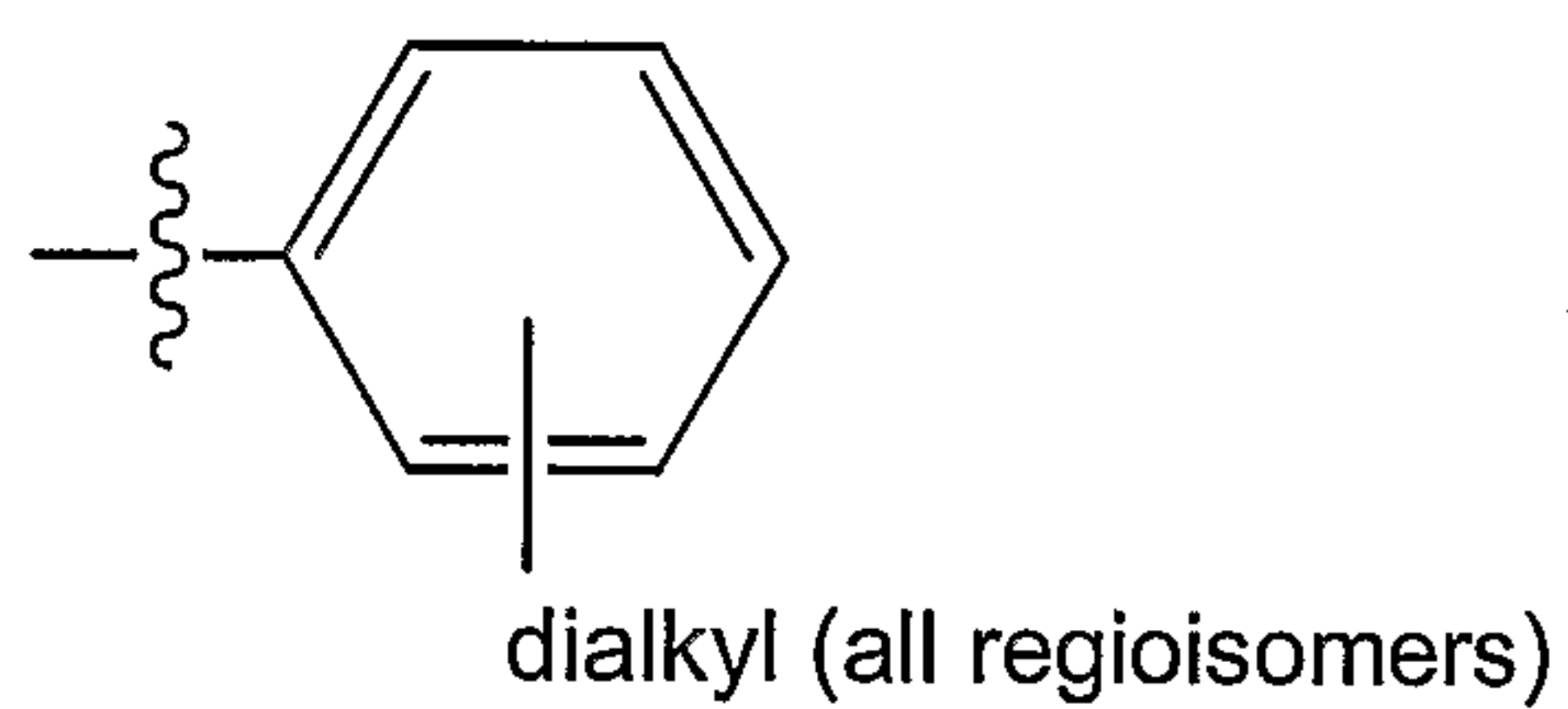
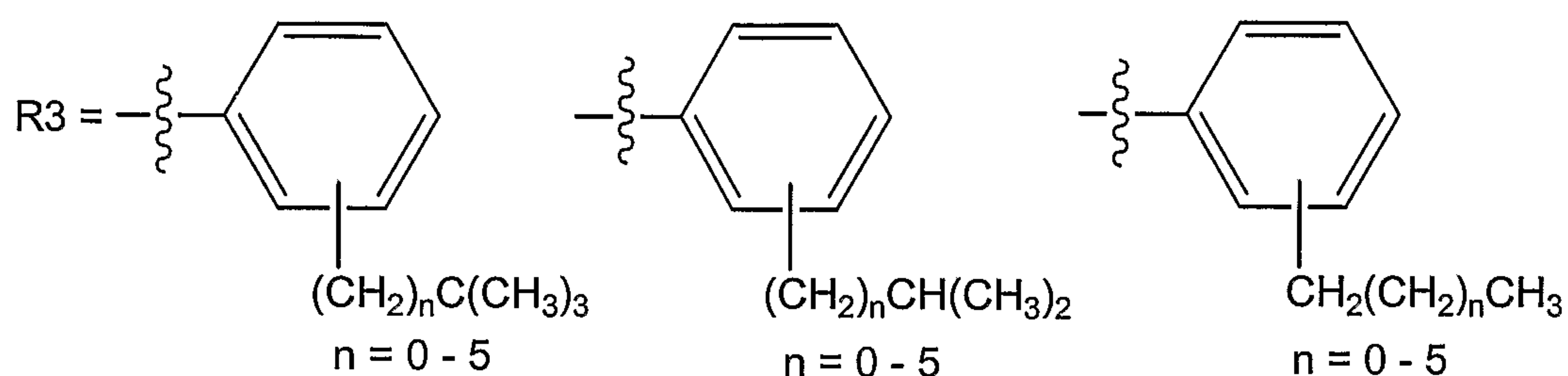
R1 = H, alkyl, or substituted alkyl
R2 = H, alkyl, or substituted alkyl

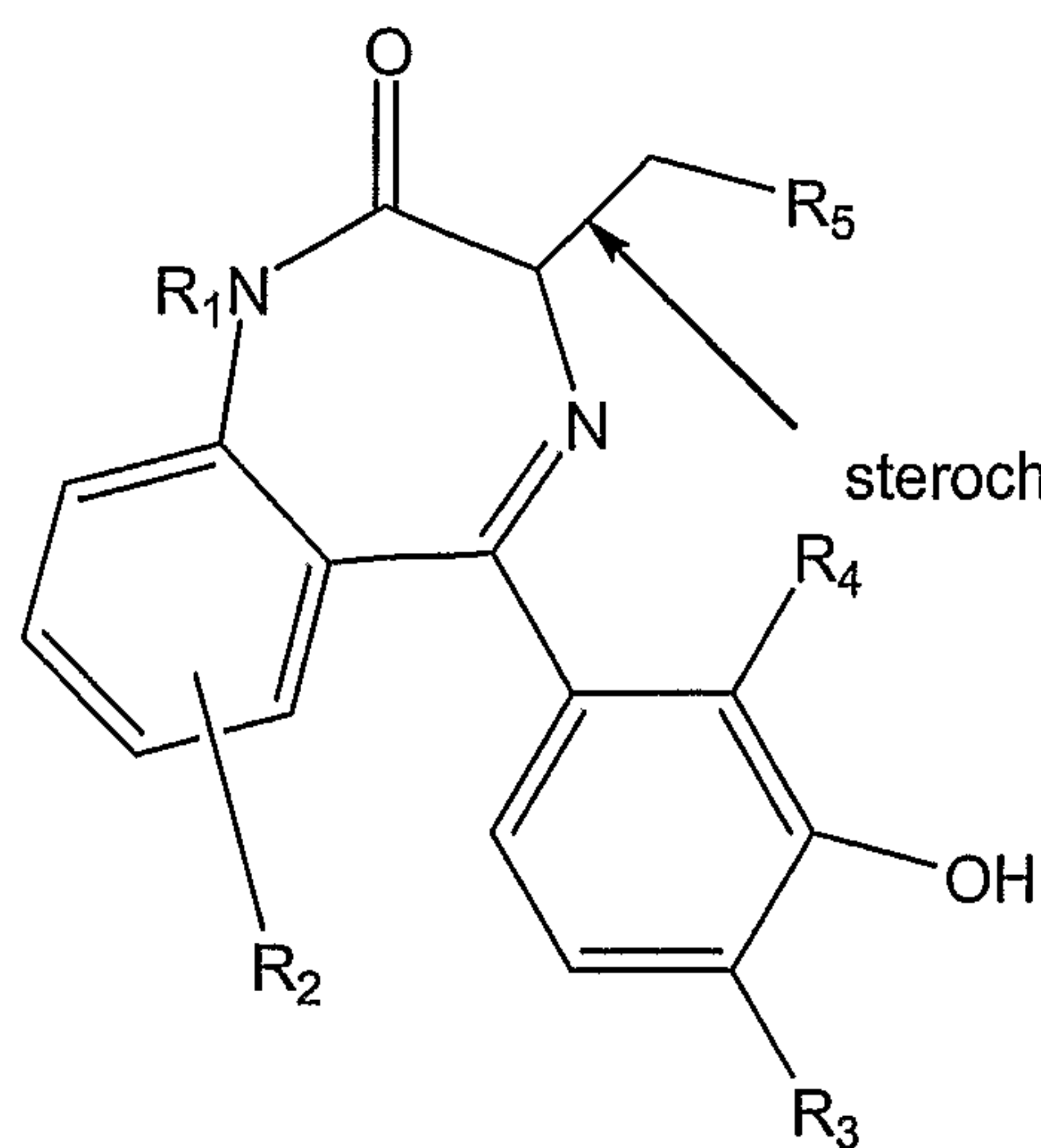




R1 = H, alkyl, or substituted alkyl
R2 = H, alkyl, or substituted alkyl

stereochemistry is R, S, or racemic





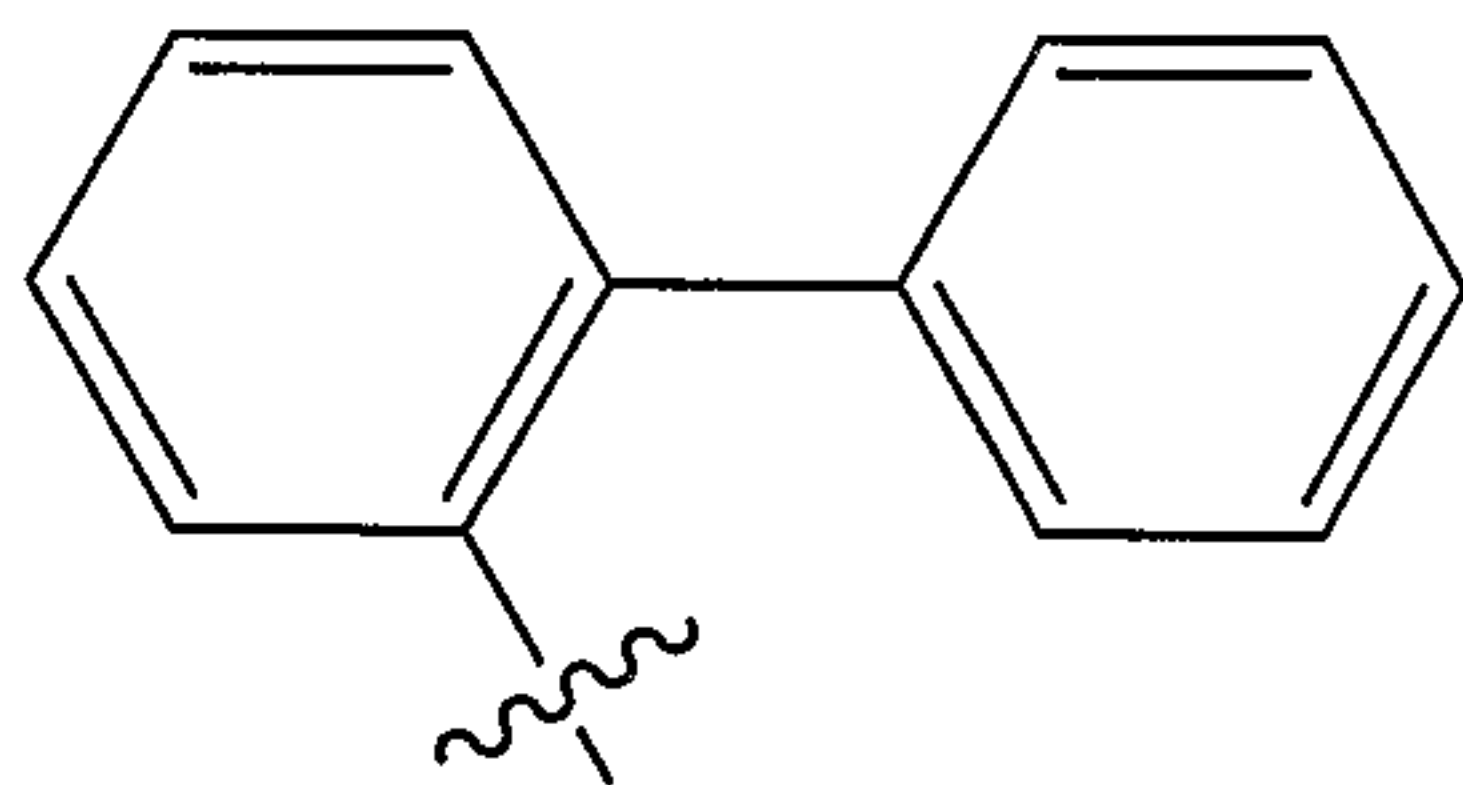
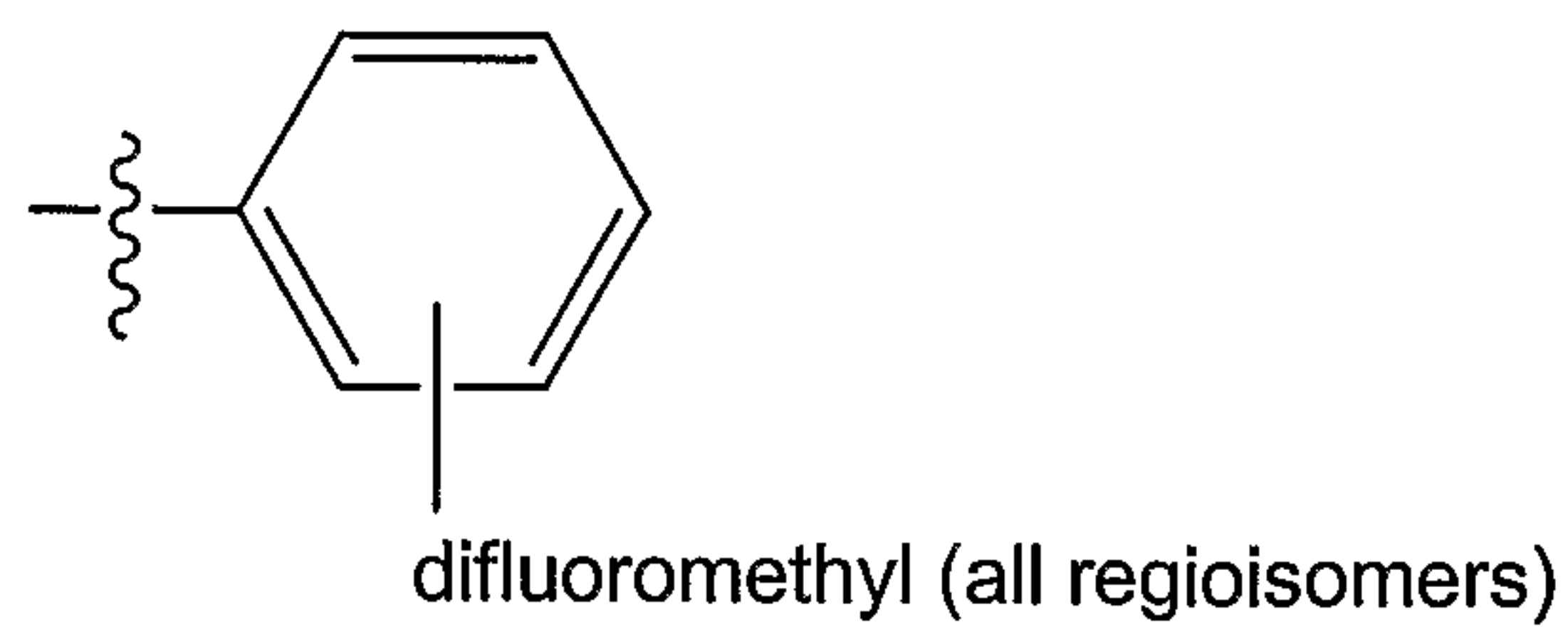
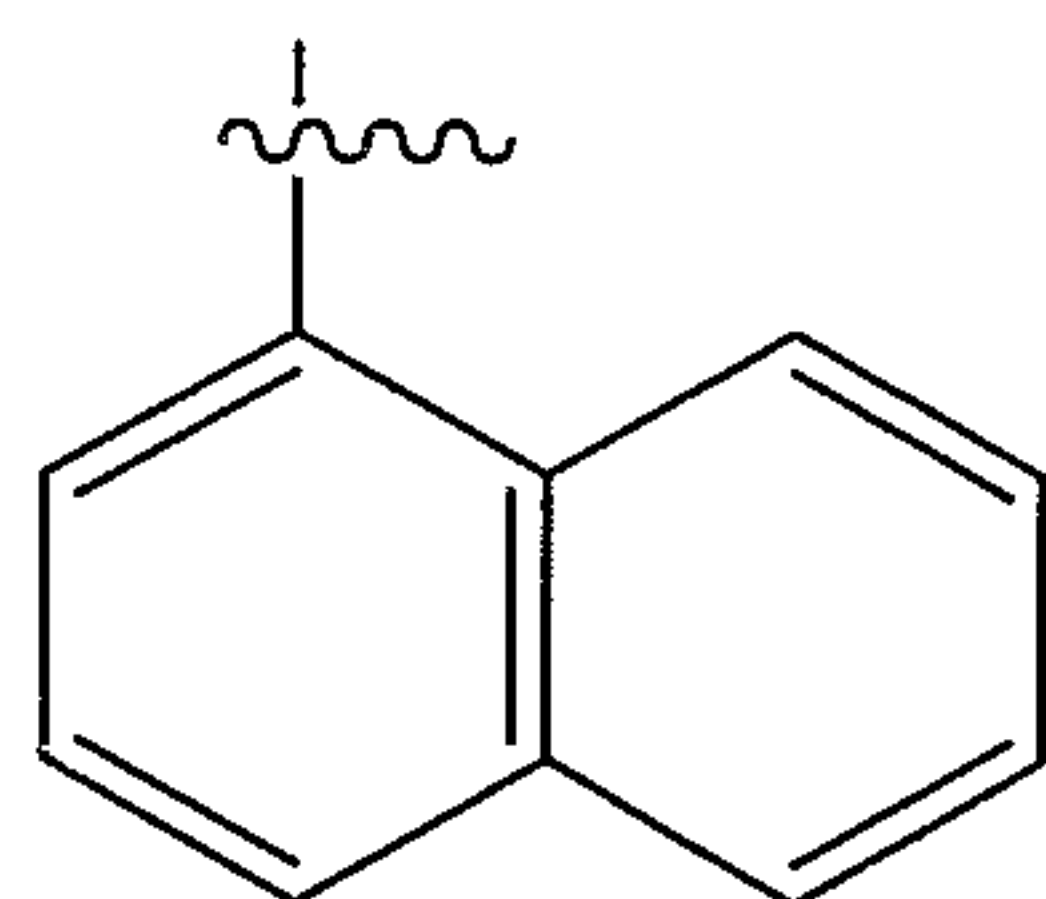
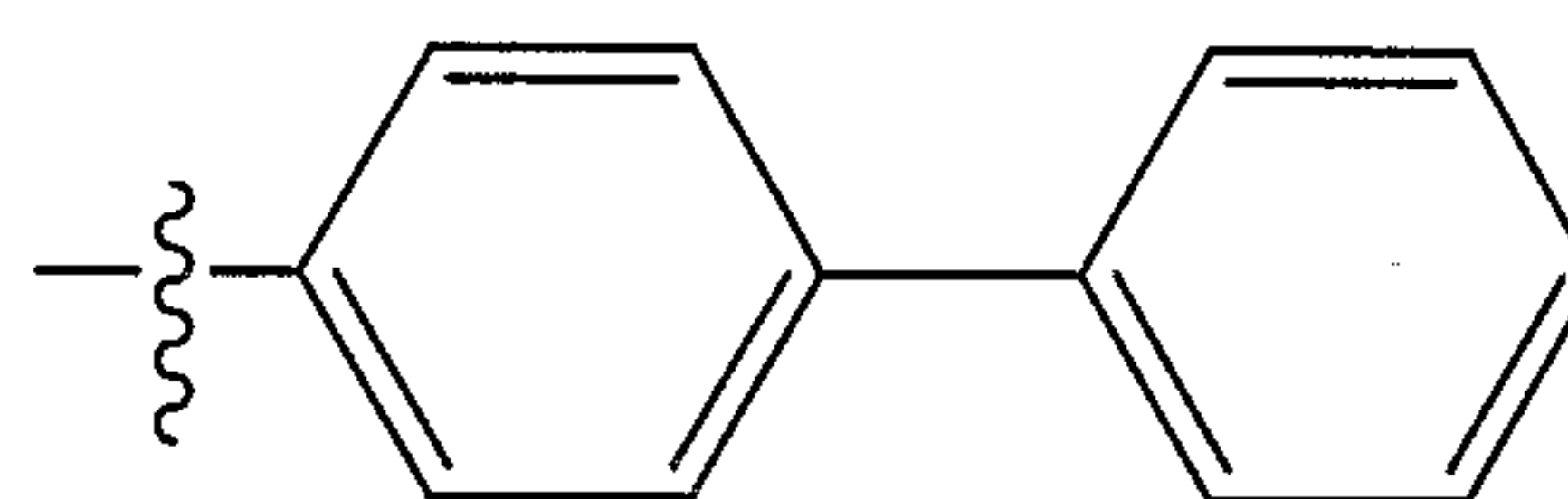
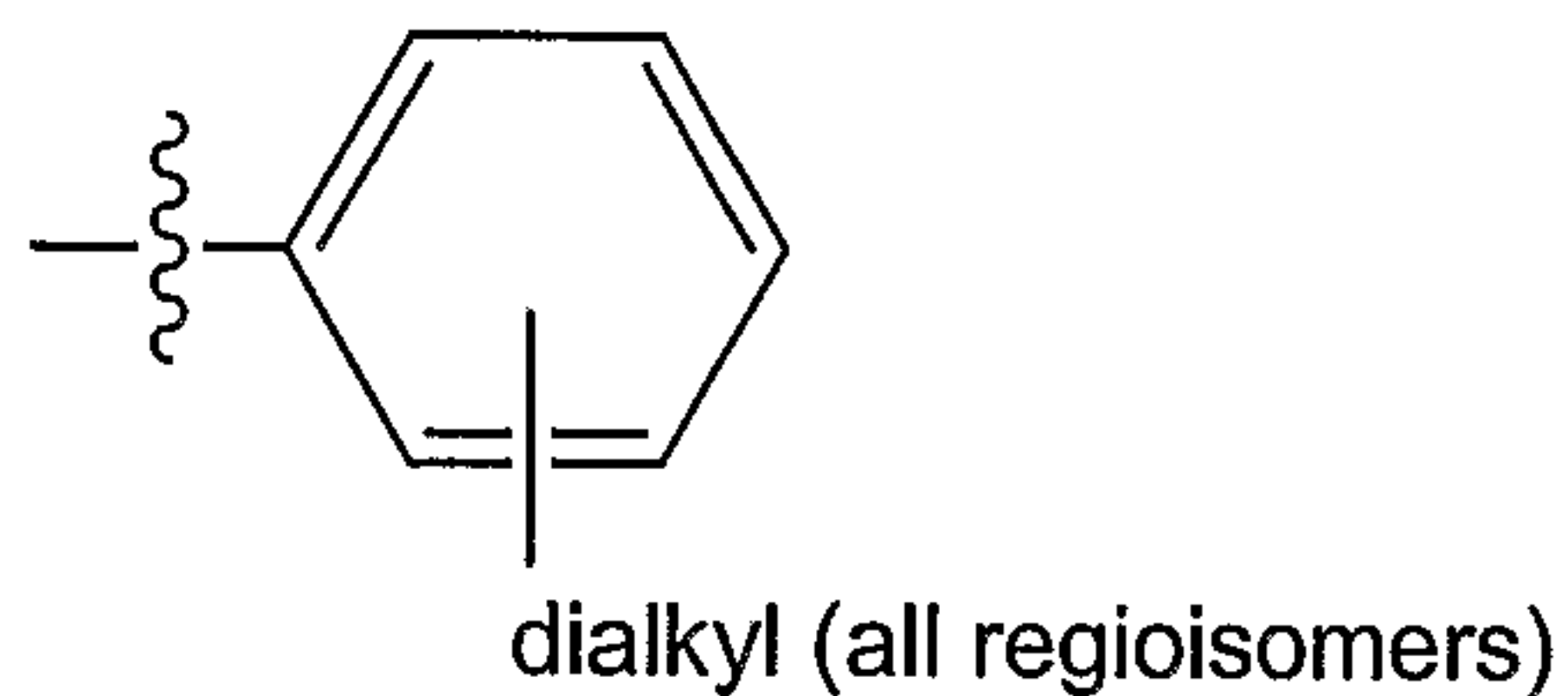
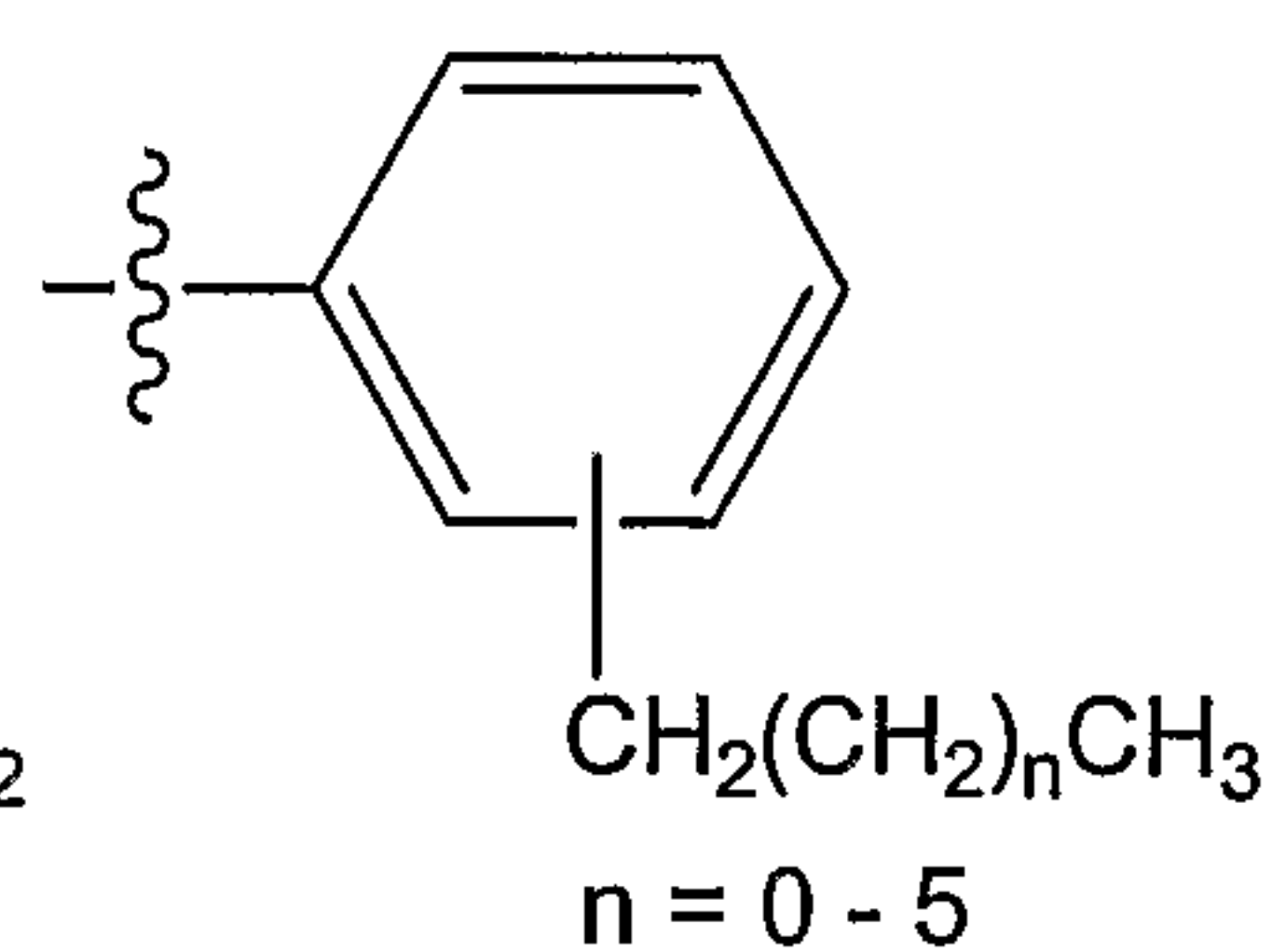
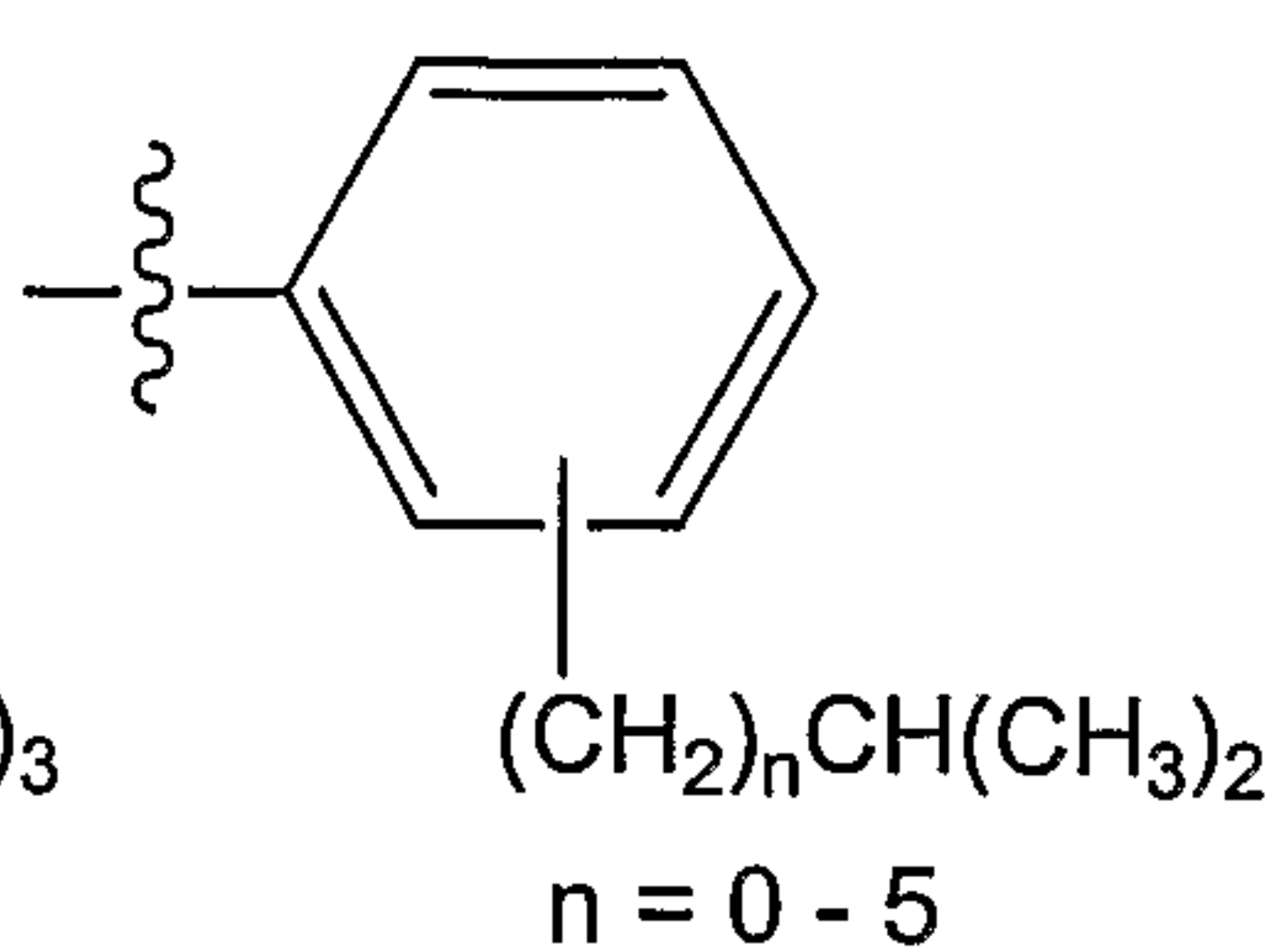
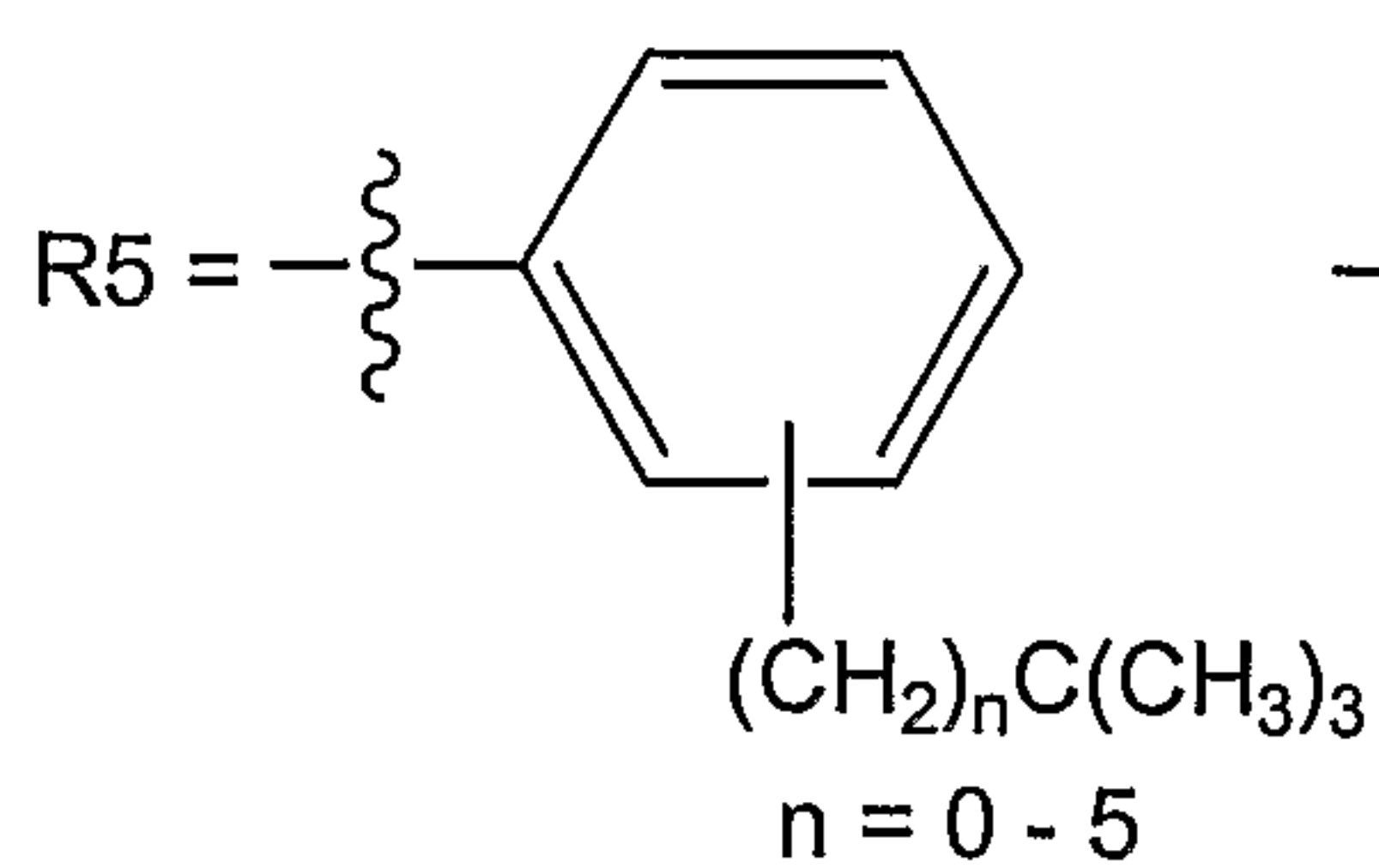
R1 = H, alkyl, or substituted alkyl

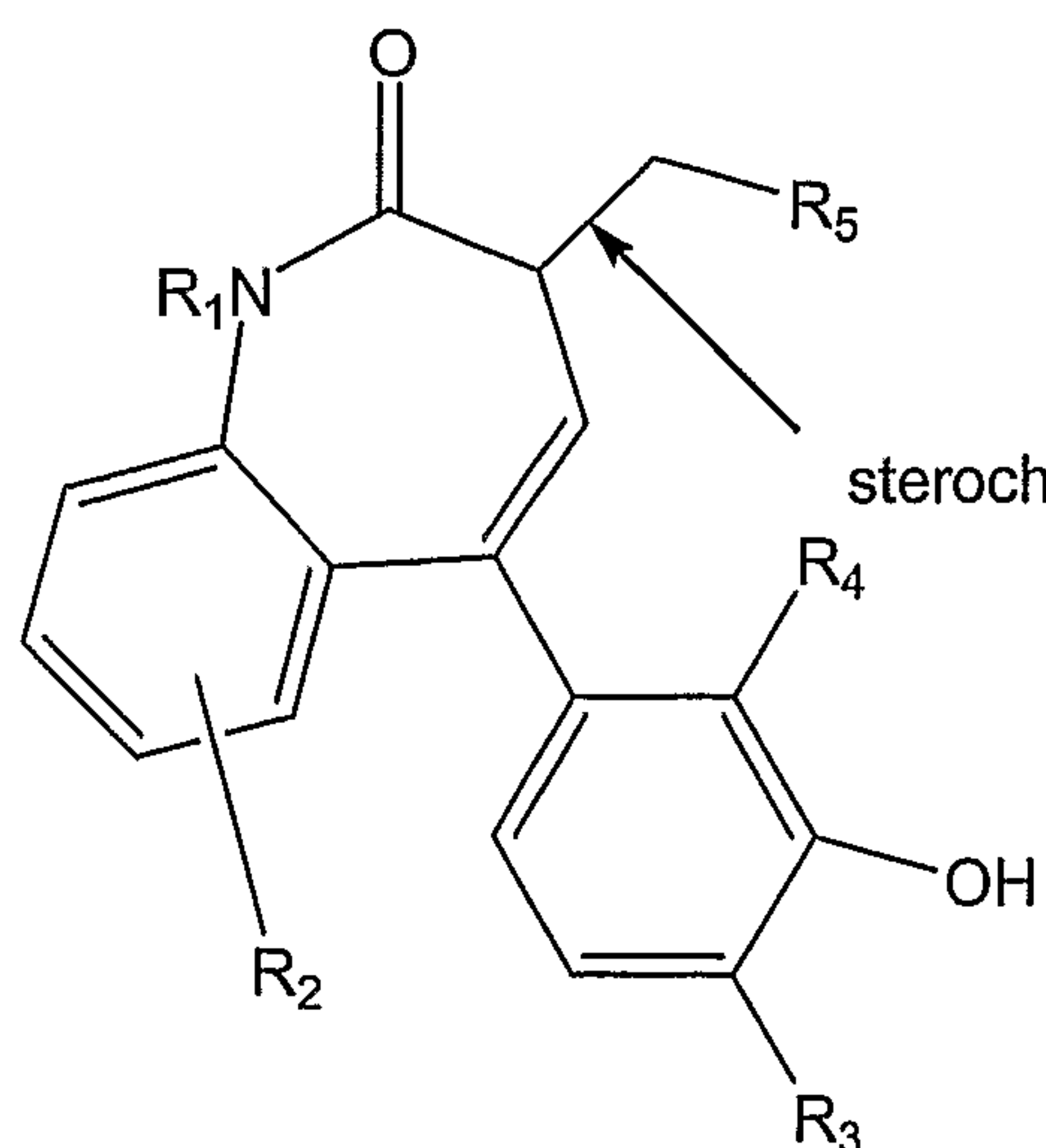
R3 = H, alkyl, or substituted alkyl

R4 = H, alkyl, or substituted alkyl

stereochemistry is R, S, or racemic

R2 is selected from hydrogen, hydroxy, an alkoxy, a halo, an amino, a lower-alkyl-a substituted-amino, an acetamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic





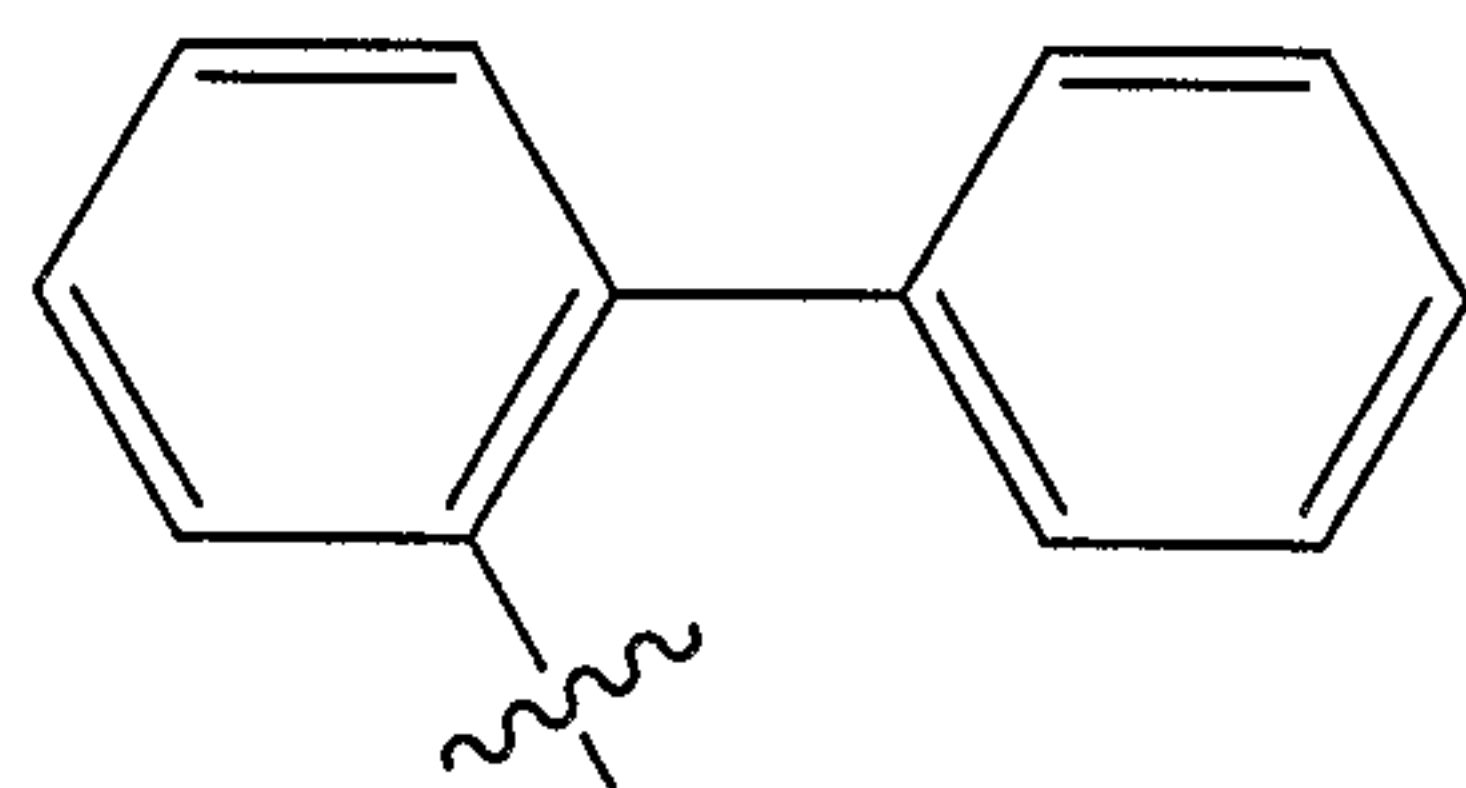
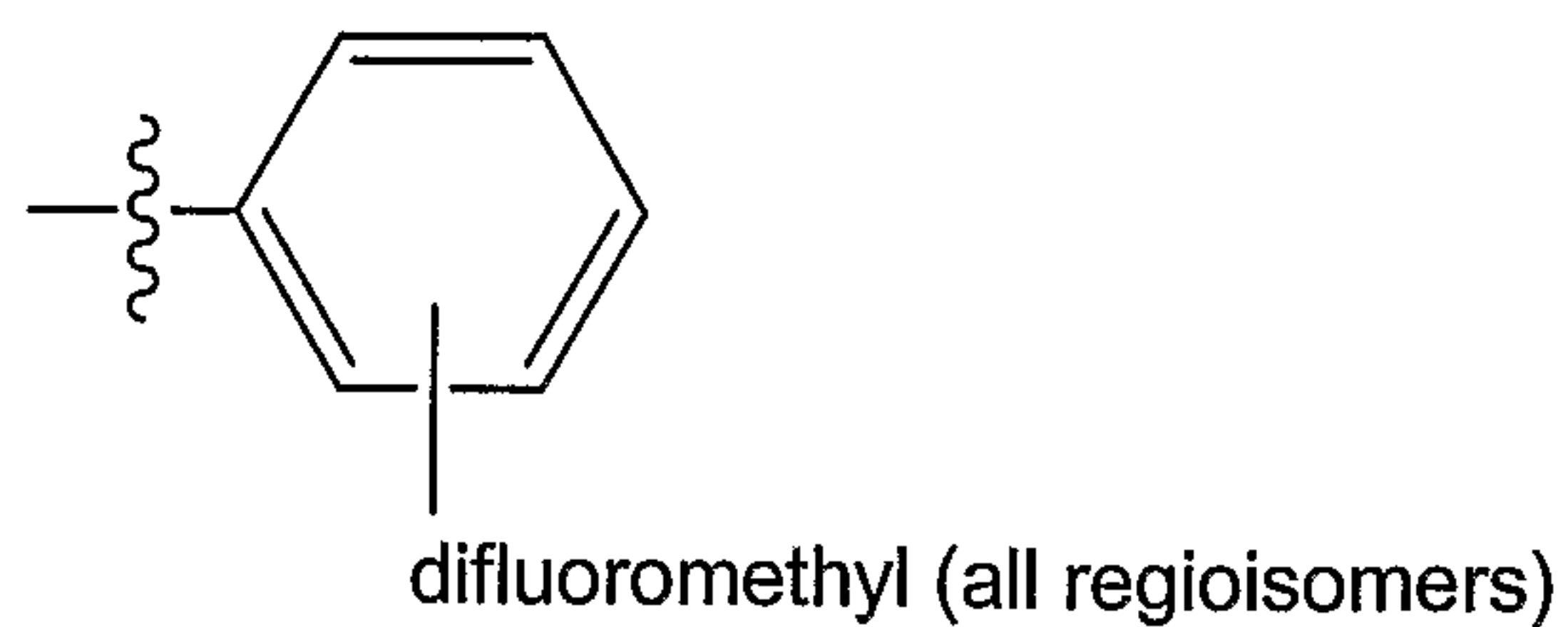
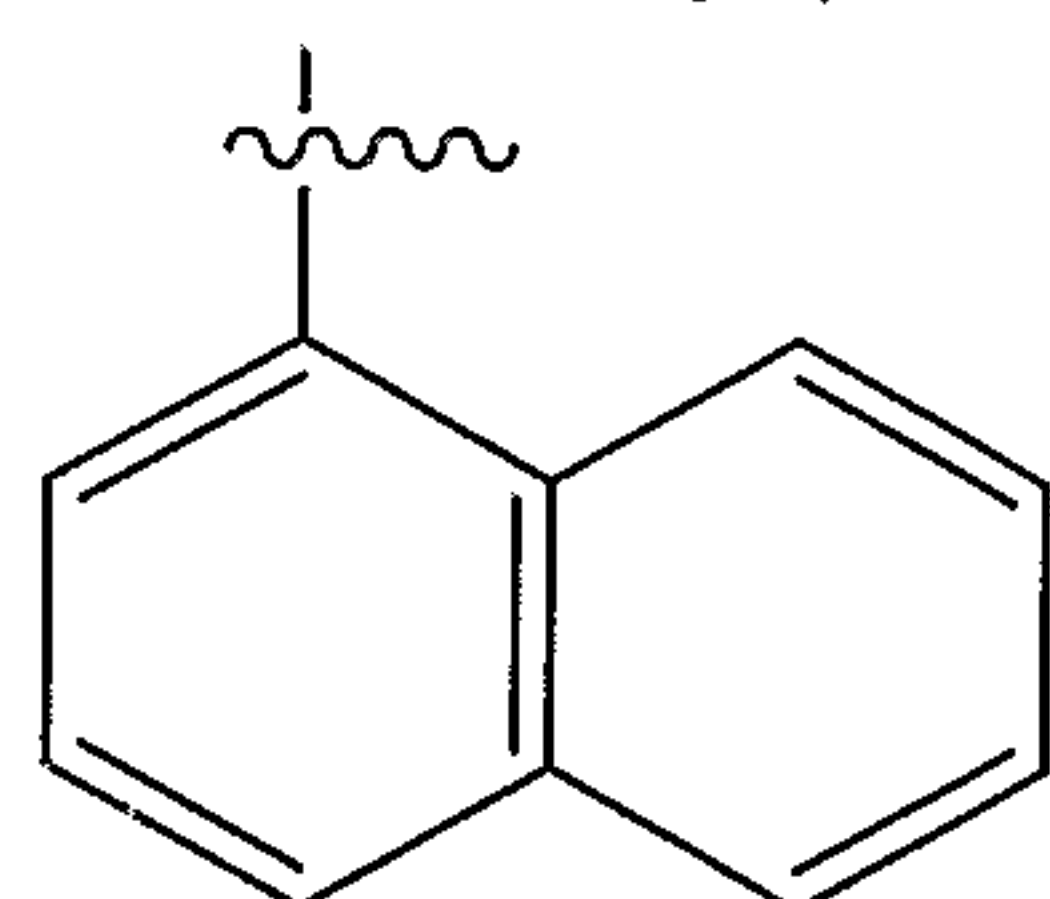
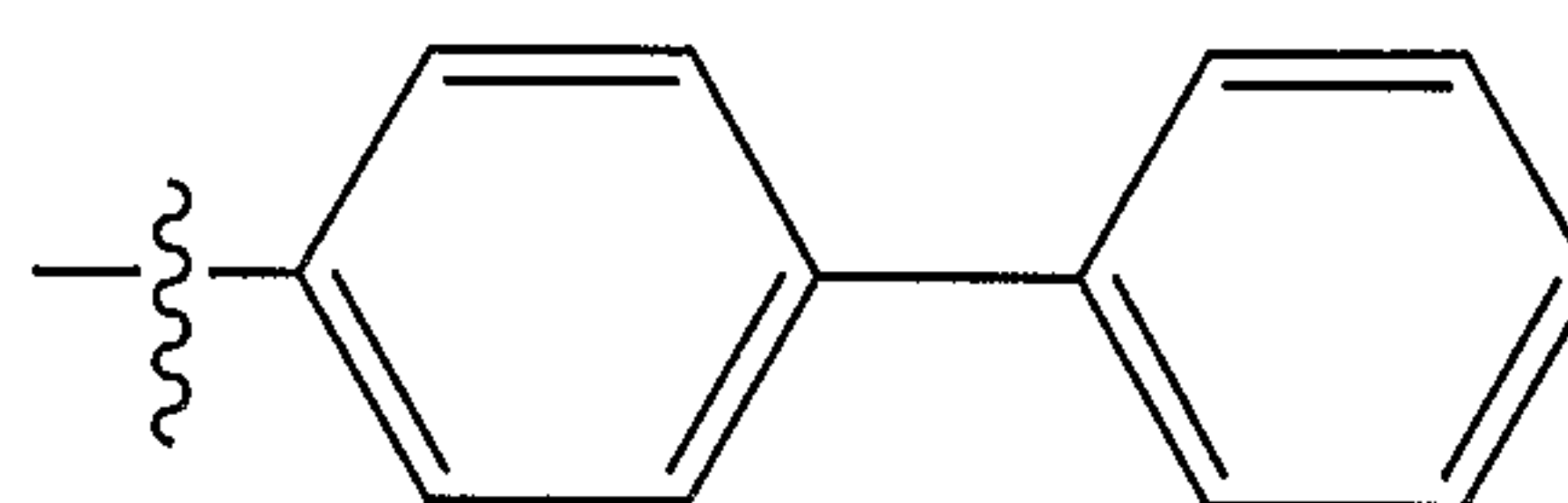
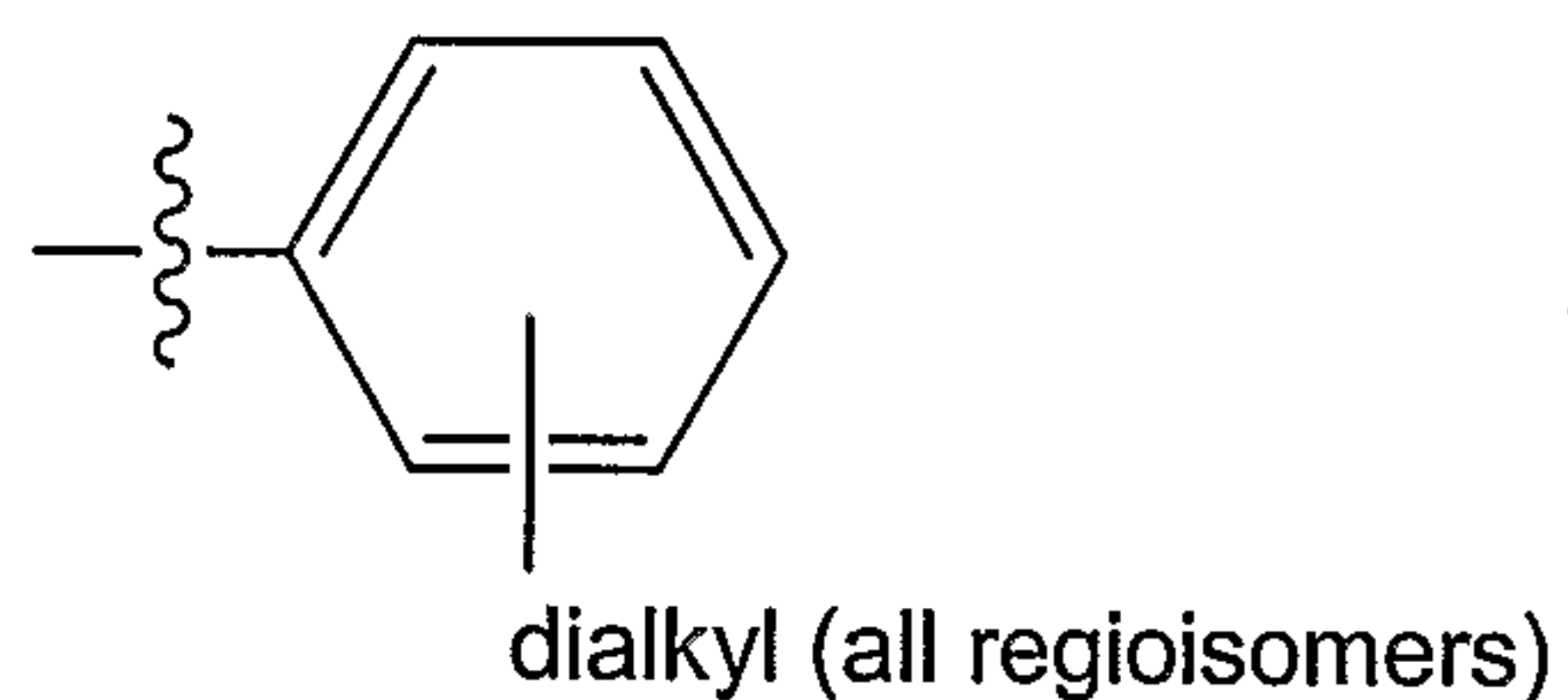
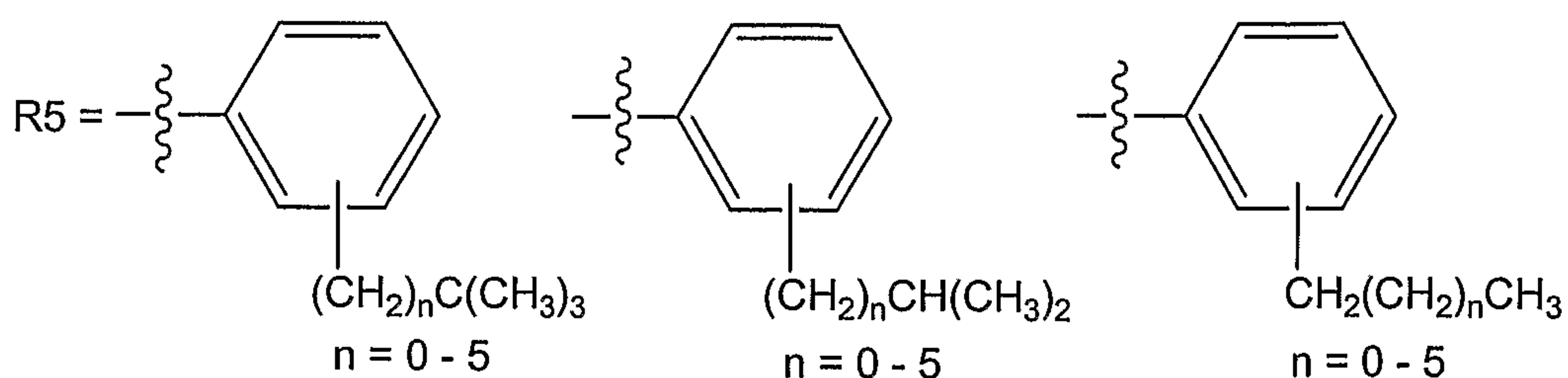
R_1 = H, alkyl, or substituted alkyl

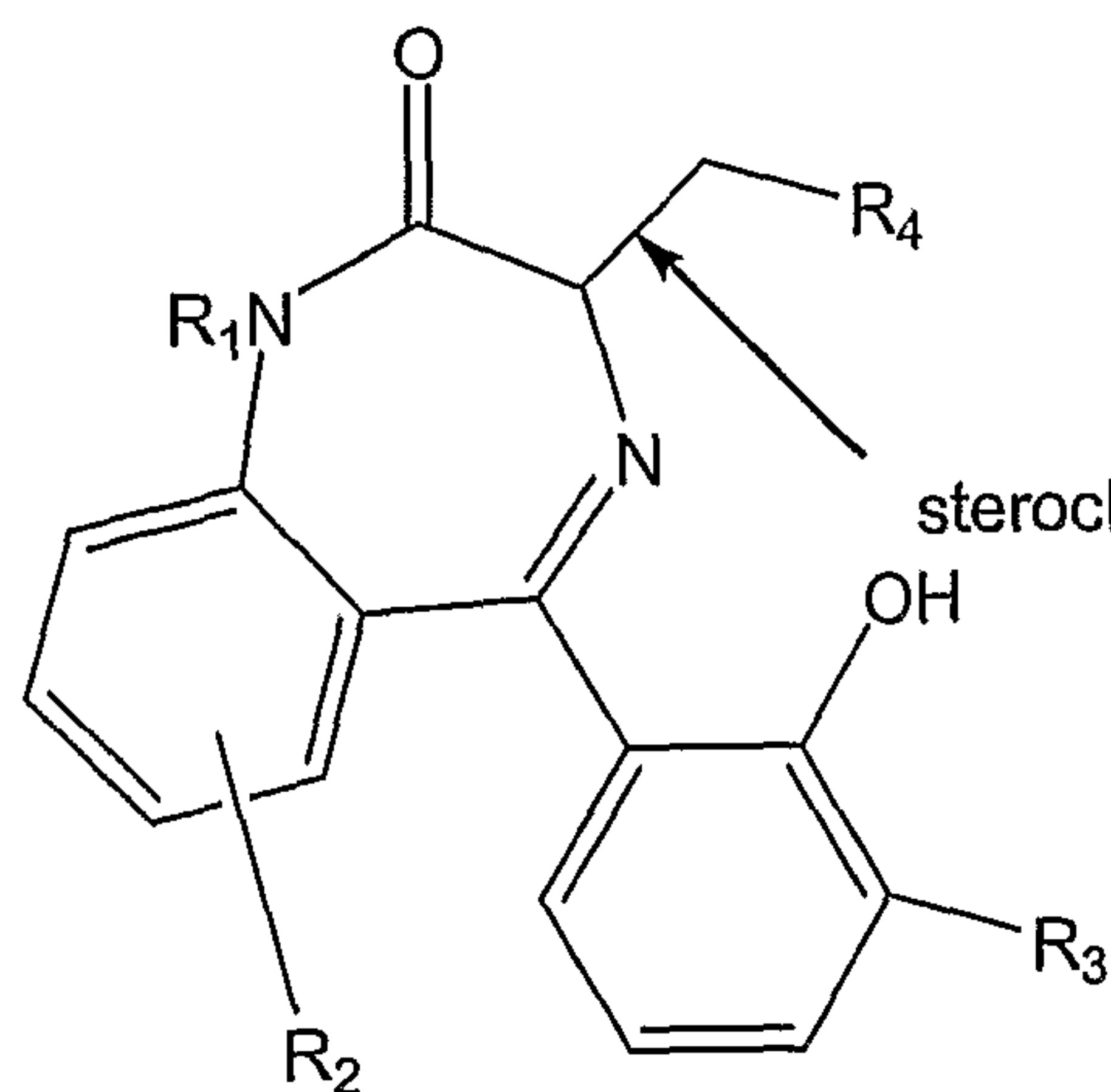
R_3 = H, alkyl, or substituted alkyl

R_4 = H, alkyl, or substituted alkyl

stereochemistry is R, S, or racemic

R_2 is selected from hydrogen, hydroxy, an alkoxy, a halo, an amino, a lower-alkyl-a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic



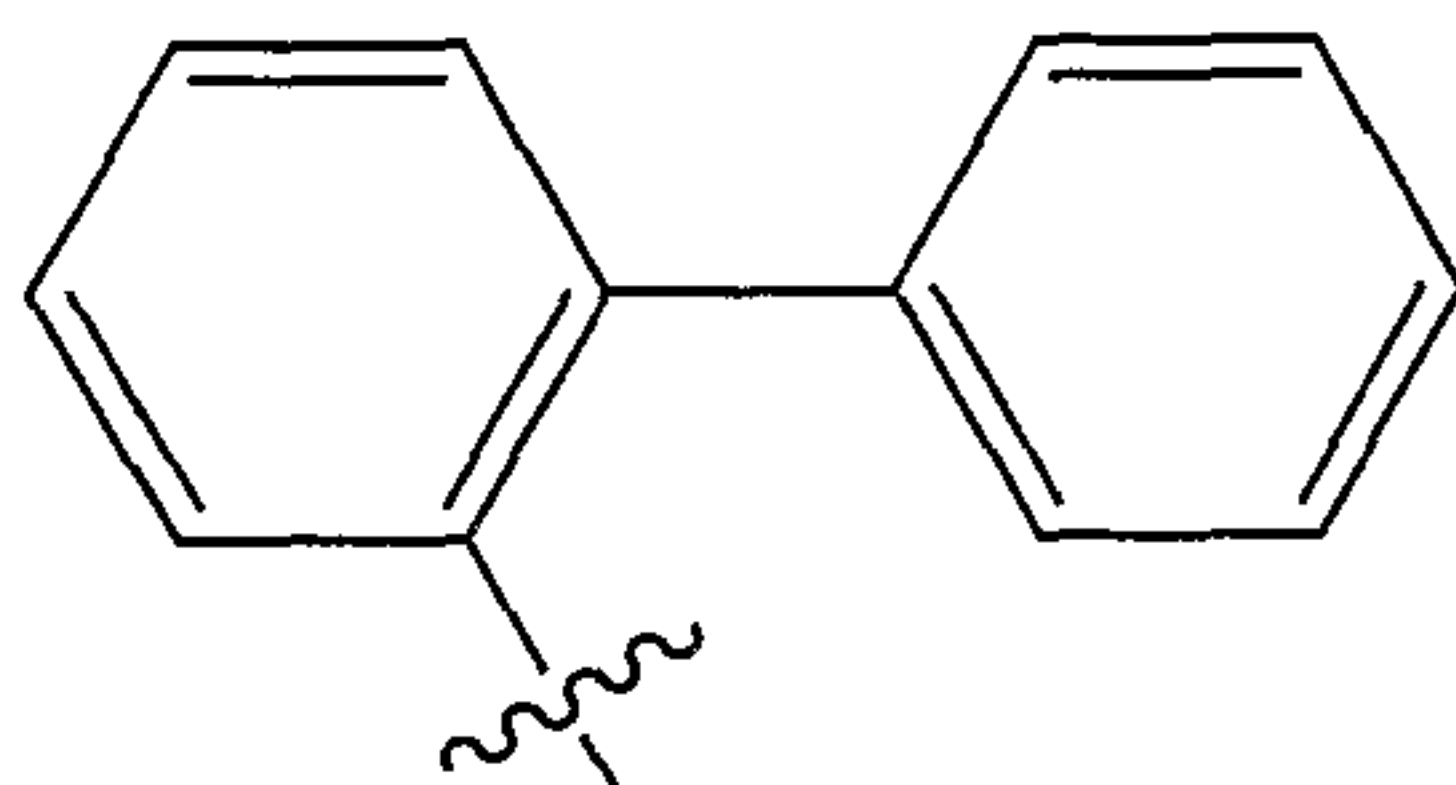
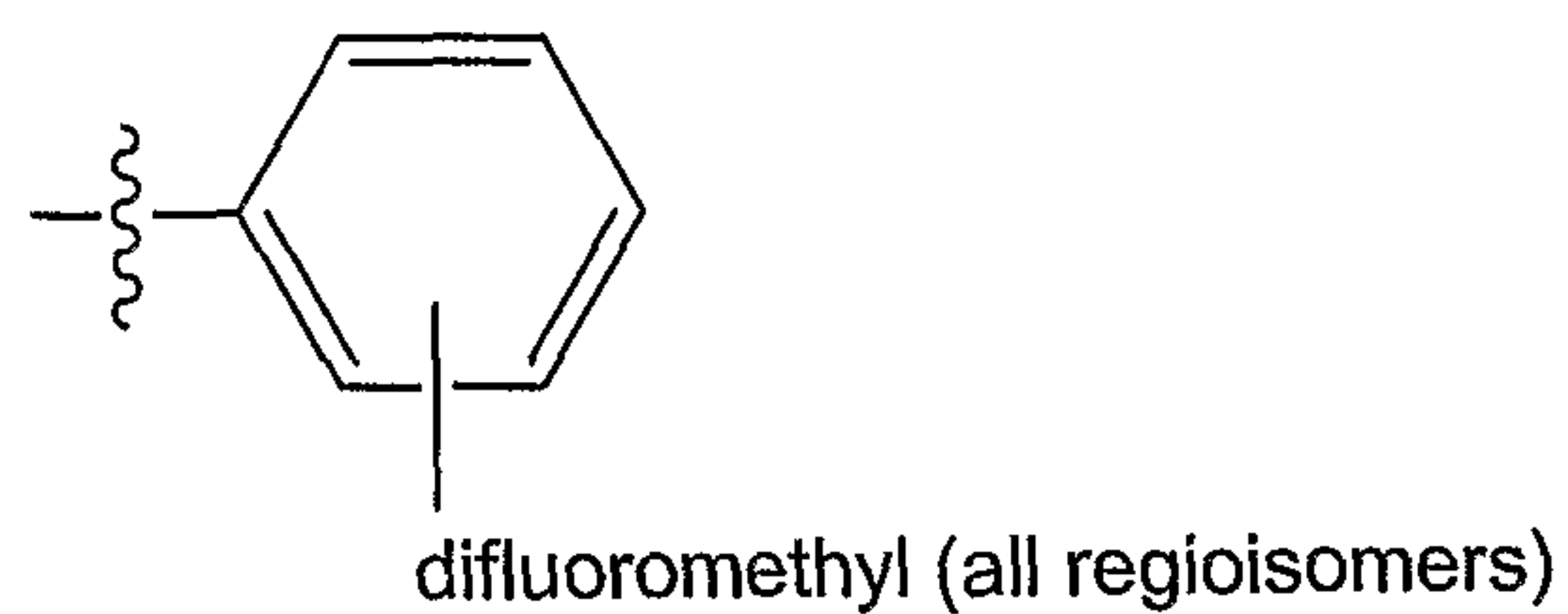
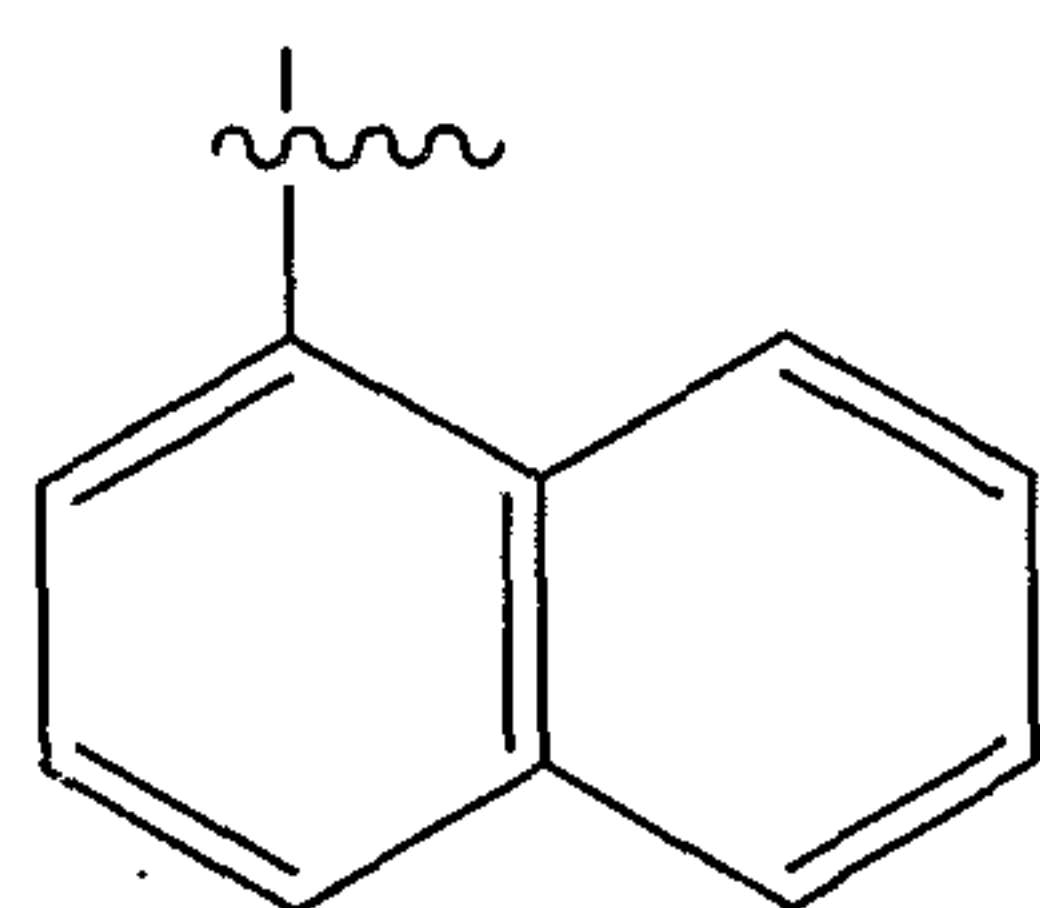
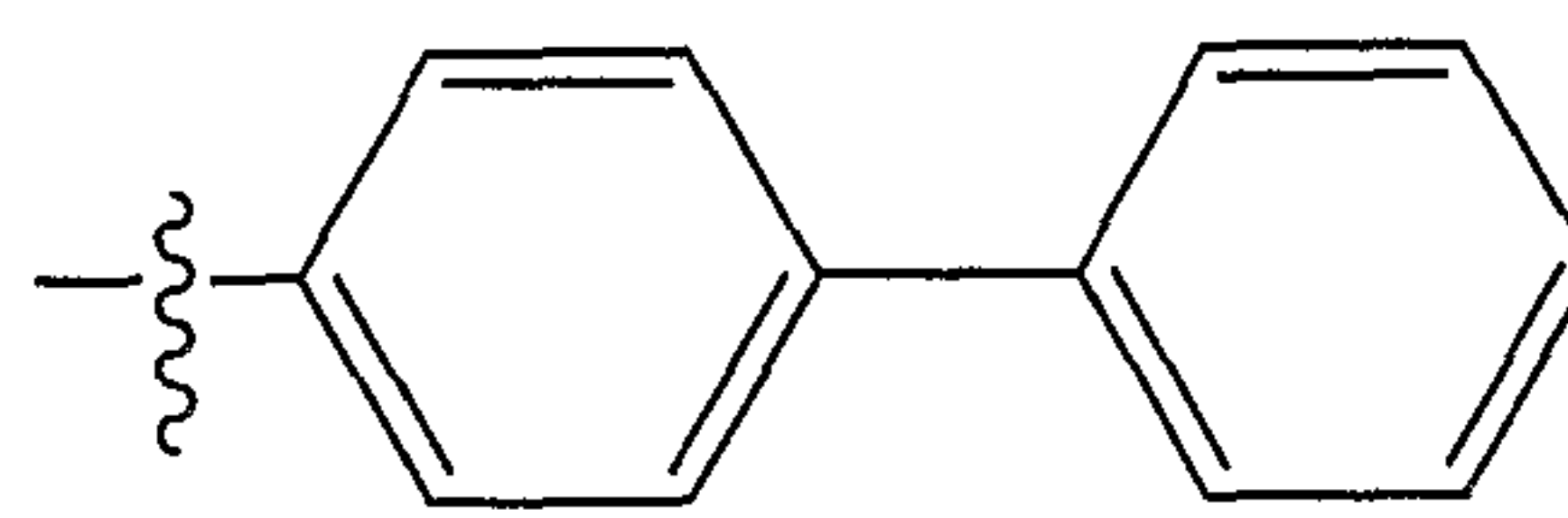
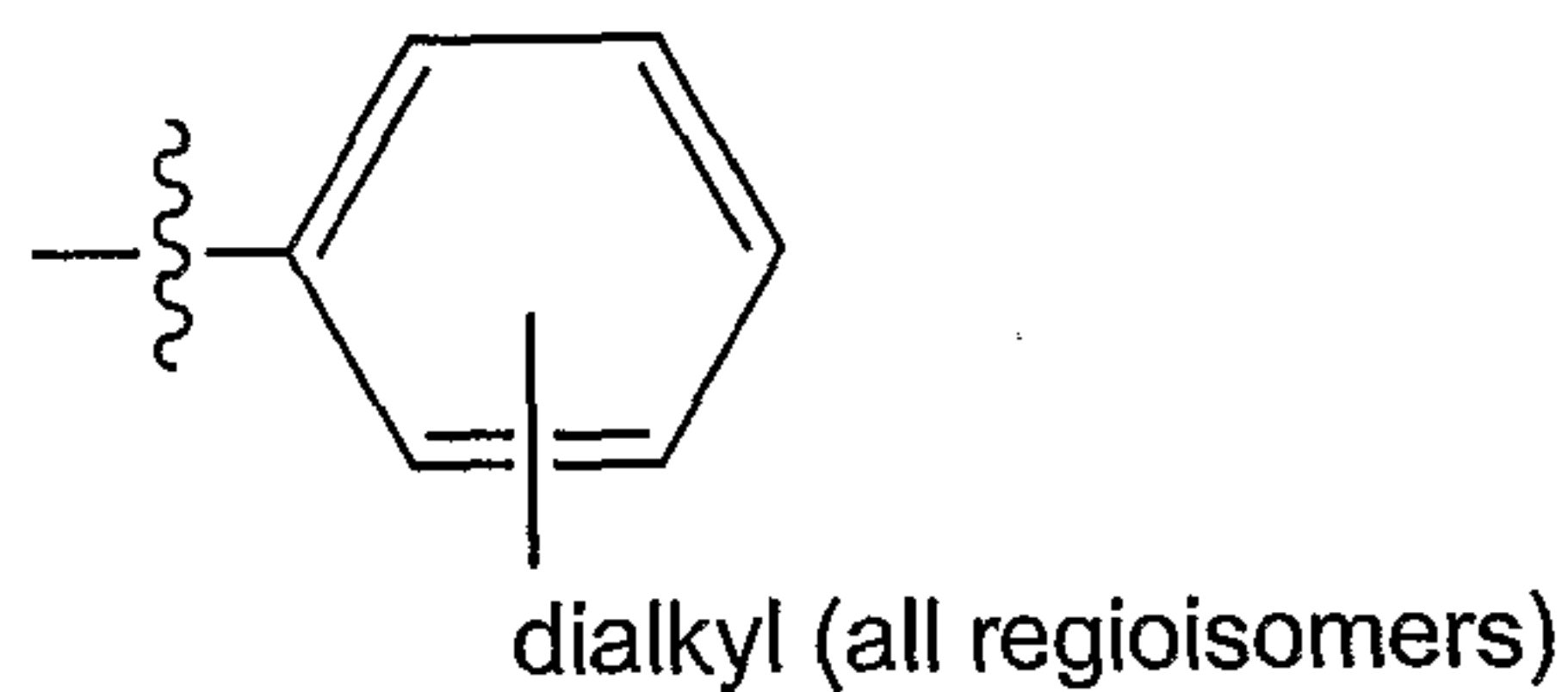
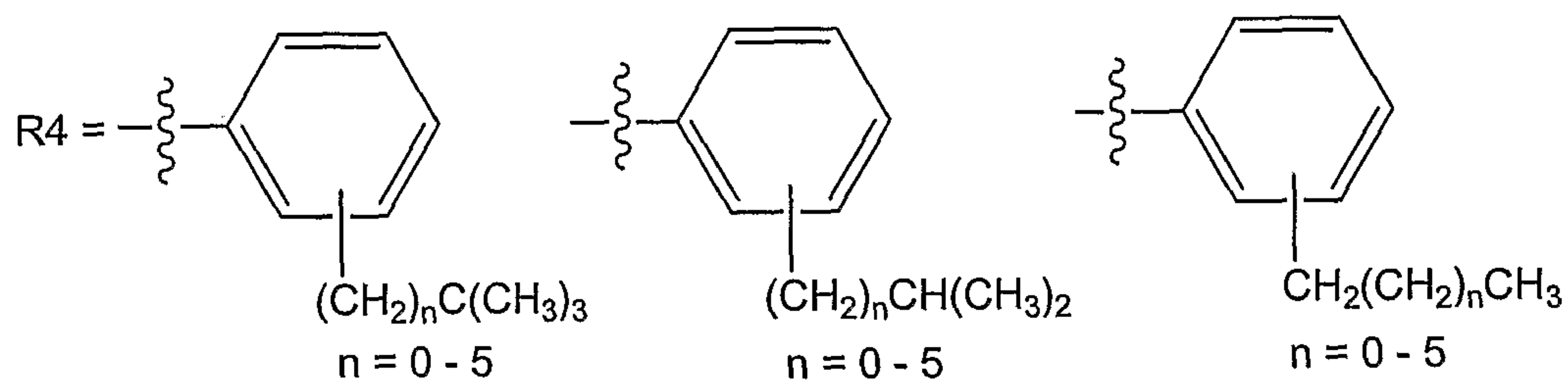


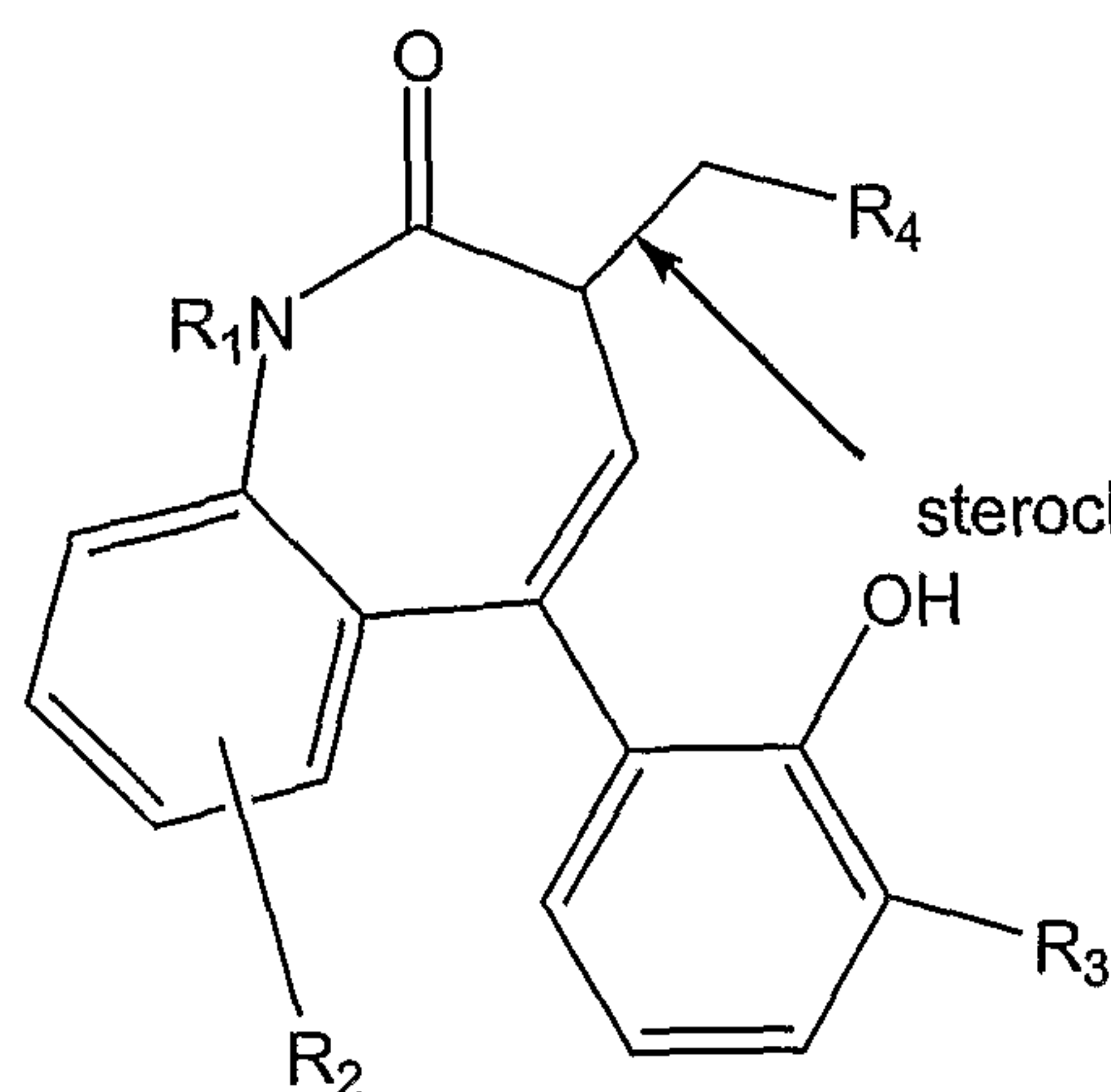
R1 = H, alkyl, or substituted alkyl

R3 = H, alkyl, or substituted alkyl

stereochemistry is R, S, or racemic

R2 is selected from hydrogen, hydroxy, an alkoxy, a halo, an amino, a lower-alkyl, a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic



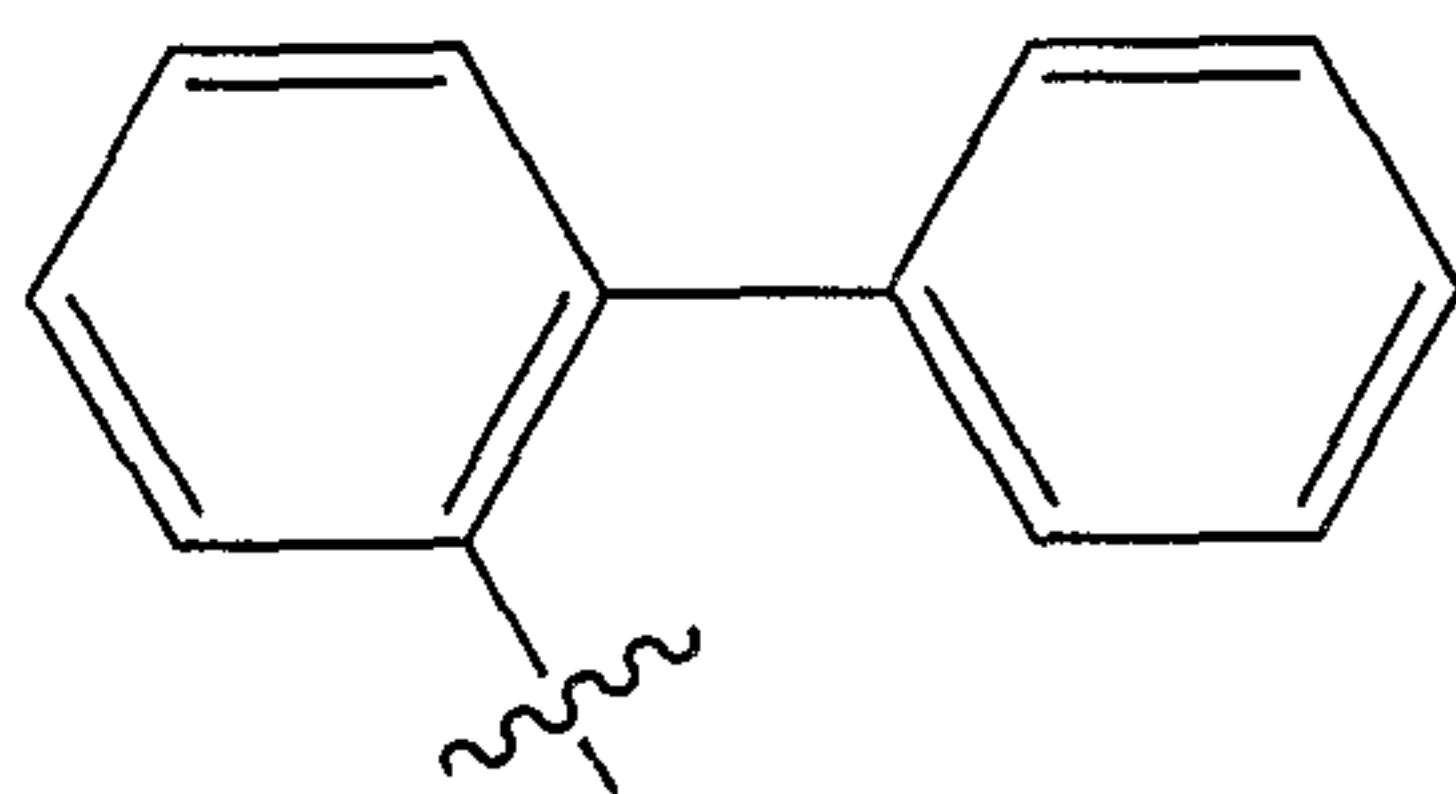
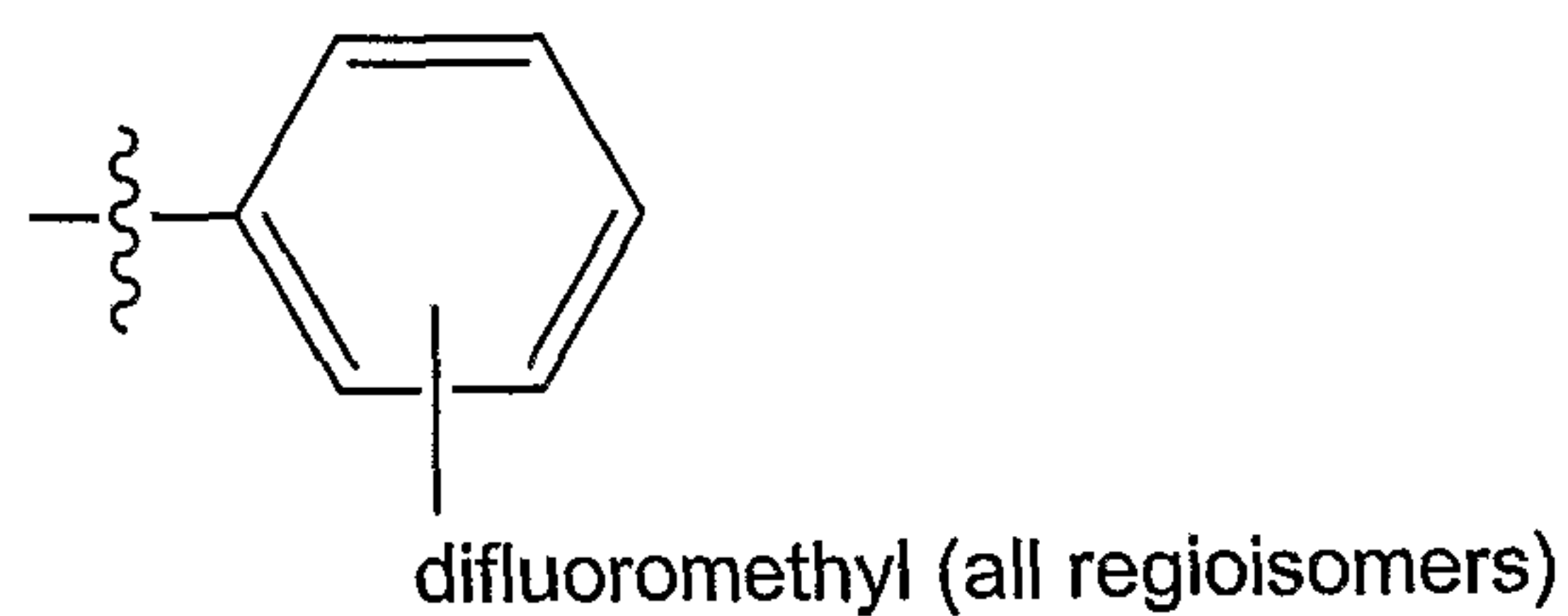
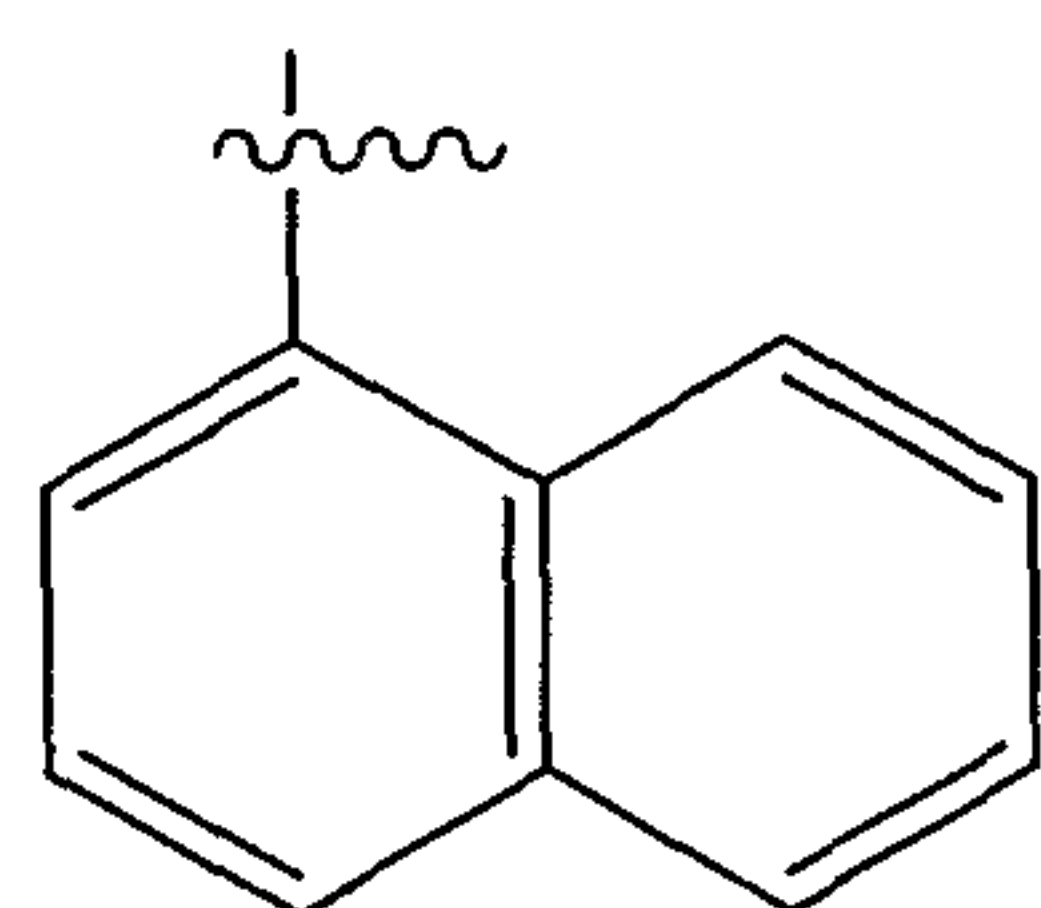
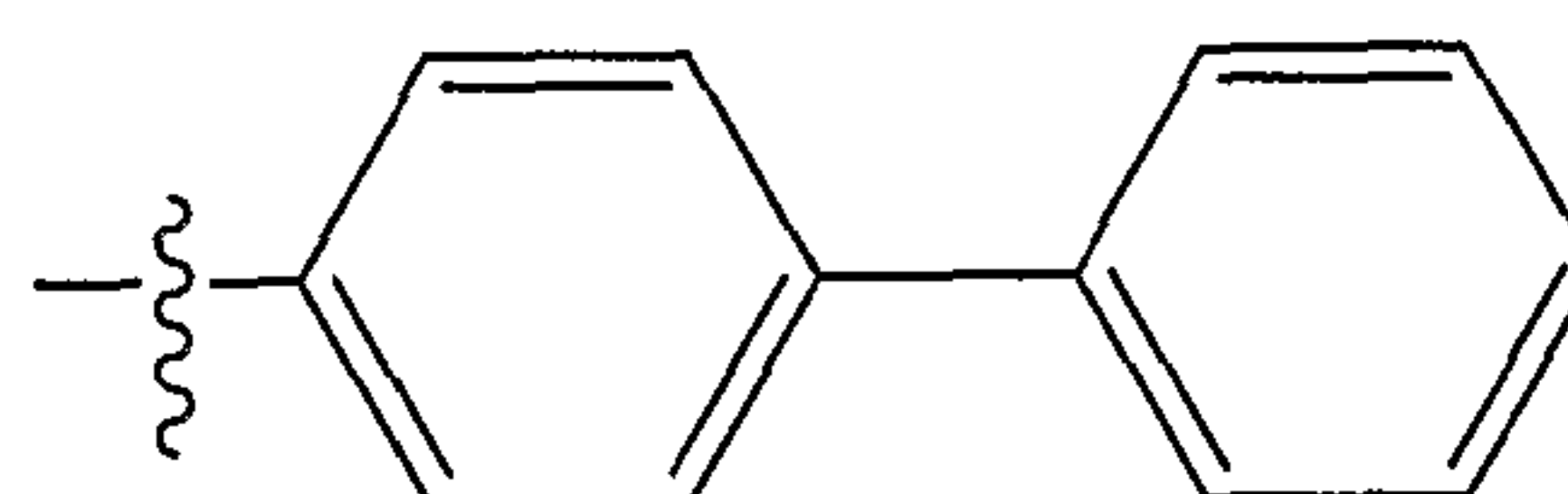
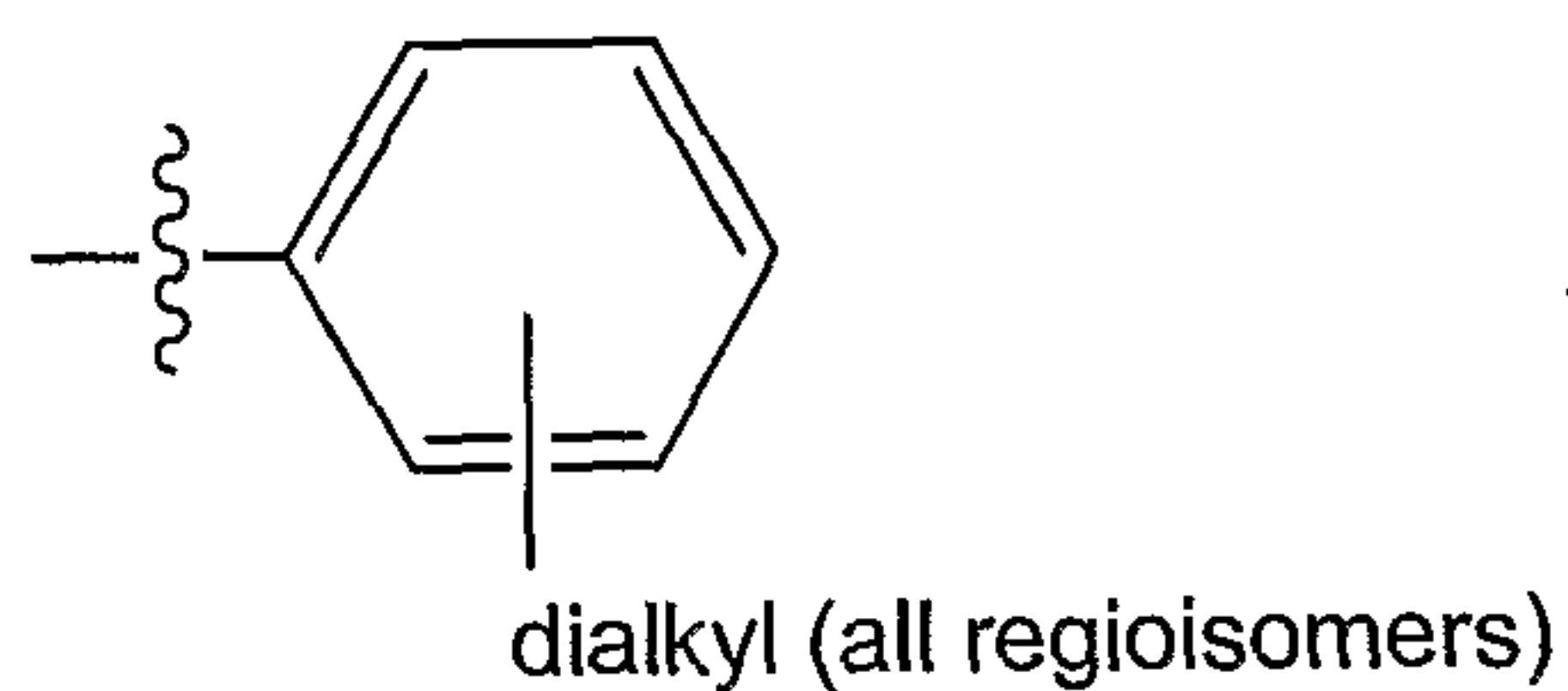
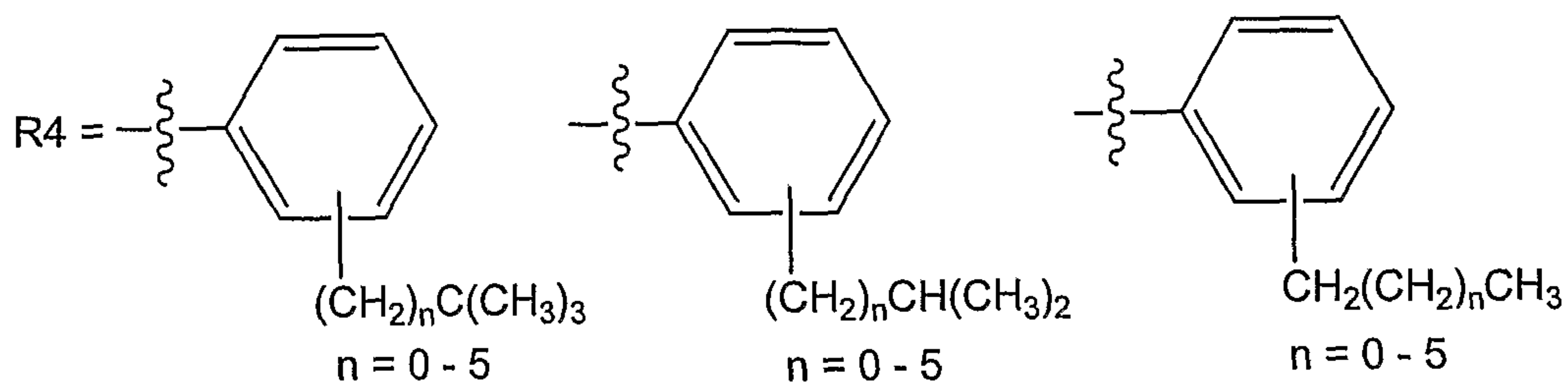


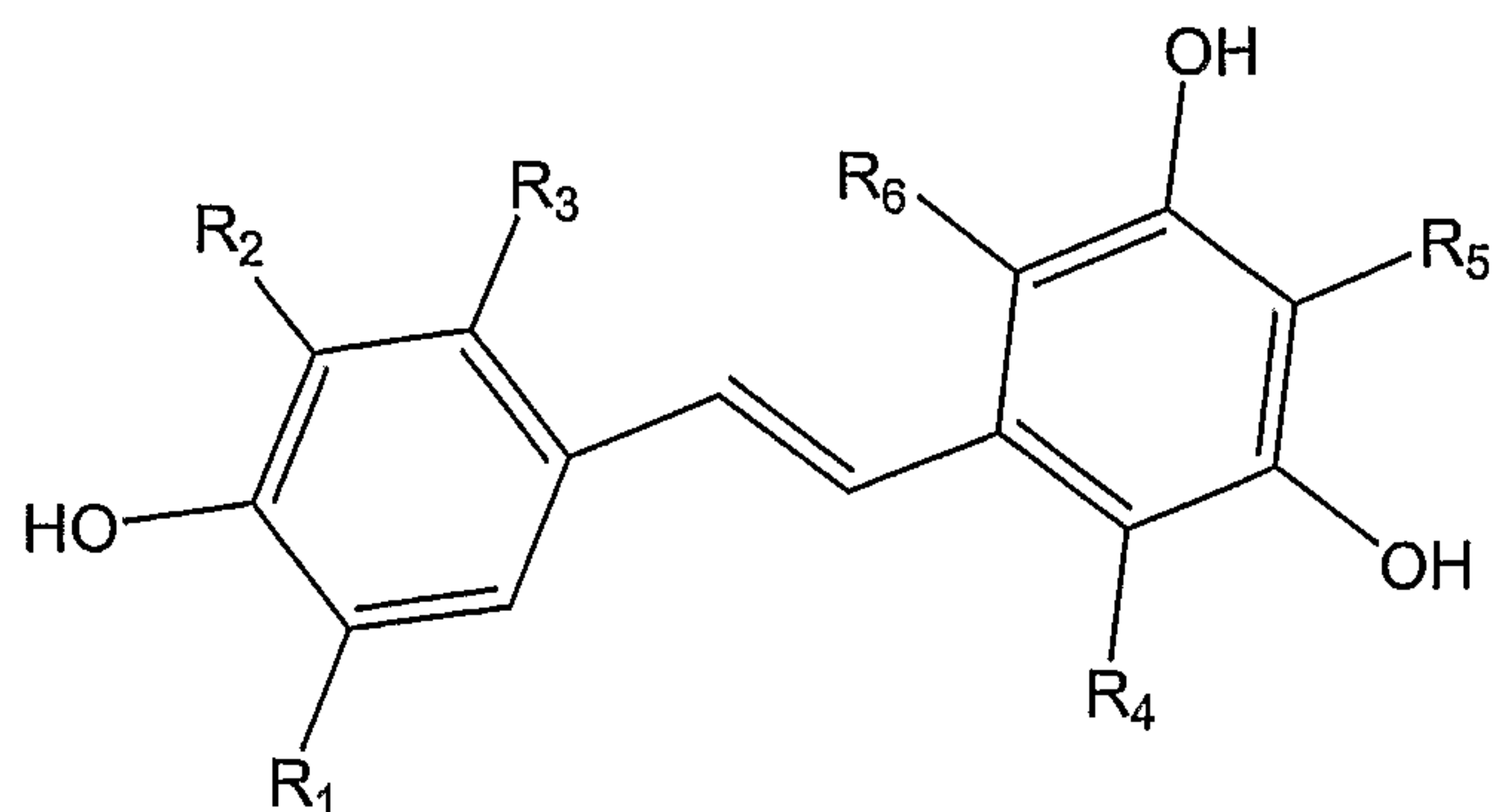
$R_1 = H$, alkyl, or substituted alkyl

$R_3 = H$, alkyl, or substituted alkyl

stereochemistry is R, S, or racemic

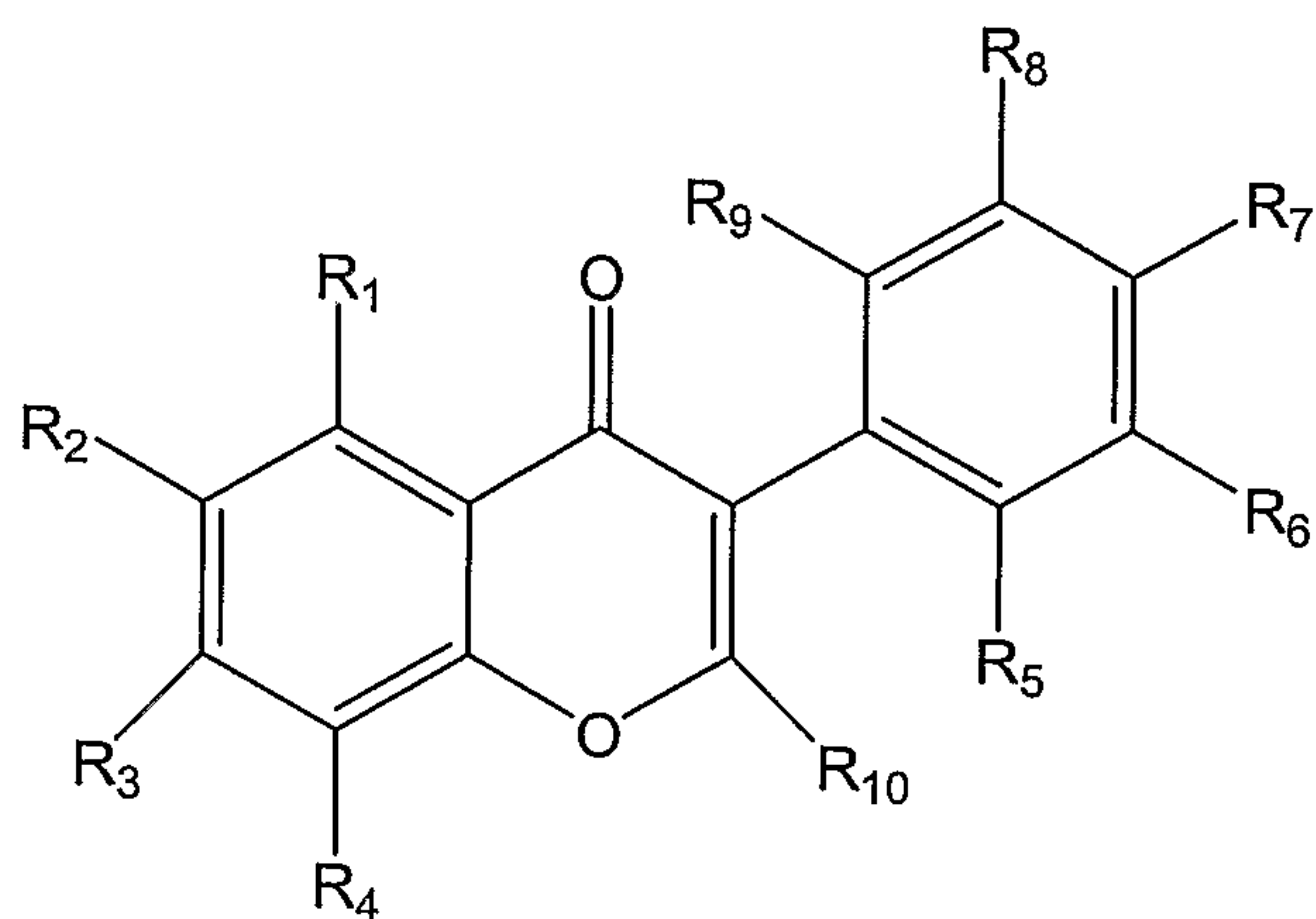
R_2 is selected from hydrogen, hydroxy, an alkoxy, a halo, an amino, a lower-alkyl, a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic



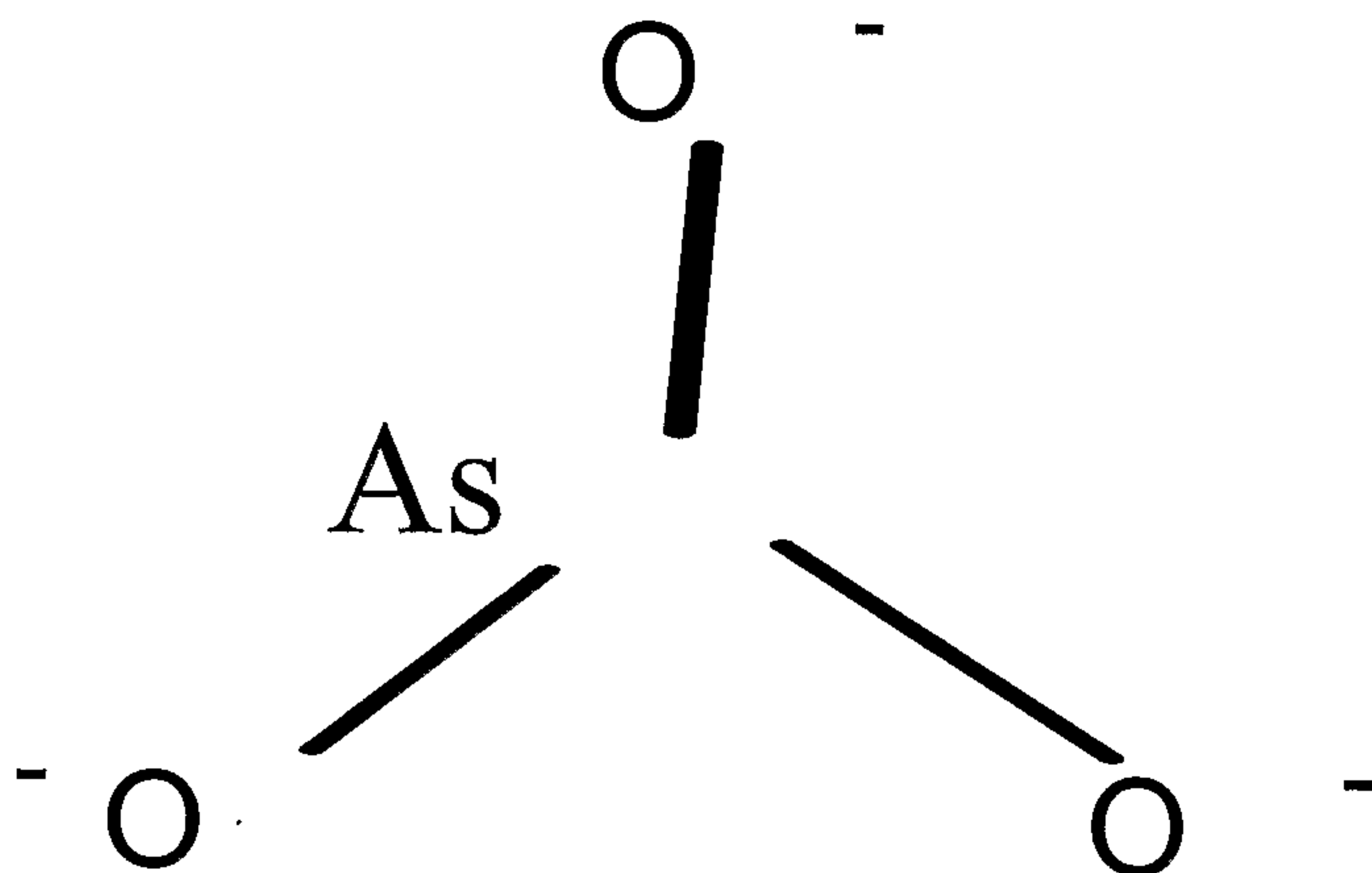


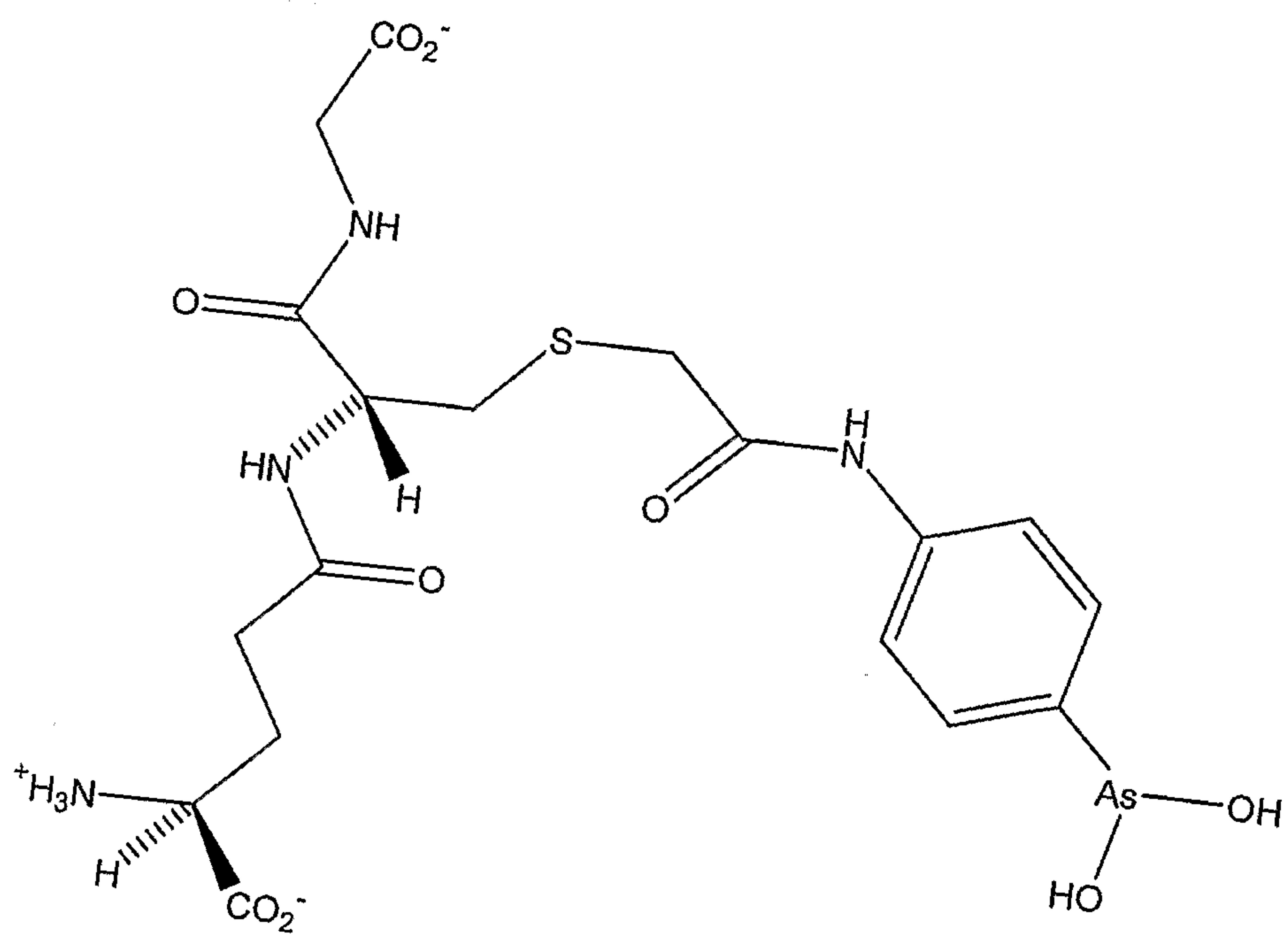
R1 is H or hydroxy

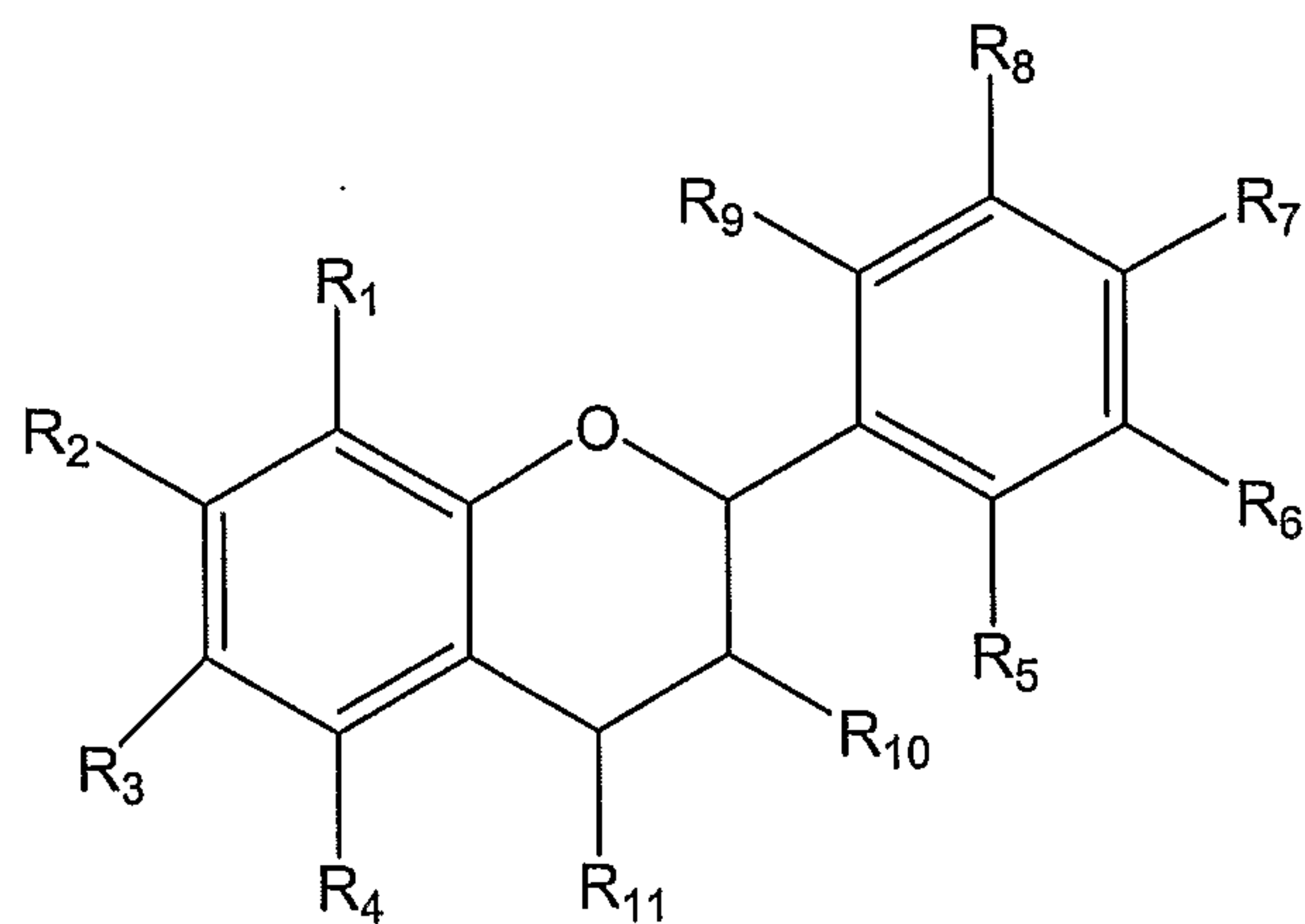
Each of R2 through R6 may be the same or different and is selected from hydrogen, a hydroxy, an alkoxy, a halo, an amino, a lower-alkyl-a substituted-amino, an acetamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic



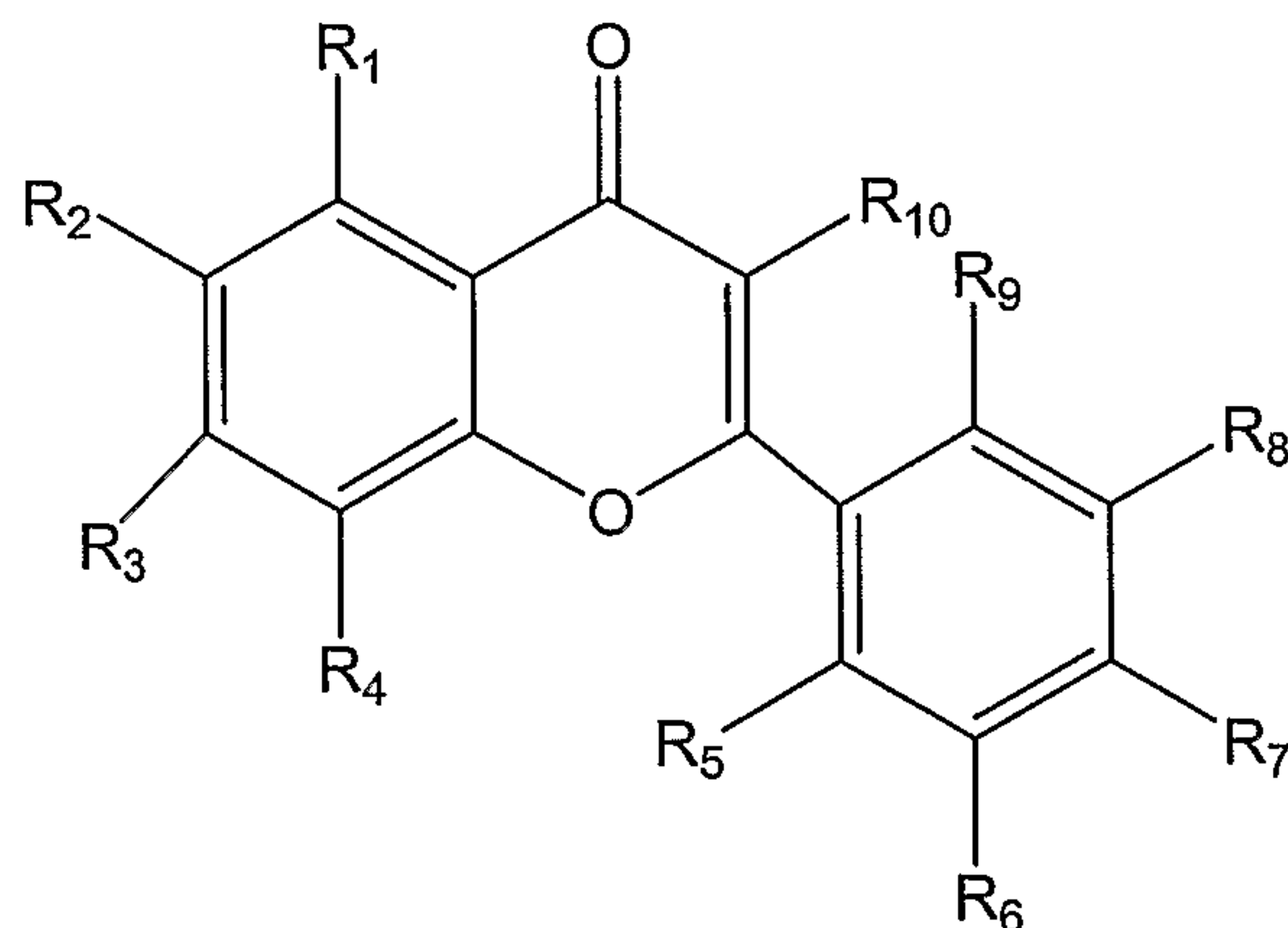
Each of R1 through R10 may be the same or different and is selected from hydrogen, a hydroxy, an alkoxy, a halo, an amino, a lower-alkyl-a substituted-amino, an acetamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic



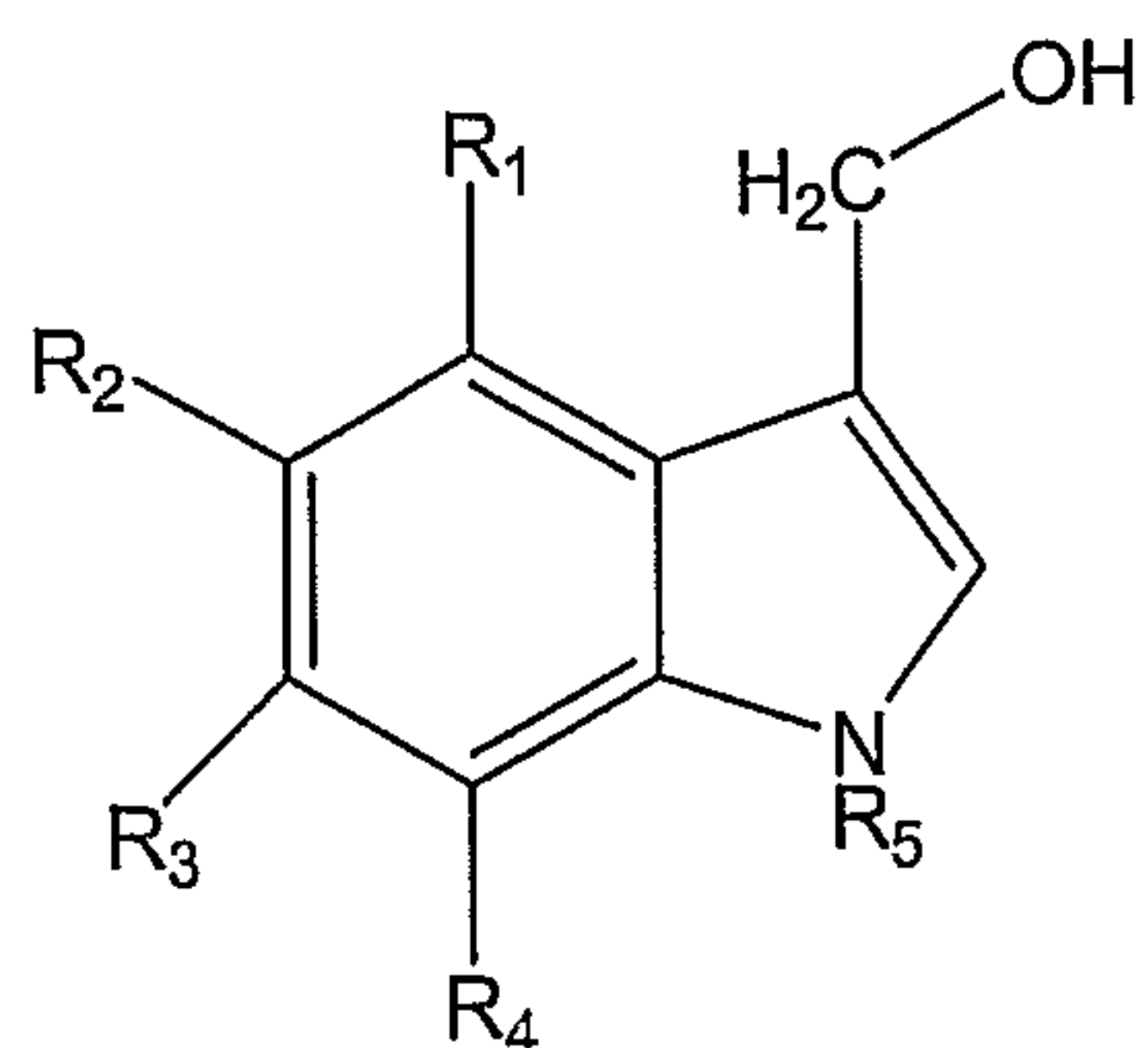




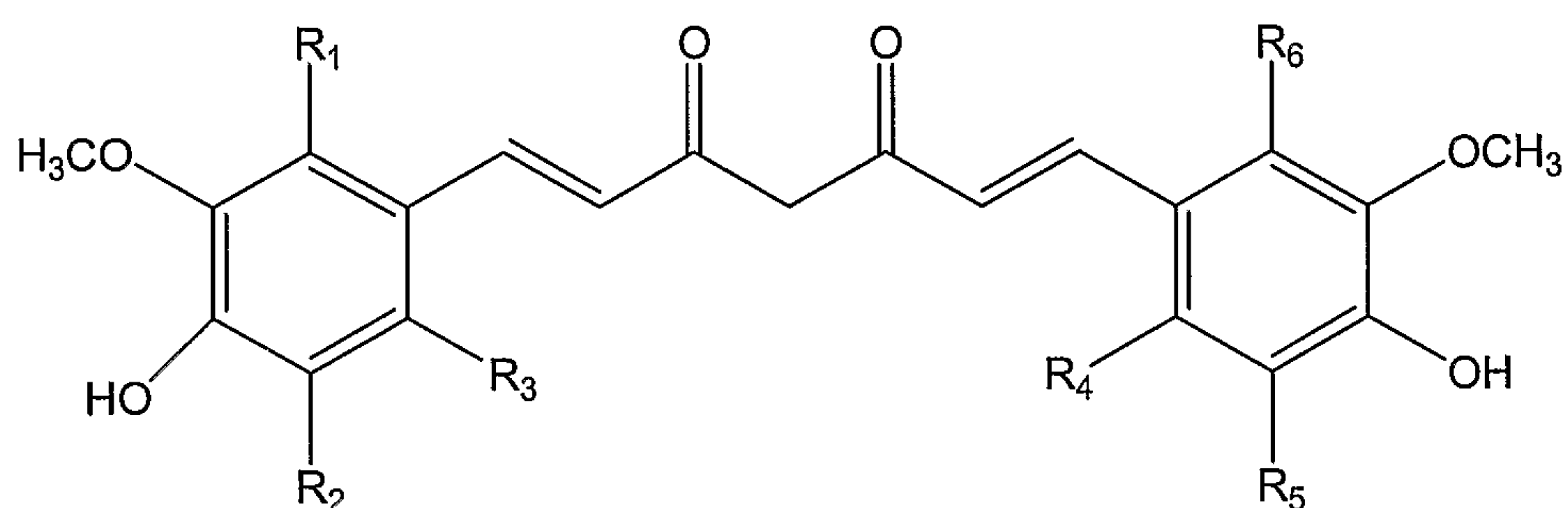
Each of R1 through R11 may be the same or different and is selected from hydrogen, a hydroxy, an alkoxy, a halo, an amino, a lower-alkyl-a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic



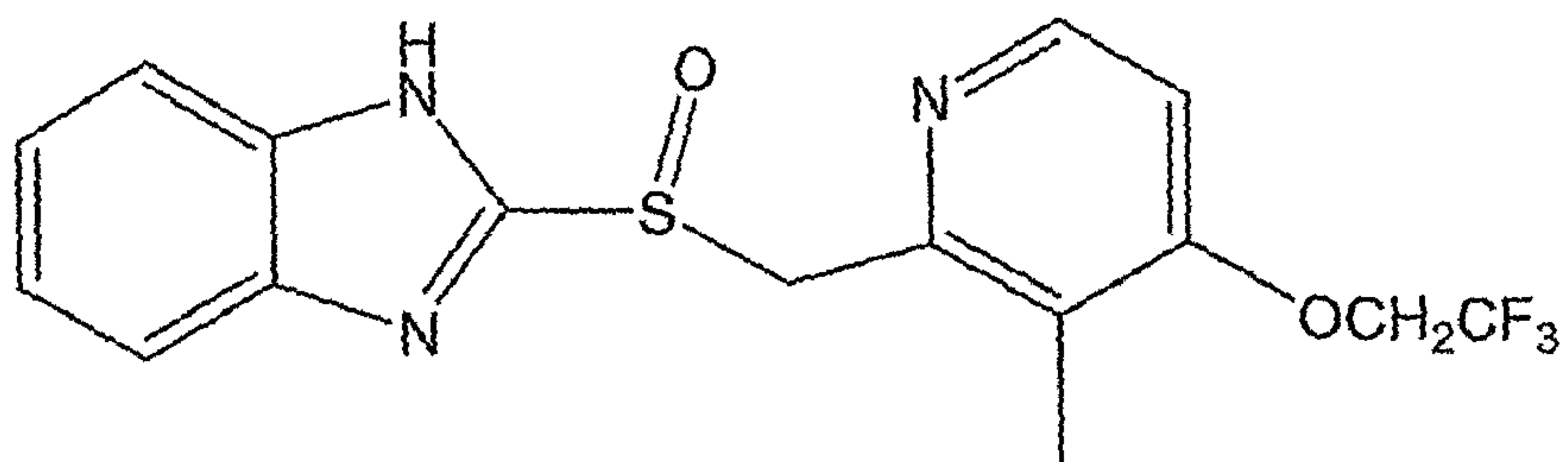
Each of R1 through R10 may be the same or different and is selected from hydrogen, a hydroxy, an alkoxy, a halo, an amino, a lower-alkyl-a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic



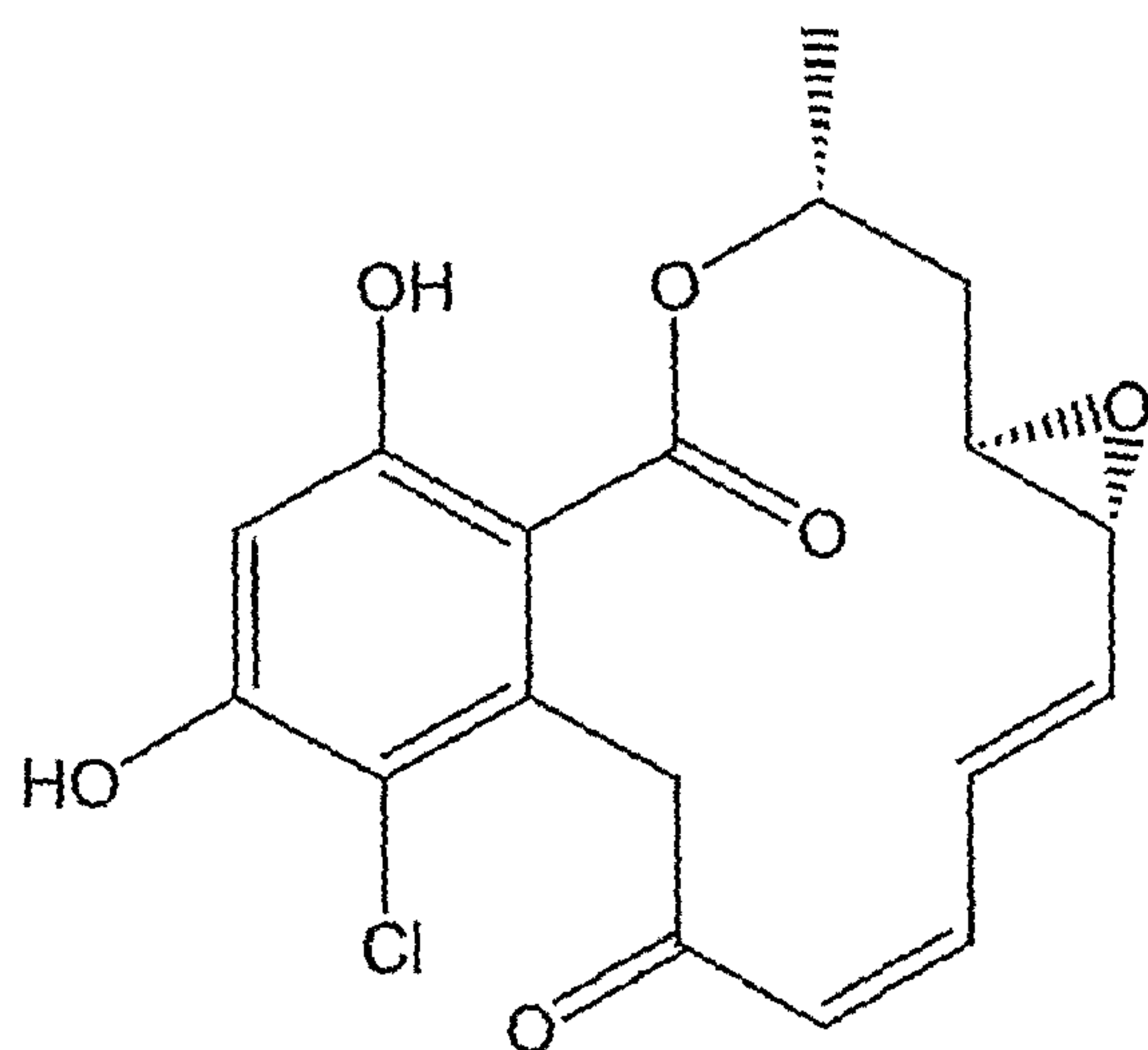
Each of R1 through R10 may be the same or different and is selected from hydrogen, a hydroxy, an alkoxy, a halo, an amino, a lower-alkyl-a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic



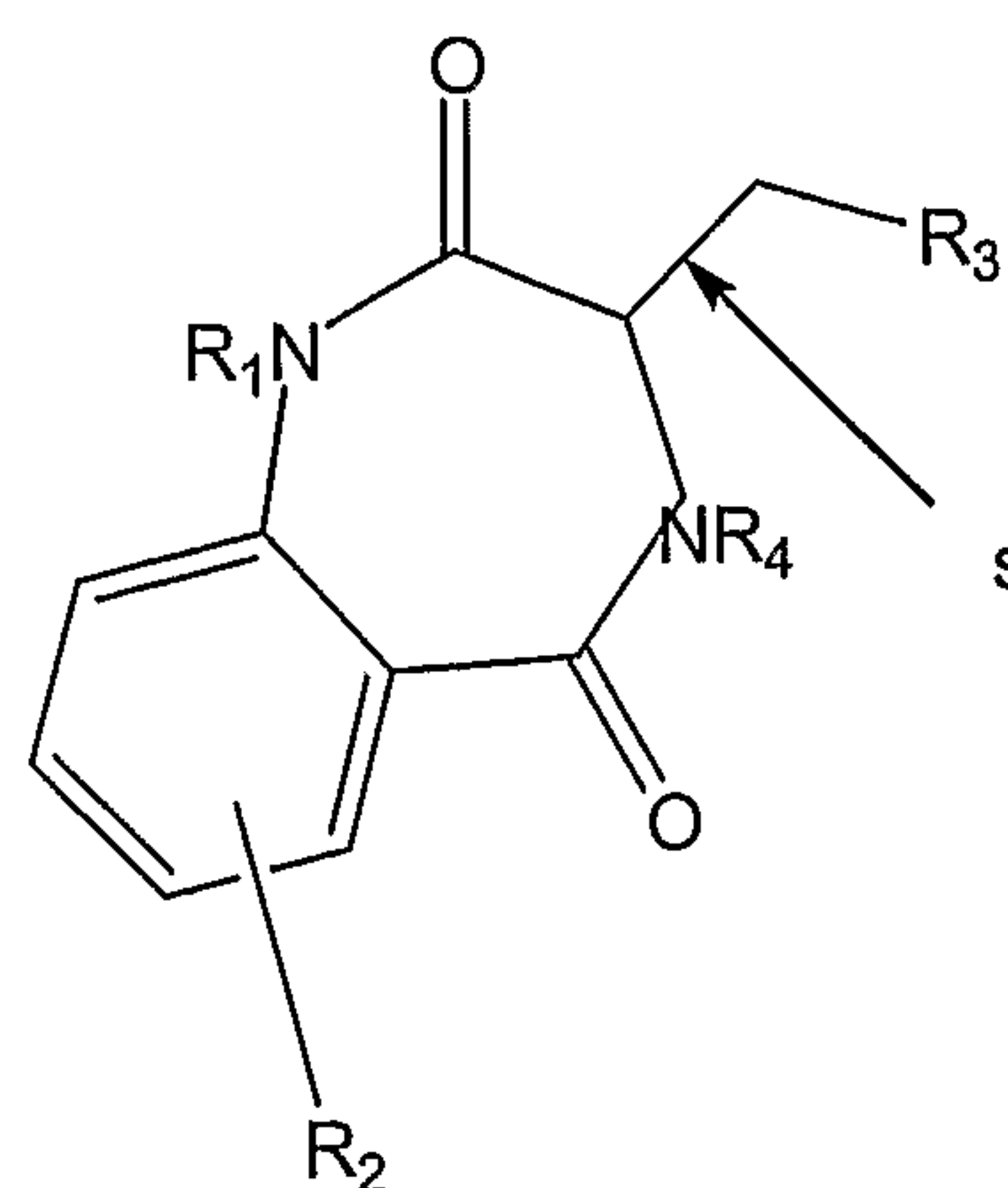
Each of R1 through R6 may be the same or different and is selected from hydrogen, a hydroxy, an alkoxy, a halo, an amino, a lower-alkyl-a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic



lansoprazole



radicicol

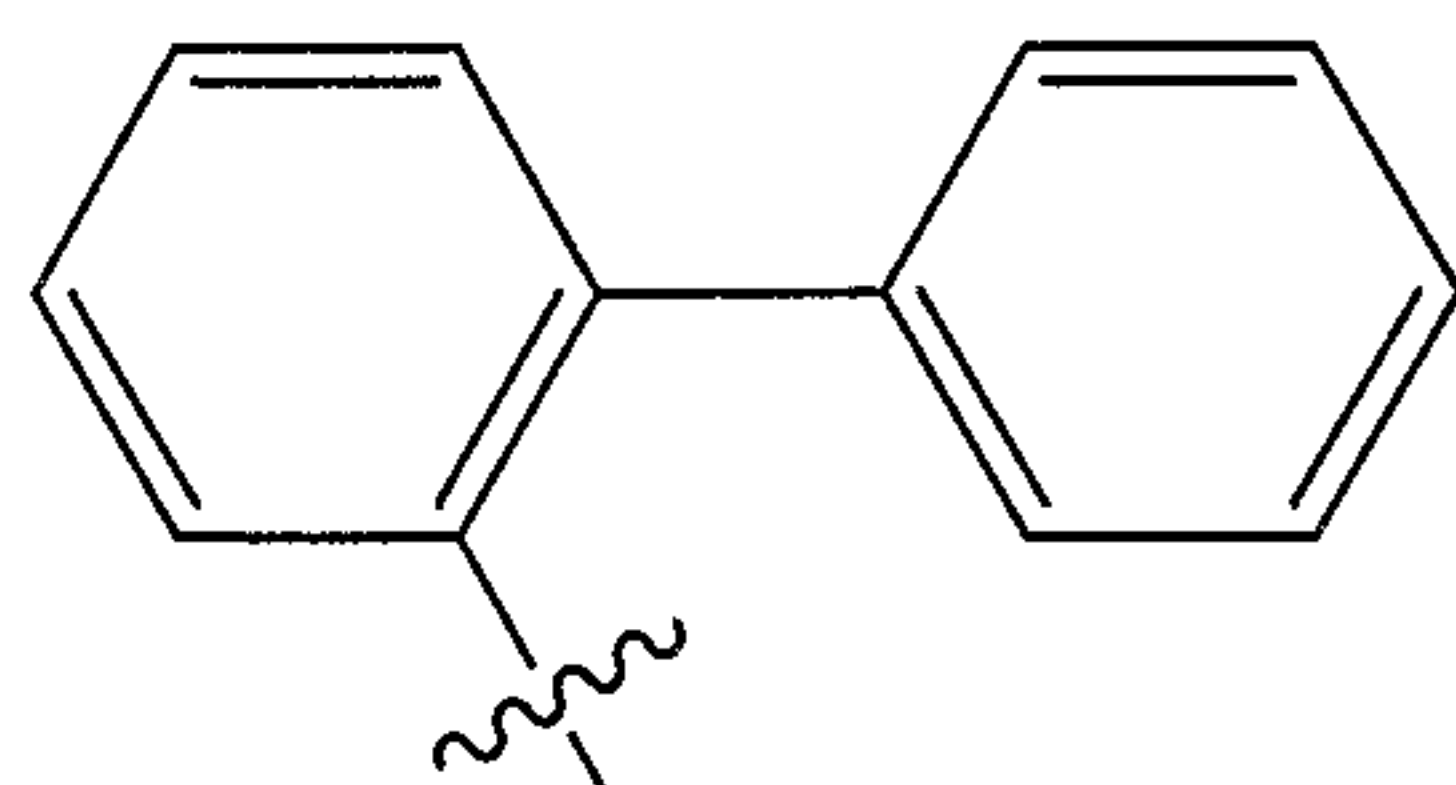
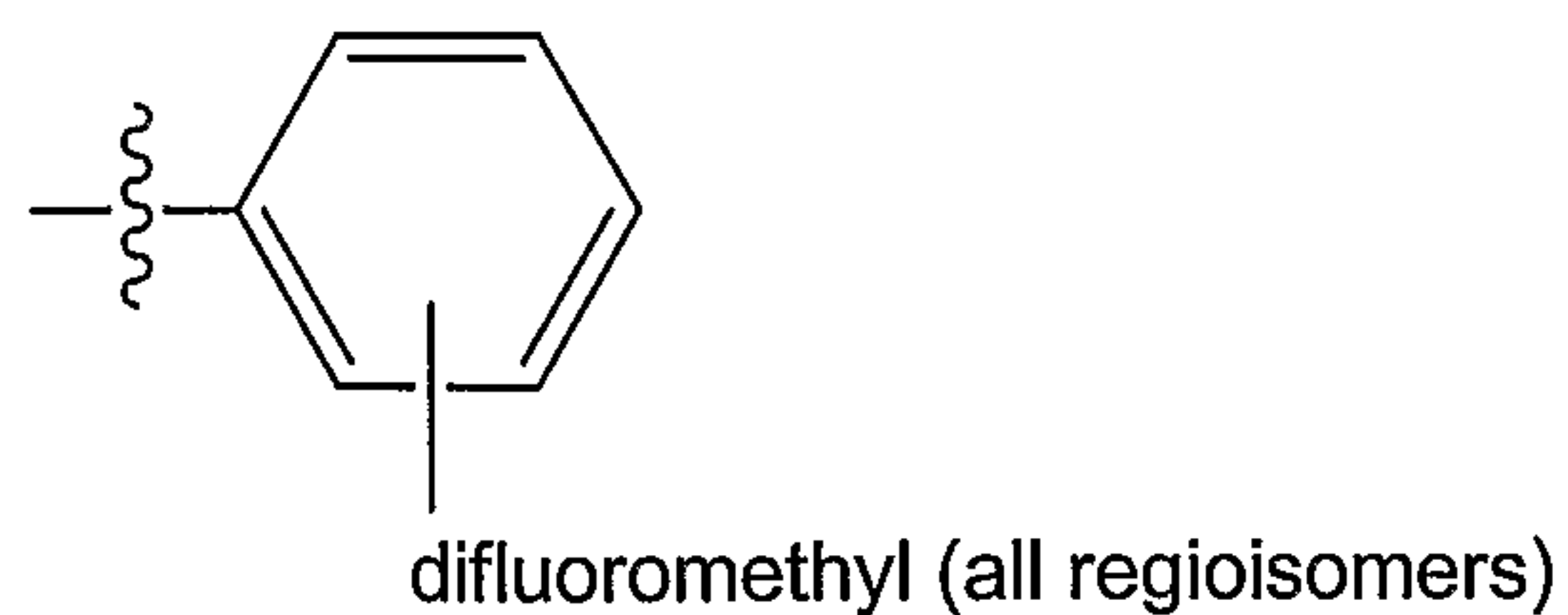
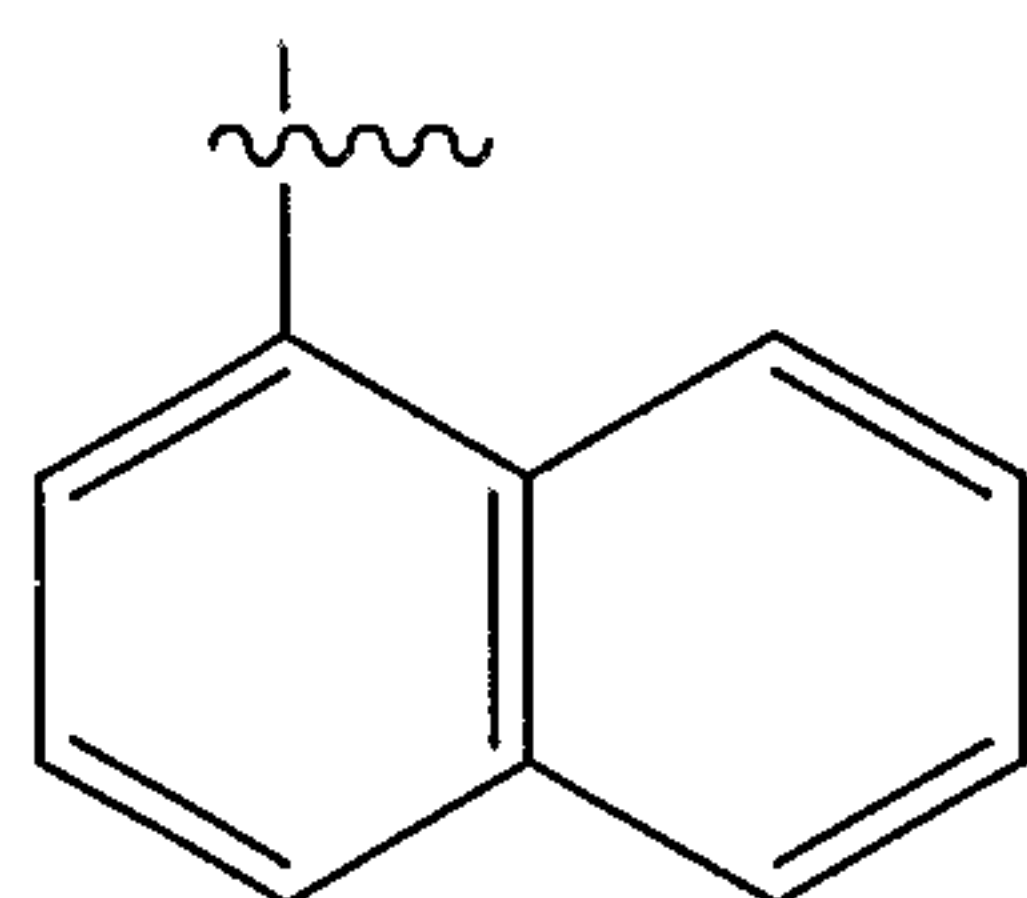
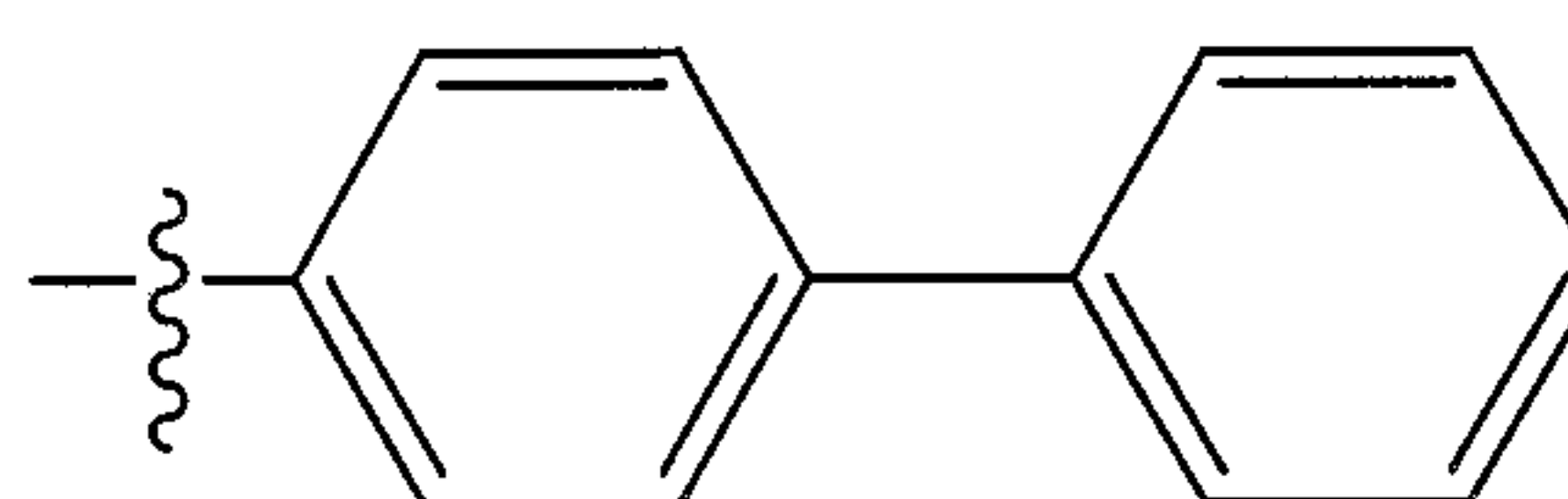
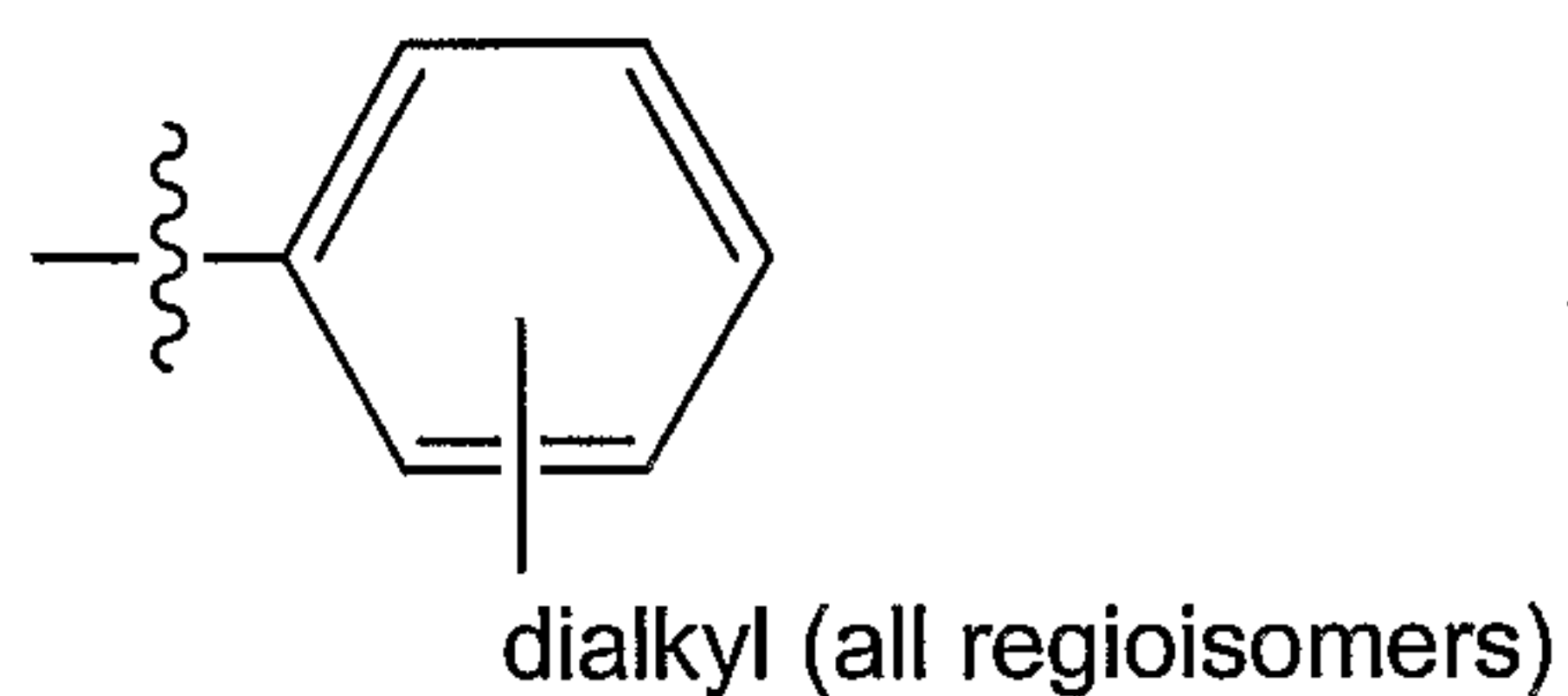
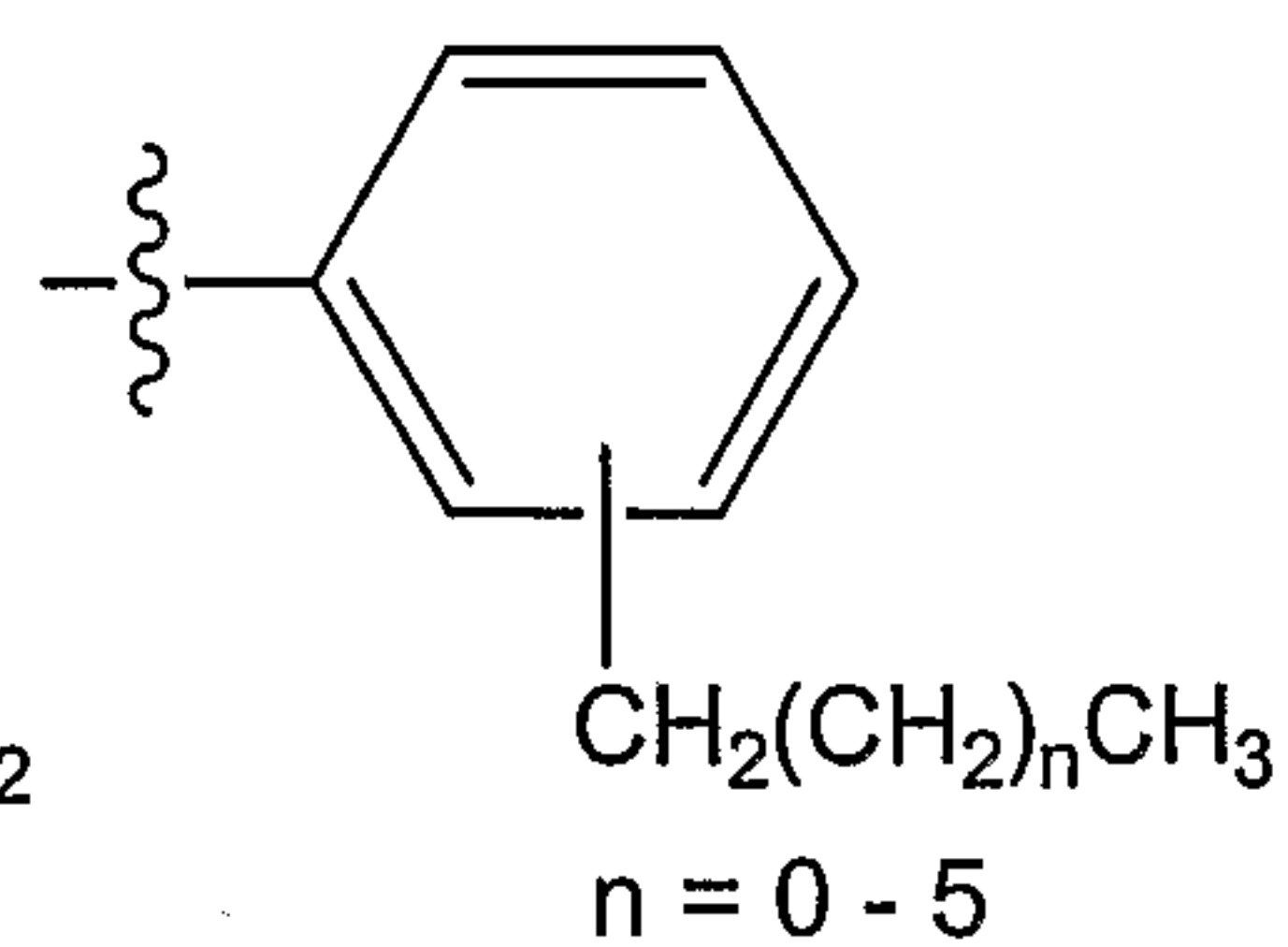
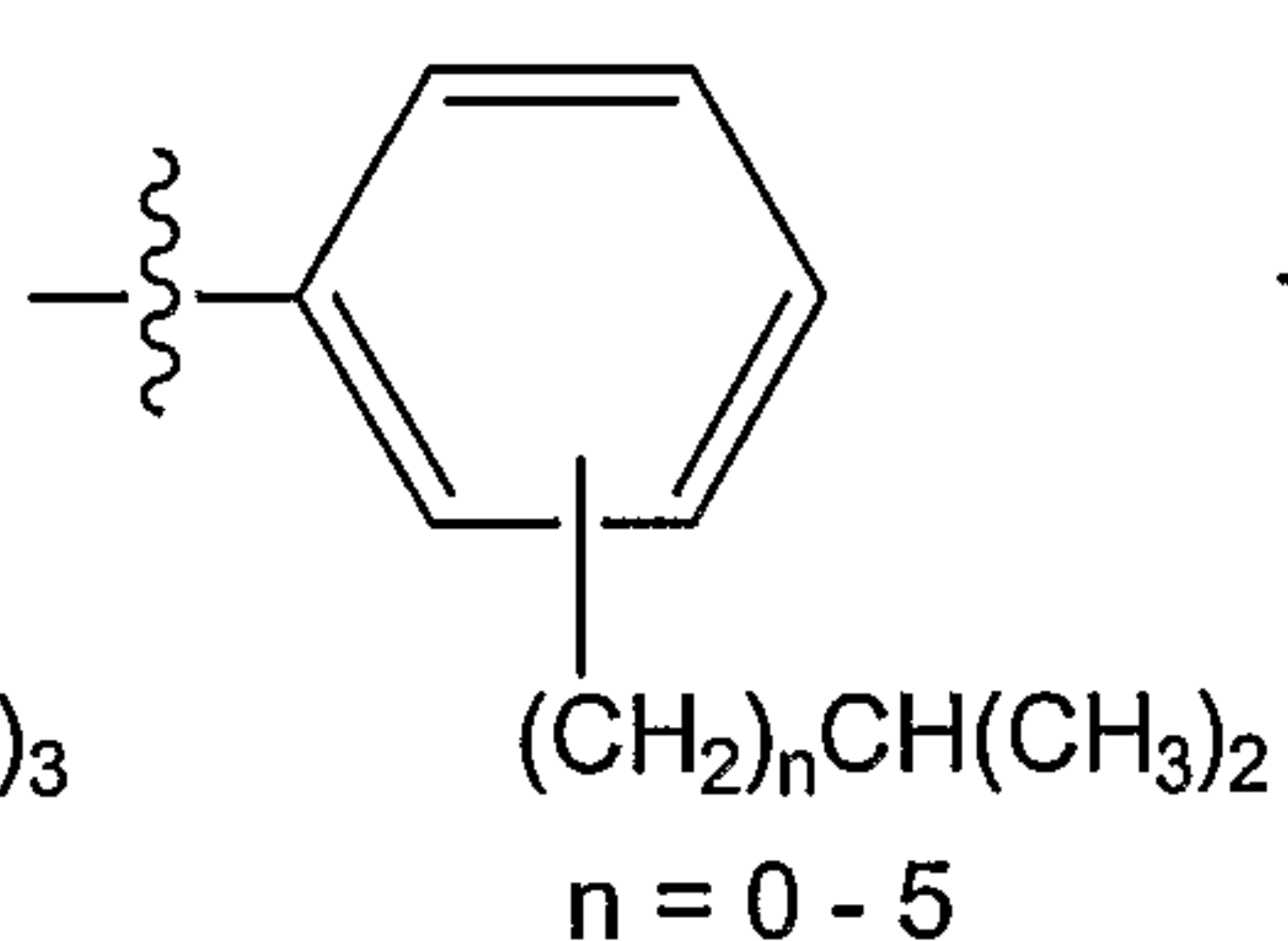
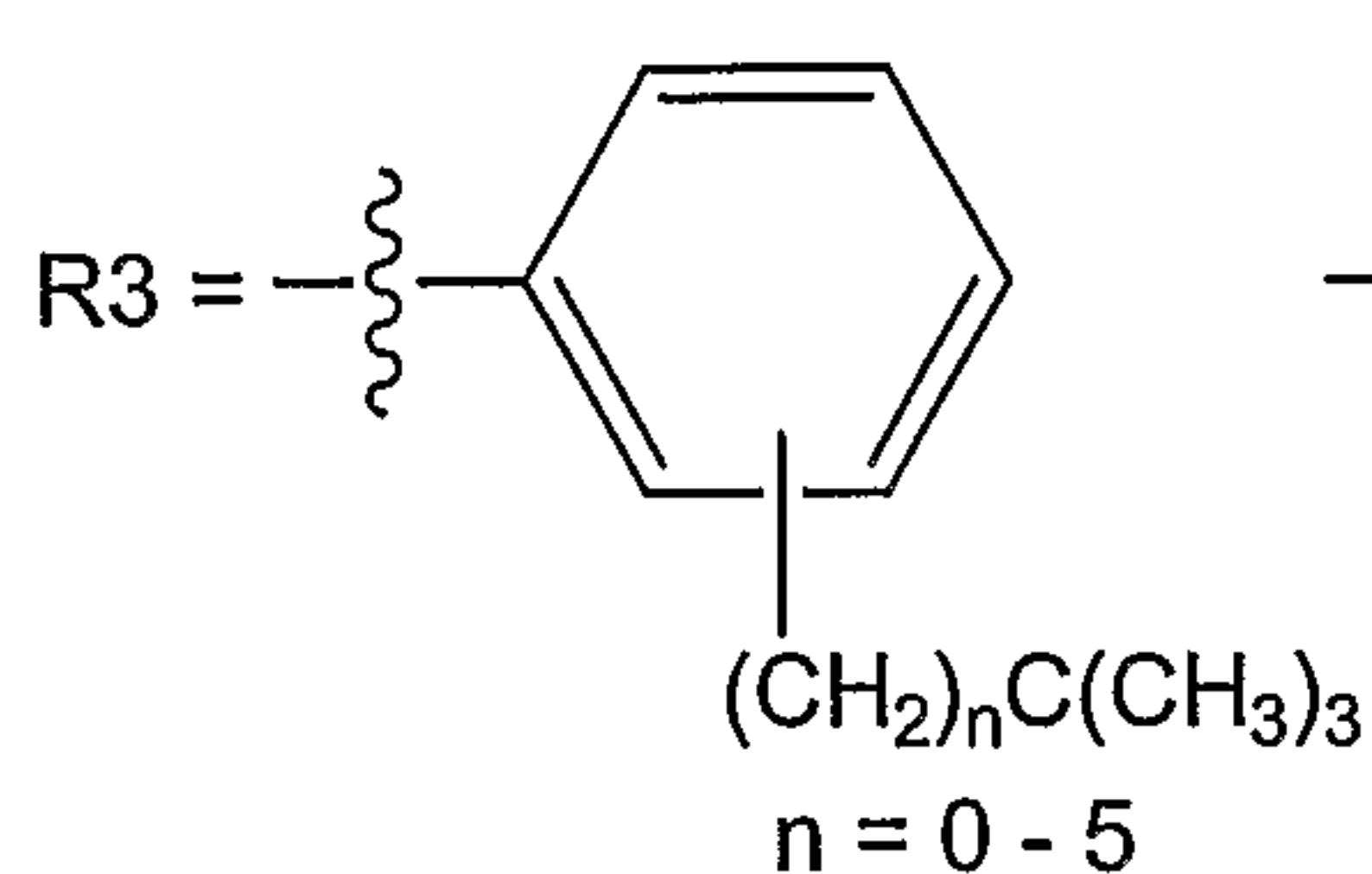


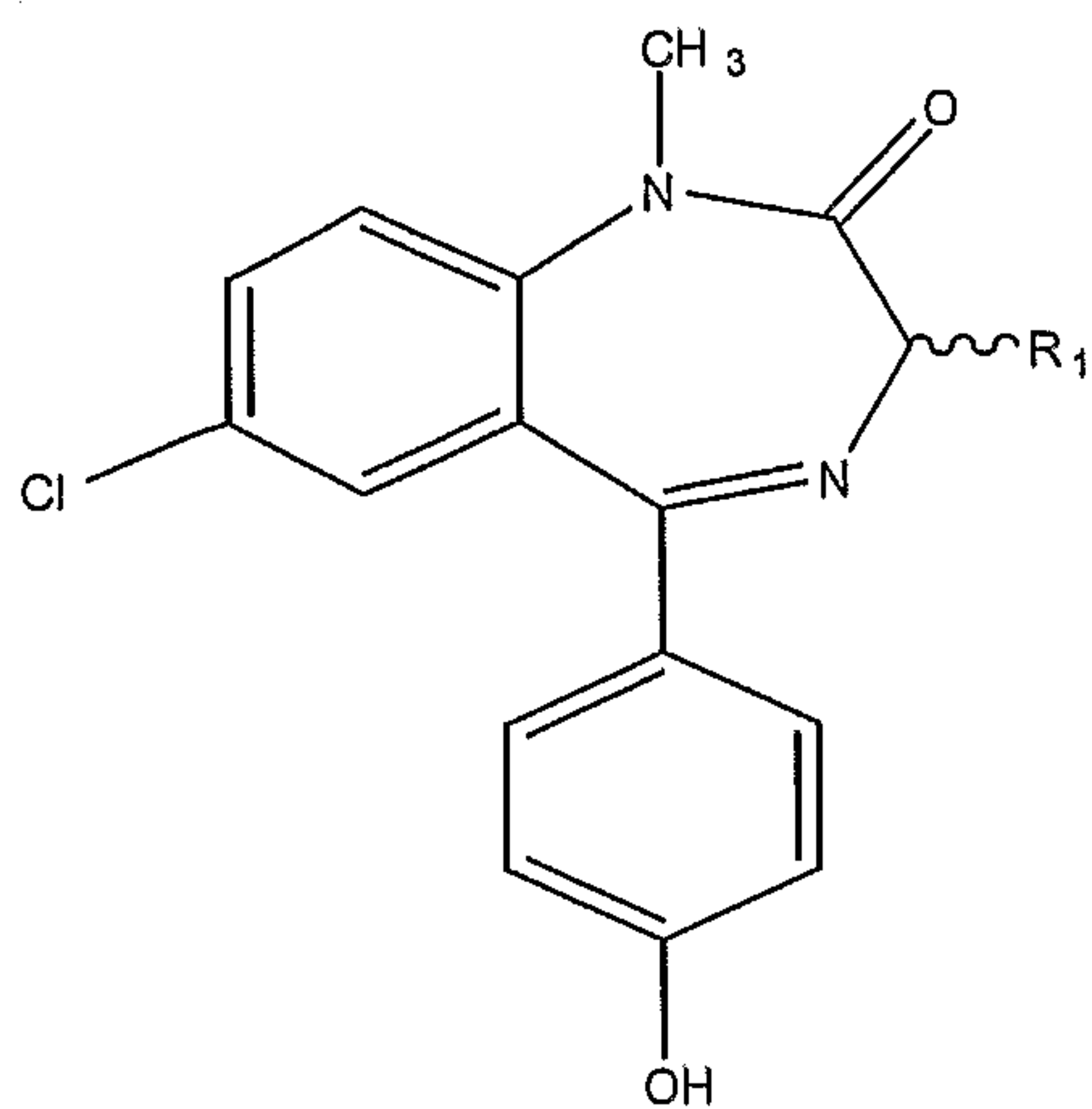
R1 = H, alkyl, or substituted alkyl

R4 = H, alkyl, or substituted alkyl

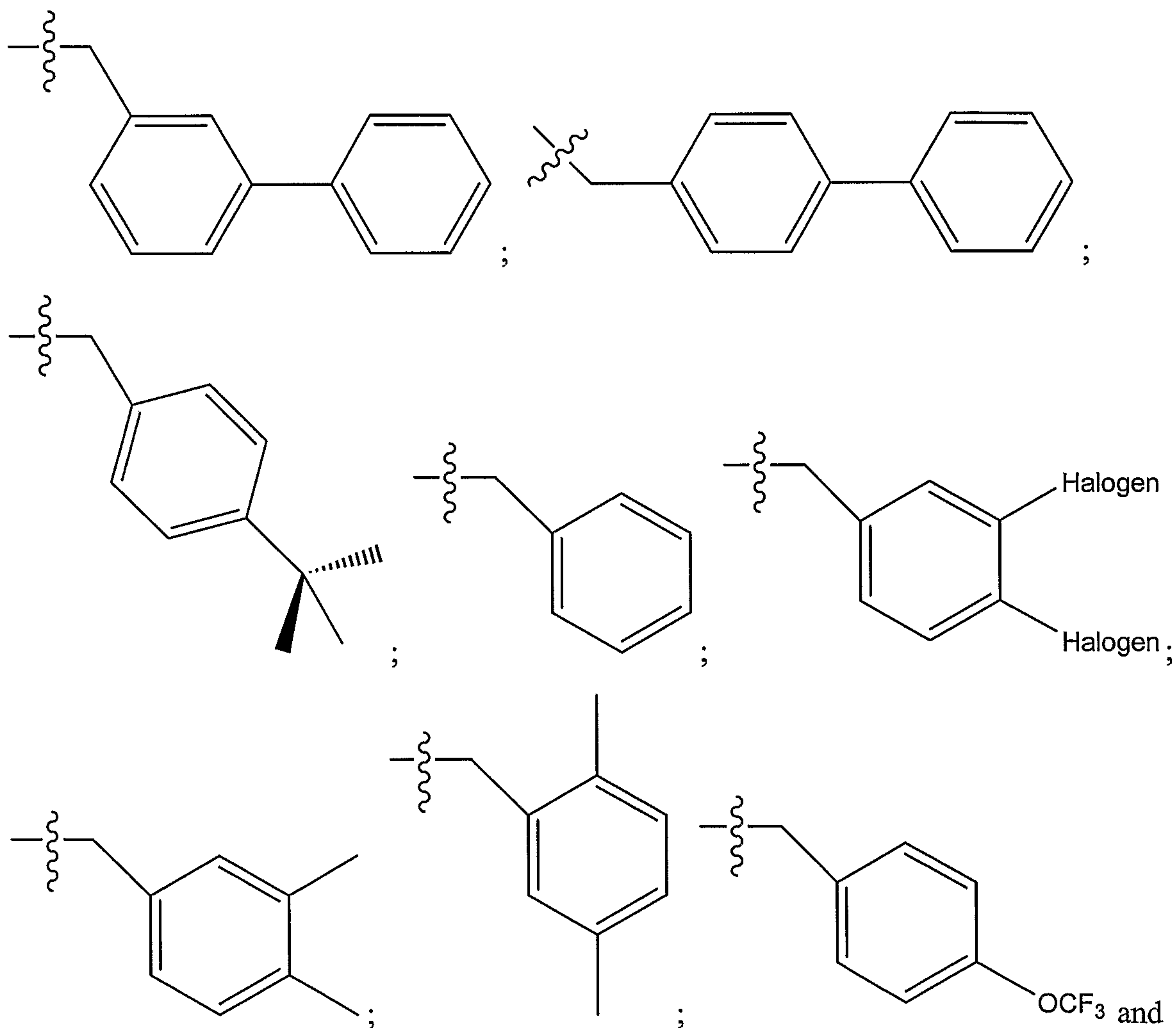
stereochemistry is R, S, or racemic

R2 is selected from hydrogen, a hydroxy, an alkoxy, a halo, an amino, a lower-alkyl-a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of < 10 carbons, a substituted cycloaliphatic group, an aryl, and a heterocyclic



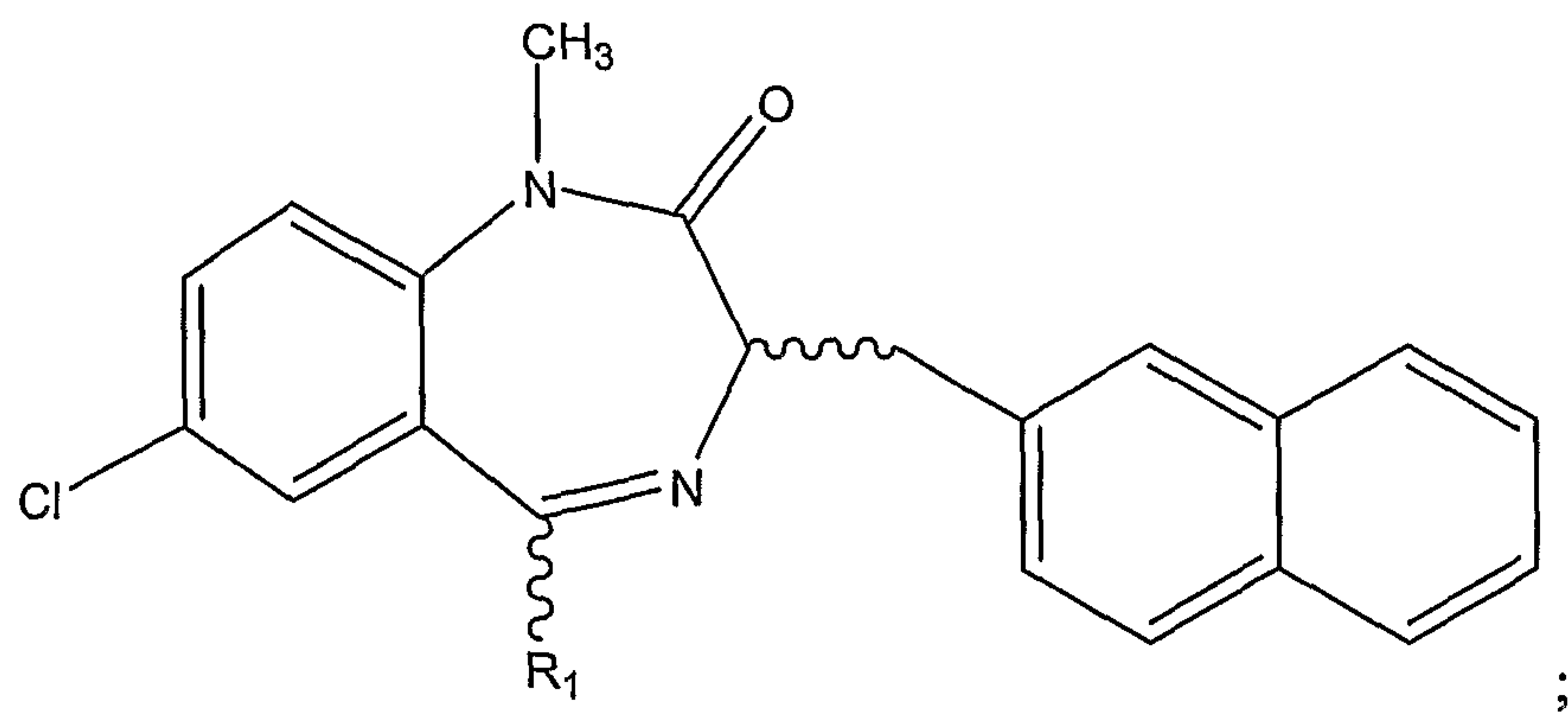


wherein R₁ is selected from naphthalanine; phenol; 1-Napthalenol; 2-Napthalenol;

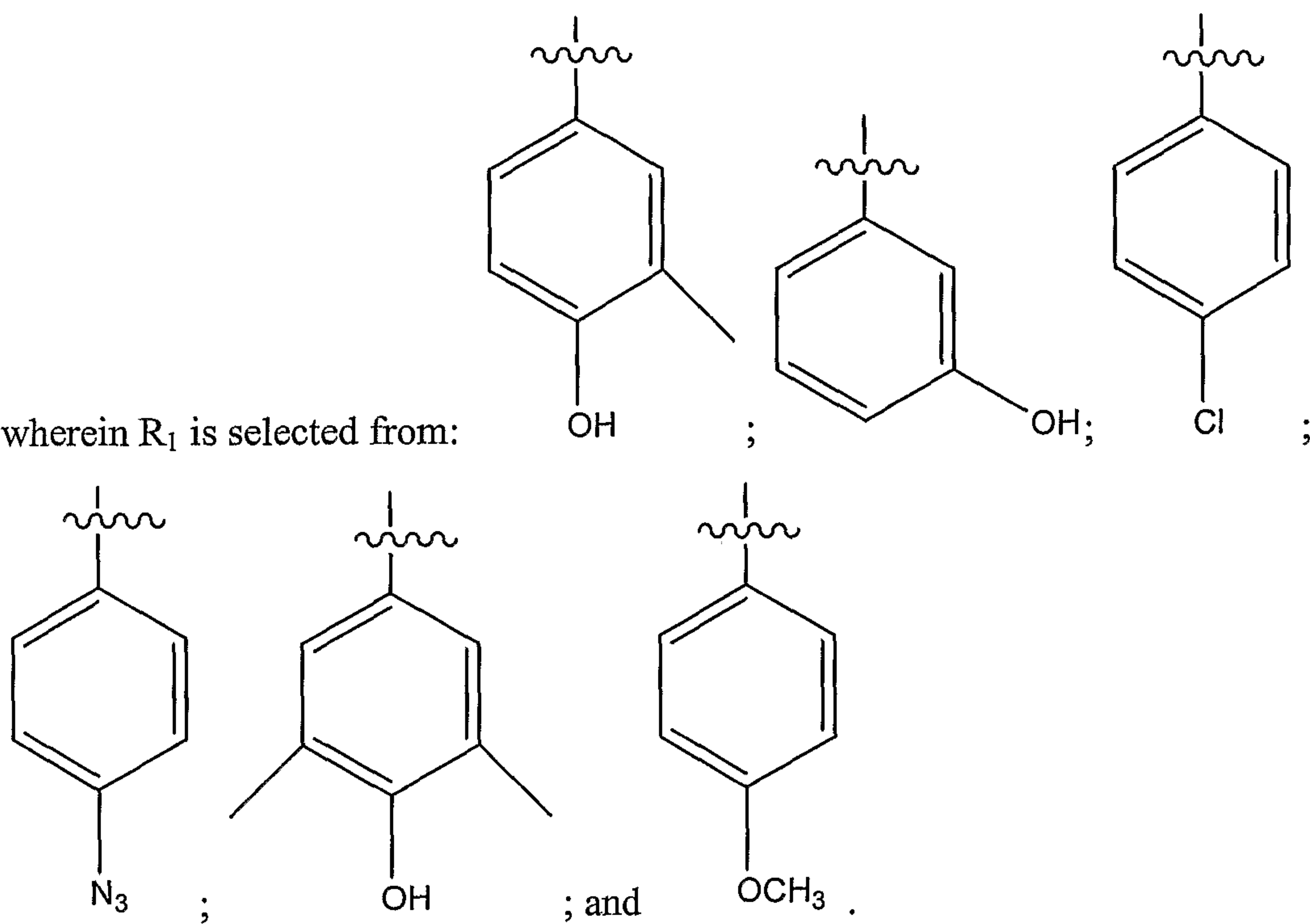


quinolines.

In some preferred embodiments, compositions comprising crystal forms and formulations of the following exemplary compounds are provided:



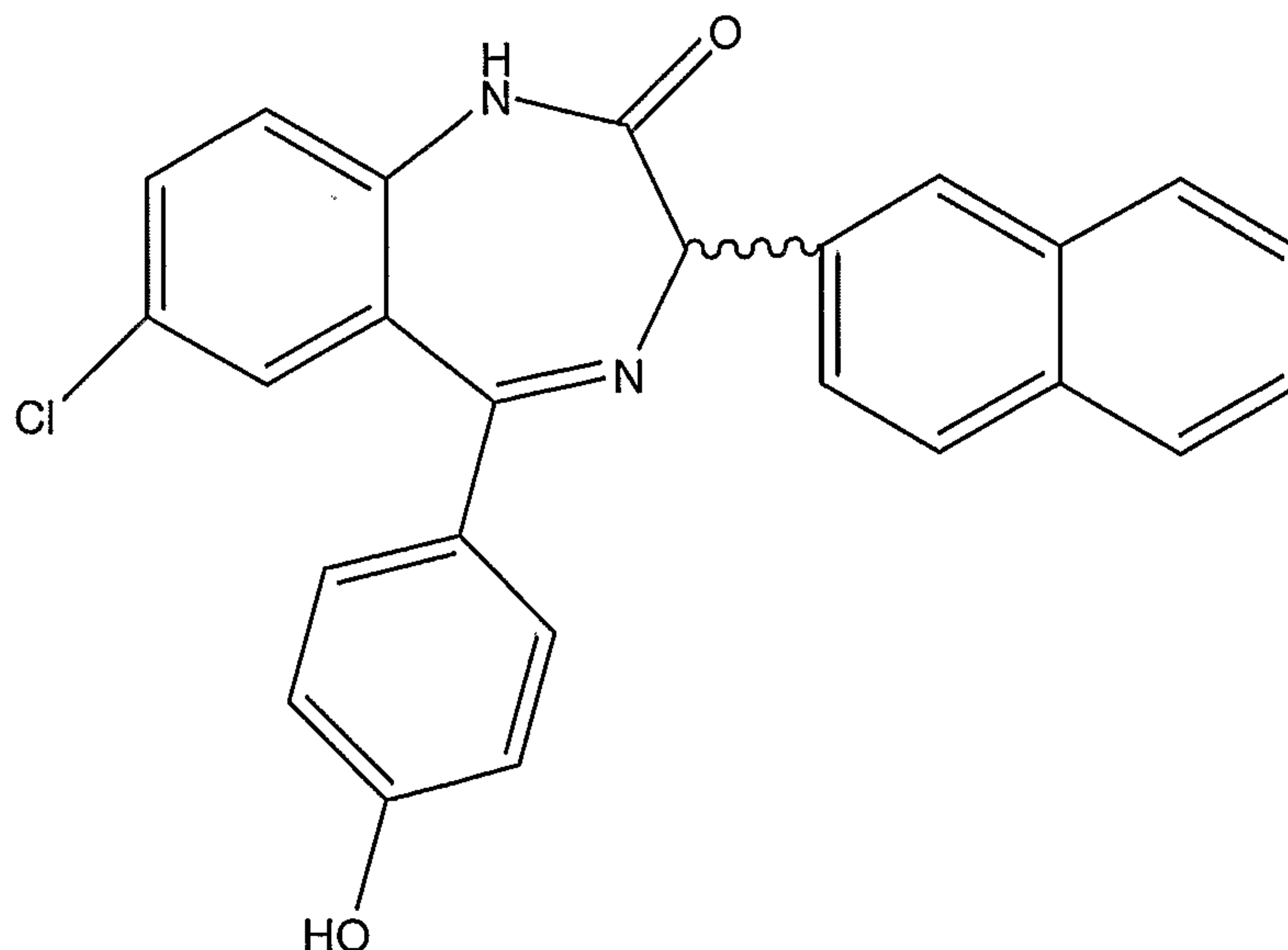
wherein R_1 is selected from:



5

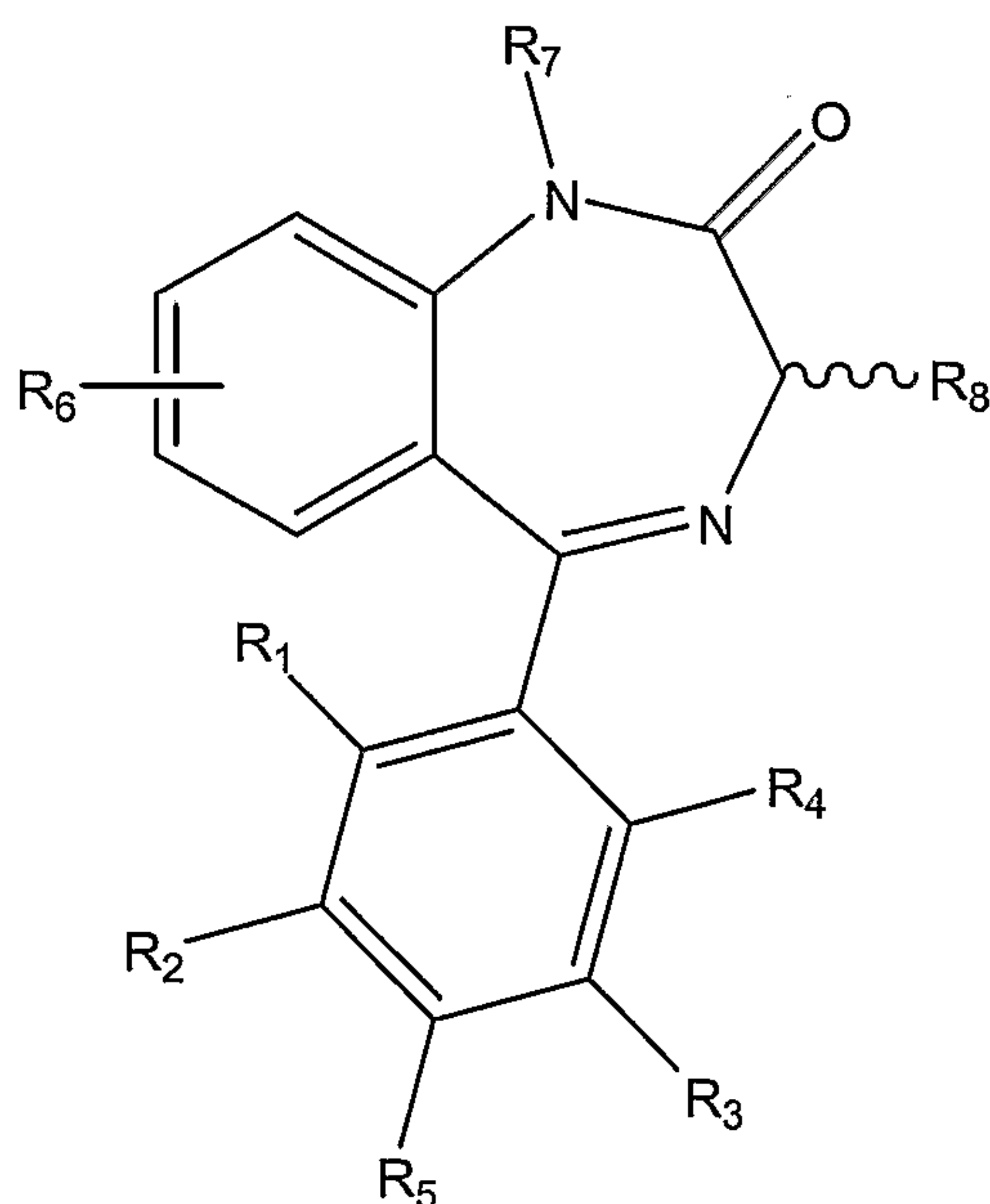
The stereochemistry of all derivatives embodied in the present invention is R, S, or racemic.

In some preferred embodiments, compositions comprising crystal forms and formulations of the following exemplary compound are provided:



In some preferred embodiments, compositions comprising crystal forms and formulations of the following exemplary compounds are provided:

5

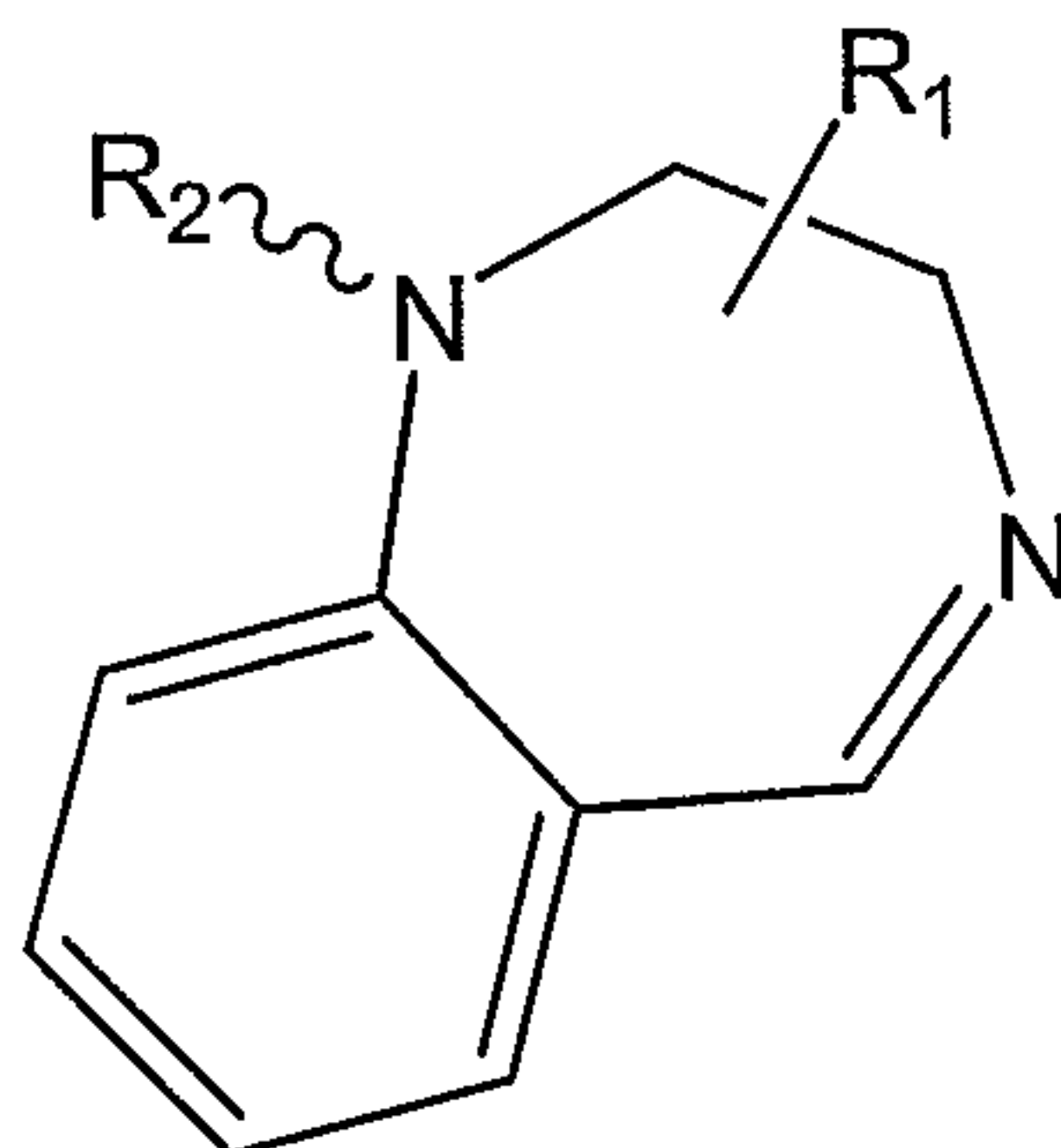


wherein R1, R2, R3 and R4 are selected from the group consisting of: hydrogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxy subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein said aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2

carbons; wherein said aliphatic chain terminates with a carboxylic acid subgroup; a linear or
 branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at
 least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain
 having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated
 5 or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen
 containing moiety (*e.g.*, nitro, nitrile, etc.); a linear or branched, saturated or unsaturated
 aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or
 branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at
 least one ether subgroup; a linear or branched, saturated or unsaturated aliphatic chain
 10 having at least 2 carbons, and having at least one halogen subgroup; a linear or branched,
 saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one
 nitronium subgroup; wherein R5 is selected from the group consisting of: OH; NO₂; NR';
 OR'; wherein R' is selected from the group consisting of: a linear or branched, saturated or
 unsaturated aliphatic chain having at least one carbon; a linear or branched, saturated or
 15 unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxyl
 subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2
 carbons, and having at least one thiol subgroup; a linear or branched, saturated or
 unsaturated aliphatic chain having at least 2 carbons, wherein said aliphatic chain terminates
 with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain
 20 having at least 2 carbons, and having at least one ketone subgroup; a linear or branched,
 saturated or unsaturated aliphatic chain having at least 2 carbons; wherein said aliphatic
 chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or
 unsaturated aliphatic chain having at least 2 carbons, and having at least one amide
 subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2
 25 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated
 aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety
 (*e.g.*, nitro, nitrile, etc.); a linear or branched, saturated or unsaturated aliphatic chain having
 at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or
 unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen
 30 subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2
 carbons, and having at least one nitronium subgroup; wherein R6 is selected from the group
 consisting of: Hydrogen; NO₂; Cl; F; Br; I; SR'; and NR'₂; wherein R' is defined as above
 in R5; wherein R7 is selected from the group consisting of: Hydrogen; a linear or branched,

saturated or unsaturated aliphatic chain having at least 2 carbons; and wherein R8 is an aliphatic cyclic group larger than benzene; wherein said larger than benzene comprises any chemical group containing 7 or more non-hydrogen atoms, and is an aryl or aliphatic cyclic group. In some embodiments, R' is any functional group that protects the oxygen of R5
 5 from metabolism in vivo, until the compound reaches its biological target (e.g., mitochondria). In some embodiments, R' protecting group(s) is metabolized at the target site, converting R5 to a hydroxyl group.

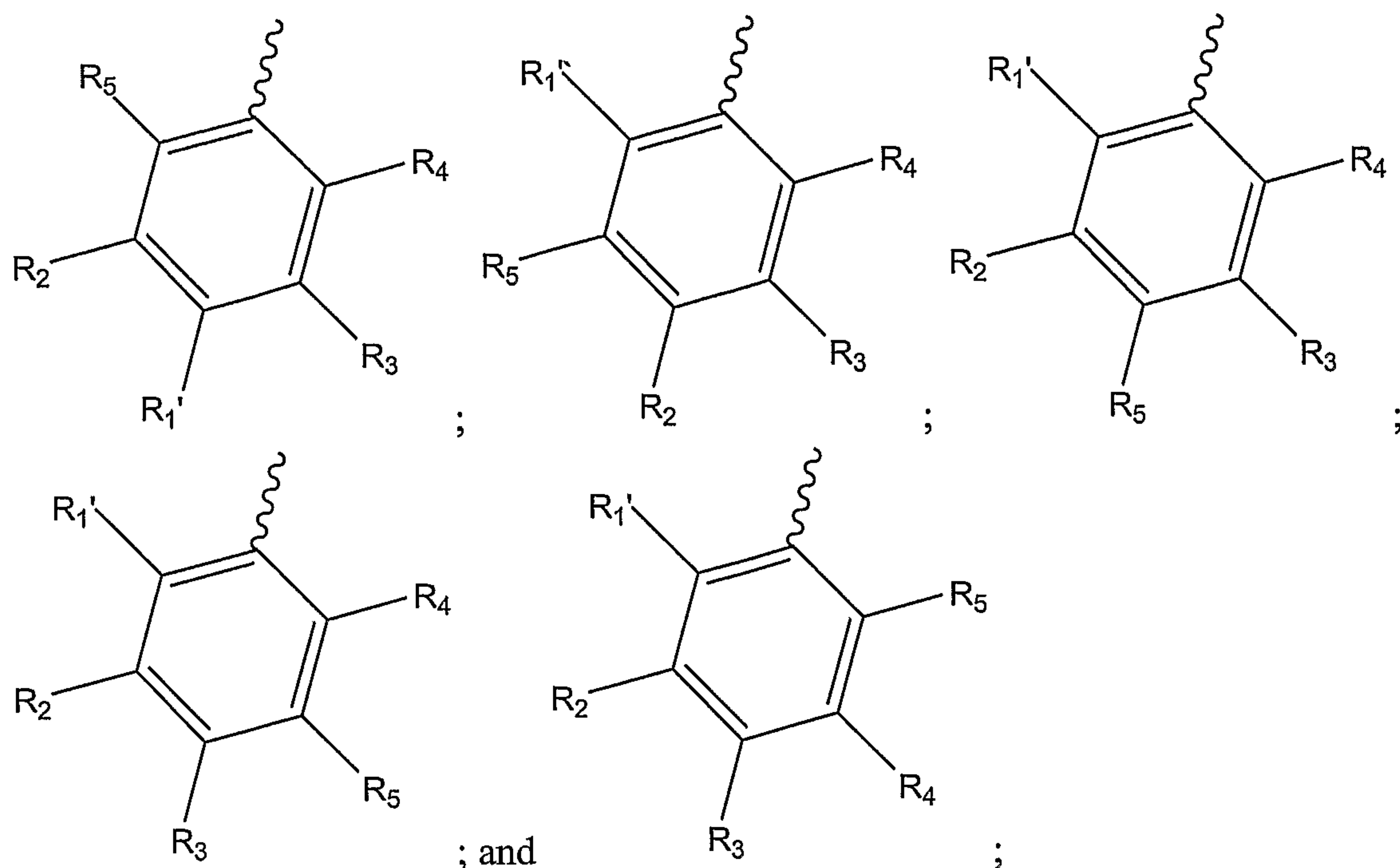
In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



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including both R and S enantiomeric forms and racemic mixtures; wherein R1 comprises a chemical moiety comprising a hydrogen bonding proton donor (e.g., a hydroxyl group, a phenol group, an amide group, a sulfonamide group, an amine group, an aniline group, a benzimidizalone group, a carbamate group, and an imidazole
 15 group); and R2 comprises a hydrophobic chemical moiety.

In preferred embodiments, R1 is selected from the group consisting of:

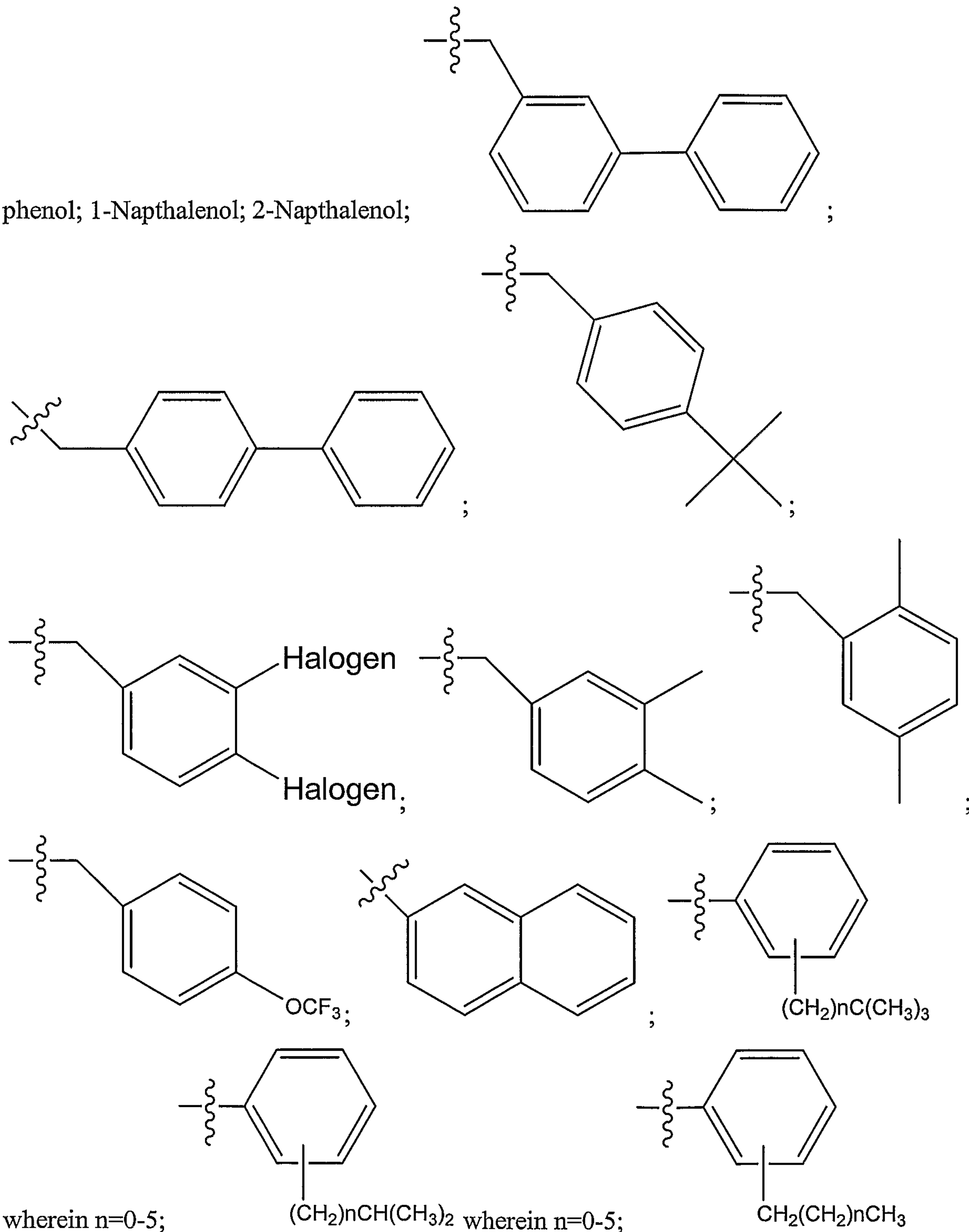


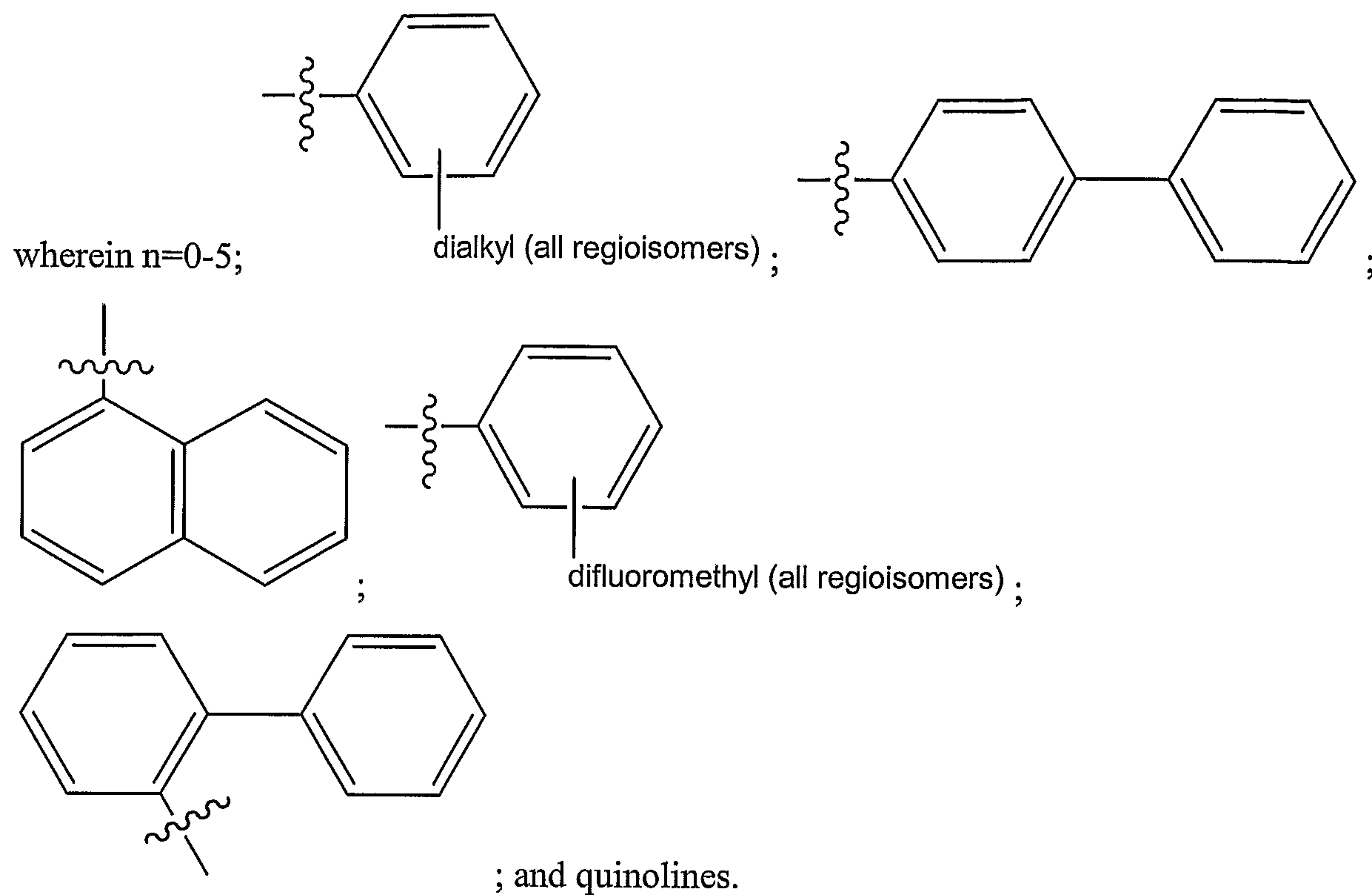
wherein R_1' , R_2 , R_3 and R_4 are selected from the group consisting of: hydrogen; CH_3 ; a linear or branched, saturated or unsaturated aliphatic chain having at least 1 carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons and at least one hydroxy subgroup; a linear or branched, saturated or unsaturated, substituted or non-substituted, aliphatic chain having at least 2 carbons and having at least one thiol subgroup; a linear or branched, saturated or unsaturated, substituted or non-substituted, aliphatic chain having at least 2 carbons wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated, substituted or non-substituted, aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated, substituted or non-substituted, aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated, substituted or non-substituted, aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated, substituted or non-substituted, aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated, substituted or non-substituted, aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated, substituted or non-substituted, aliphatic chain having at least 2 carbons, and having at least one ether subgroup; a linear or branched, saturated or unsaturated, substituted or non-substituted, aliphatic chain having at

least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated, substituted or non-substituted, aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; and R5 is OH.

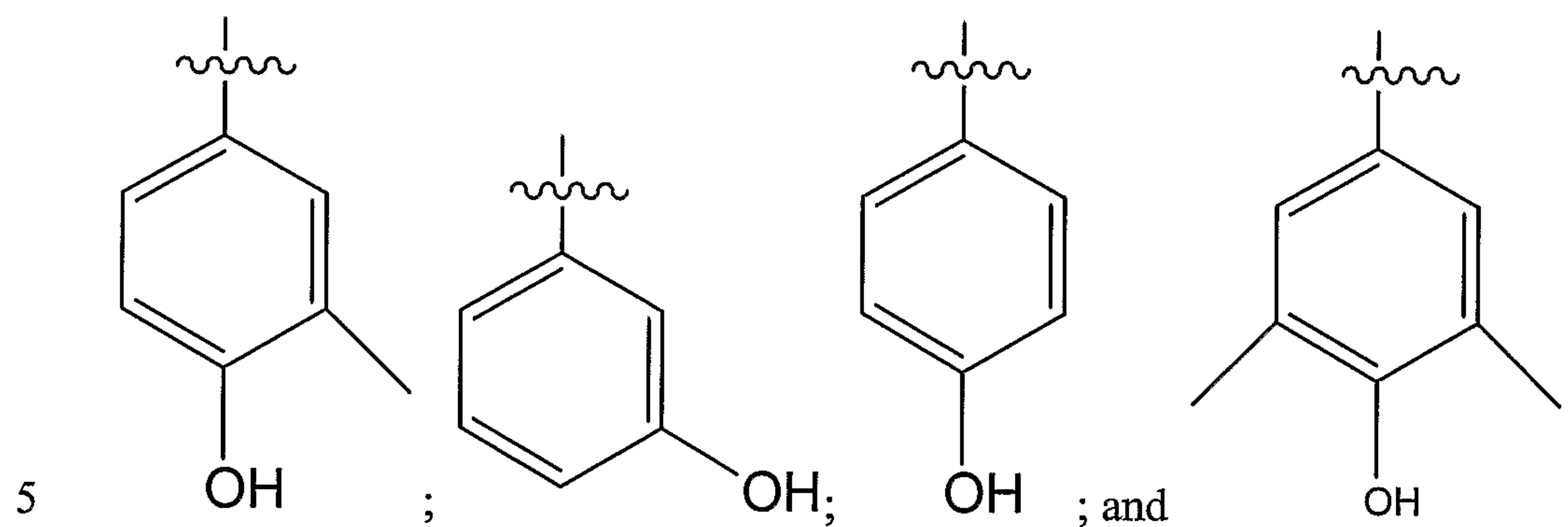
In preferred embodiments, R2 is selected from group consisting of: naphthalanine;

5 phenol; 1-Napthalenol; 2-Napthalenol;

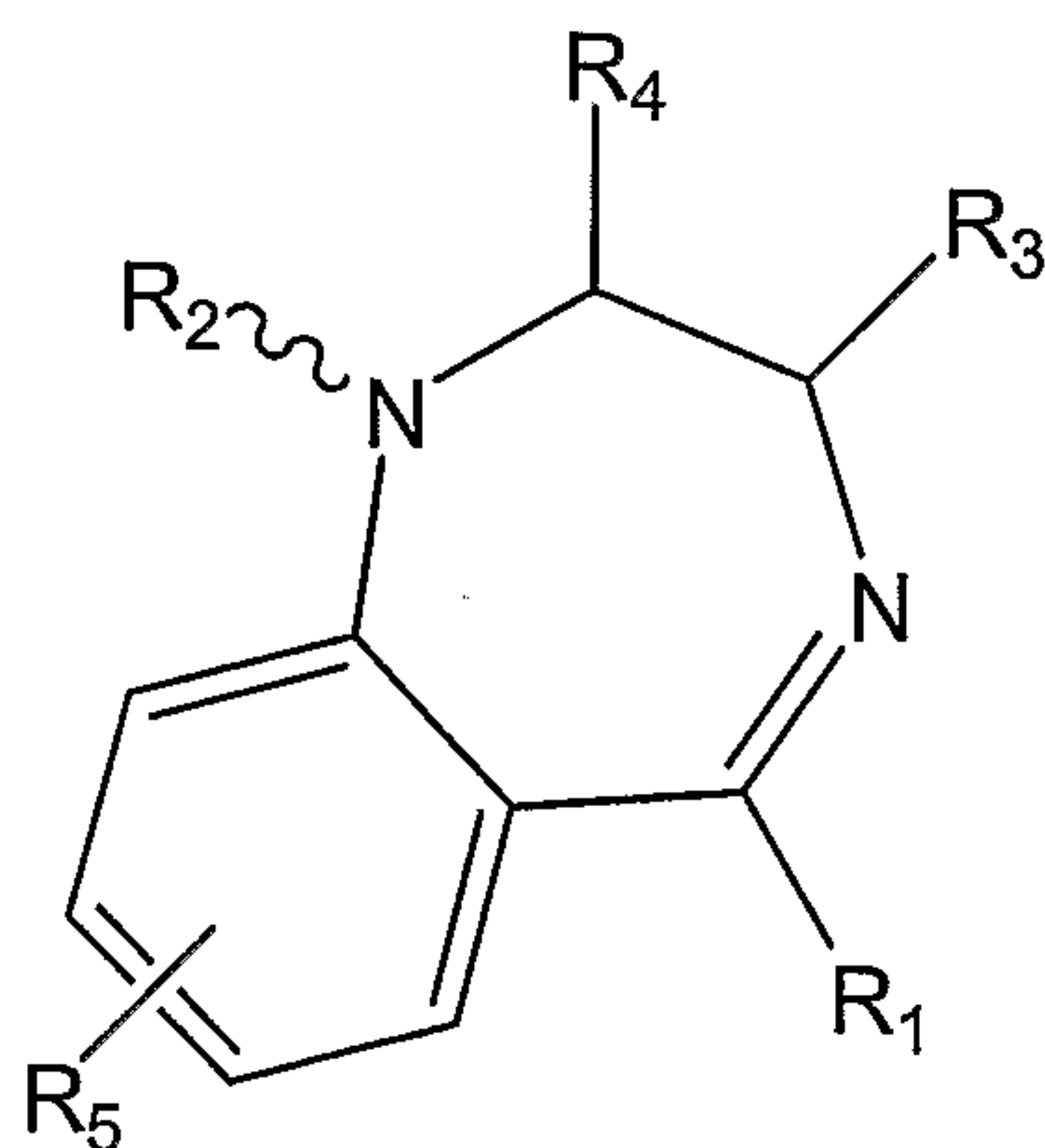




In some preferred embodiments, R1 is selected from the group consisting of:



In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



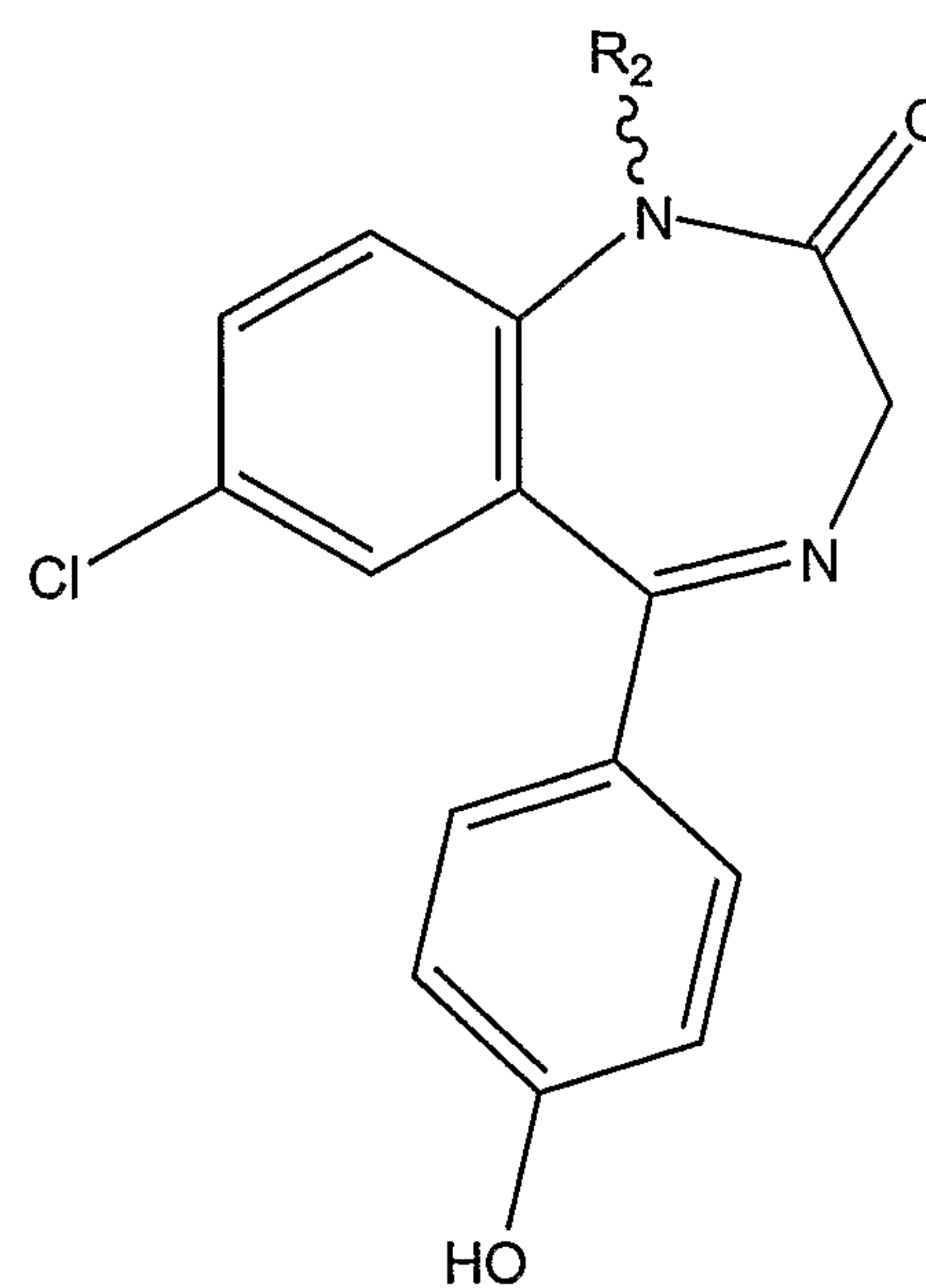
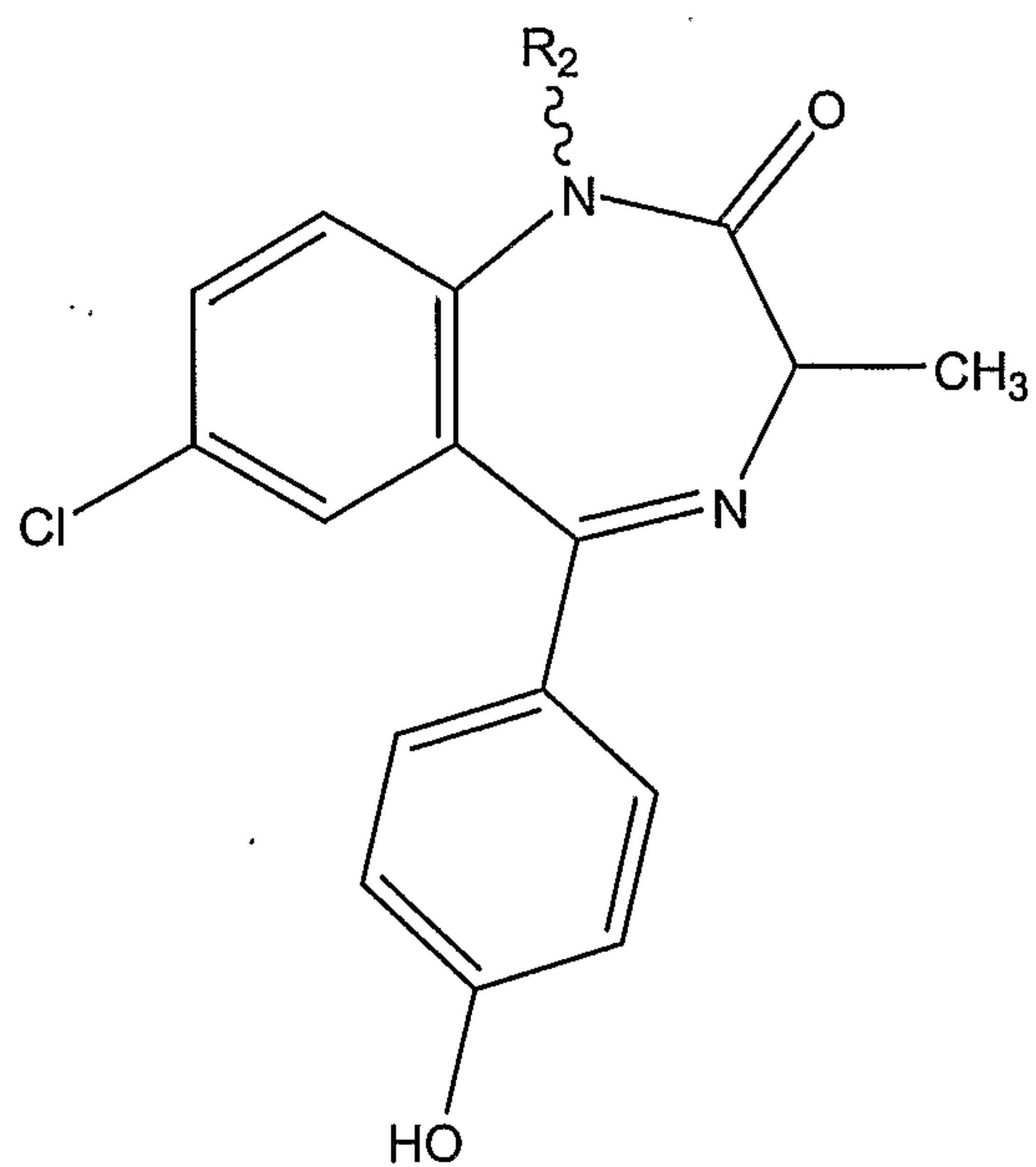
wherein R3 is selected from the group consisting of Hydrogen; amino; and a linear or branched, saturated or unsaturated, substituted (e.g., substituted with amines, esters, amides or phosphatases) or non-substituted, aliphatic chain having at least 2 carbons;

R4 is selected from the group consisting of H, a ketone, and a nitrogen; and

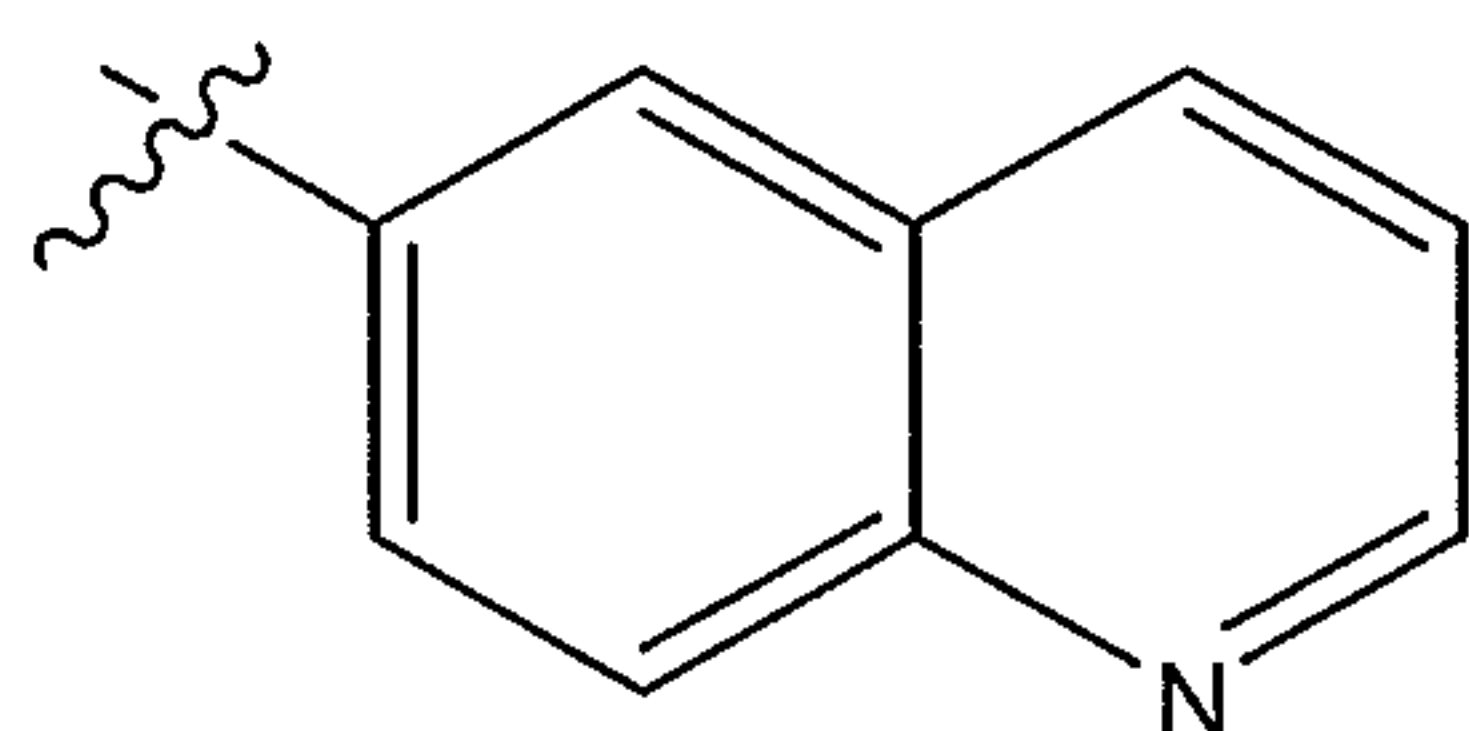
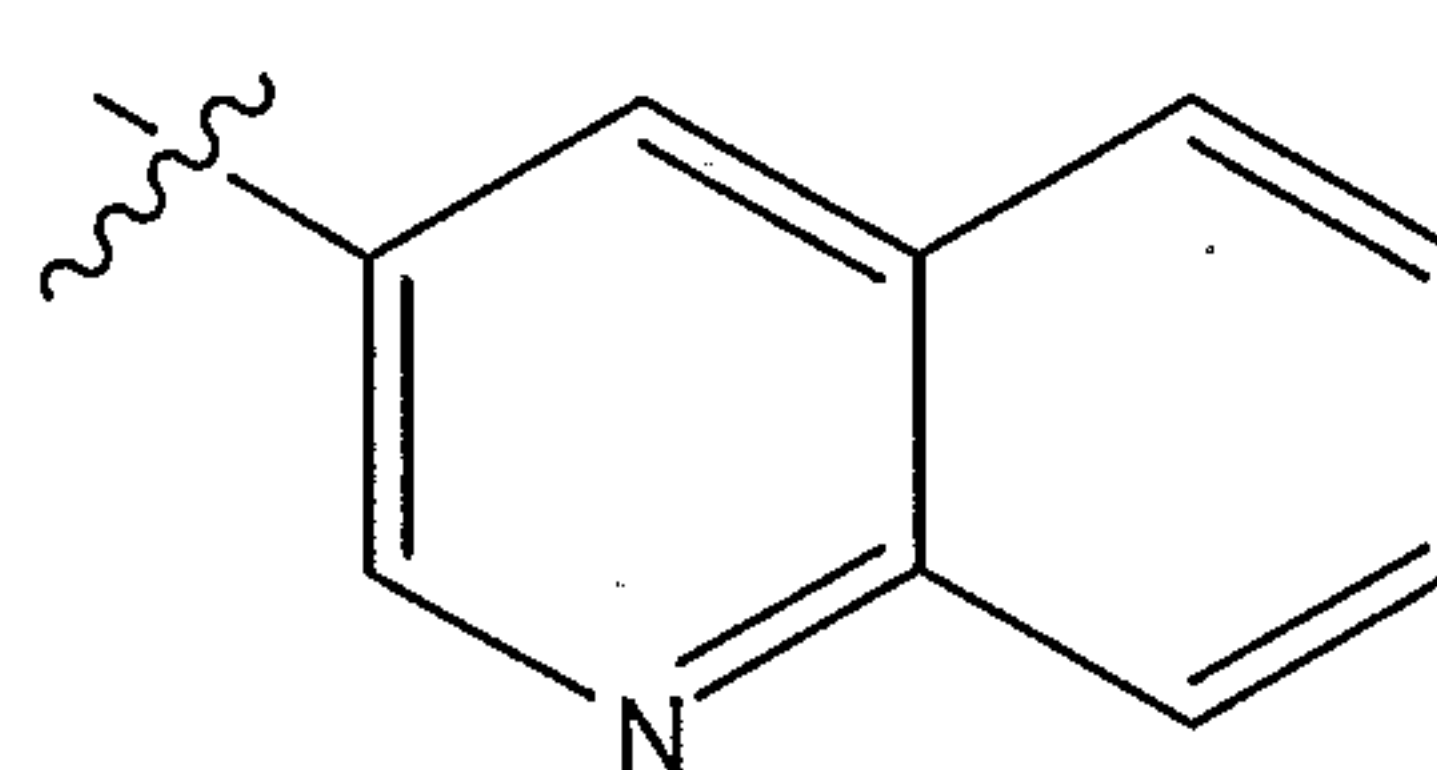
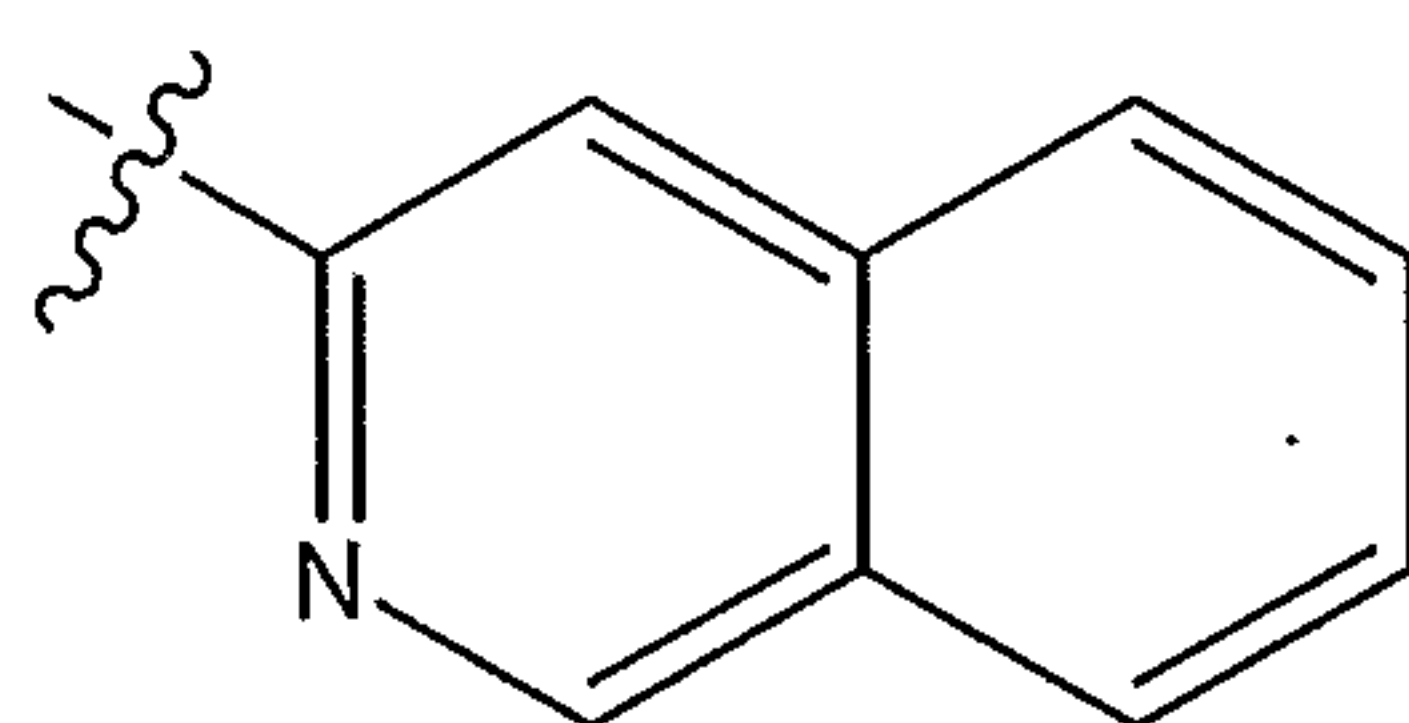
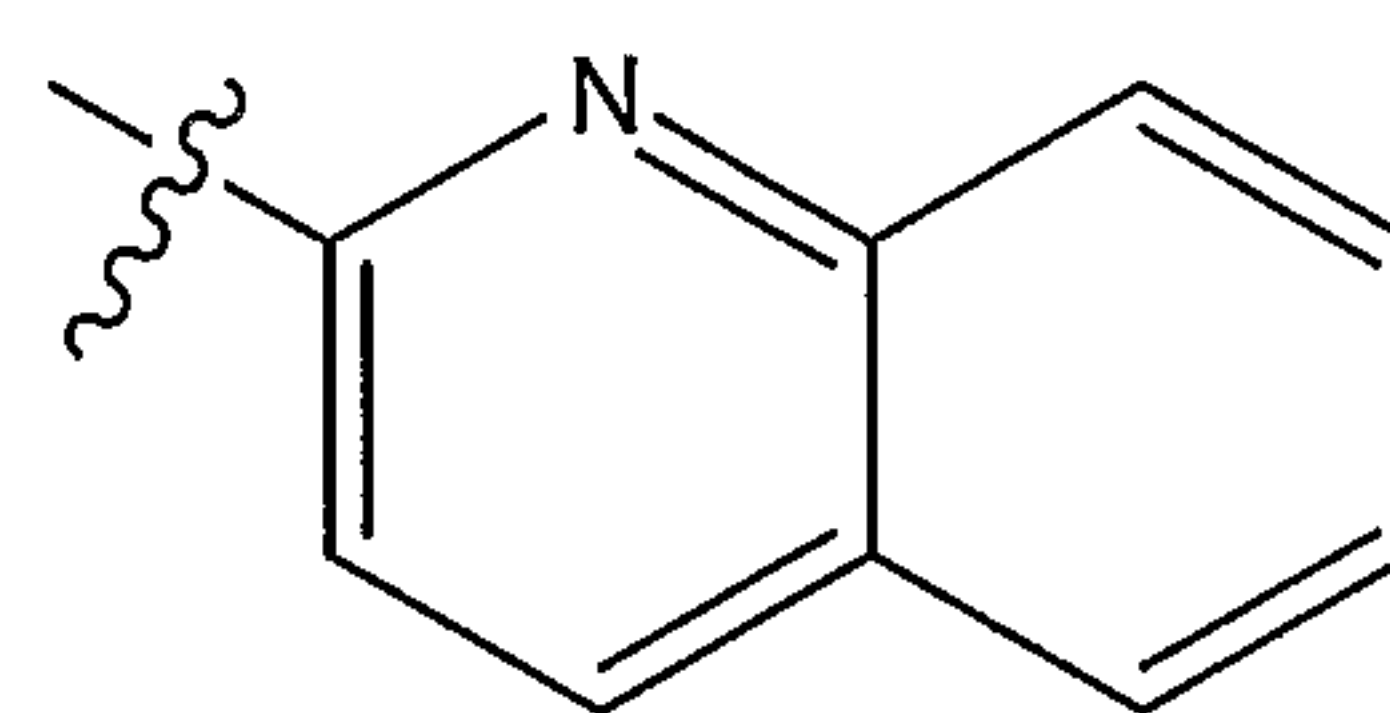
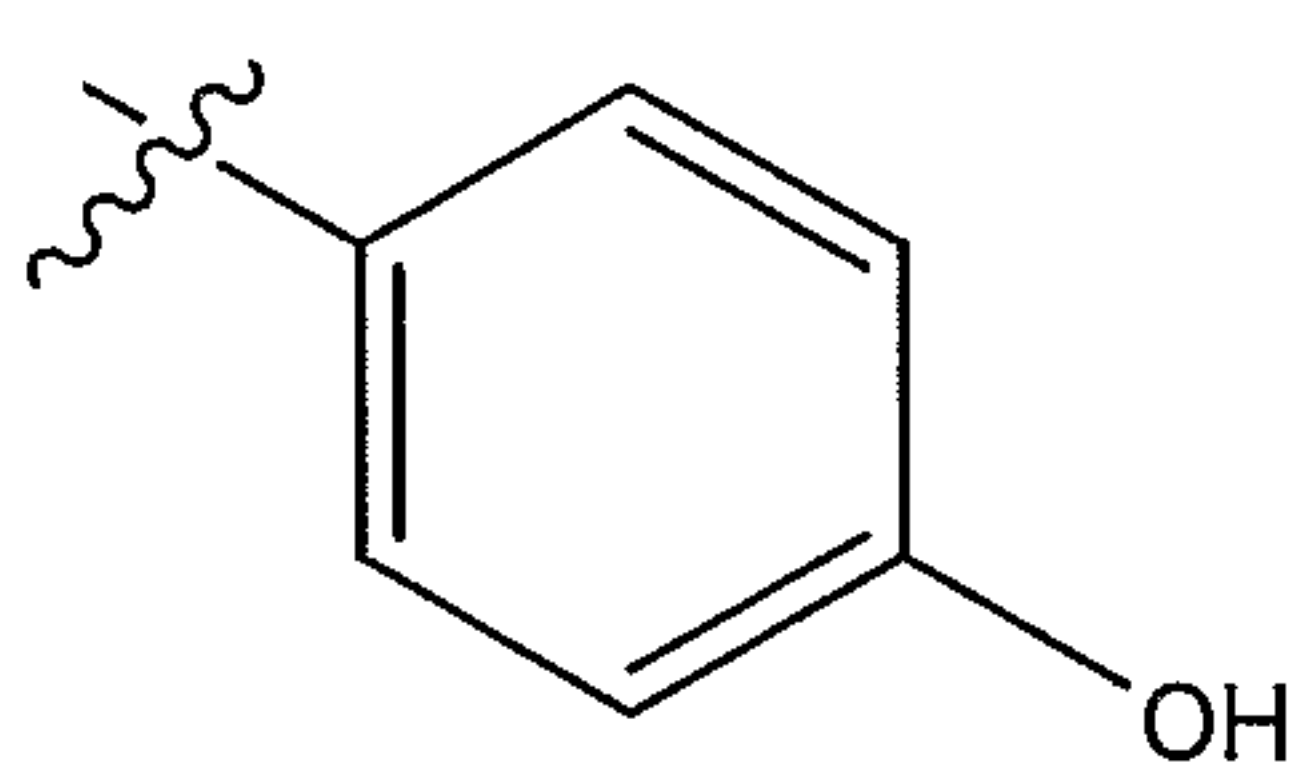
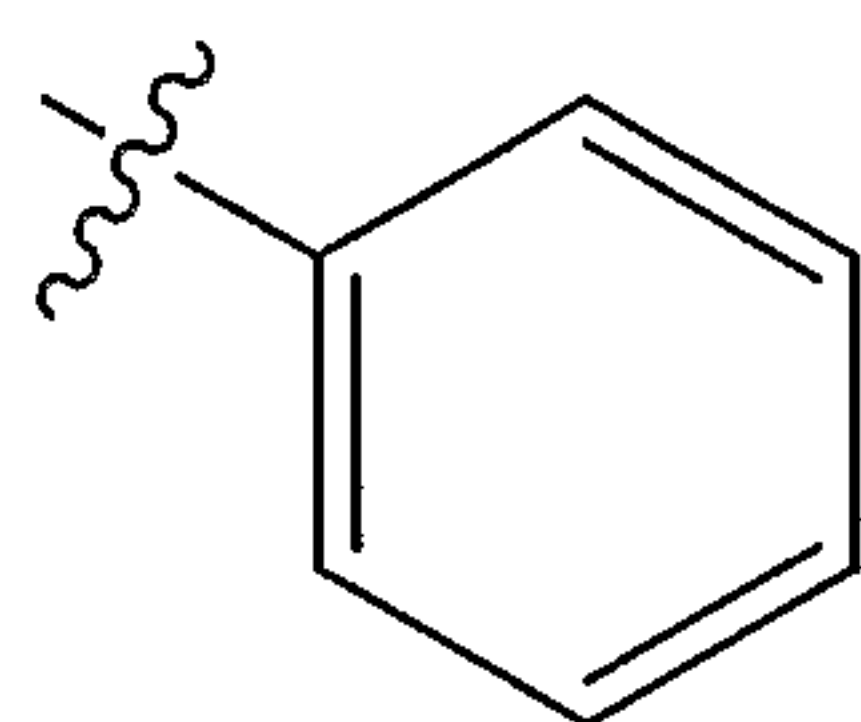
5 R5 is selected from H, a hydroxy, an alkoxy, a carboxylic acid, a carboxylic ester, a halogen, a nitro, a sulfonamide, an amide, a carbamate, an amino, a lower-alkyl, a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of less than 10 carbons, a substituted cycloaliphatic group, an aryl, a heterocyclic,
10 NO₂; SR'; and NR'₂, wherein R' is defined as a linear or branched, saturated or unsaturated aliphatic chain having at least one carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxyl subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain
15 having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least
20 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine
25 subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup.

In other preferred embodiments R2 is any chemical group that permits the
30 compound to bind to OSCP. In some such embodiments, R2 comprises a hydrophobic aromatic group. In preferred embodiments R2 comprises a hydrophobic aromatic group larger than benzene (e.g., a benzene ring with non-hydrogen substituents, a moiety having two or more aromatic rings, a moiety with 7 or more carbon atoms, etc.).

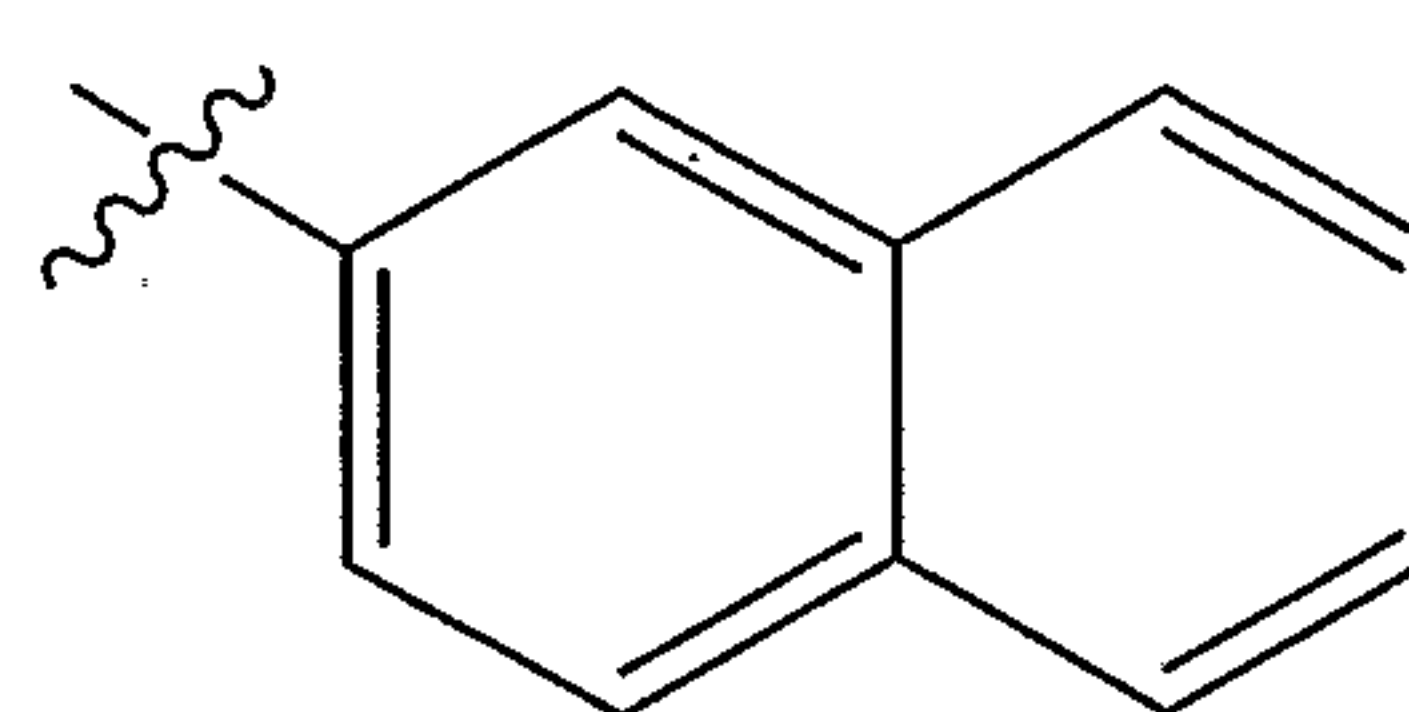
In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



5 wherein R_2 is

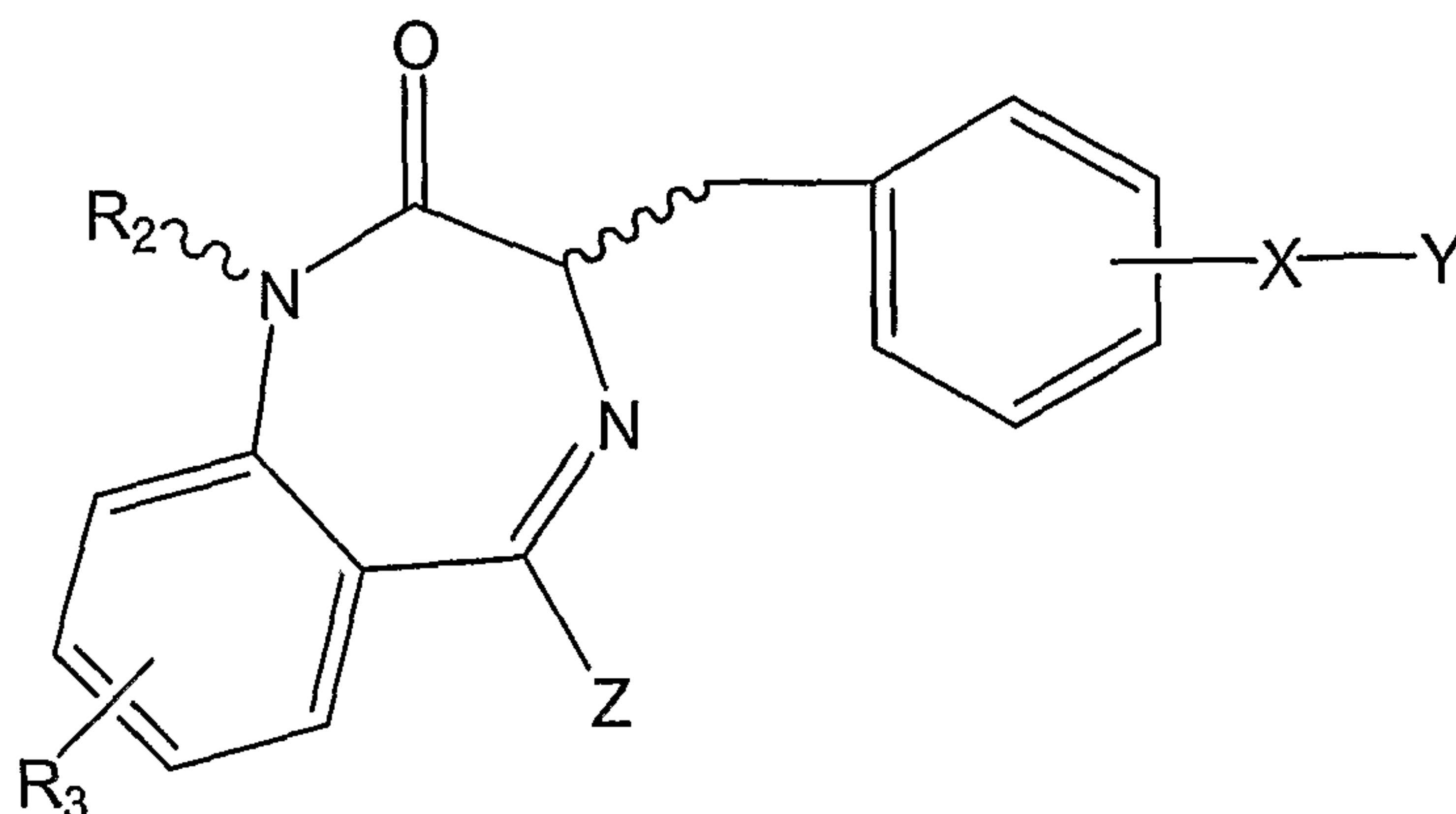


and



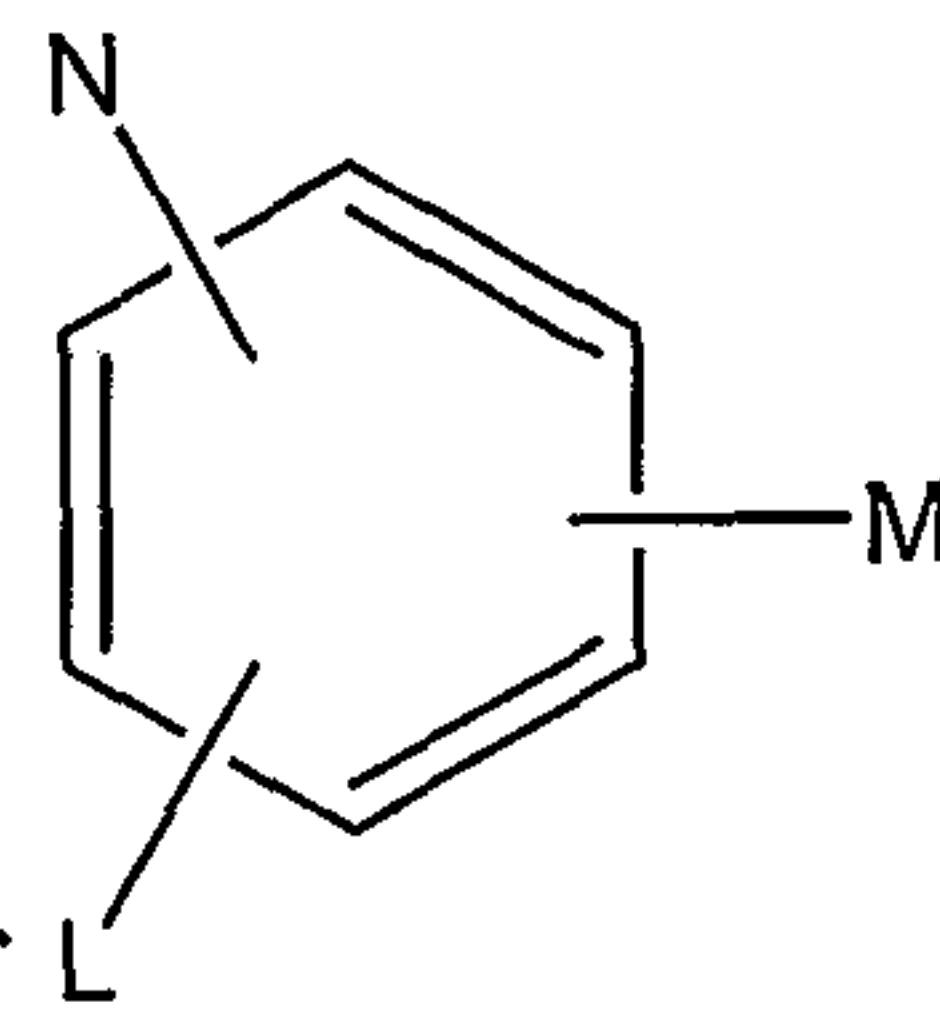
and dimethylphenyl (all isomers) and ditrifluoromethyl (all isomers).

In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:

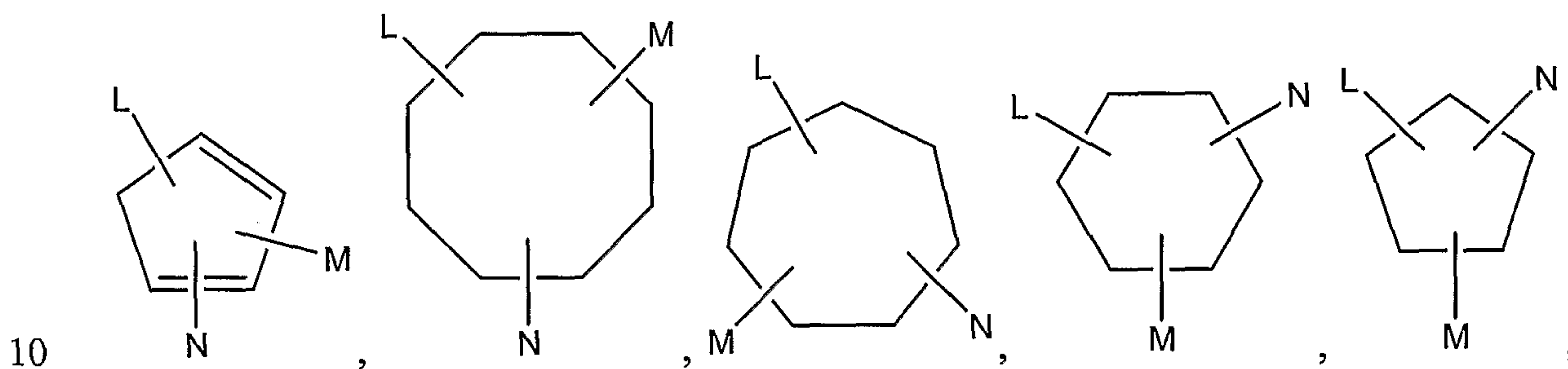


5 wherein R2 is selected from the group consisting of Hydrogen, alkyl, substituted alkyl, and $(CH_2)_n$ wherein $n=1-6$;

wherein R3 is selected from the group consisting of hydrogen, halogen, alkyl, substituted alkyl, carboxylic acid, amide SO_2NH_2 , $NHSO_2$ alkyl, and NO_2 ;



wherein X is selected from the group consisting of L ,



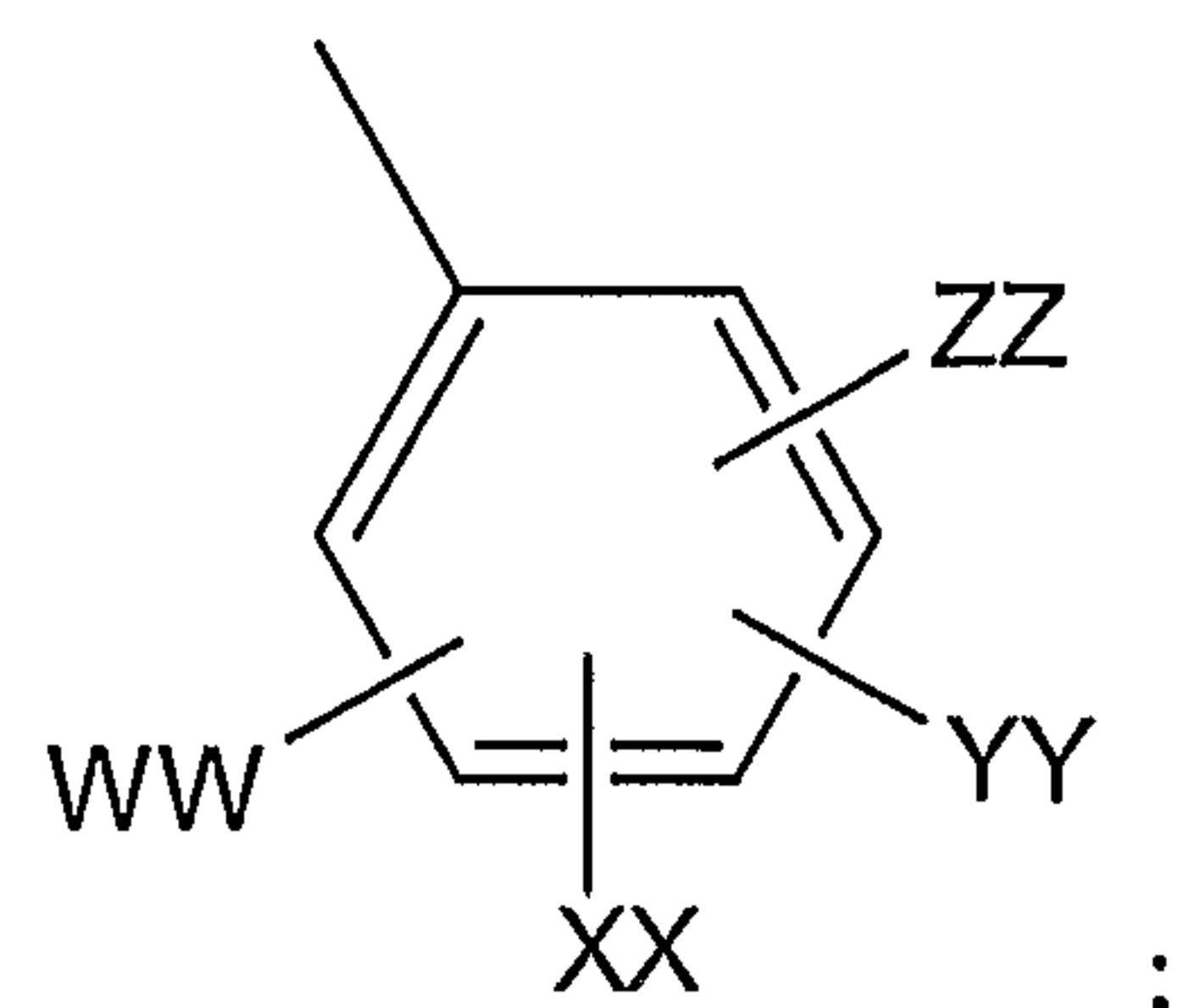
10

alkyl, substituted alkyl, sulfamide, SO_2 alkyl, $NHSO_2$, CH_2 , CH_2CH_2 , SO_2 , CH_2SO_2 , SO_2CH_2 , OCH_2CH_2O , SO , CH_2CH_2SO , $SOCH_2CH_2$; and

15 wherein L, M and N are present or absent, and are selected from the group consisting of alkyl, NO_2 , halogen, OH, O-Alkyl, methyl ester, propyl ester, ethyl ester, CO_2H , CF_3 , aniline, nitro, heterocycle, mono-substituted alkyl, di-substituted alkyl, and tri-substituted alkyl, hydrogen, SO_2NH_2 , SO_2NH -alkyl, SO alkyl, $NHSO_2$ alkyl; and

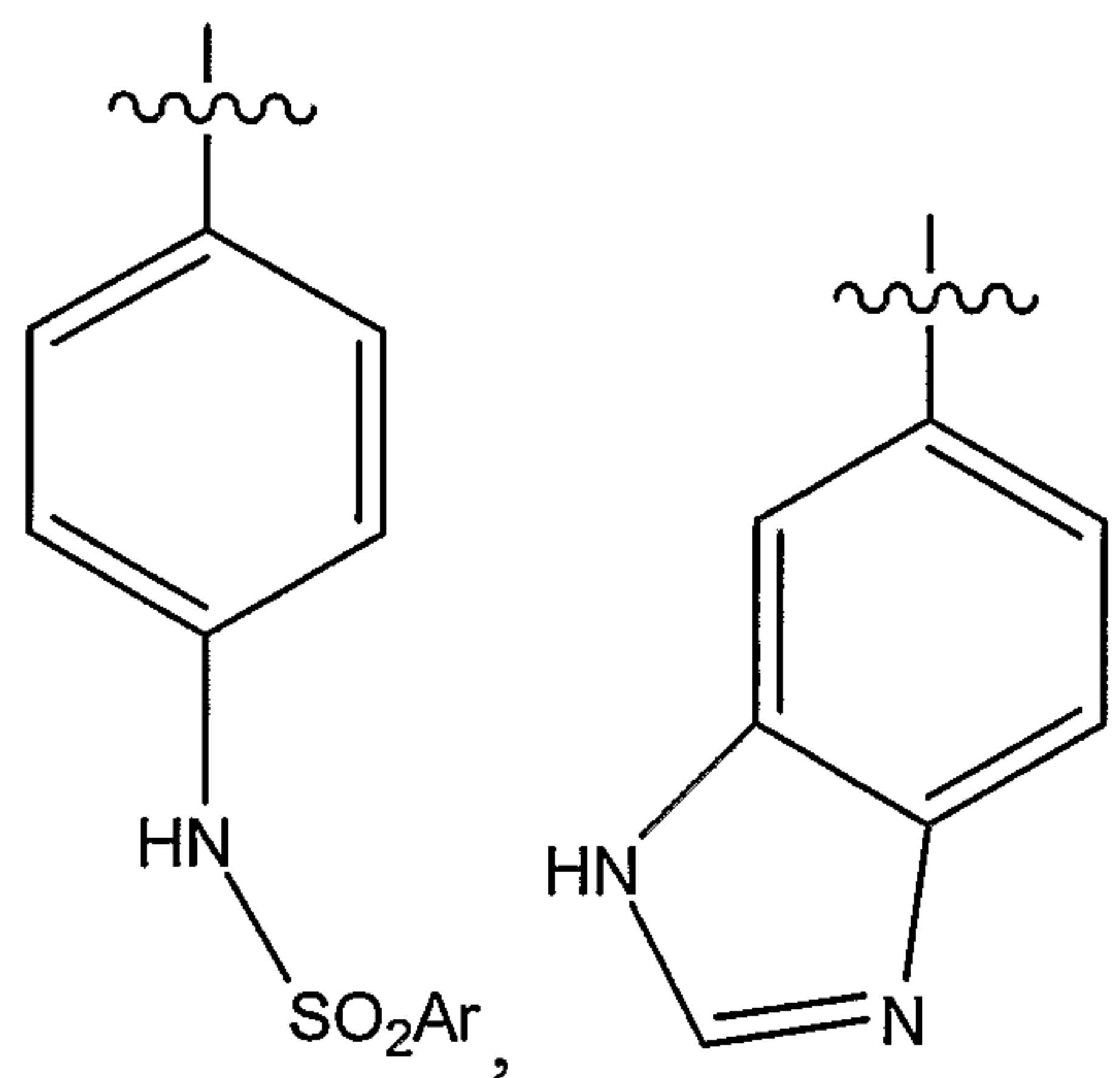
wherein Y is selected from the group consisting of hydrogen, alkyl, substituted alkyl, halogen, OH, O-Alkyl, methyl ester, propyl ester, ethyl ester, CO_2H , nitro,

heterocycle, mono-substituted alkyl, di-substituted alkyl, and tri-substituted alkyl,

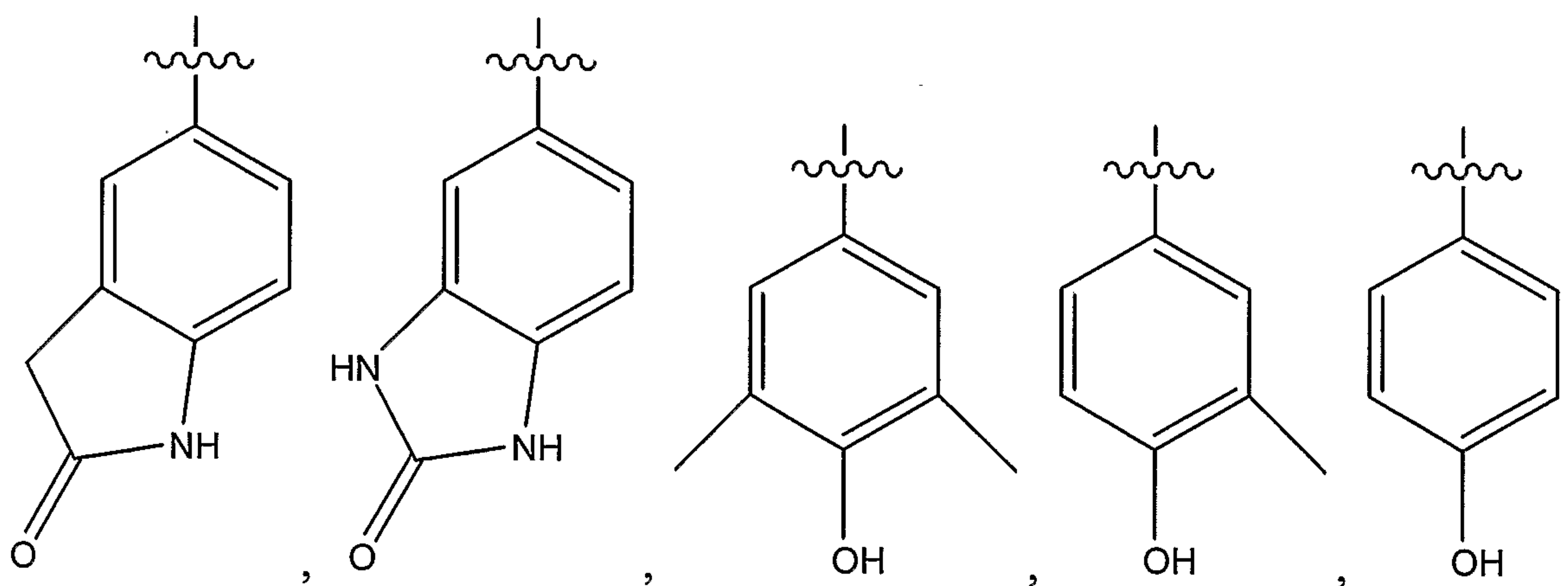


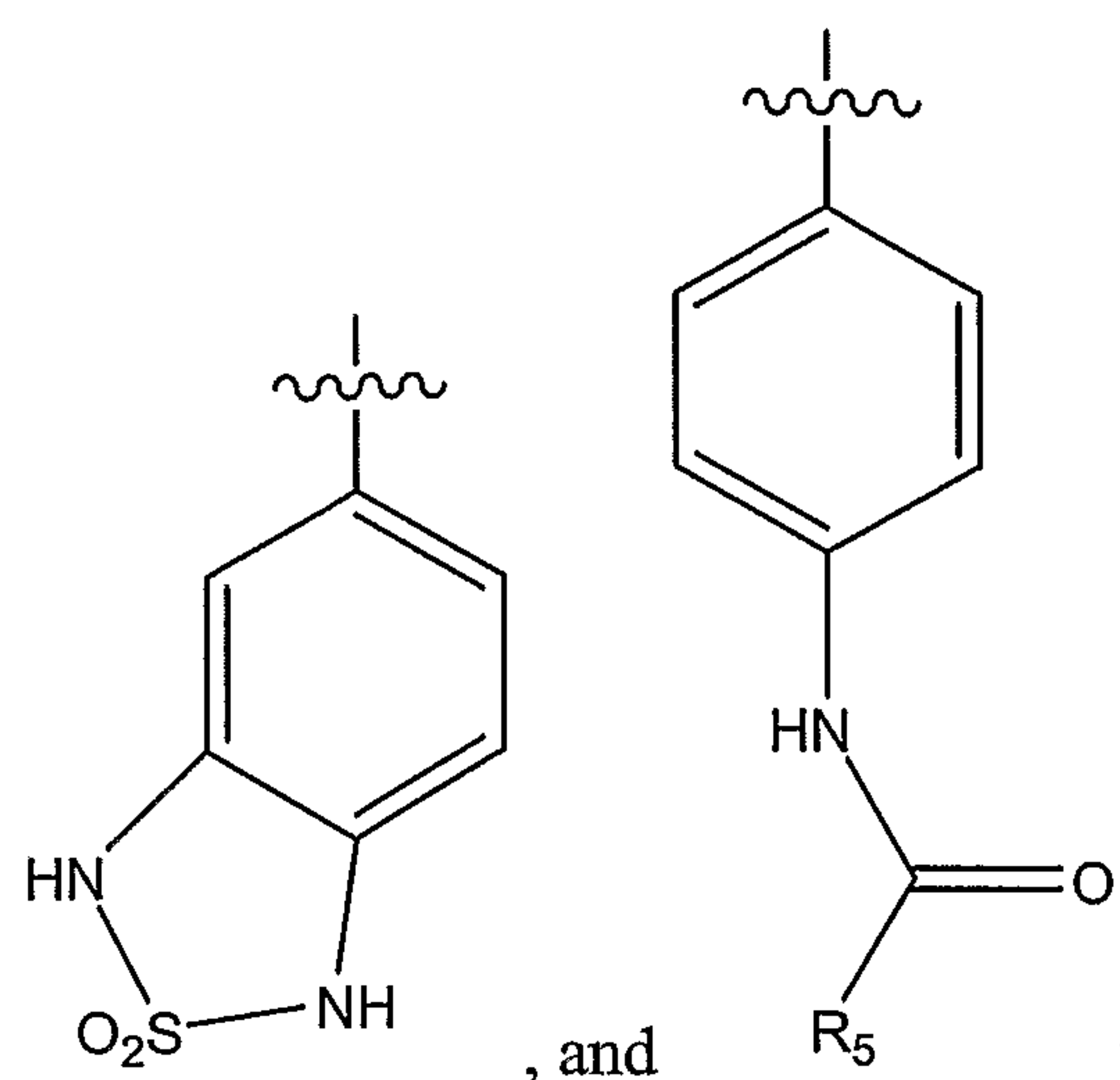
hydrogen, SOalkyl, SO₂NH₂, SO₂NH-alkyl, NHSO₂alkyl, and

wherein WW, XX, YY and ZZ are present or absent, and are selected from the group consisting of alkyl, halogen, OH, O-Alkyl, methyl ester, propyl ester, ethyl ester, CO₂H, aniline, nitro, heterocycle, mono-substituted alkyl, di-substituted alkyl, and tri-substituted alkyl, hydrogen, SO₂NH₂, SO₂NH-alkyl, NHSO₂alkyl; and



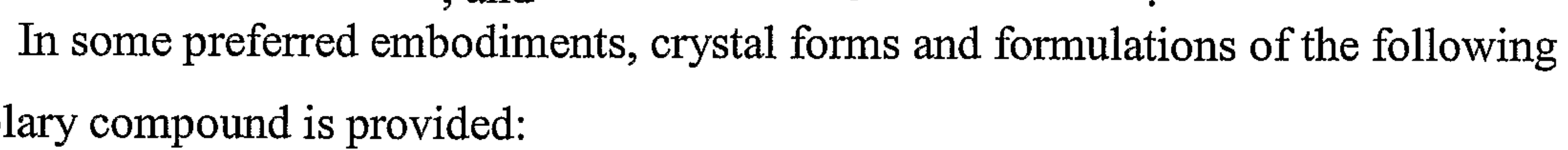
wherein Z is selected from the group consisting of

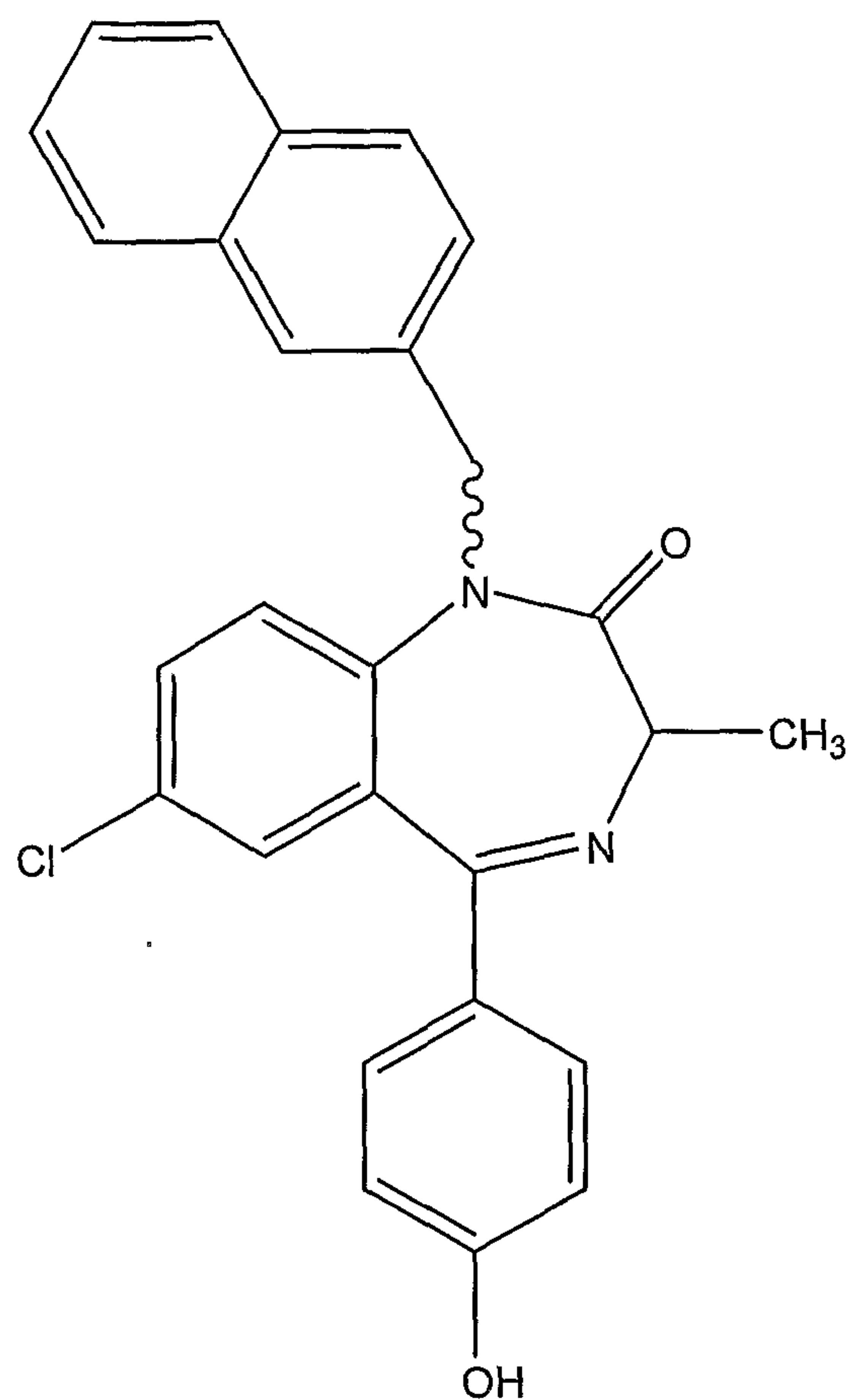




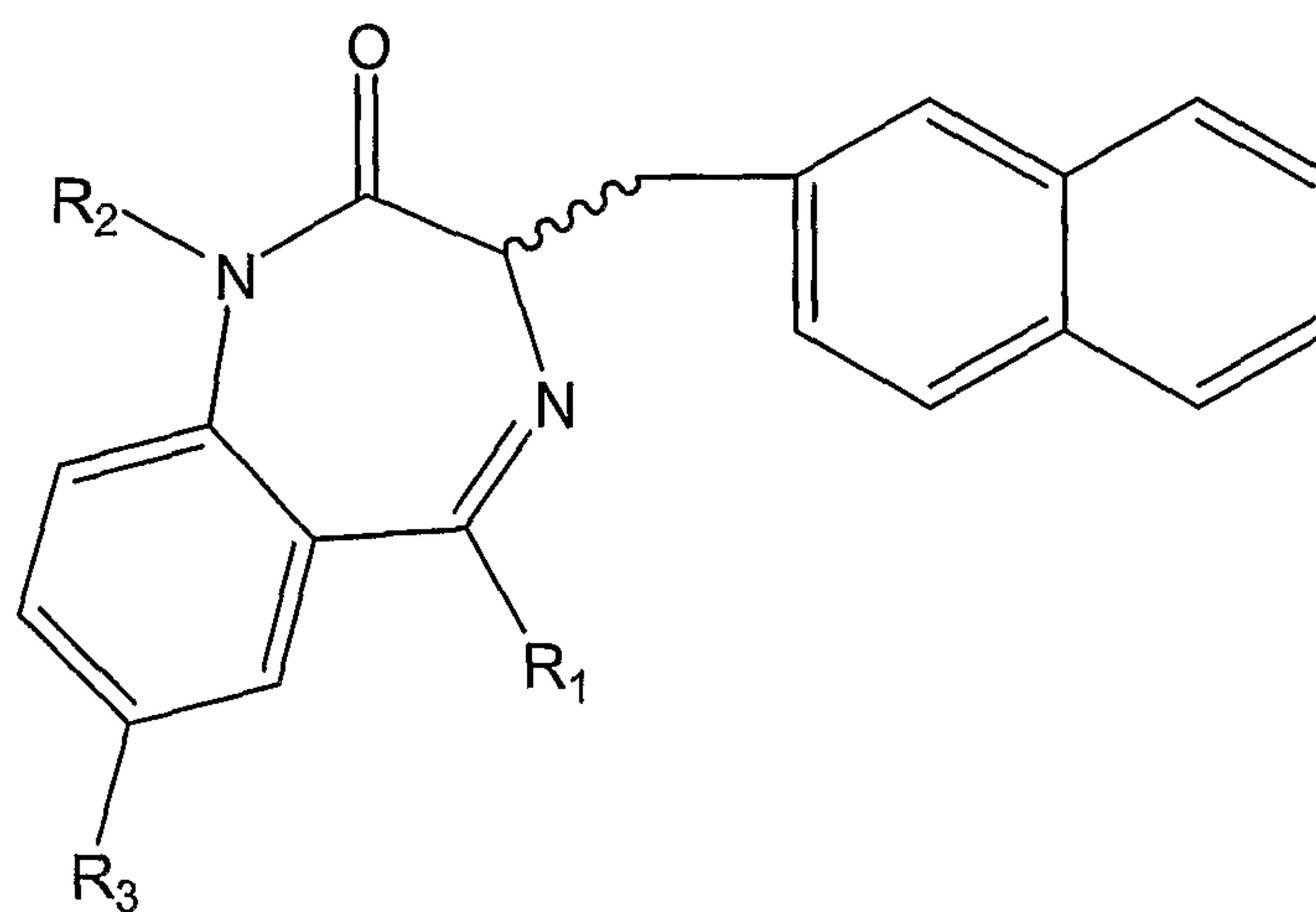
, and wherein R₅ is selected from the group consisting of alkyl, mono-substituted alkyl, di-substituted alkyl, and tri-substituted alkyl.

In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:

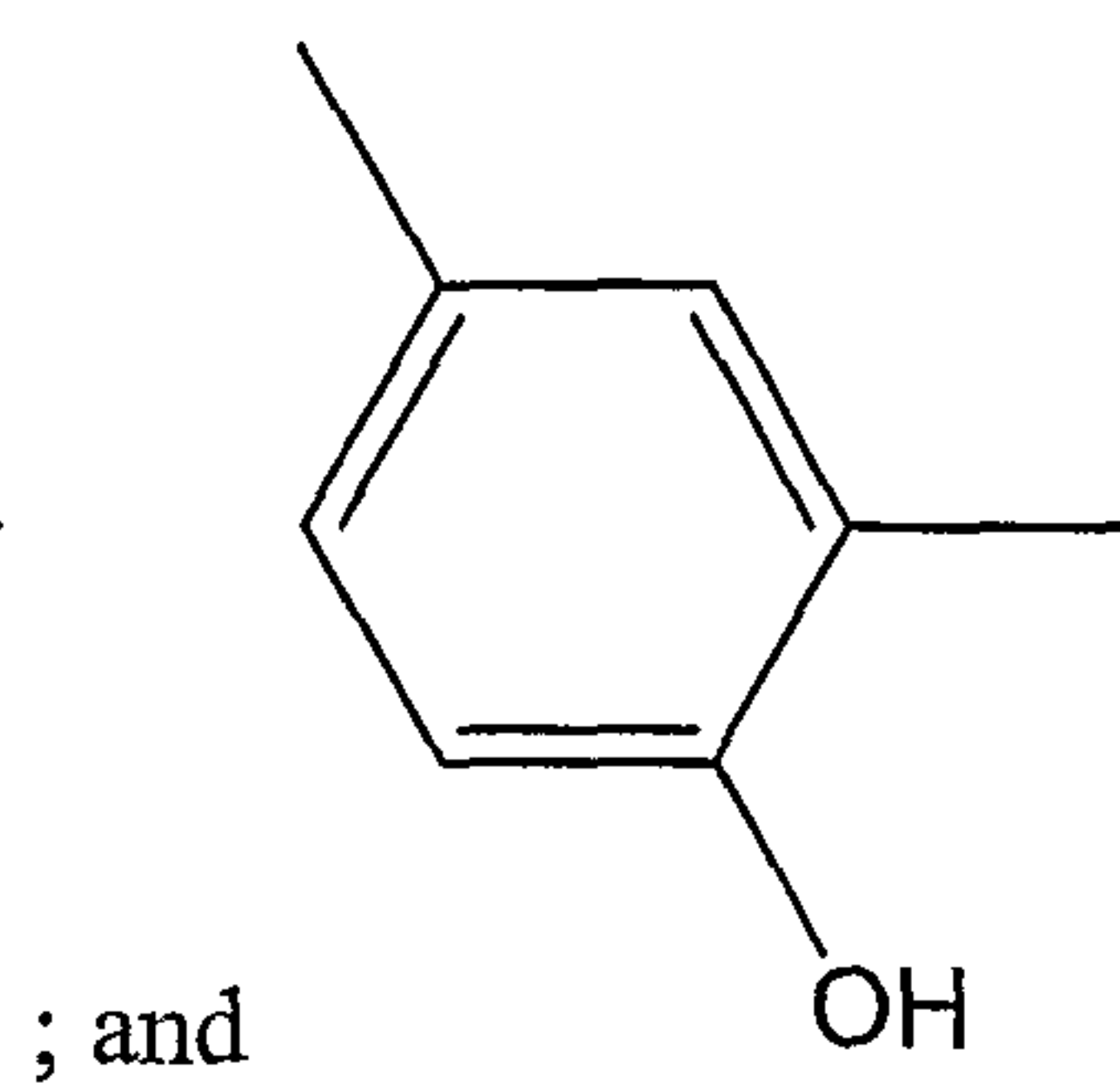
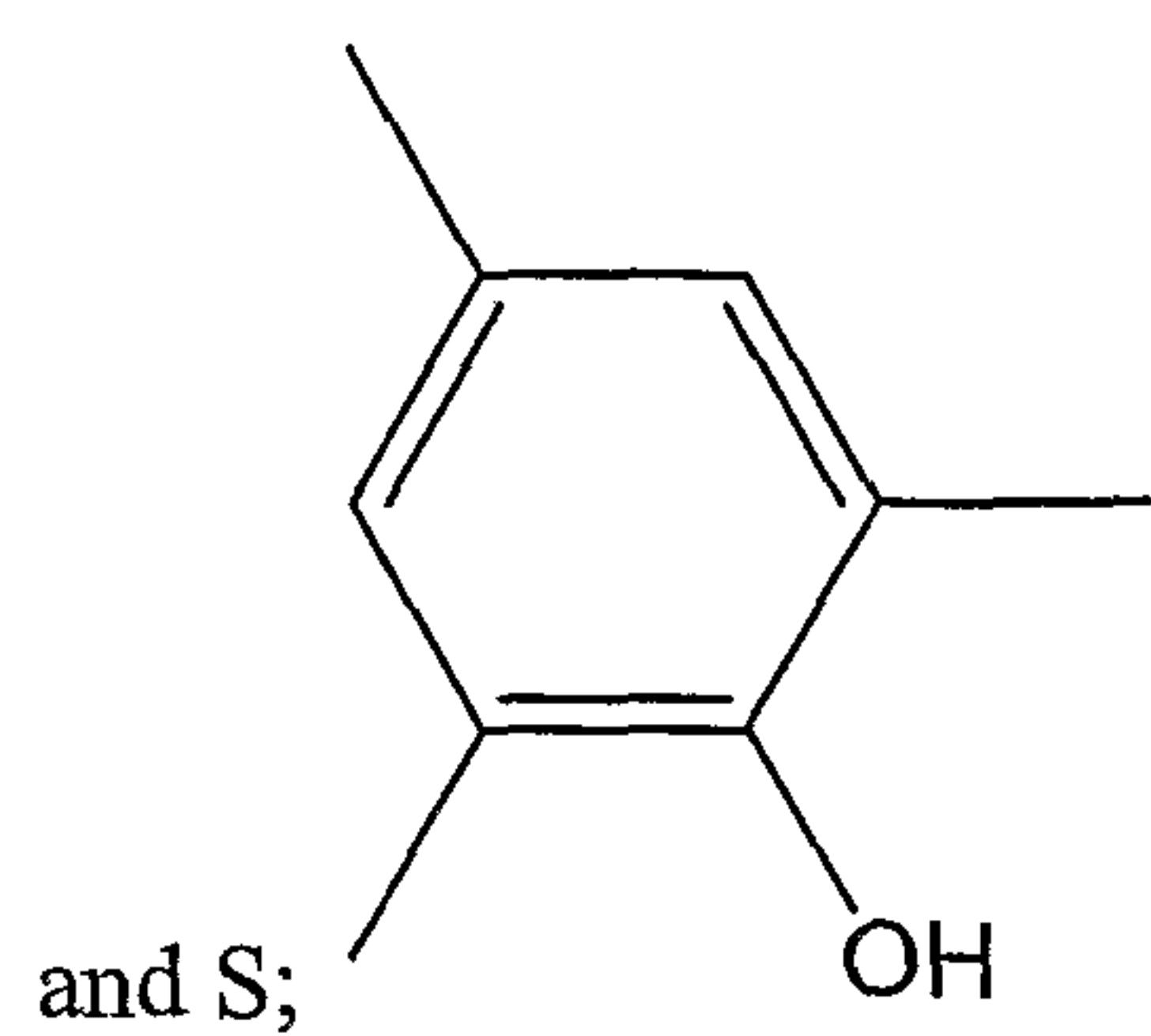
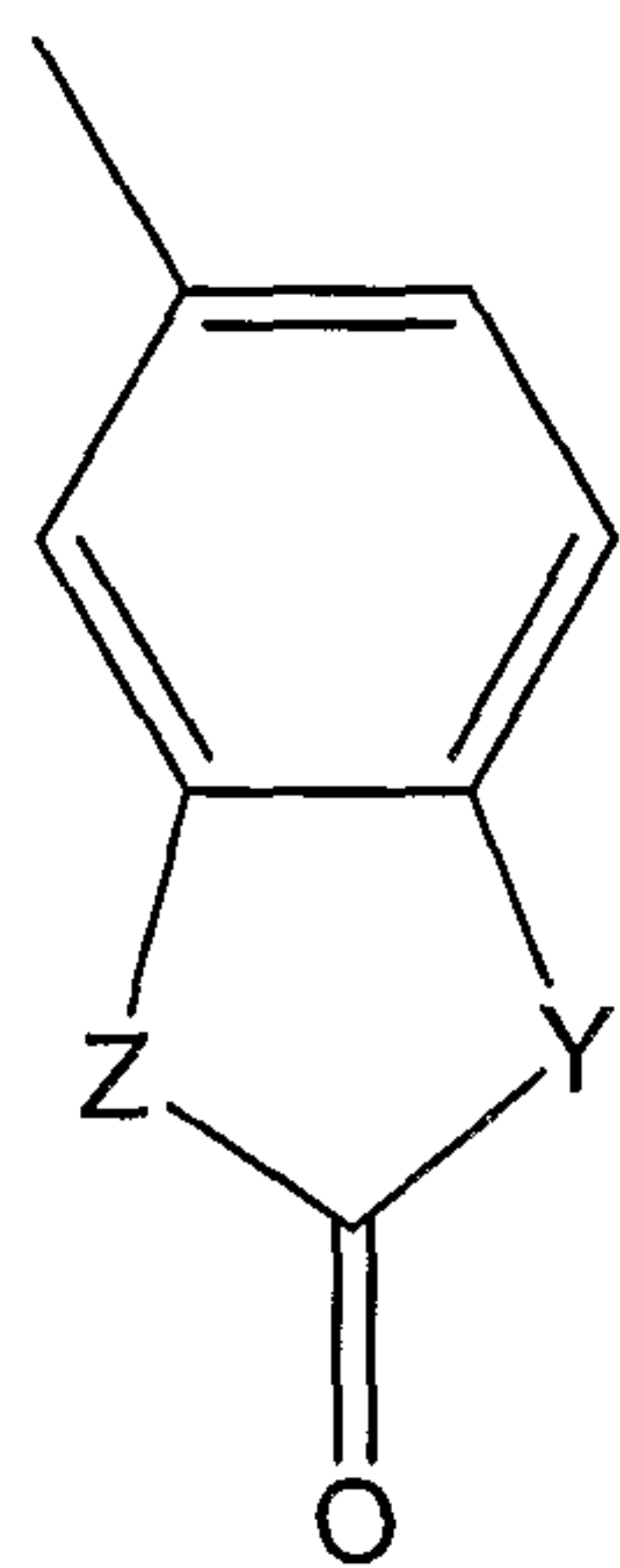
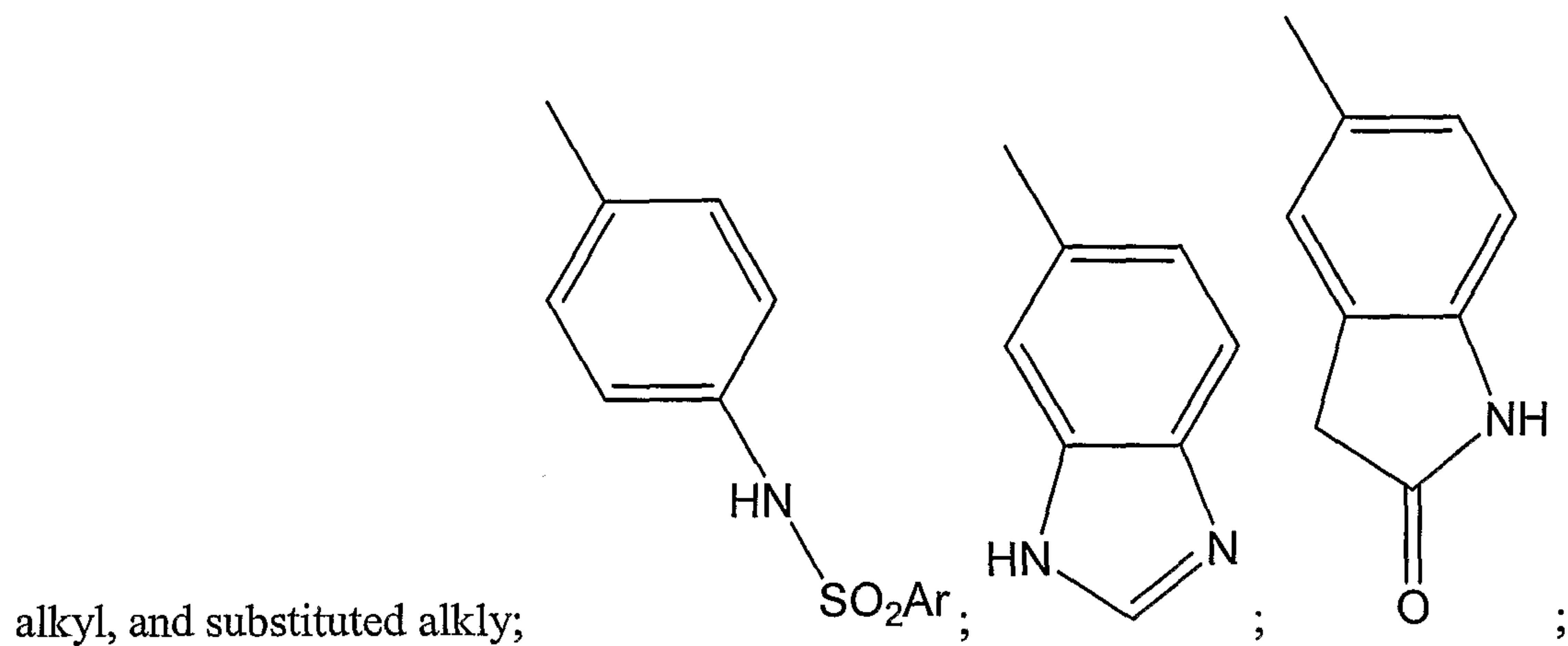
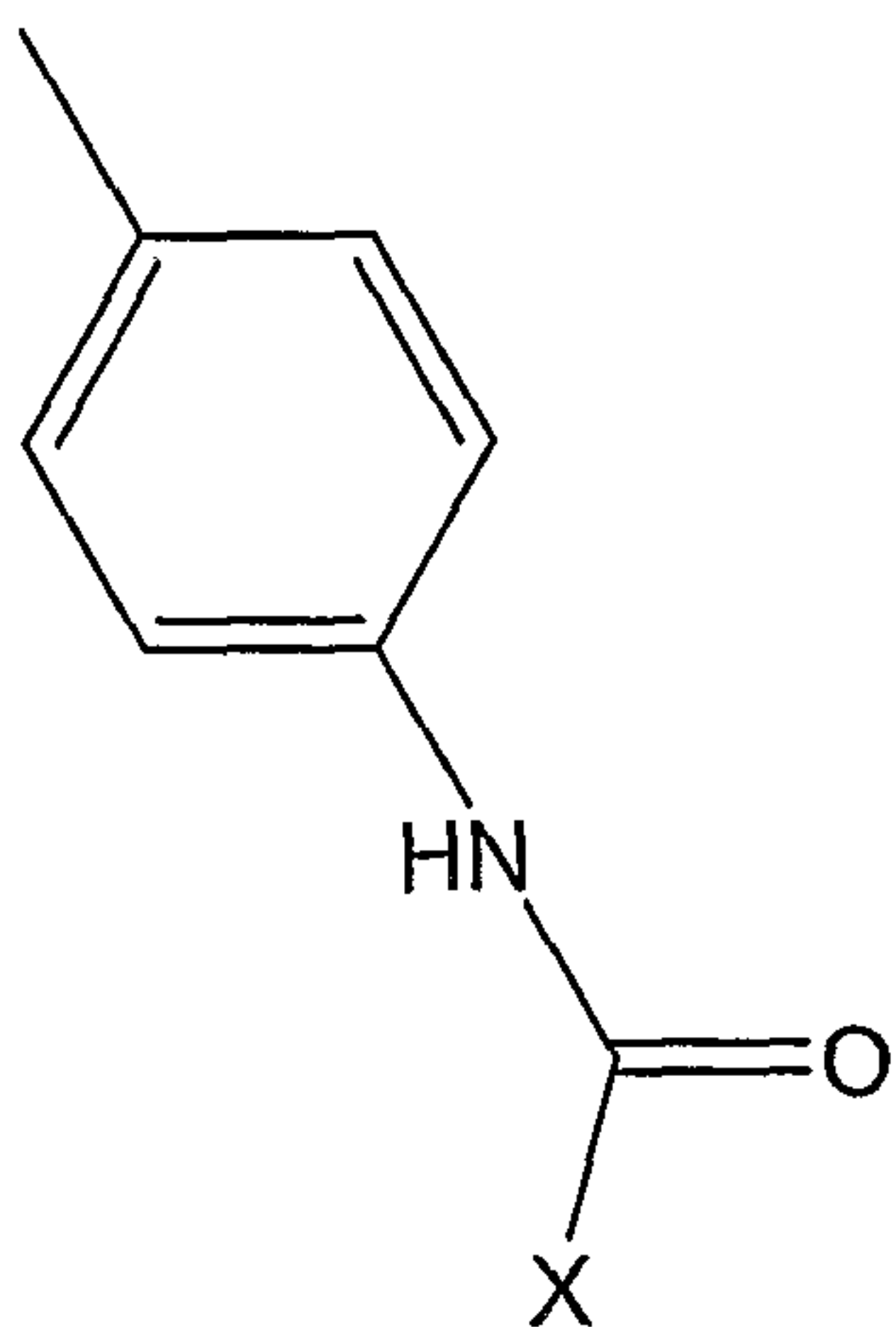




In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



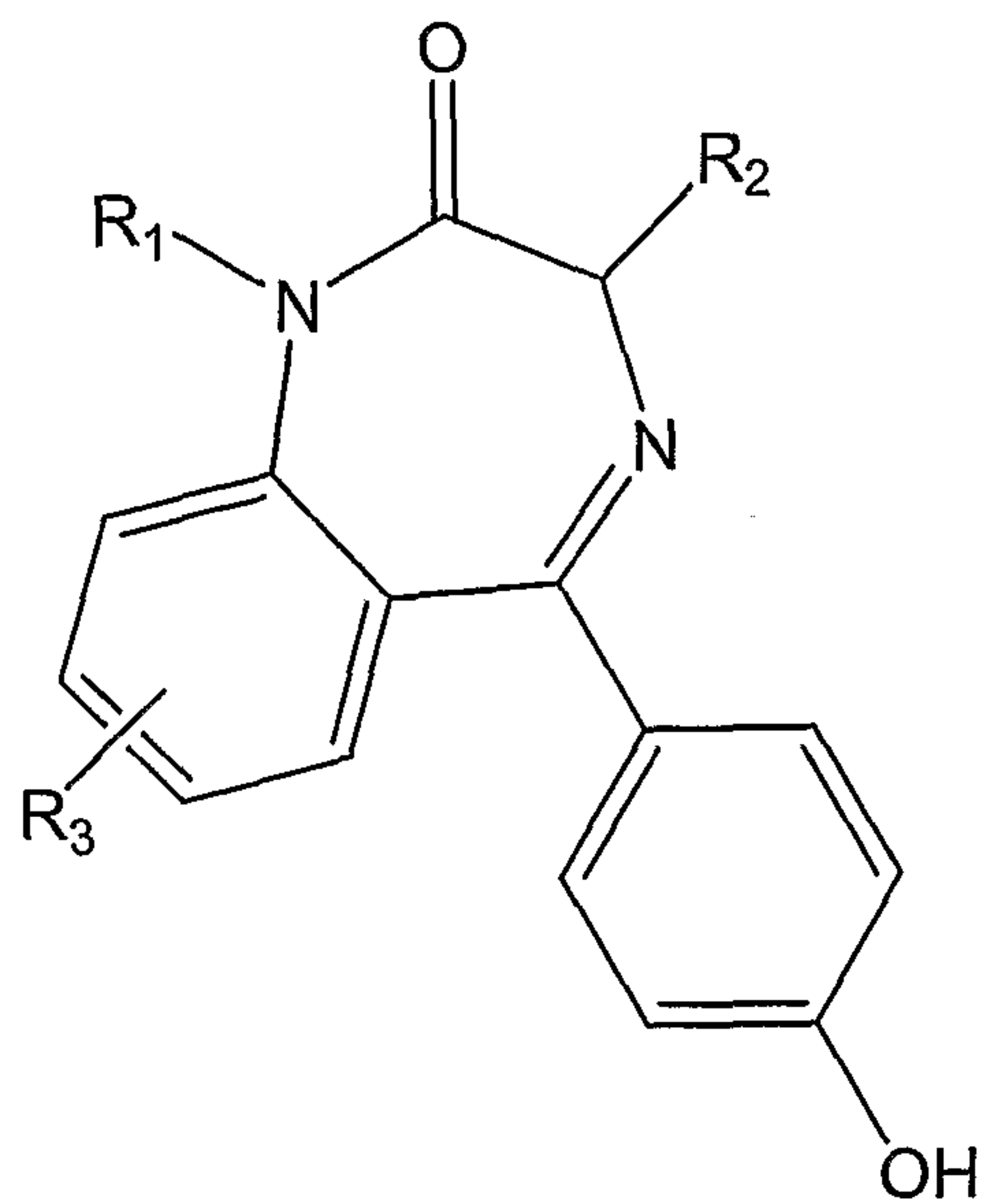
- 5 including both R and S enantiomeric forms and racemic mixtures; wherein R1 is selected from the group consisting of:



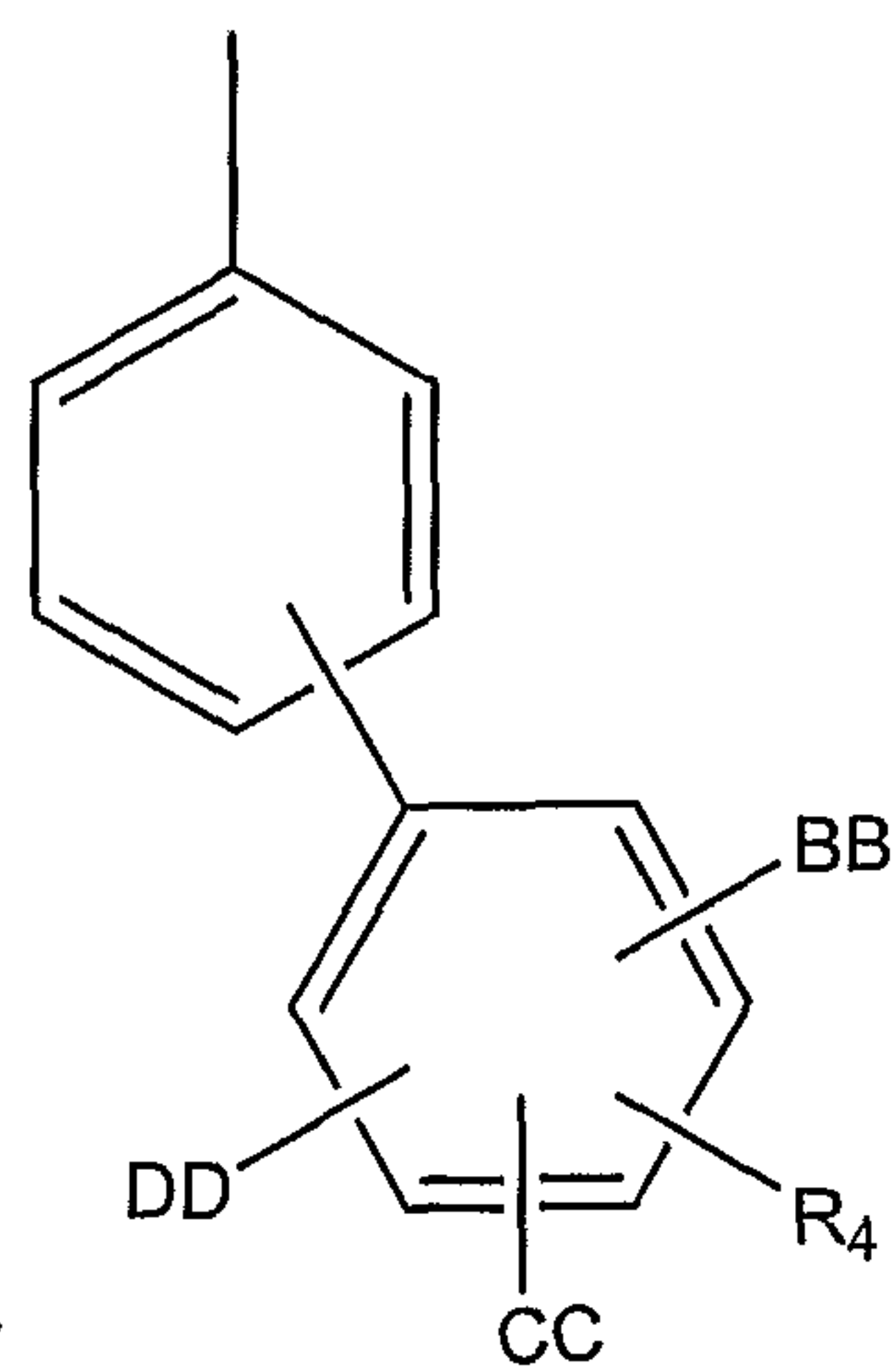
5 wherein R2 is selected from the group consisting of methyl, H, alkyl, and $(\text{CH}_2)_n$ -morpholino wherein $n=1-6$; and wherein R3 is

selected from the group consisting of hydrogen, halogen, alkyl, substituted alkyl, carboxylic acid, amide SO_2NH_2 , $\text{NHSO}_2\text{alkyl}$, and NO_2 .

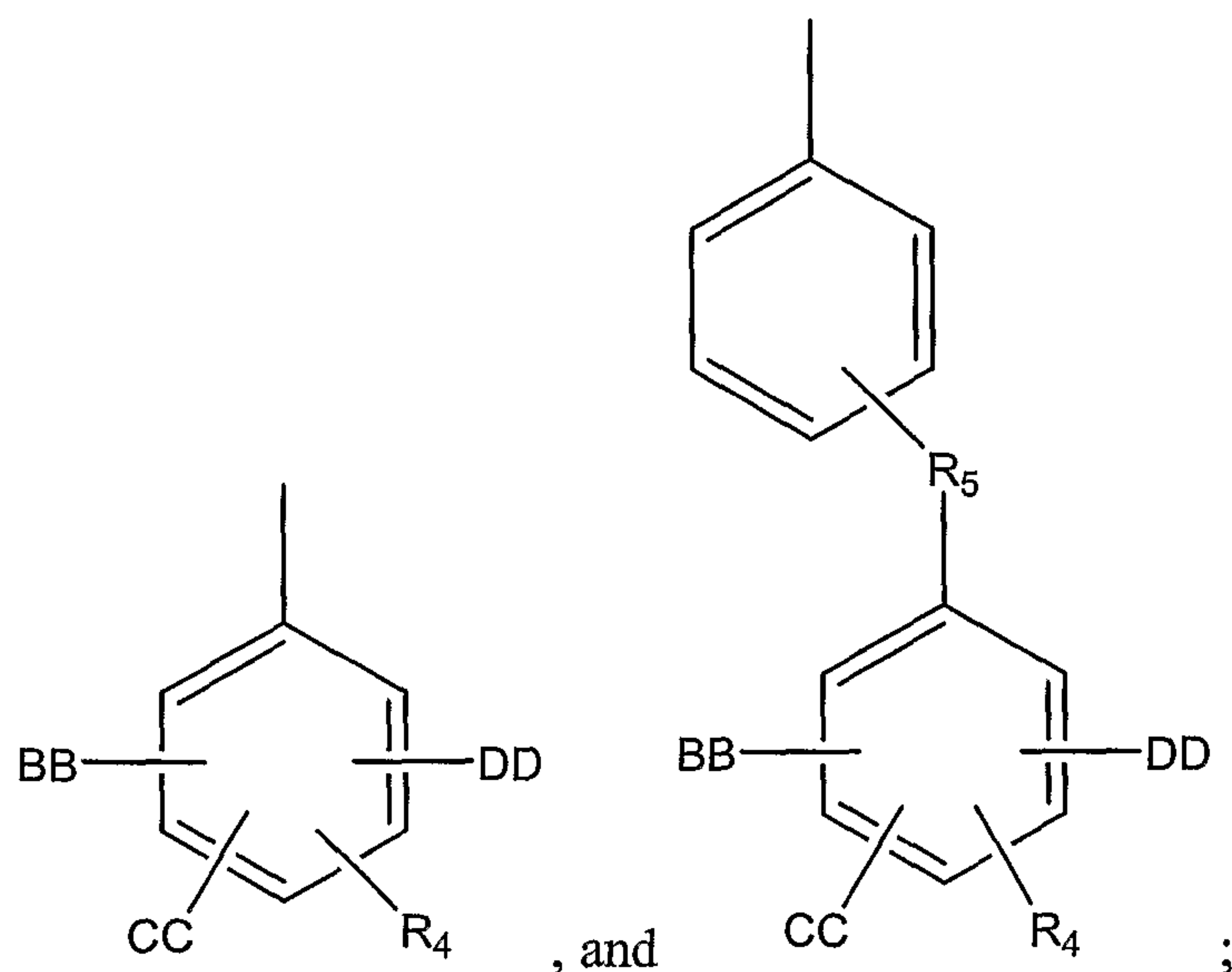
In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



5 wherein R1 is selected from the group consisting of methyl, hydrogen, alkyl, and $(\text{CH}_2)_n$ -morpholino wherein $n=1-6$;

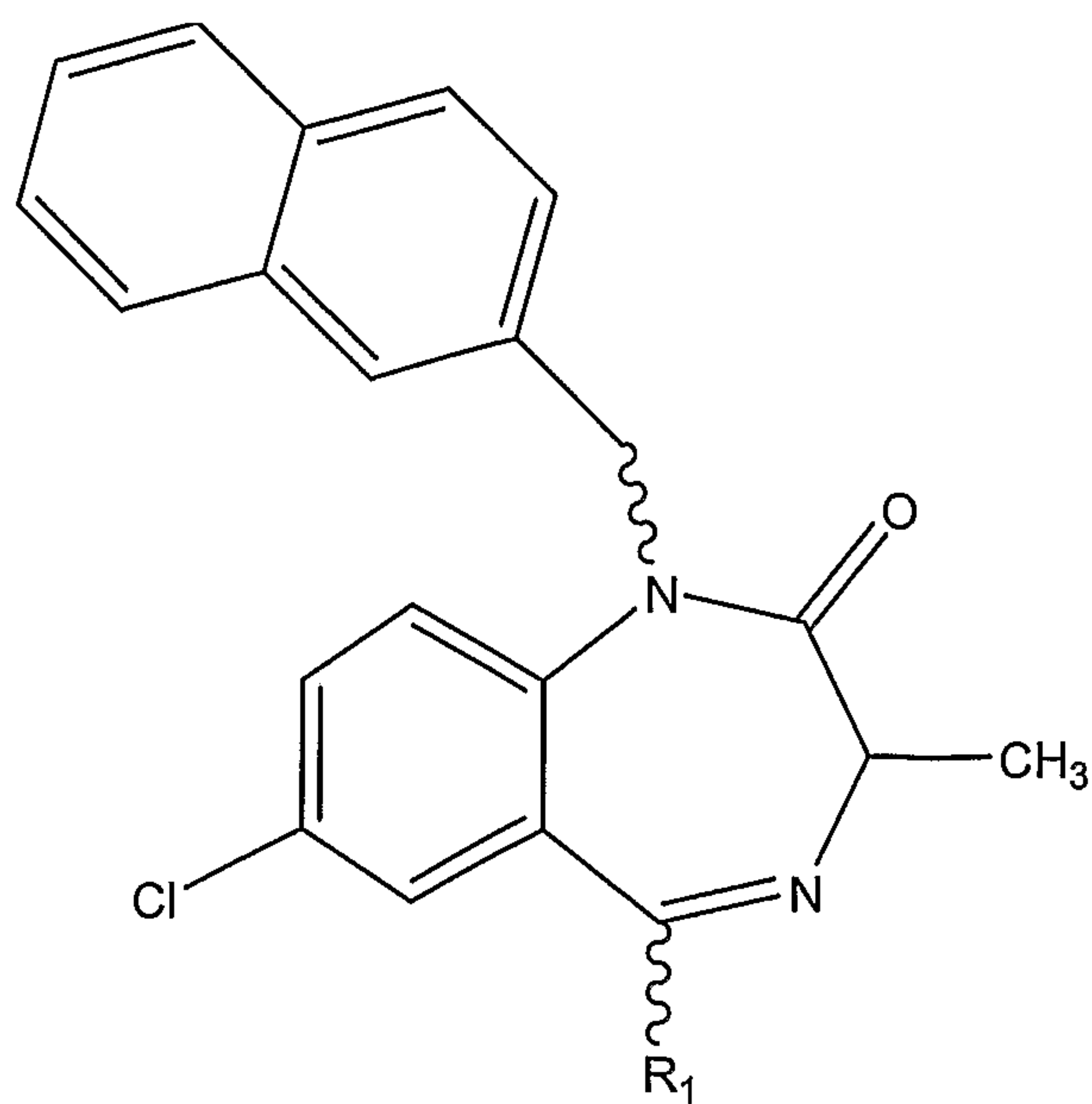


wherein R2 is selected from the group consisting of

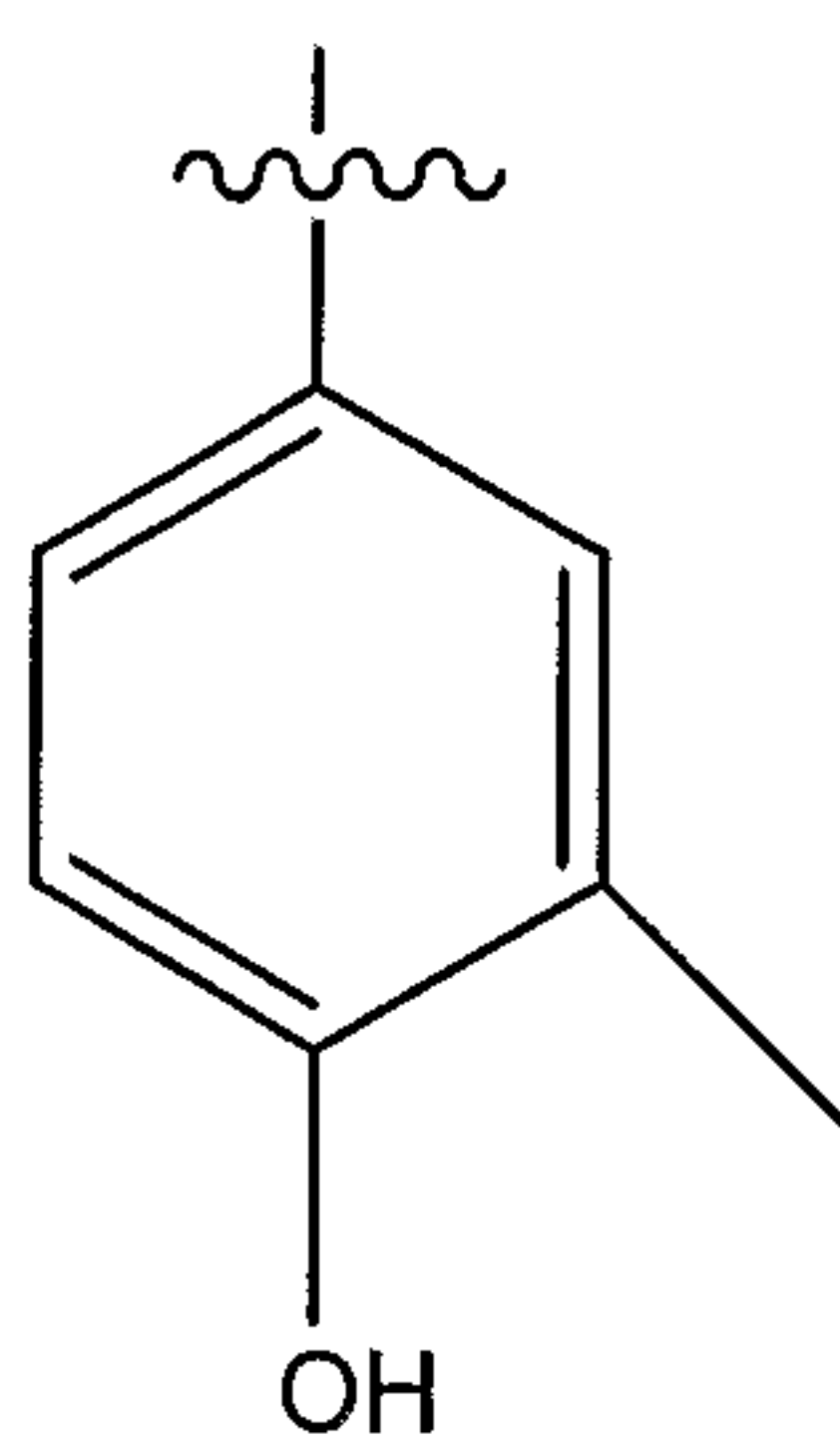


wherein R3 is selected from the group consisting of hydrogen, halogen, alkyl, substituted alkyl, carboxylic acid, amide, SO₂NH₂, NHSO₂alkyl, and NO₂; wherein BB, CC, DD, and R4 are present or absent, and are selected from the group consisting of hydrogen, CF₃, NO₂, alkyl, halogen, OH, O-alkyl, nitro, OCH₂CH₂OH, SO₂H, mono-substituted alkyl, di-substituted alkyl, tri-substituted alkyl, CO₂H, heterocycle, SO₂NH₂, SO₂NH-alkyl, NHSO₂alkyl, methyl ester, propyl ester, and ethyl ester; and wherein R5 is selected from the group consisting of NHSO₂, CH₂NHSO₂, CH₂CH₂NHSO₂, CH₂CH₂CH₂NHSO₂, SO₂NH, SO₂NHCH₂, SO₂NHCH₂CH₂, SO₂NHCH₂CH₂CH₂, CH₂, CH₂CH₂, CH₂CH₂CH₂, SO₂, CH₂SO, SOCH₂, OCH₂CH₂O, SO, CH₂CH₂SO, and SOCH₂CH₂.

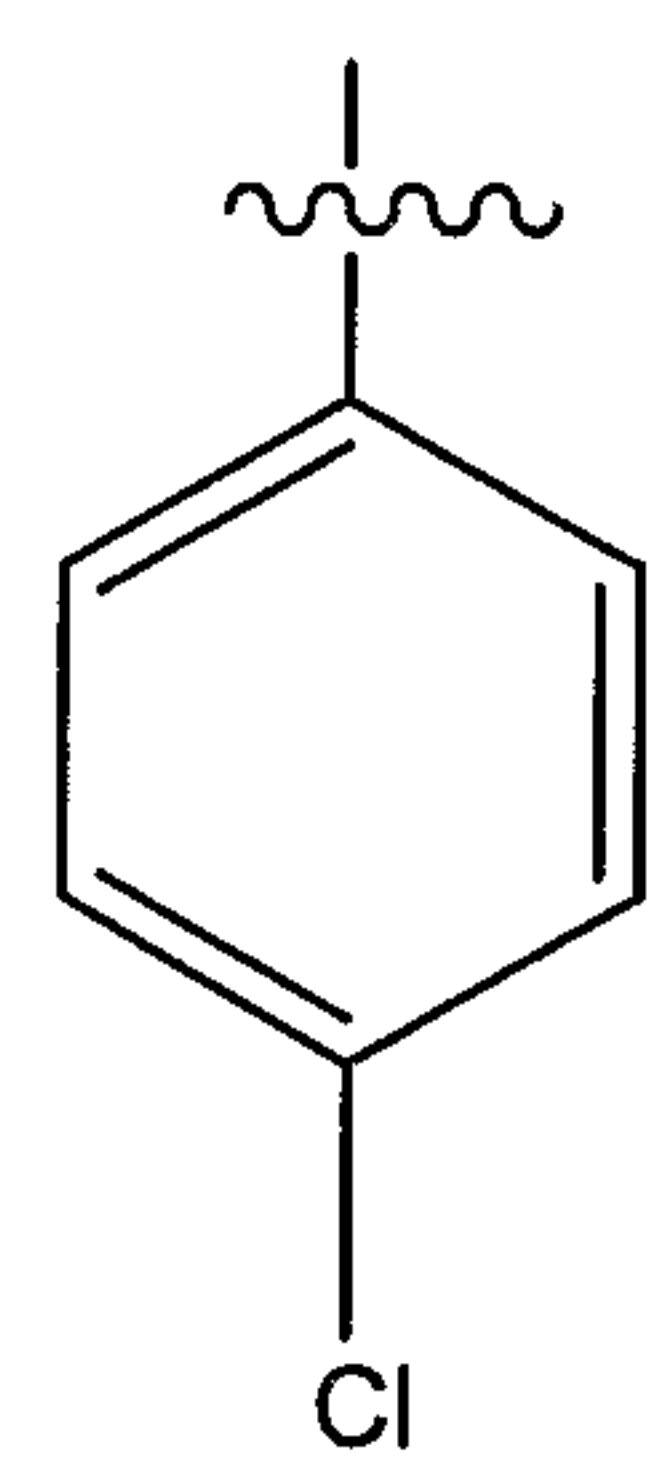
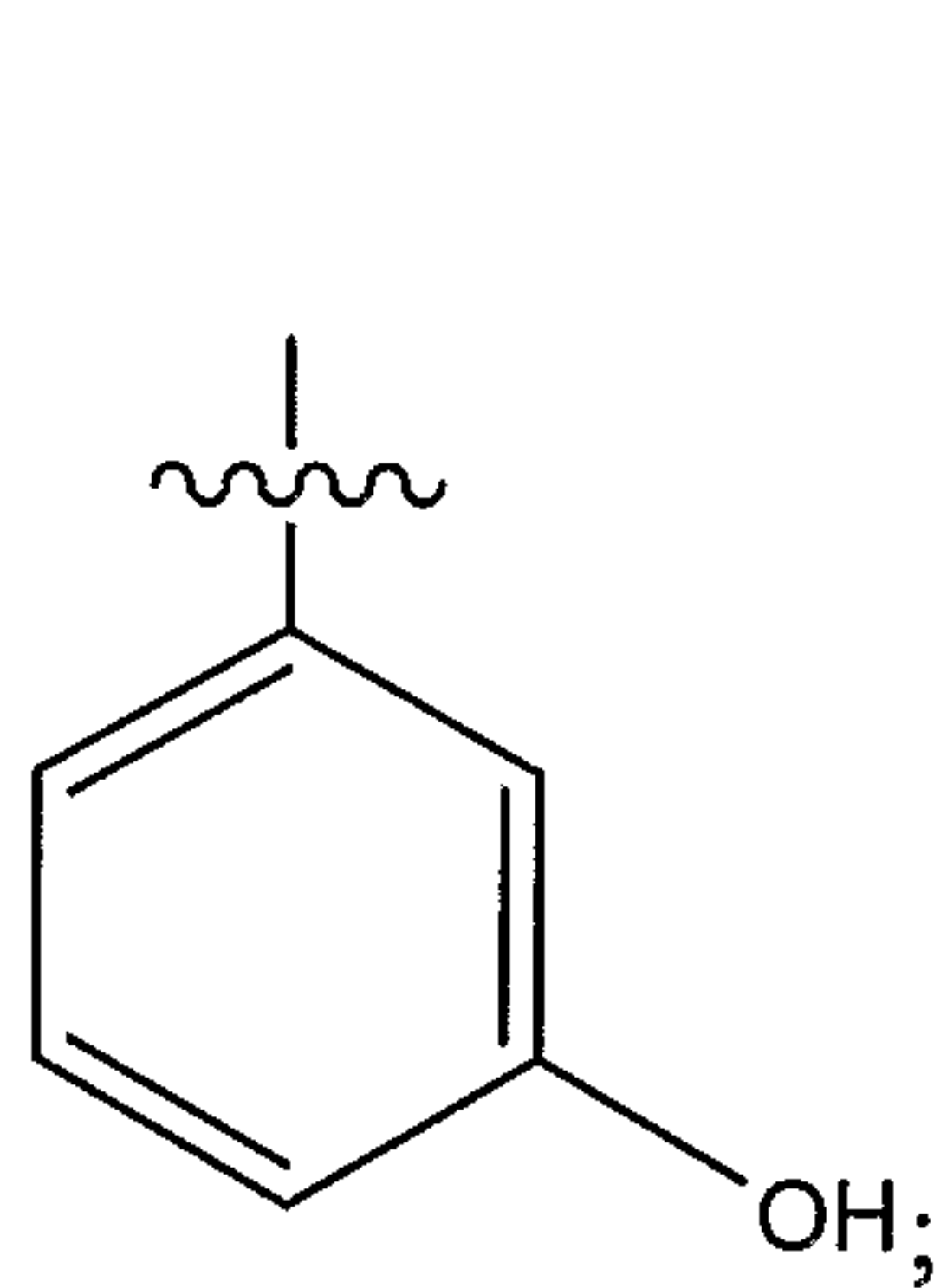
In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



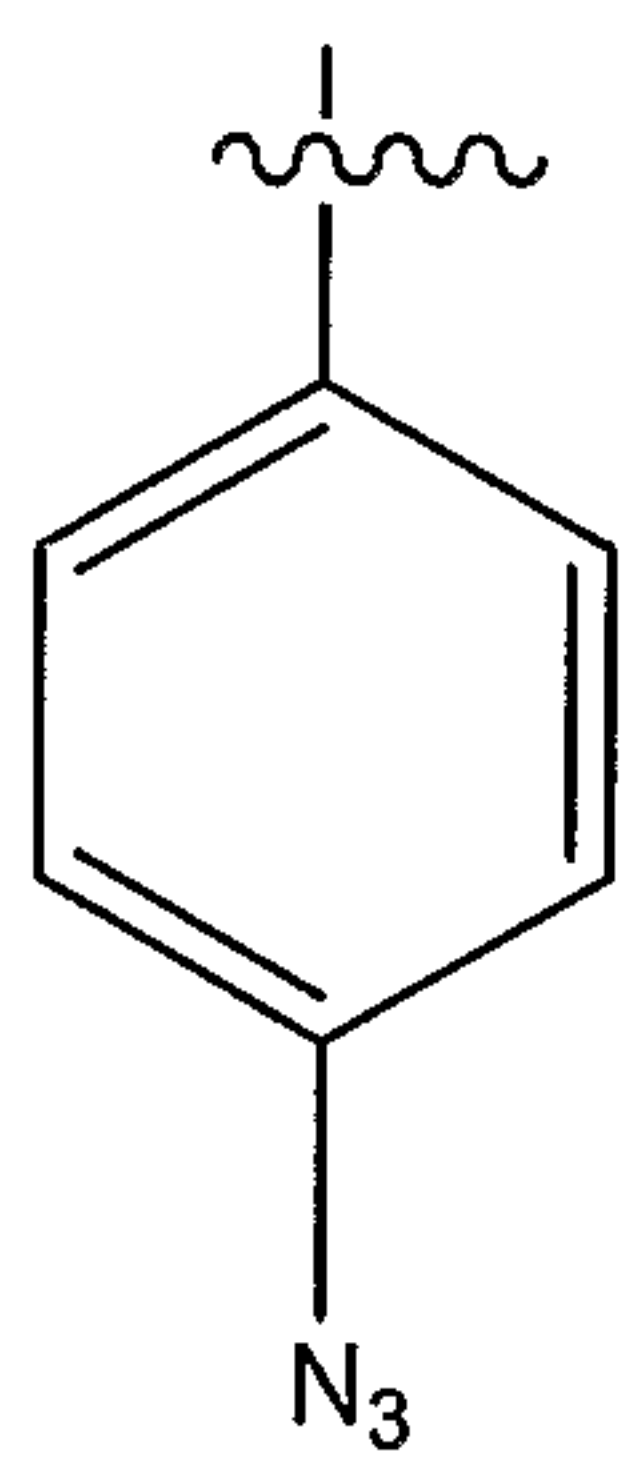
;



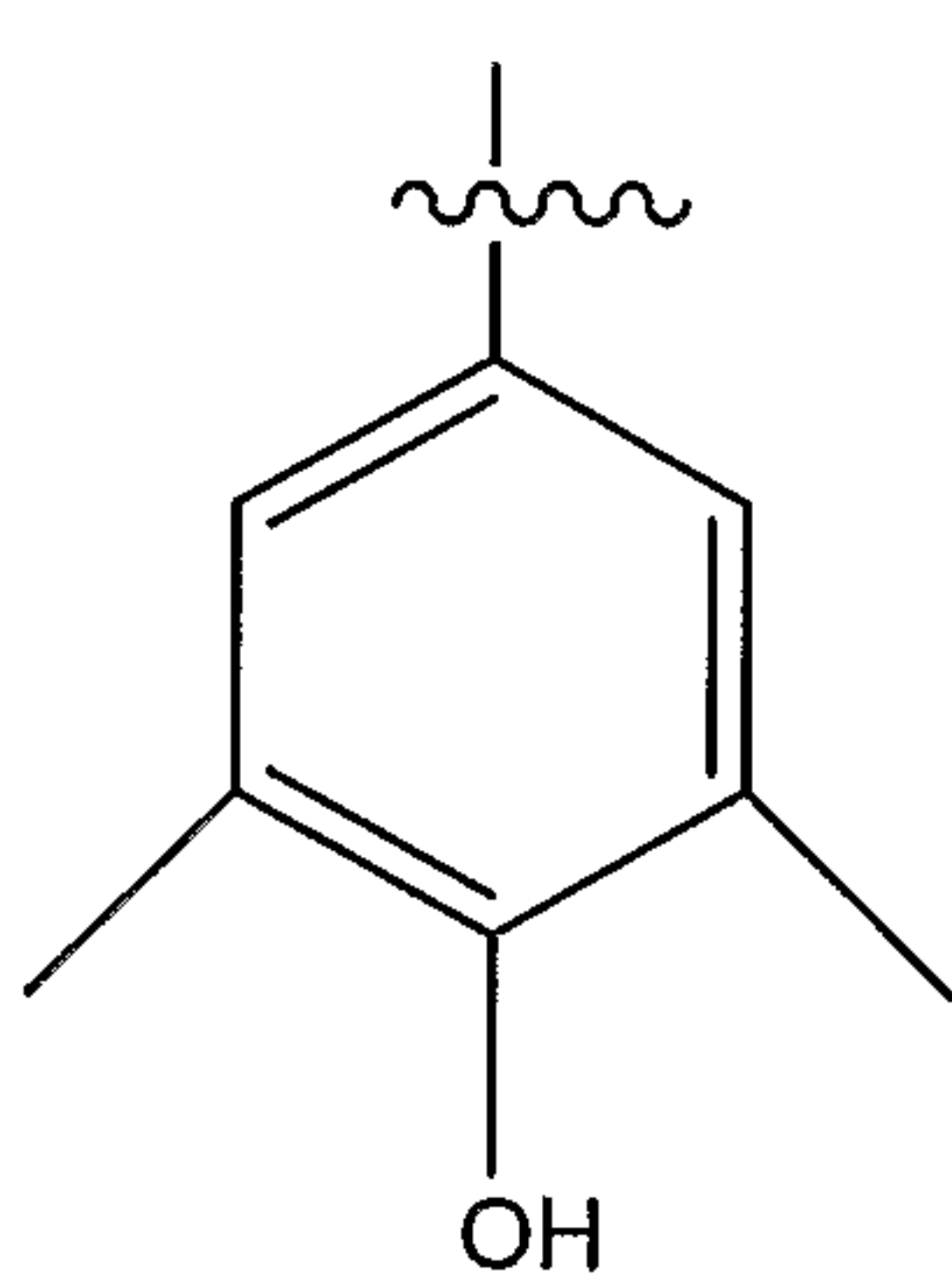
;



;

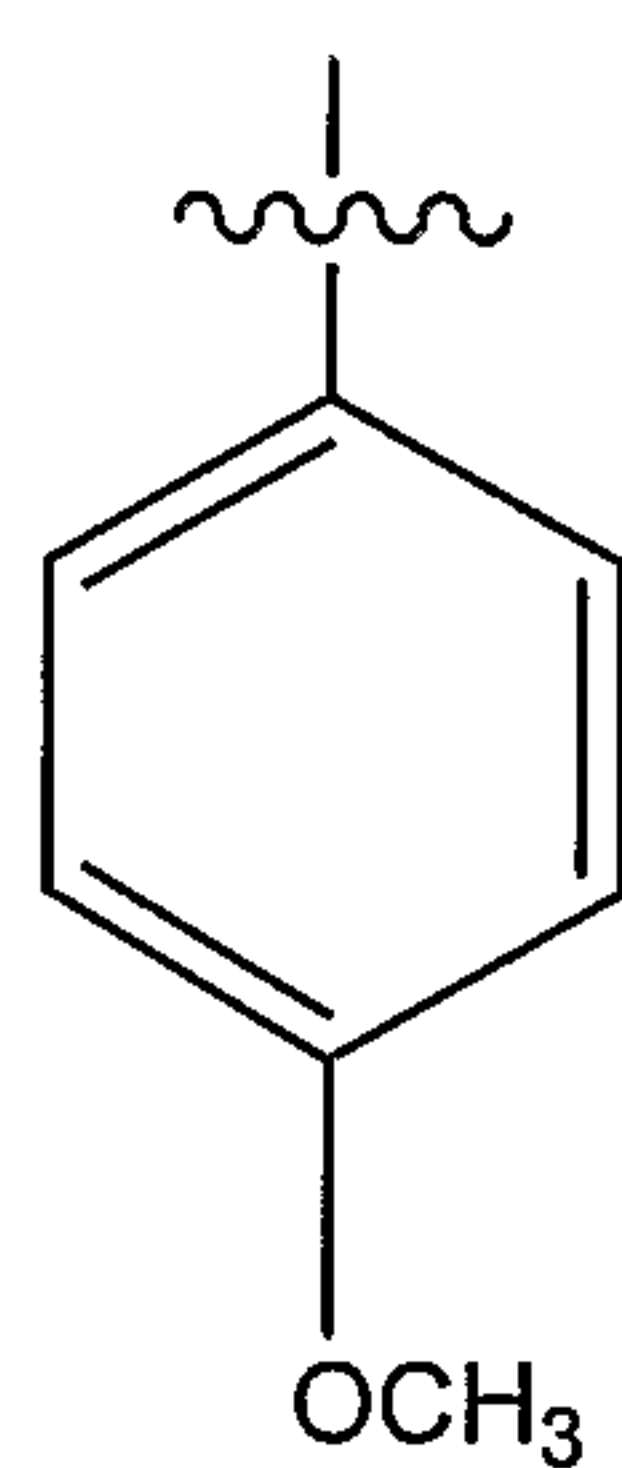
wherein R₁ is selected from:

;



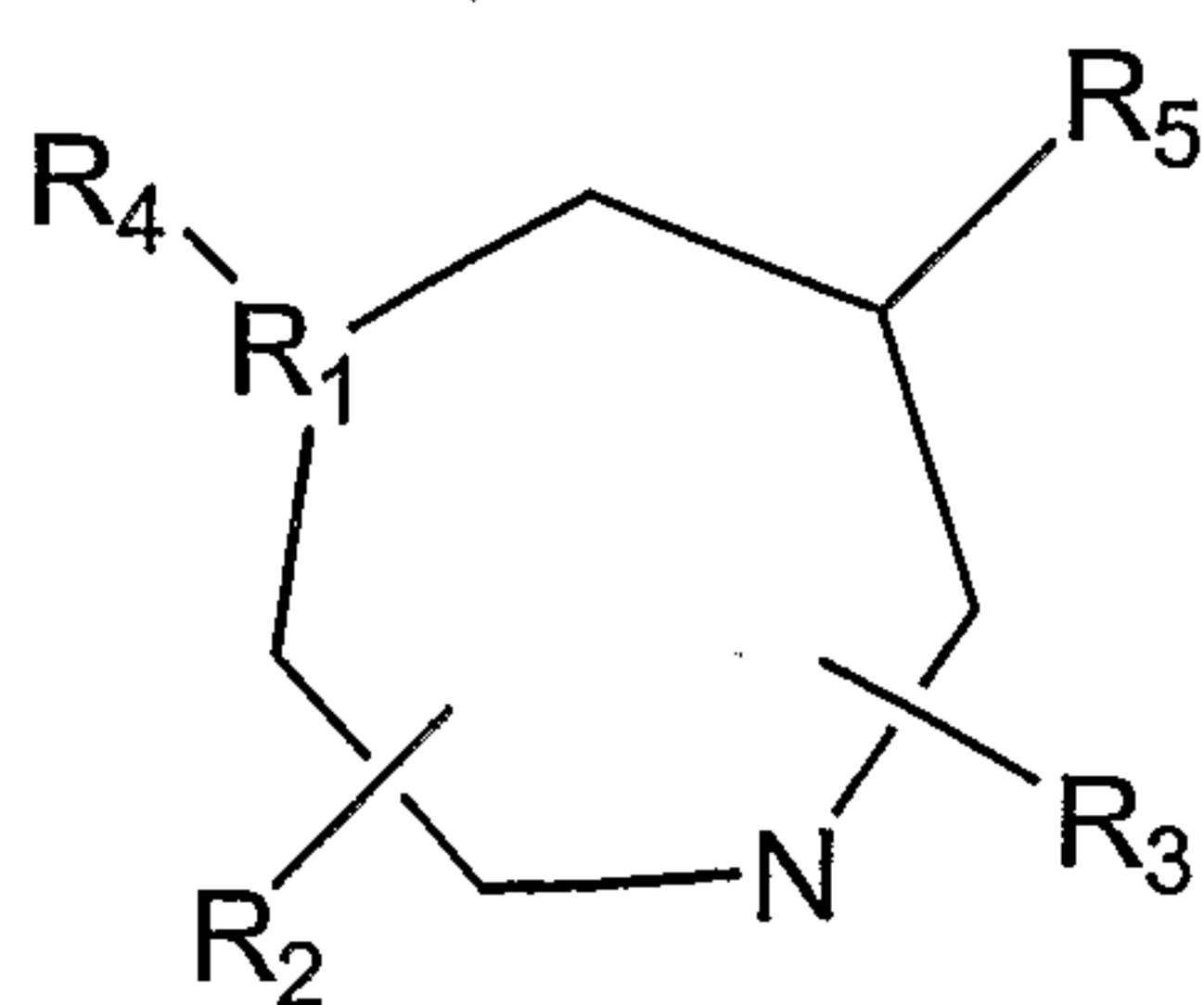
;

and



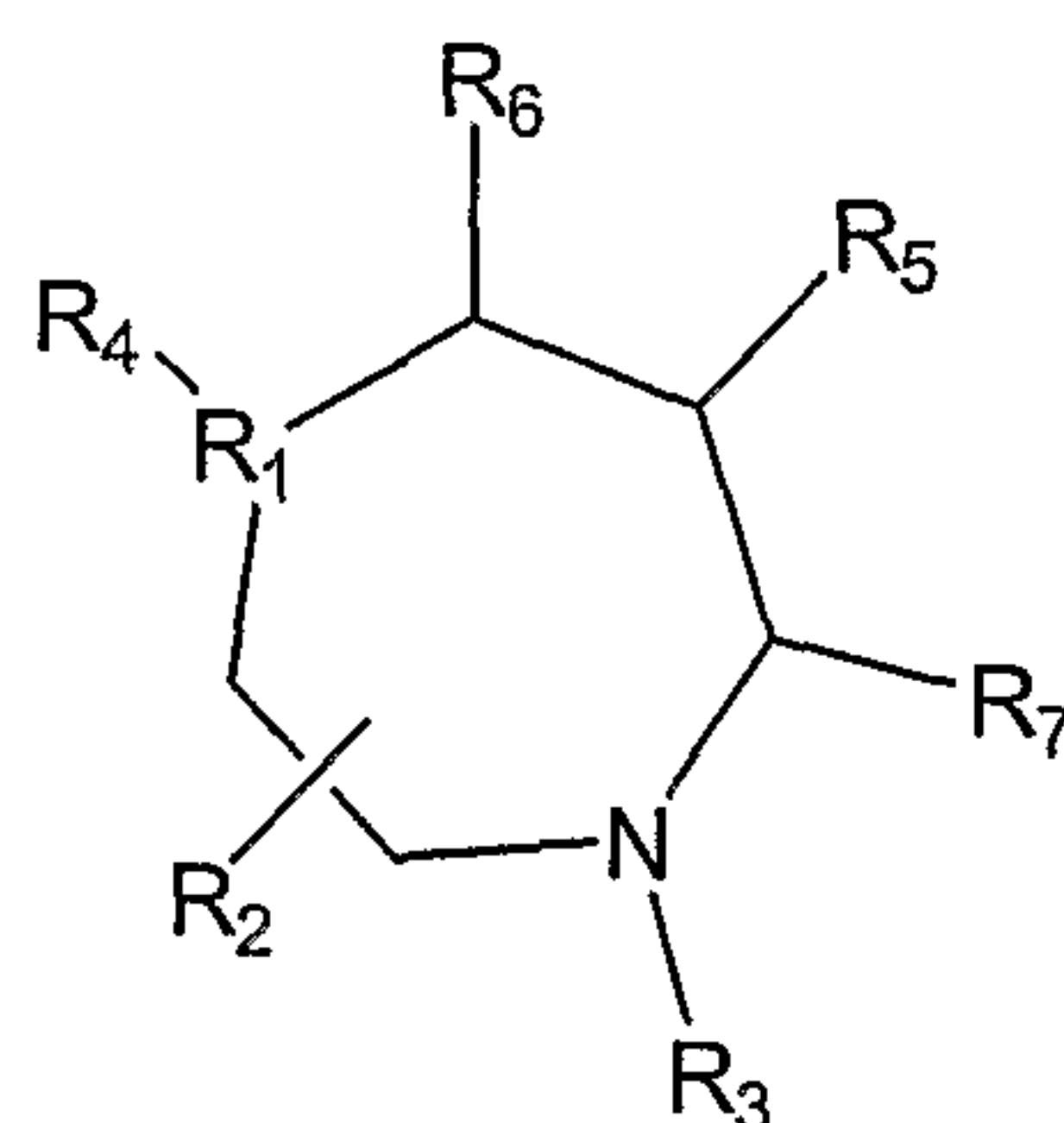
.

In certain preferred embodiments, crystal forms and formulations of the following
 5 exemplary compounds are provided:



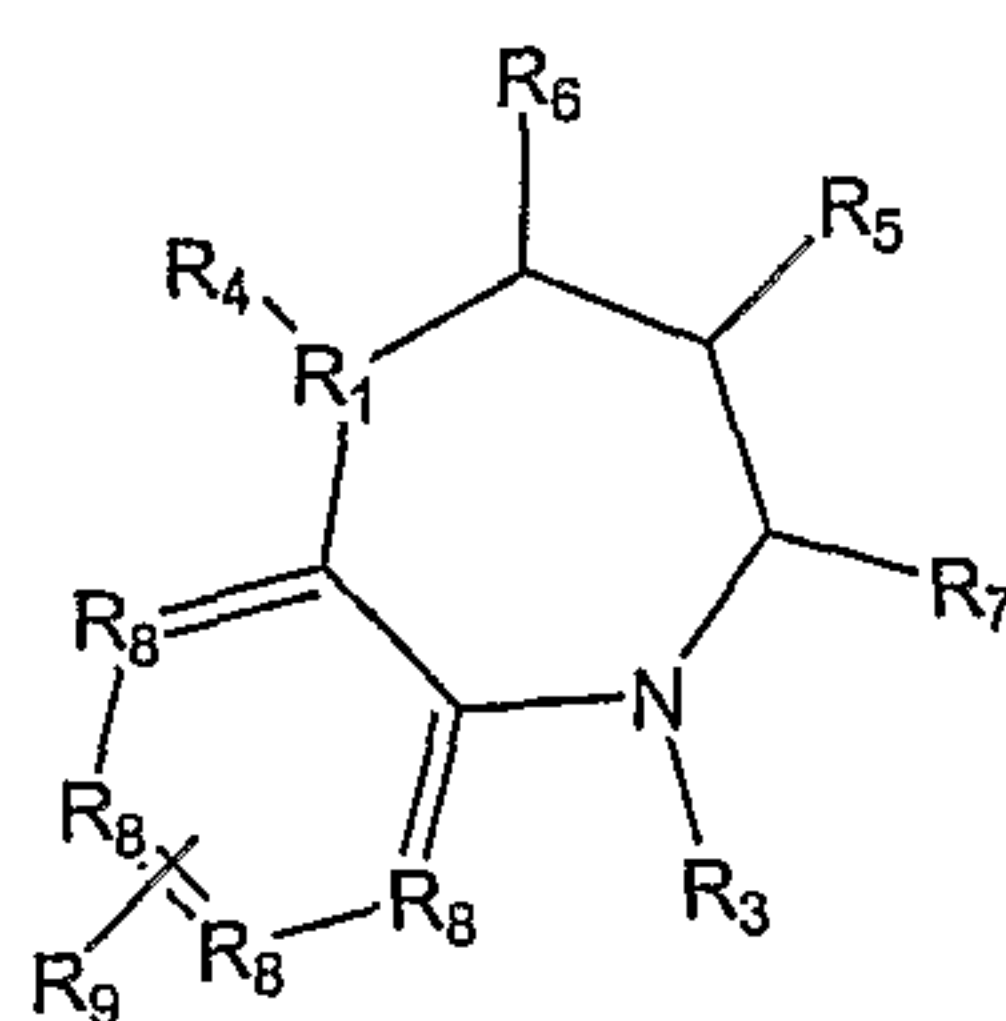
including both R and S enantiomeric forms and racemic mixtures; wherein R1 is a nitrogen atom or a carbon atom; wherein R2 is comprises a chemical moiety comprising a heterocyclic group containing 3 or more carbon atoms; wherein R3 comprises a chemical moiety comprising a heterocyclic group containing 3 or more carbon atoms; and wherein
 5 R4 and R5 are separately selected from the group consisting of: hydrogen; halogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; a chemical moiety comprising a halogen; a chemical moiety comprising Sulfur; a chemical moiety comprising Nitrogen; an aromatic chemical moiety; a hydrophilic chemical moiety; and a hydrophobic chemical moiety.

10 In preferred embodiments, the compound comprises the formula:



wherein R6 is selected from the group consisting of H and a ketone; and wherein R7 is selected from the group consisting of H and a ketone.

In preferred embodiments, the compound comprises the formula:

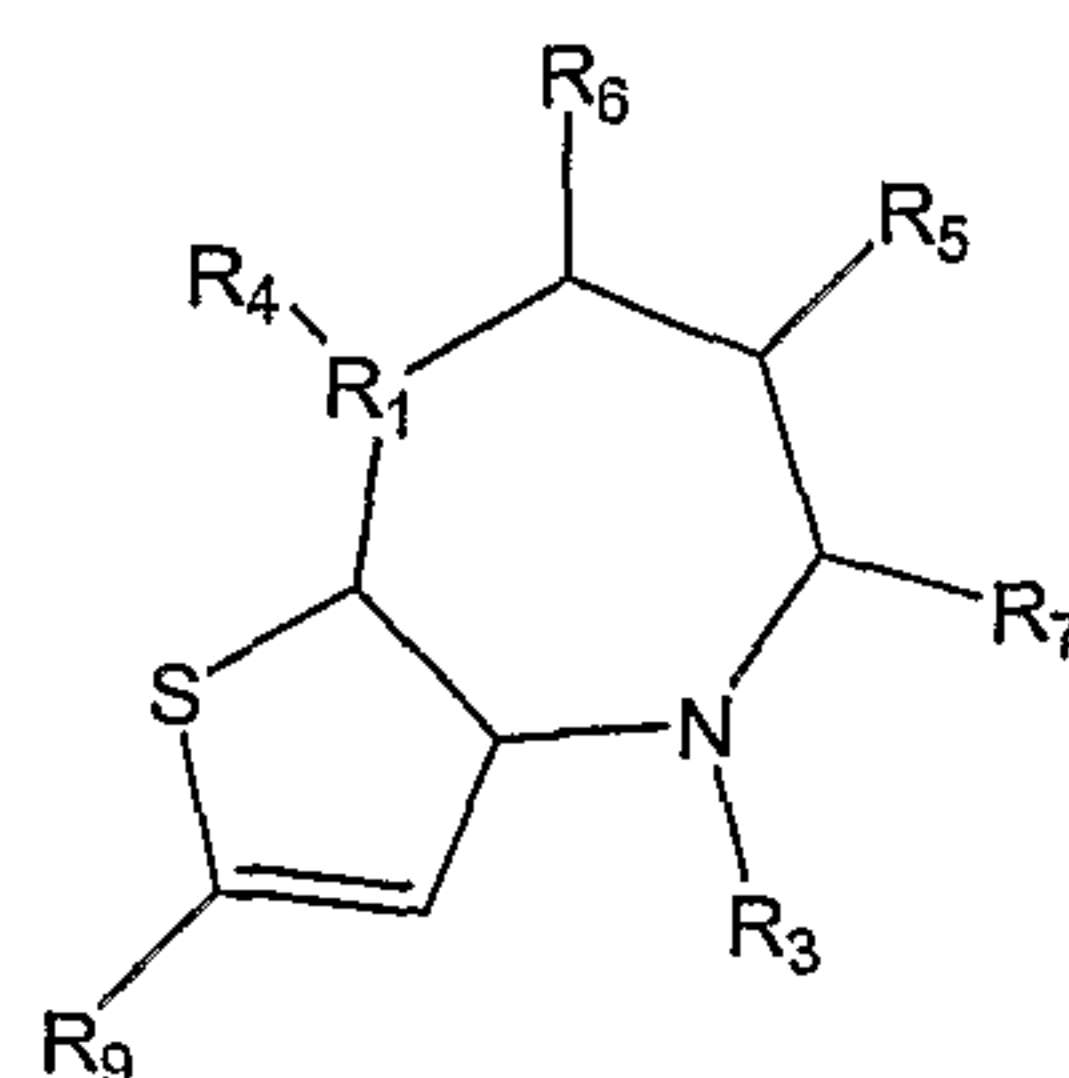


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In such preferred embodiments, R8 is carbon or nitrogen and R9 is selected from H, a hydroxy, an alkoxy, a halogen, an amino, a lower-alkyl, a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of less than 10
 20 carbons, a substituted cycloaliphatic group, an aryl, a heterocyclic, NO₂; SR'; and NR'₂, wherein R' is defined as a linear or branched, saturated or unsaturated aliphatic chain having at least one carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxyl subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one
 25 thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least

2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup.

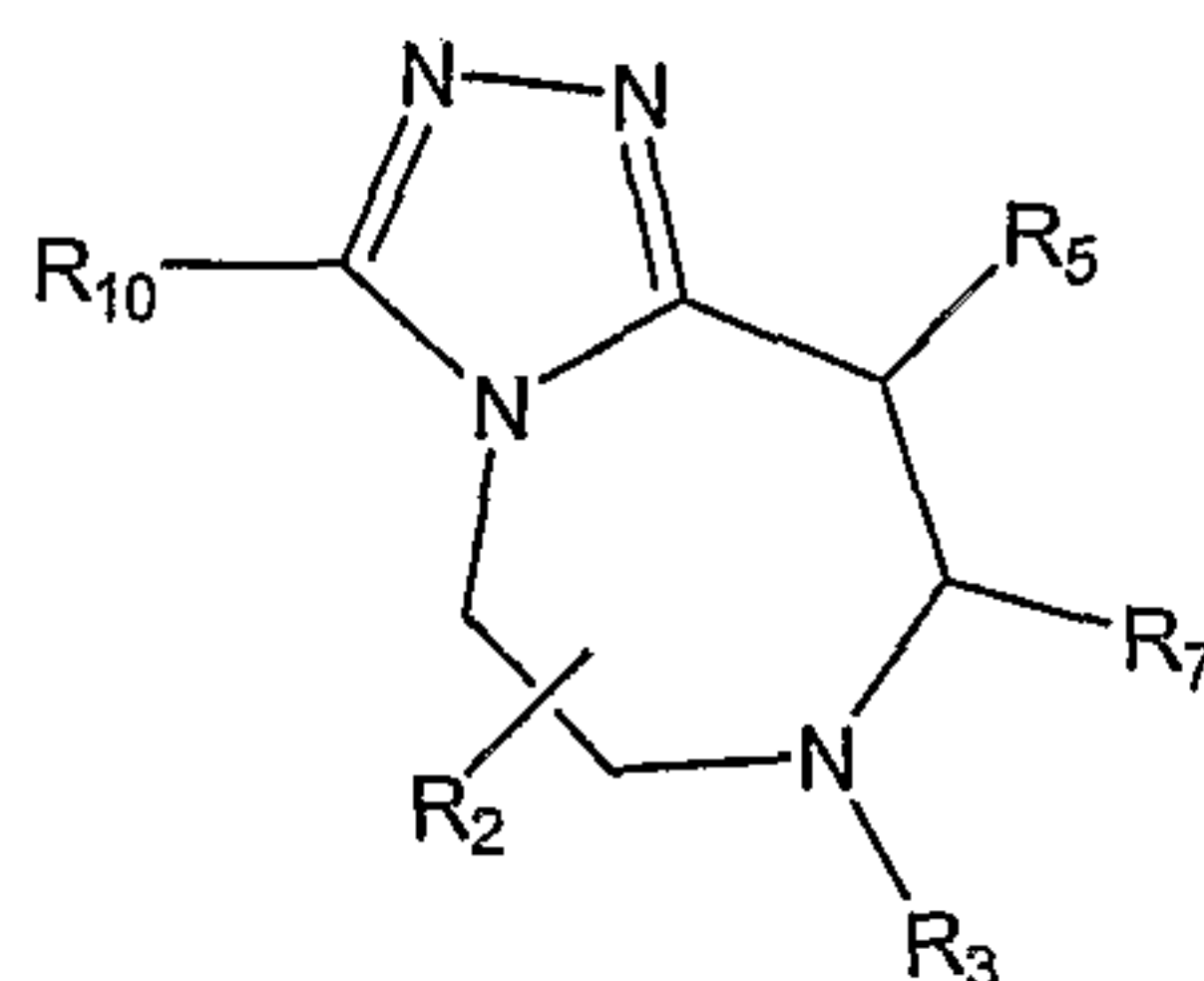
15 In preferred embodiments, the compound comprises the formula:



wherein R9 is selected from H, a hydroxy, an alkoxy, a halo, an amino, a lower-alkyl, a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of less than 10 carbons, a substituted cycloaliphatic group, an aryl, a heterocyclic, NO₂; SR'; and NR'₂, wherein R' is defined as a linear or branched, saturated or unsaturated aliphatic chain having at least one carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxyl subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic

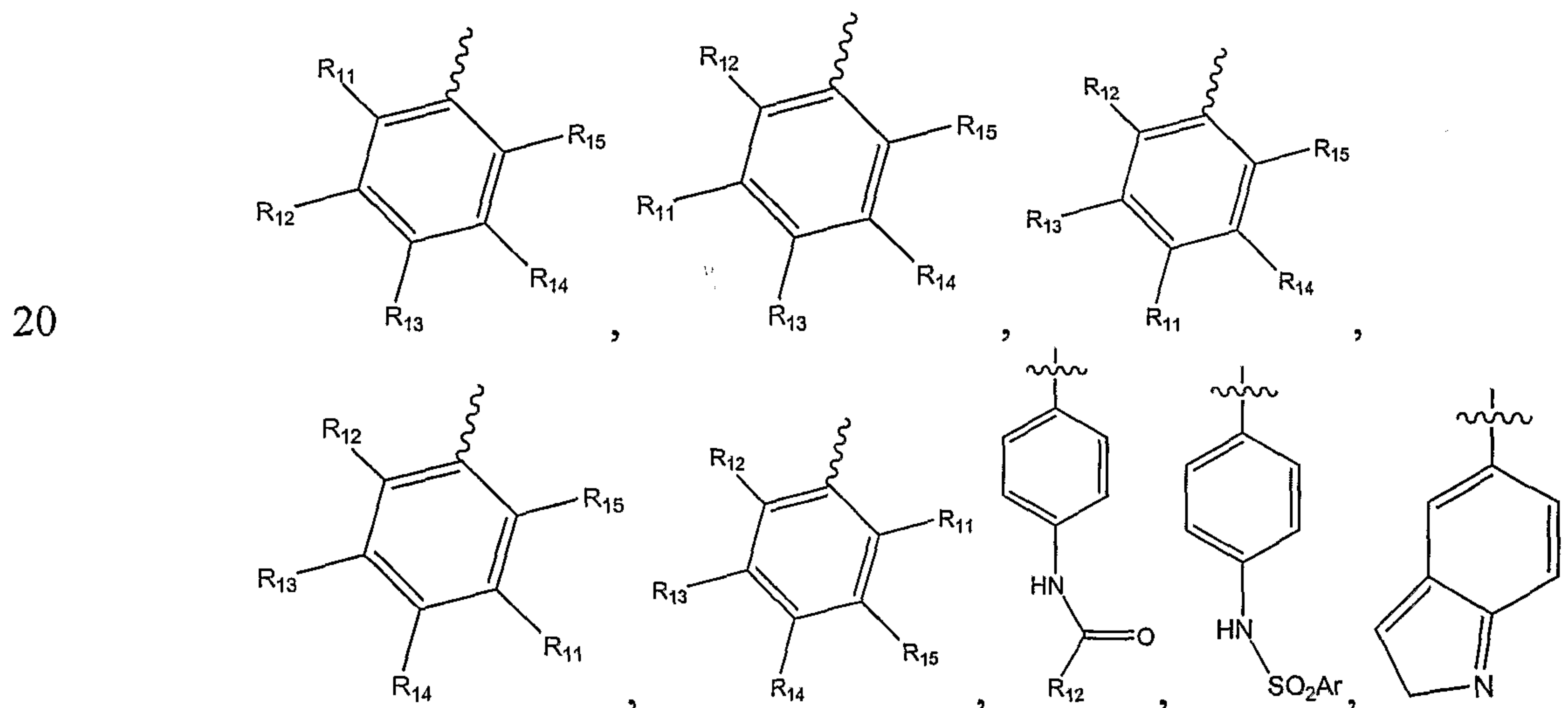
chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup.

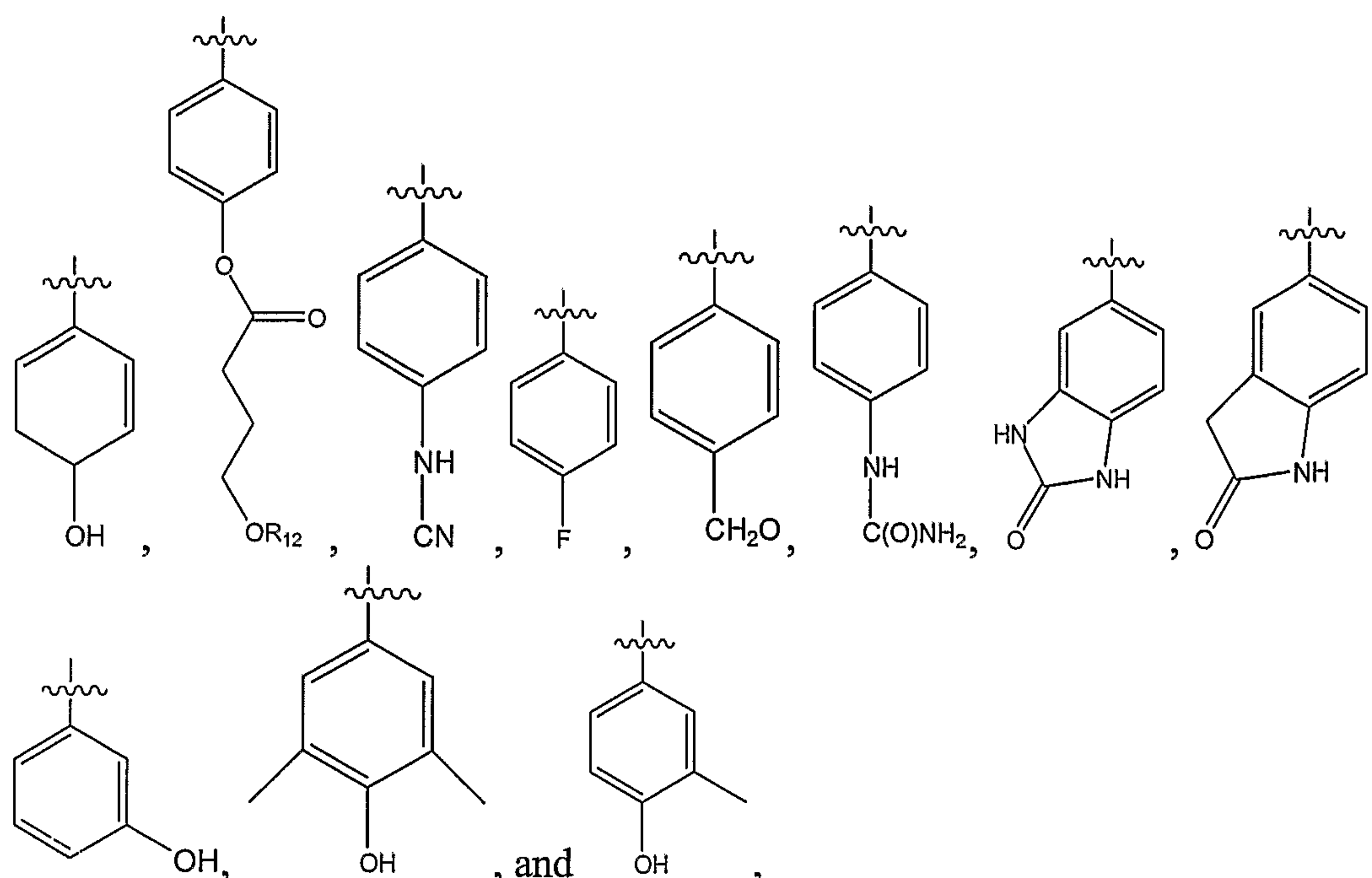
In preferred embodiments, the compound comprises the formula:



wherein R10 is selected from the group consisting of: hydrogen; halogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; a chemical moiety comprising a halogen; a chemical moiety comprising Sulfur; a chemical moiety comprising Nitrogen; an aromatic chemical moiety; a hydrophilic chemical moiety; and a hydrophobic chemical moiety; and wherein R7 is selected from the group consisting of H and a ketone.

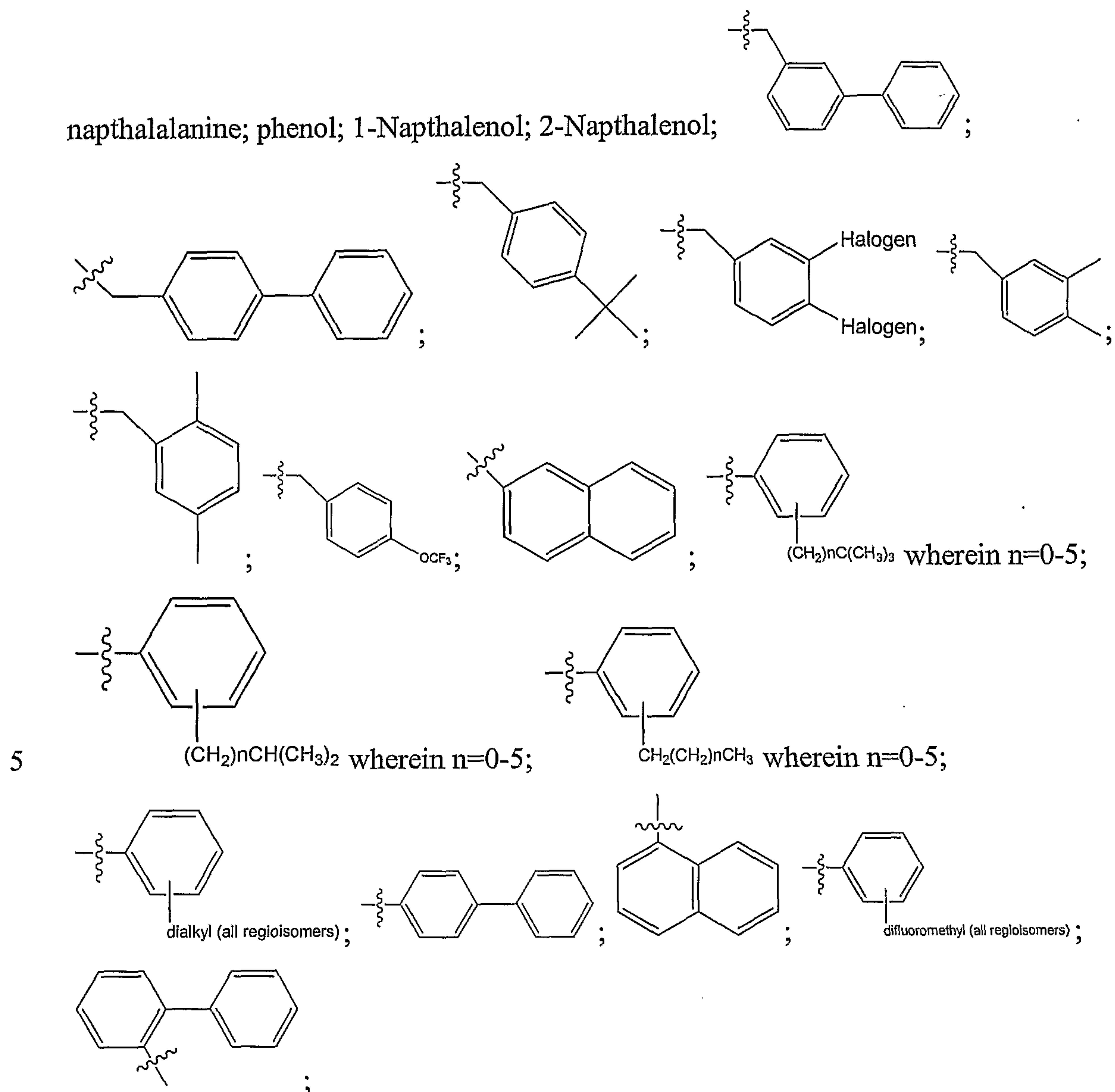
In other preferred embodiments, R3 is selected from the group consisting of:





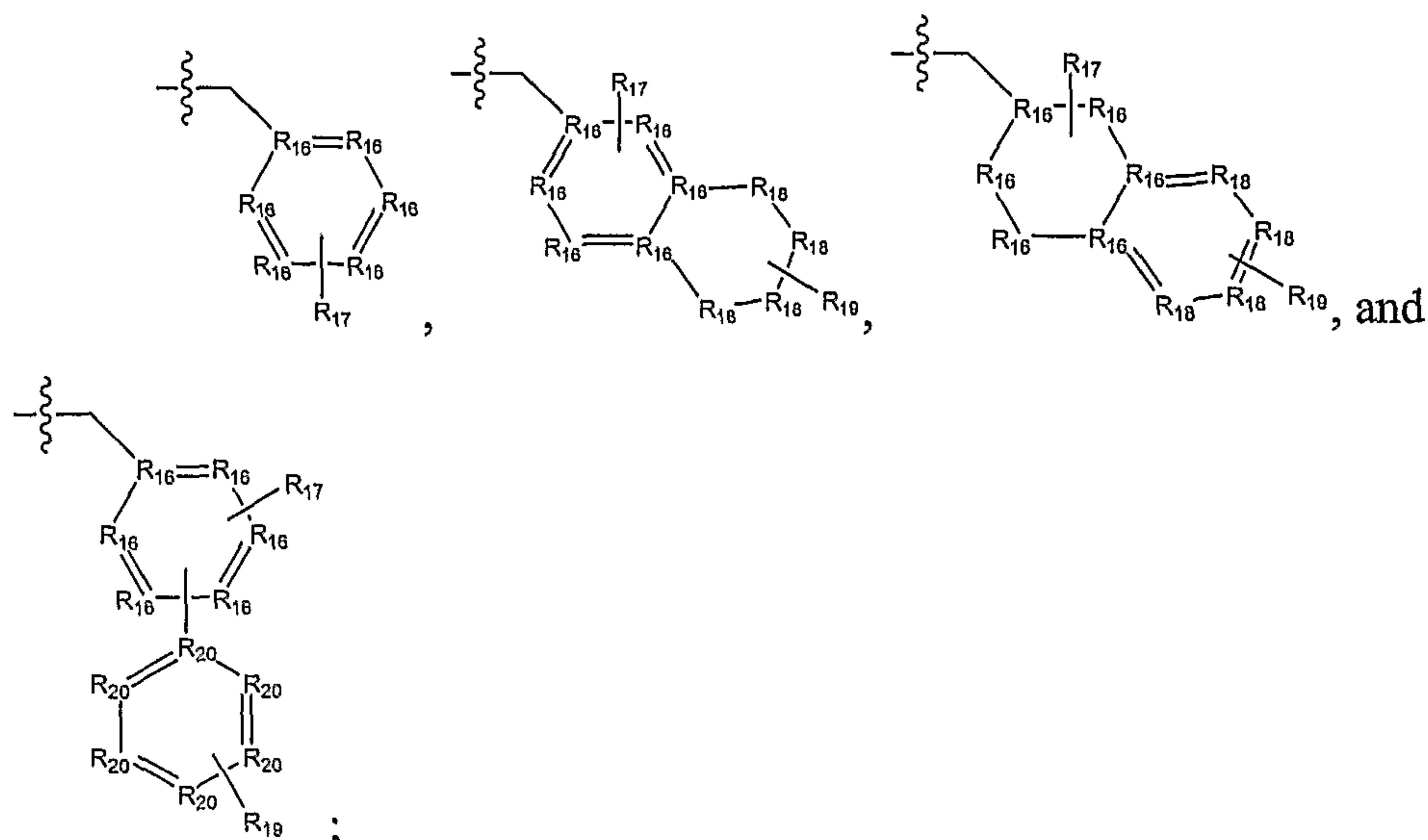
wherein R12, R13, R14 and R15 are selected from the group consisting of: hydrogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 1 carbon; a
 5 linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxy subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or
 10 unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain
 15 having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ether subgroup;
 20 a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; and R11 is OH.

In yet other preferred embodiments, R4 or R5 are selected from group consisting of:



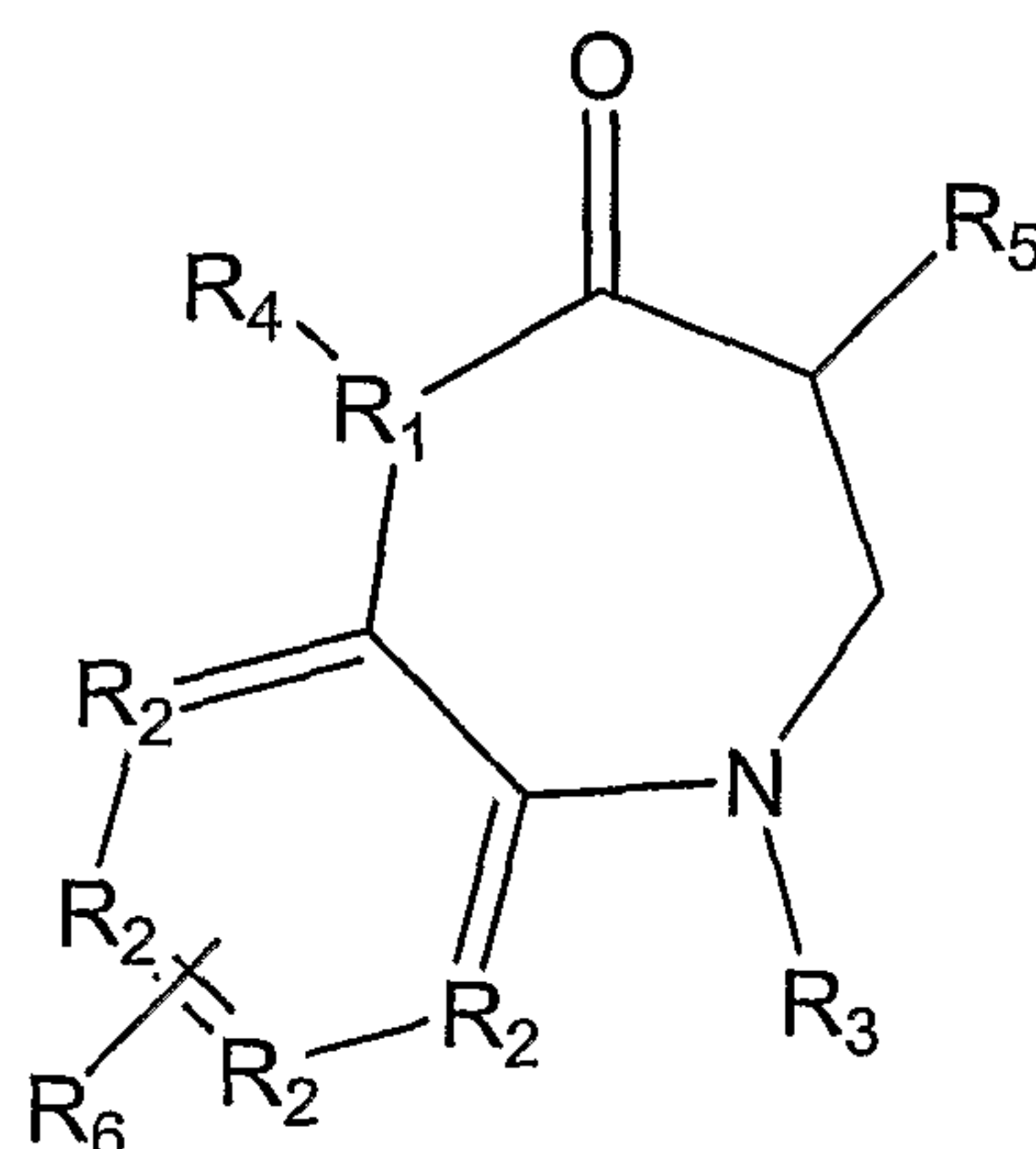
quinolines, and all aromatic regioisomers.

In other preferred embodiments, R4 or R5 is selected from the group consisting of:



- wherein R16 is carbon or nitrogen; wherein R17 is selected from the group consisting of hydrogen; halogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; a chemical moiety comprising a halogen; a chemical moiety comprising Sulfur; a chemical moiety comprising Nitrogen; an aromatic chemical moiety; a hydrophilic chemical moiety; and a hydrophobic chemical moiety; wherein R18 is carbon or nitrogen; wherein R19 is selected from the group consisting of hydrogen; halogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; a chemical moiety comprising a halogen; a chemical moiety comprising Sulfur; a chemical moiety comprising Nitrogen; an aromatic chemical moiety; a hydrophilic chemical moiety; and a hydrophobic chemical moiety; and wherein R20 is carbon or nitrogen.

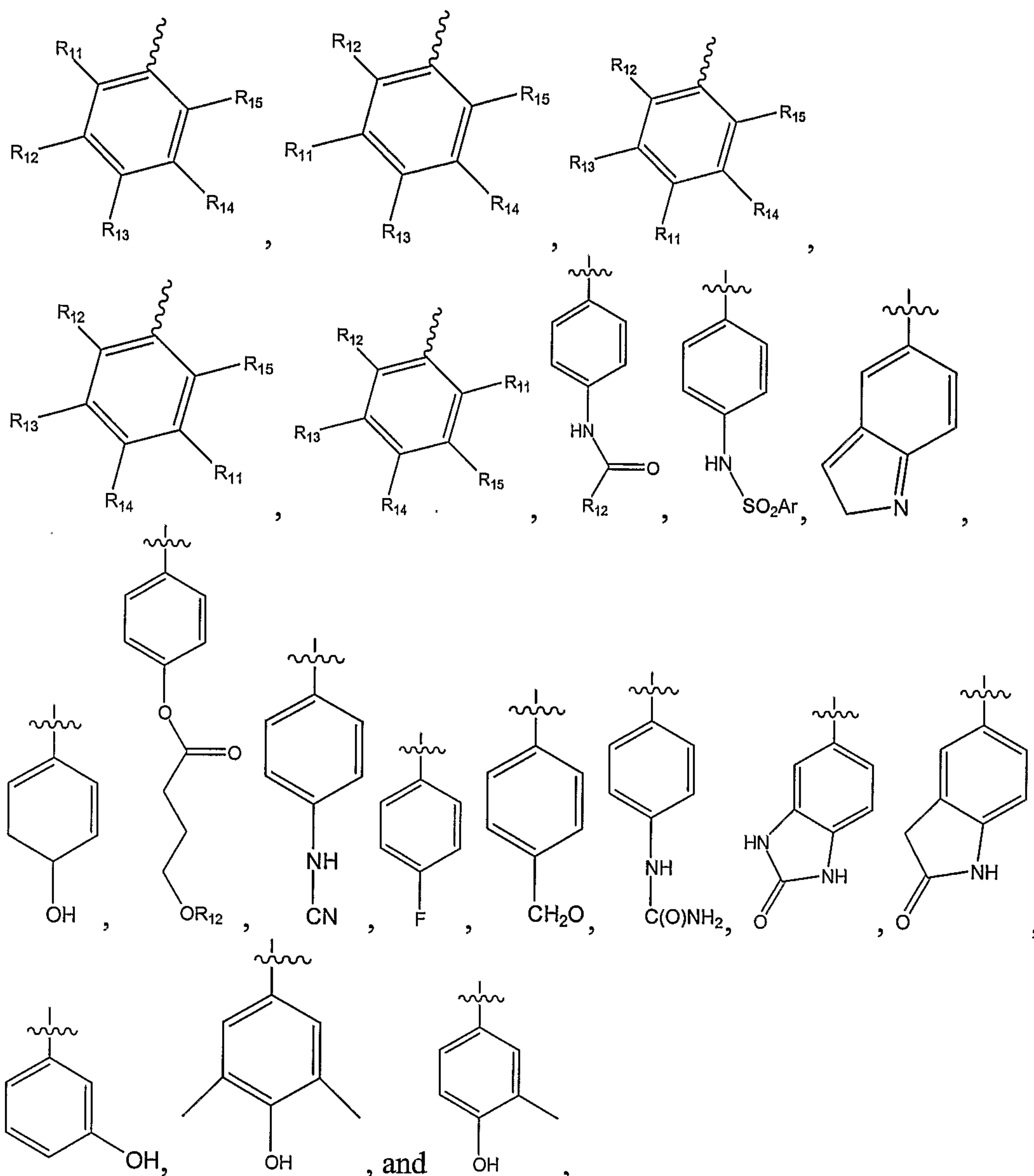
- In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



including both R and S enantiomeric forms and racemic mixtures. In such preferred embodiments, R1 is a nitrogen atom or a carbon atom; R2 is carbon or nitrogen; R3

comprises a chemical moiety comprising a heterocyclic group containing 3 or more carbon atoms; R4 and R5 are separately selected from the group consisting of: hydrogen; halogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; a chemical moiety comprising a halogen; a chemical moiety comprising Sulfur; a chemical moiety comprising Nitrogen; an aromatic chemical moiety; a hydrophilic chemical moiety; and a hydrophobic chemical moiety; and R6 is selected from H, a hydroxy, an alkoxy, a halogen, an amino, a lower-alkyl, a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of less than 10 carbons, a substituted cycloaliphatic group, an aryl, a heterocyclic, NO₂; SR'; and NR'₂, wherein R' is defined as a linear or branched, saturated or unsaturated aliphatic chain having at least one carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxyl subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup.

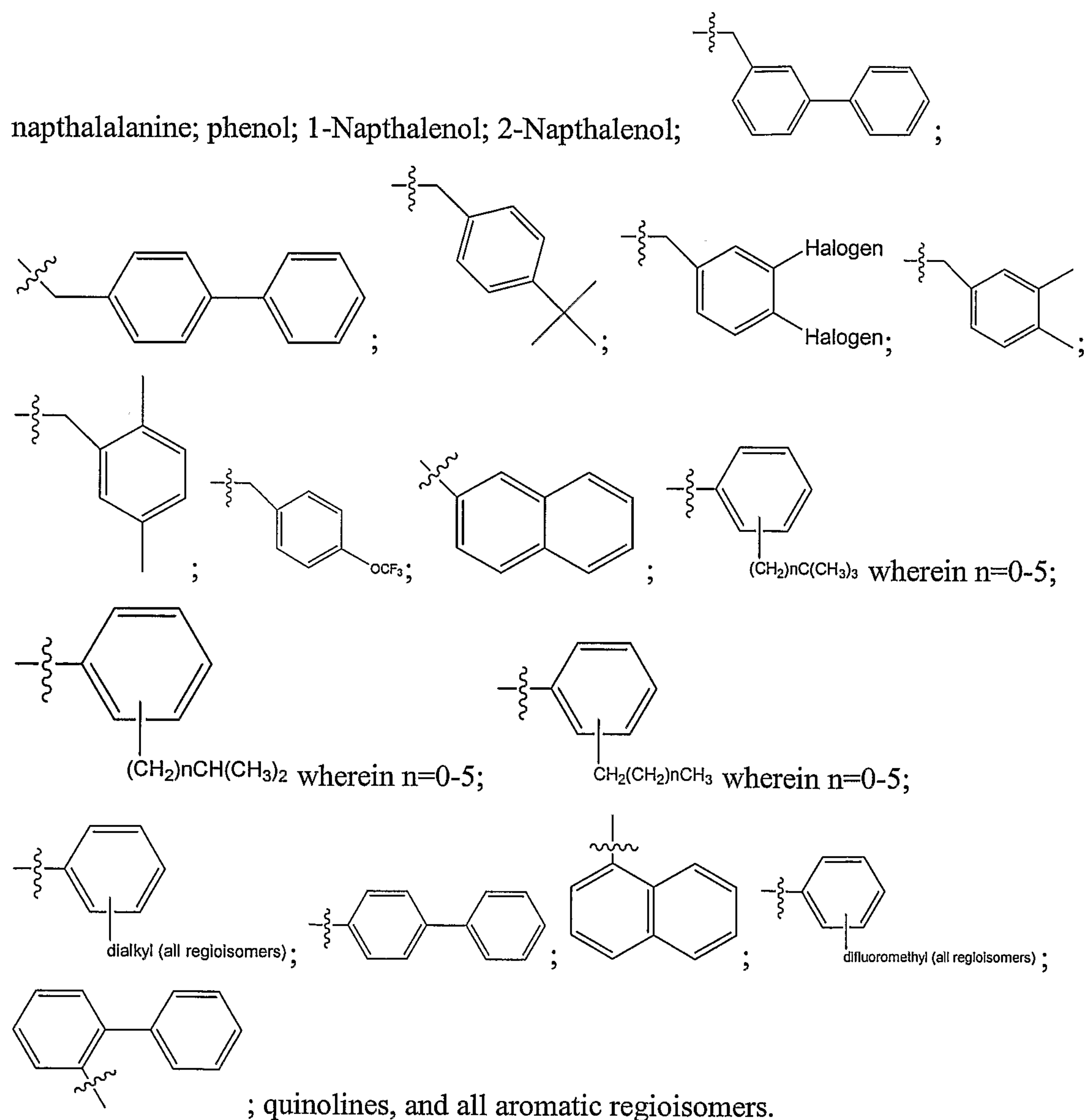
In preferred embodiments, R3 is selected from the group consisting of:



- 5 wherein R12, R13, R14 and R15 are selected from the group consisting of: hydrogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 1 carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxy subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or
- 10 branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or
- 15 branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain

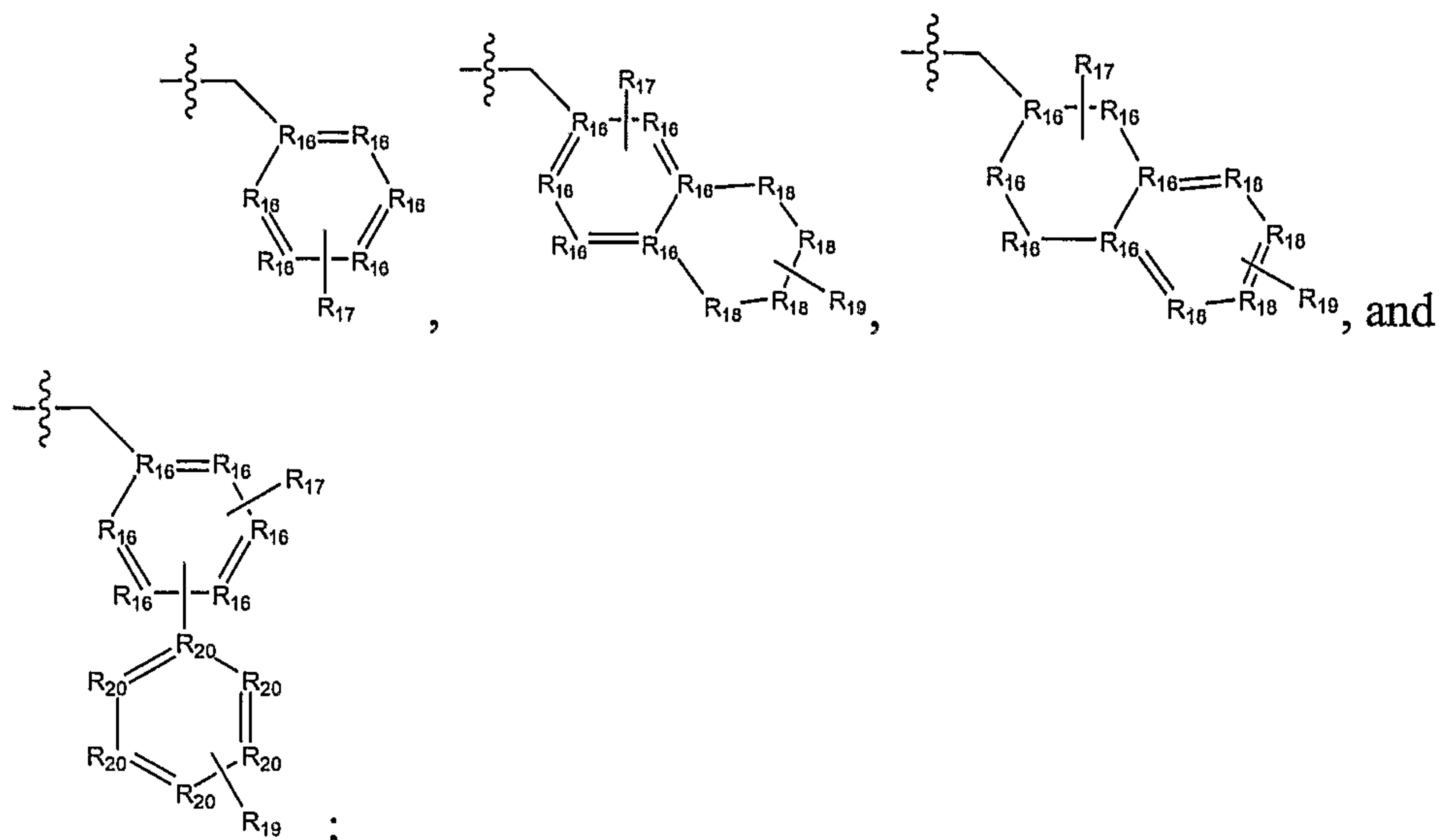
having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ether subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; and R11 is OH.

10 In preferred embodiments, R4 or R5 are selected from group consisting of:



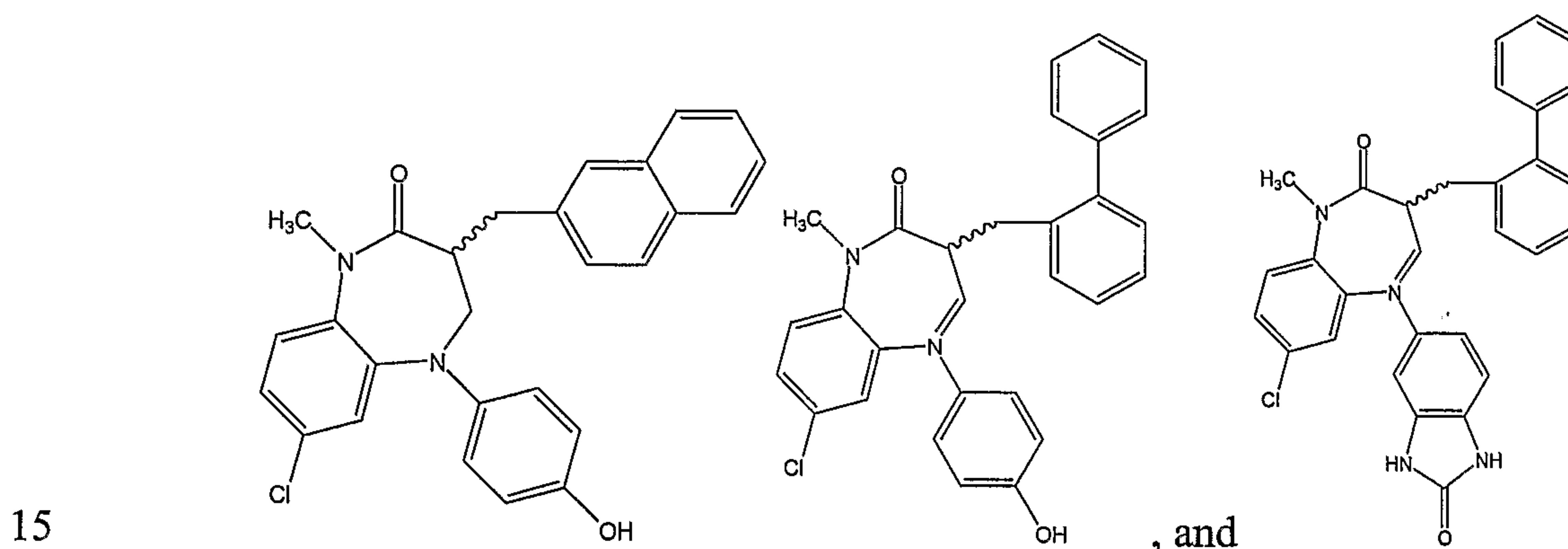
15

In other preferred embodiments, R4 or R5 is selected from the group consisting of:

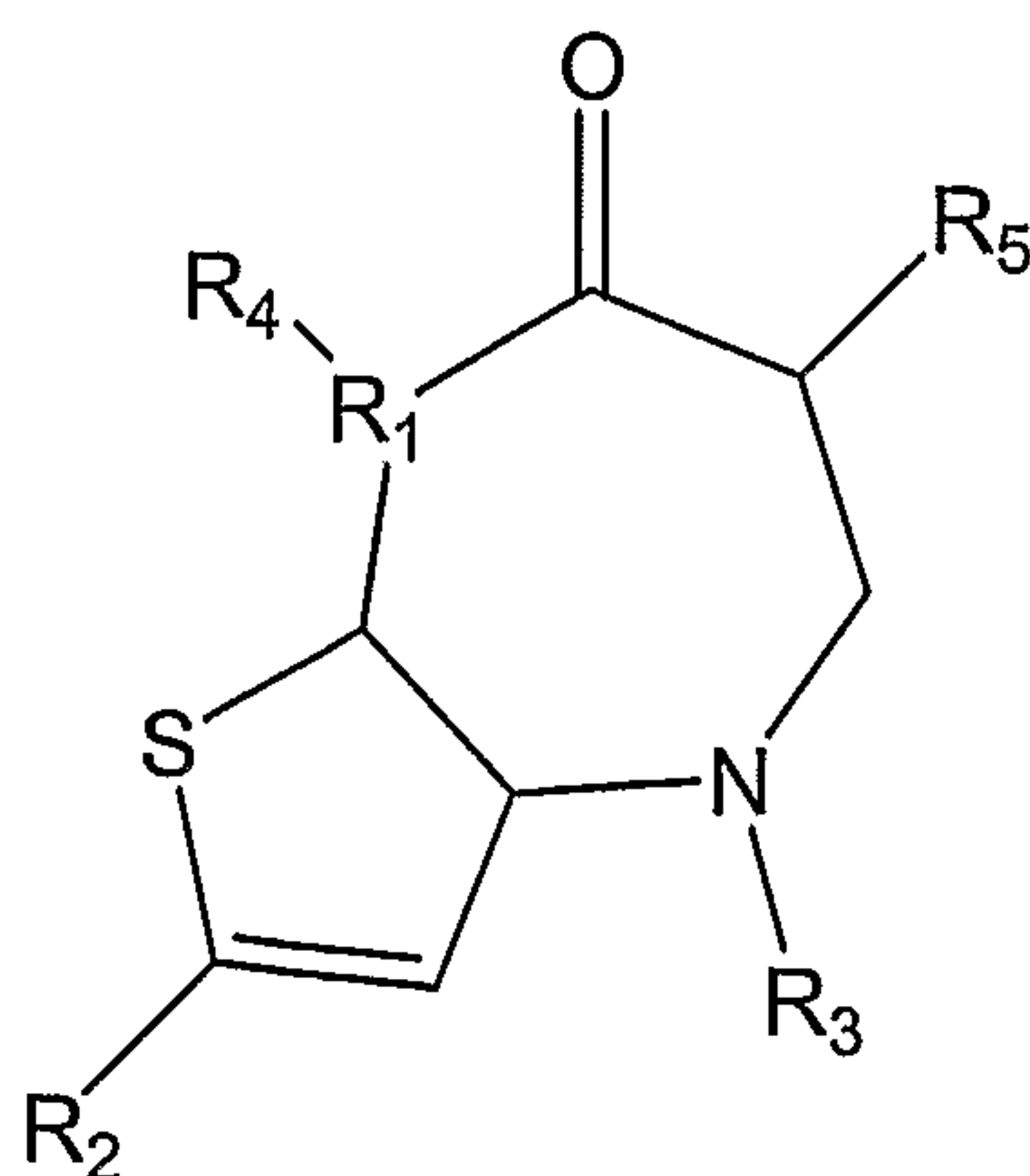


wherein R16 is carbon or nitrogen; wherein R17 is selected from the group consisting of hydrogen; halogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; a chemical moiety comprising a halogen; a chemical moiety comprising Sulfur; a chemical moiety comprising Nitrogen; an aromatic chemical moiety; a hydrophilic chemical moiety; and a hydrophobic chemical moiety; wherein R18 is carbon or nitrogen; wherein R19 is selected from the group consisting of hydrogen; halogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; a chemical moiety comprising a halogen; a chemical moiety comprising Sulfur; a chemical moiety comprising Nitrogen; an aromatic chemical moiety; a hydrophilic chemical moiety; and a hydrophobic chemical moiety; and wherein R20 is carbon or nitrogen.

In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



including both R and S enantiomeric forms and racemic mixtures.

In such preferred embodiments, R1 is a nitrogen atom or a carbon atom; R2 is selected from H, a hydroxy, an alkoxy, a halo, an amino, a lower-alkyl, a substituted-amino, an acetilamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of less than 10 carbons, a substituted cycloaliphatic group, an aryl, a heterocyclic, NO₂; SR'; and NR'₂, wherein R' is defined as a linear or branched, saturated or unsaturated aliphatic chain having at least one carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxyl subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; R3 comprises a chemical moiety comprising a heterocyclic group containing 3 or

In preferred embodiments, R3 is selected from the group consisting of:



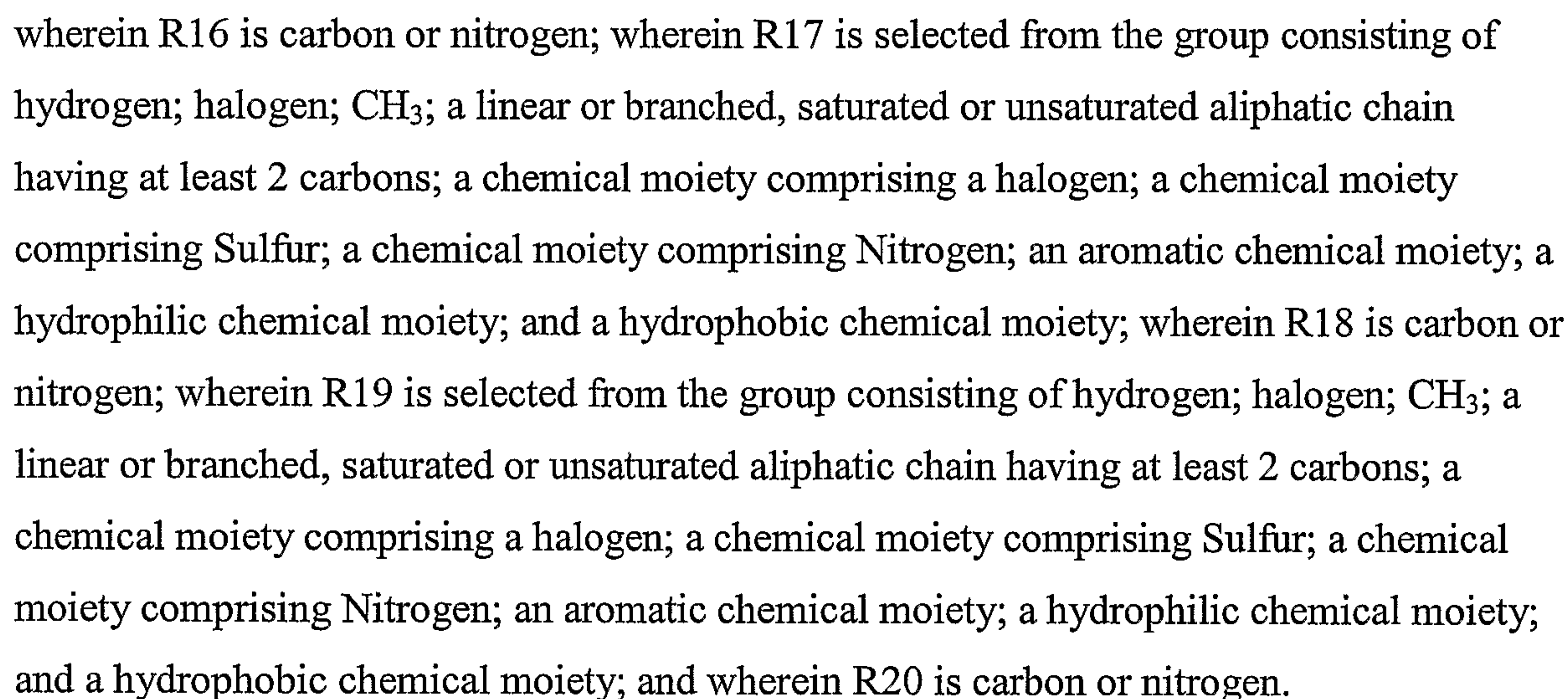
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In preferred embodiments, R4 or R5 are selected from group consisting of:

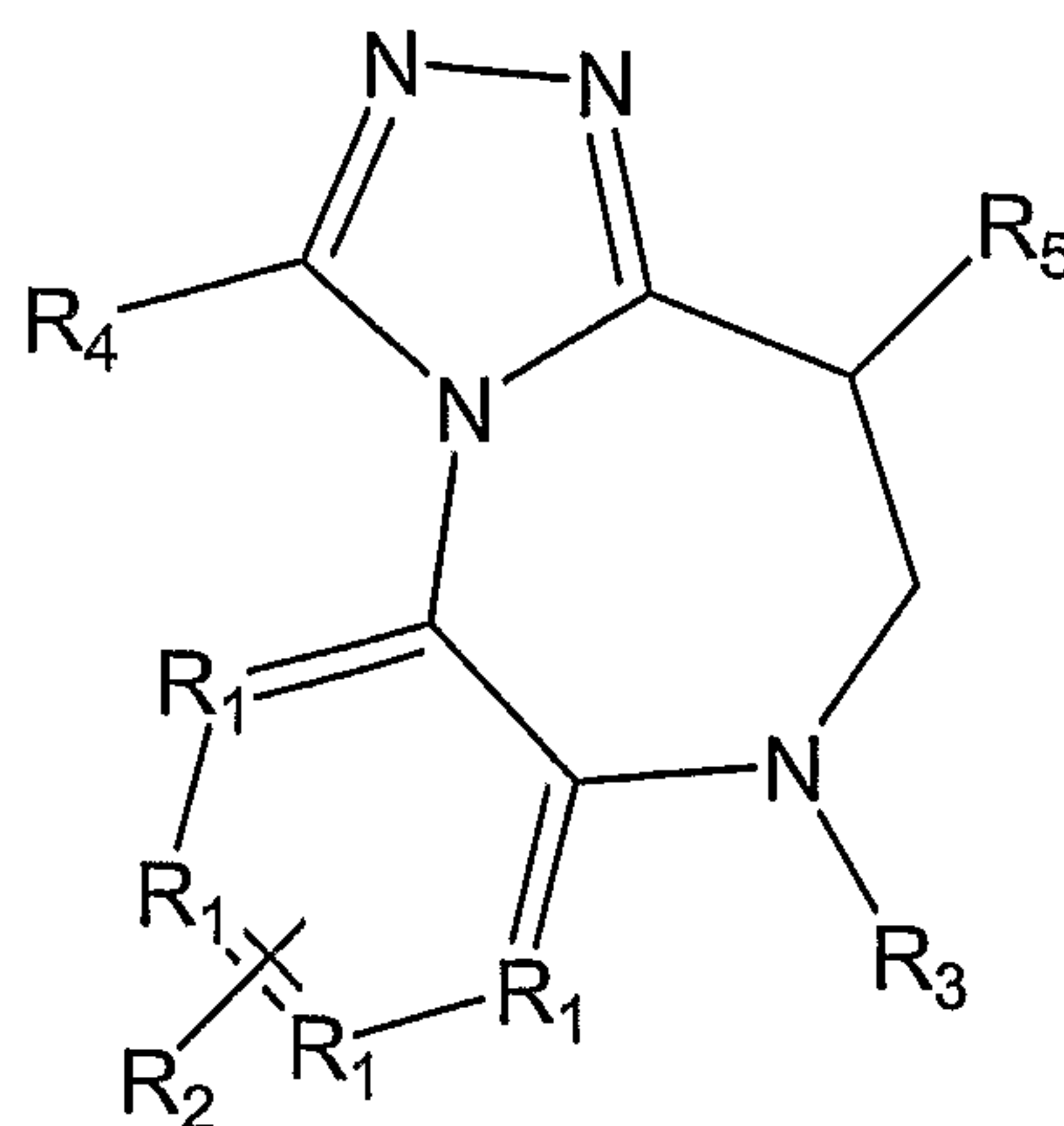




In other preferred embodiments, R4 or R5 is selected from the group consisting of:



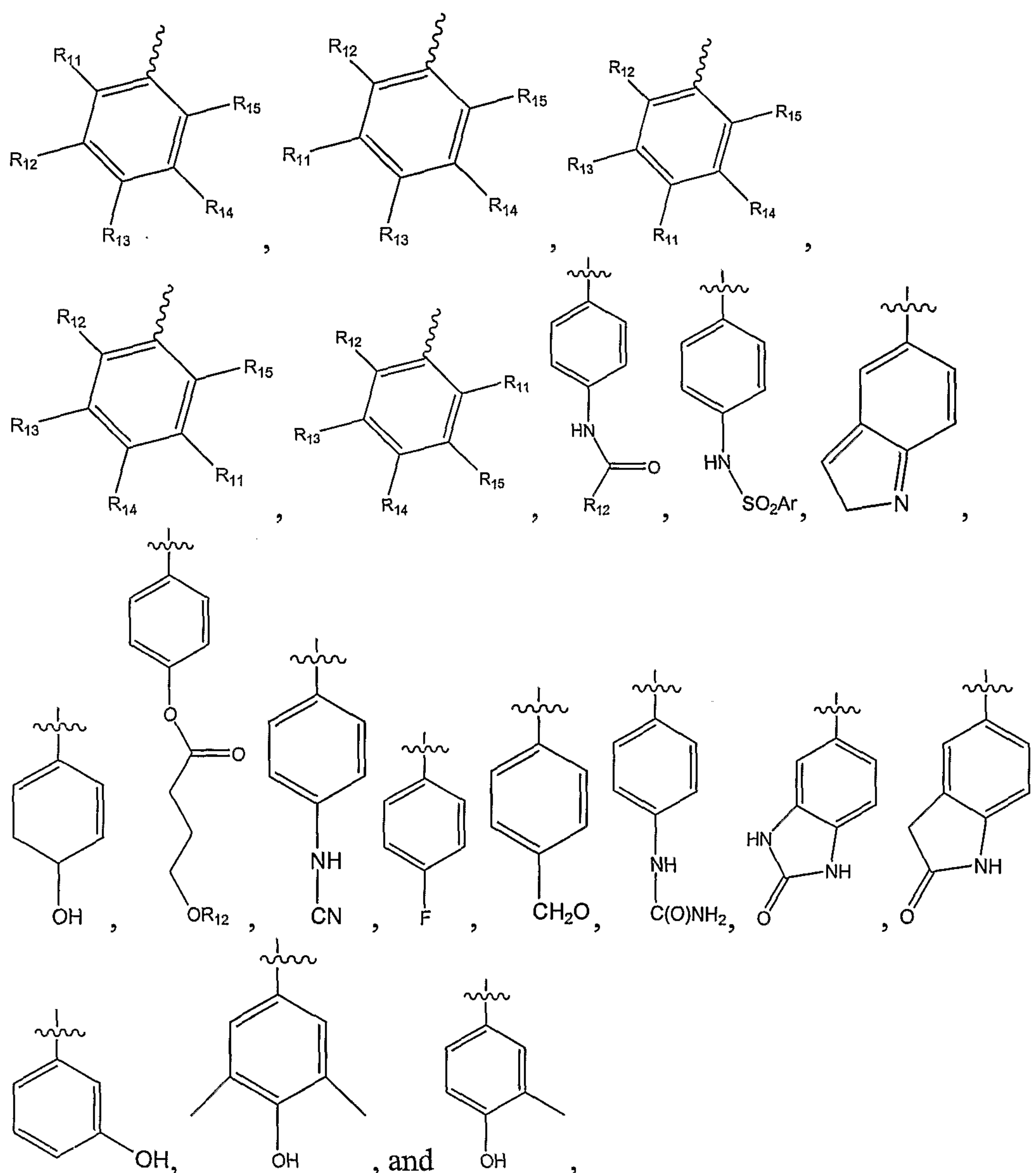
In some preferred embodiments, crystal forms and formulations of the following exemplary compounds are provided:



including both R and S enantiomeric forms and racemic mixtures.

In such preferred embodiments, R1 is carbon or nitrogen; R2 is selected from H, a hydroxy, an alkoxy, a halogen, an amino, a lower-alkyl, a substituted-amino, an acetylamino, a hydroxyamino, an aliphatic group having 1-8 carbons and 1-20 hydrogens, a substituted aliphatic group of similar size, a cycloaliphatic group consisting of less than 10 carbons, a substituted cycloaliphatic group, an aryl, a heterocyclic, NO₂; SR'; and NR'₂, wherein R' is defined as a linear or branched, saturated or unsaturated aliphatic chain having at least one carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxyl subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; R3 comprises a chemical moiety comprising a heterocyclic group containing 3 or

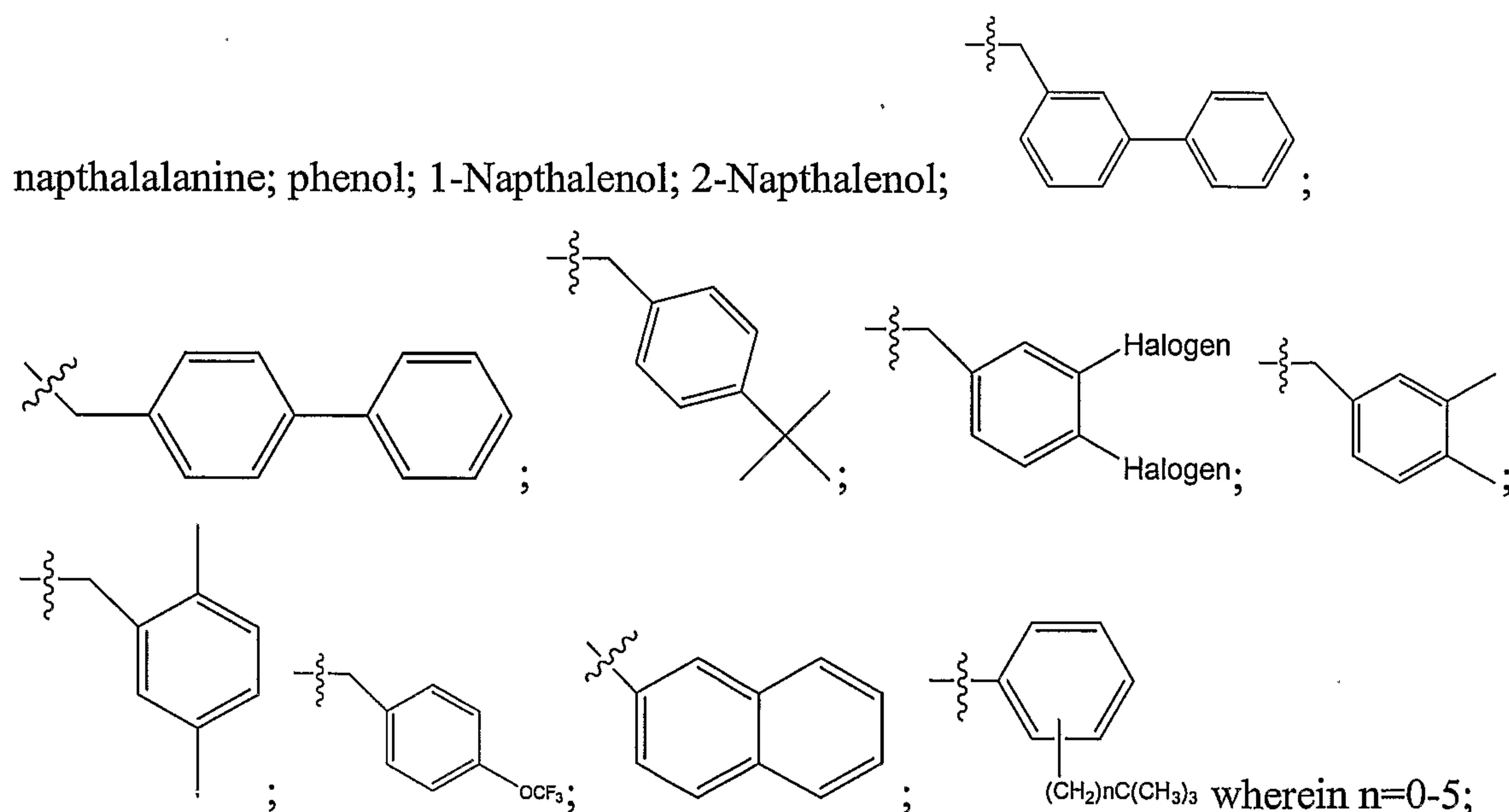
10 In preferred embodiments, R3 is selected from the group consisting of:

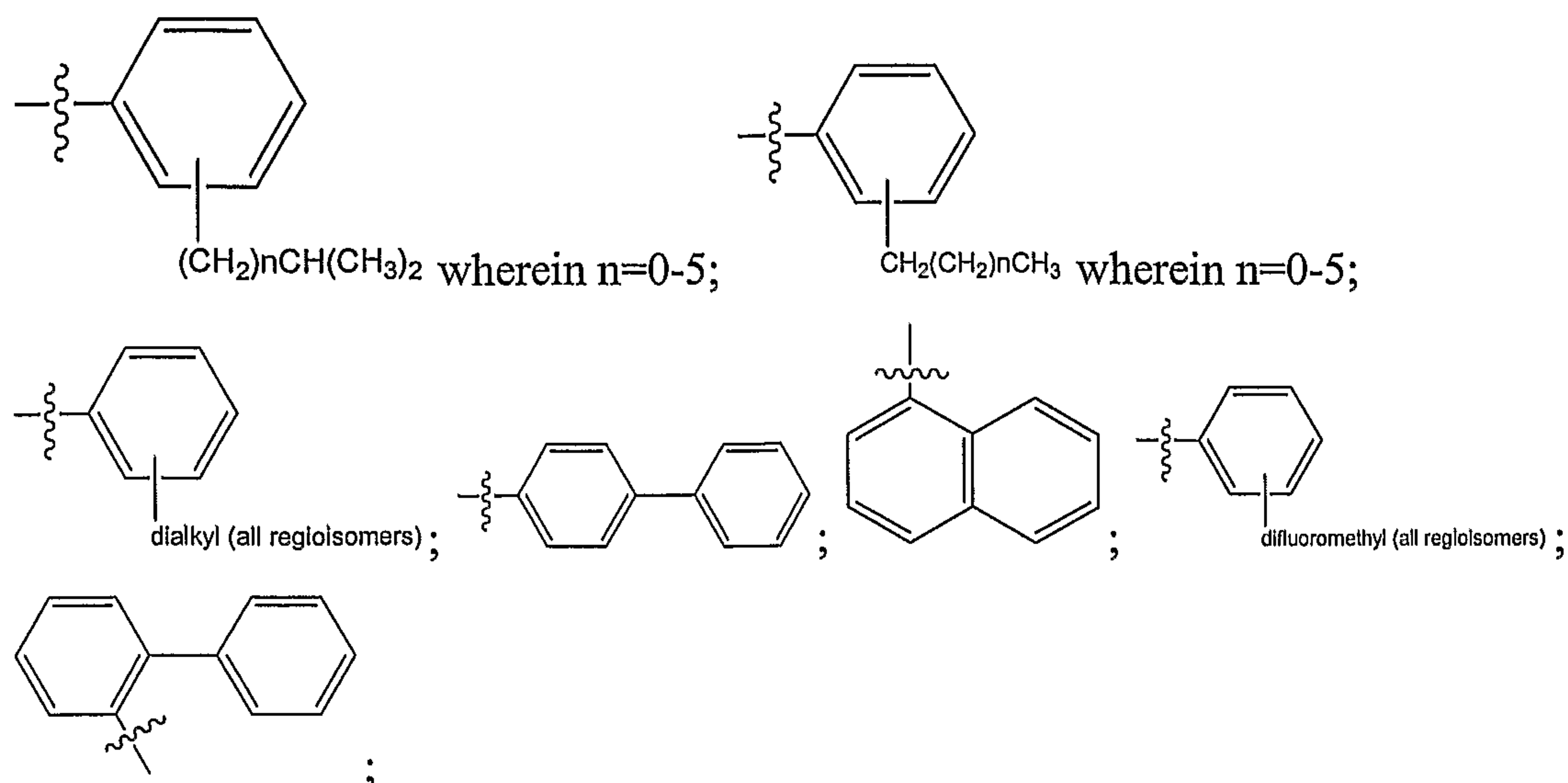


15 wherein R12, R13, R14 and R15 are selected from the group consisting of: hydrogen; CH₃;
a linear or branched, saturated or unsaturated aliphatic chain having at least 1 carbon; a

linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxy subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ether subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; and R11 is OH.

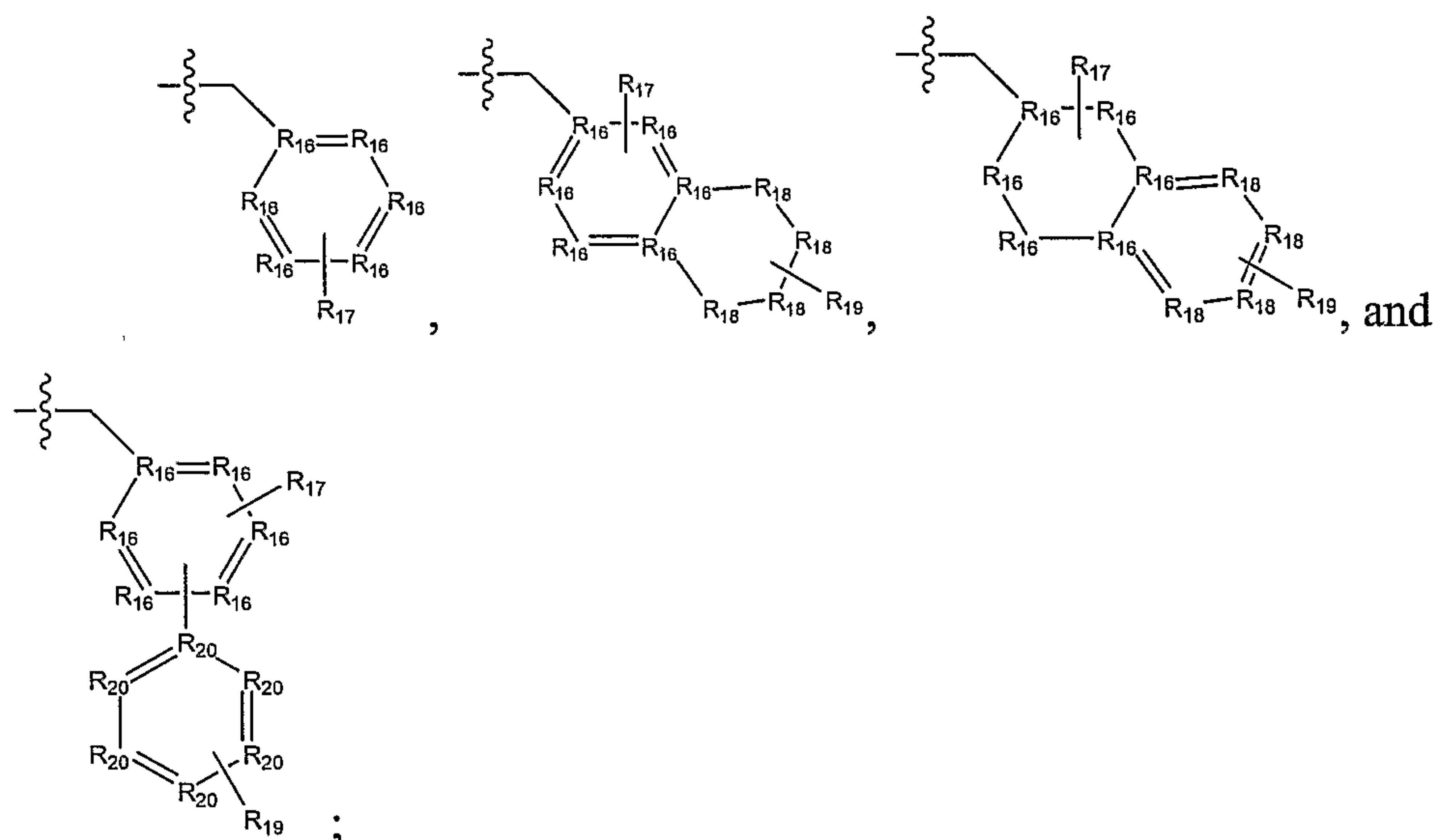
In preferred embodiments, R4 or R5 are selected from group consisting of:





quinolines, and all aromatic regioisomers.

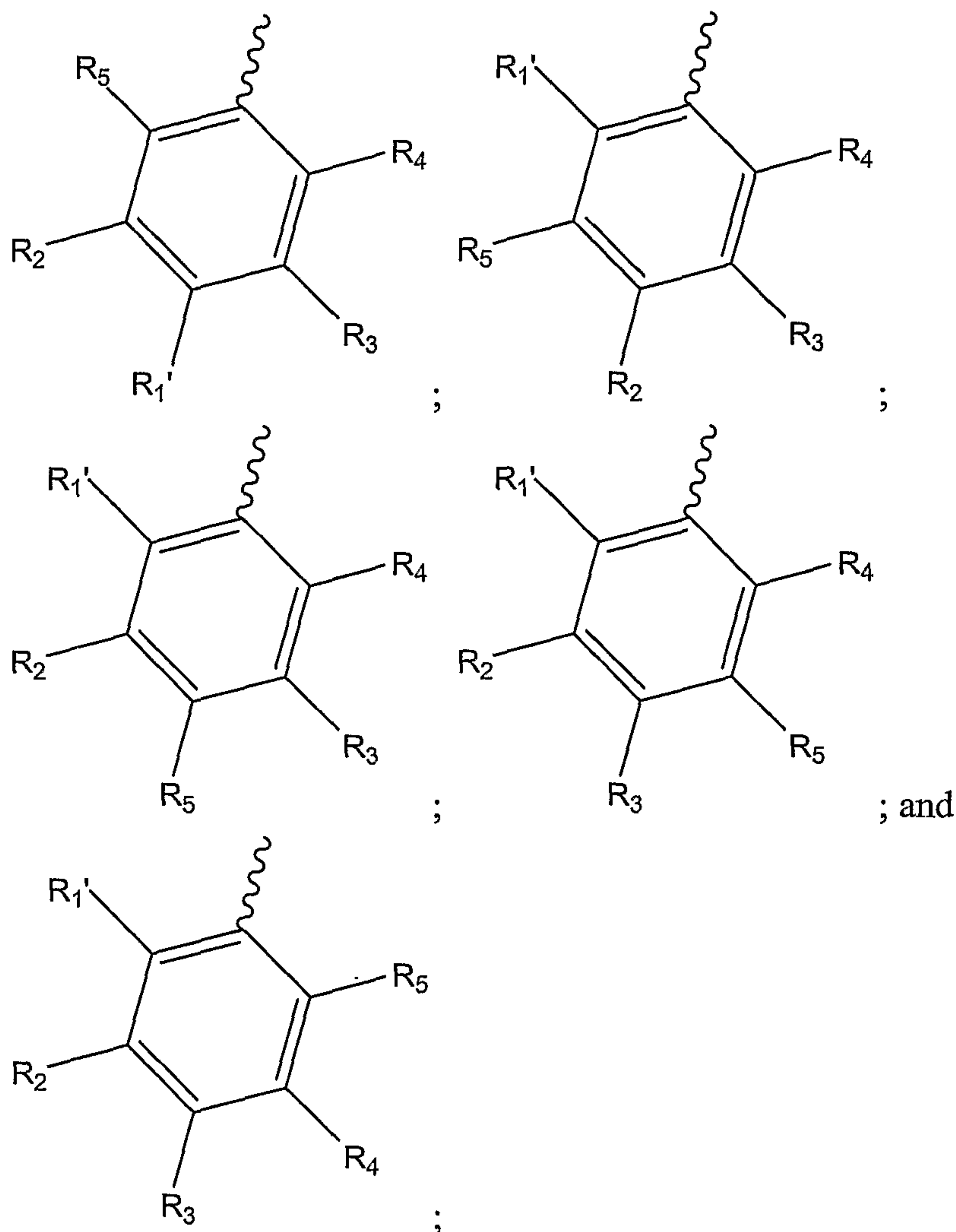
5 In other preferred embodiments, R4 or R5 is selected from the group consisting of:



wherein R16 is carbon or nitrogen; wherein R17 is selected from the group consisting of hydrogen; halogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; a chemical moiety comprising a halogen; a chemical moiety comprising Sulfur; a chemical moiety comprising Nitrogen; an aromatic chemical moiety; a hydrophilic chemical moiety; and a hydrophobic chemical moiety; wherein R18 is carbon or nitrogen; wherein R19 is selected from the group consisting of hydrogen; halogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; a chemical moiety comprising a halogen; a chemical moiety comprising Sulfur; a chemical moiety comprising Nitrogen; an aromatic chemical moiety; a hydrophilic chemical moiety; and a hydrophobic chemical moiety; and wherein R20 is carbon or nitrogen.

In certain embodiments, crystal forms and formulations of the following exemplary compounds comprising the following formula are provided: A – B – C; wherein A is a chemical moiety comprising a hydroxyl group (e.g., a phenolic ring); wherein B is a chemical moiety (e.g., scaffold molecule) separating A and C by at least 1 atom; and
 5 wherein C is a hydrophobic chemical moiety (e.g., naphyl group).

In some embodiments, A is selected from the group consisting of: is selected from the group consisting of:



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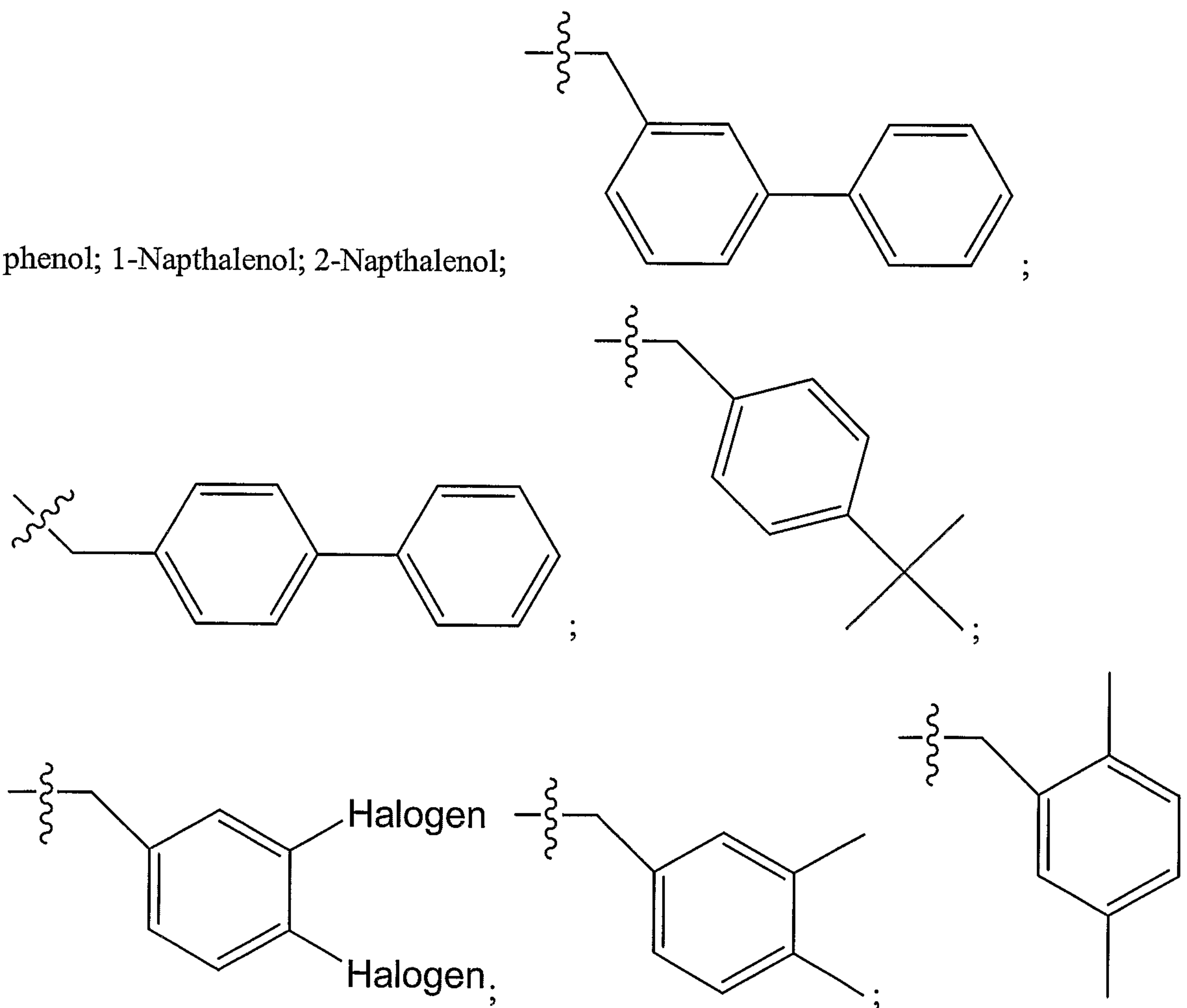
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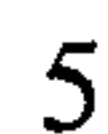
wherein R1', R2, R3 and R4 are selected from the group consisting of: hydrogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 1 carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxy subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein the aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated

aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein the aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide
 5 subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic
 10 chain having at least 2 carbons, and having at least one ether subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; and R5 is OH.

In some embodiments, C is selected from group consisting of: naphthalalanine;

15 phenol; 1-Napthalenol; 2-Napthalenol;





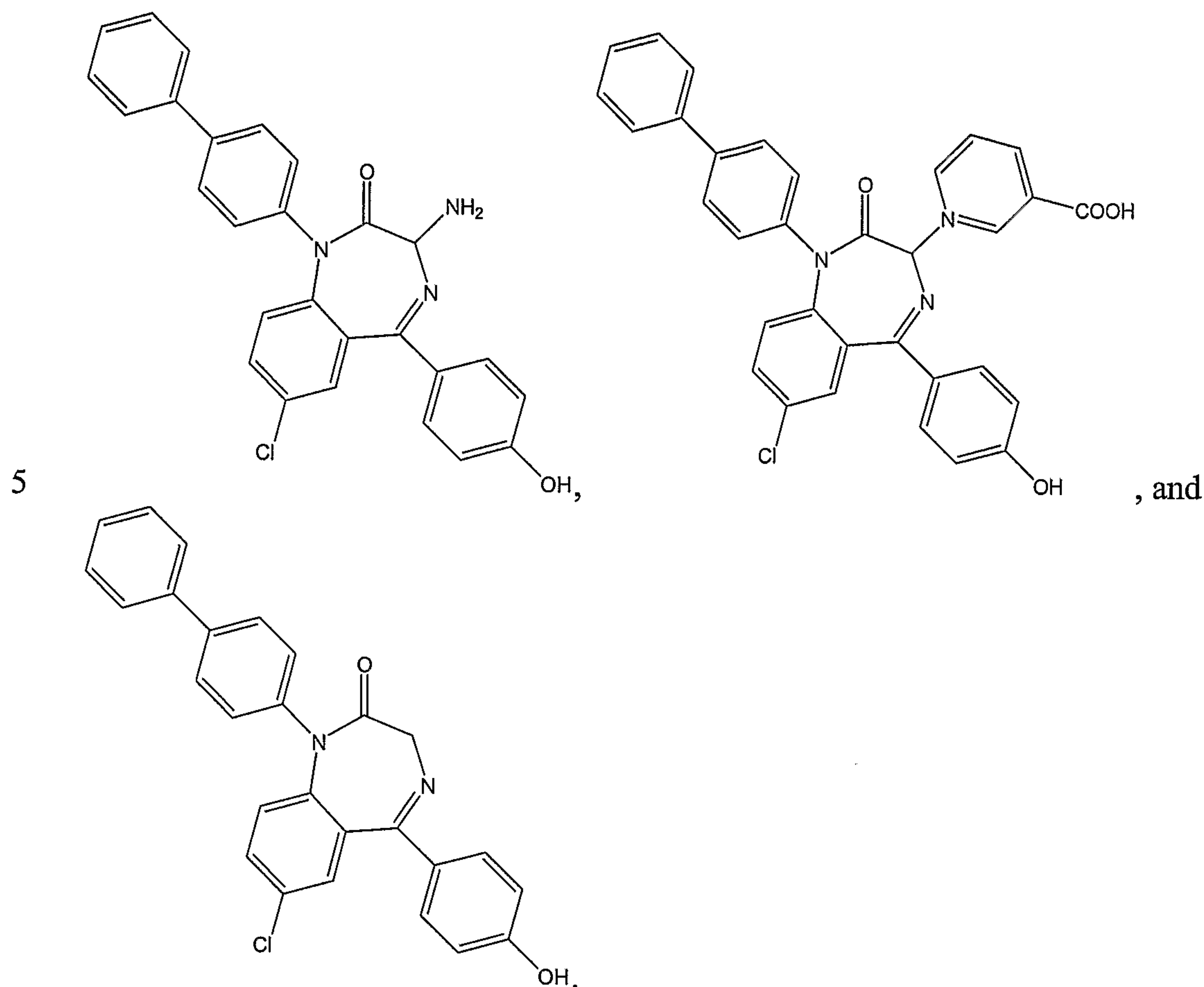
In some embodiments, B is a benzodiazepine structure described by the following formula:



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selected from the group consisting of position 1, position 2, position 3, position 4, position 5, position 6, position 7, position 8, position 9, and position 10.

In some embodiments, crystal forms and formulations of the following exemplary compounds are provided:

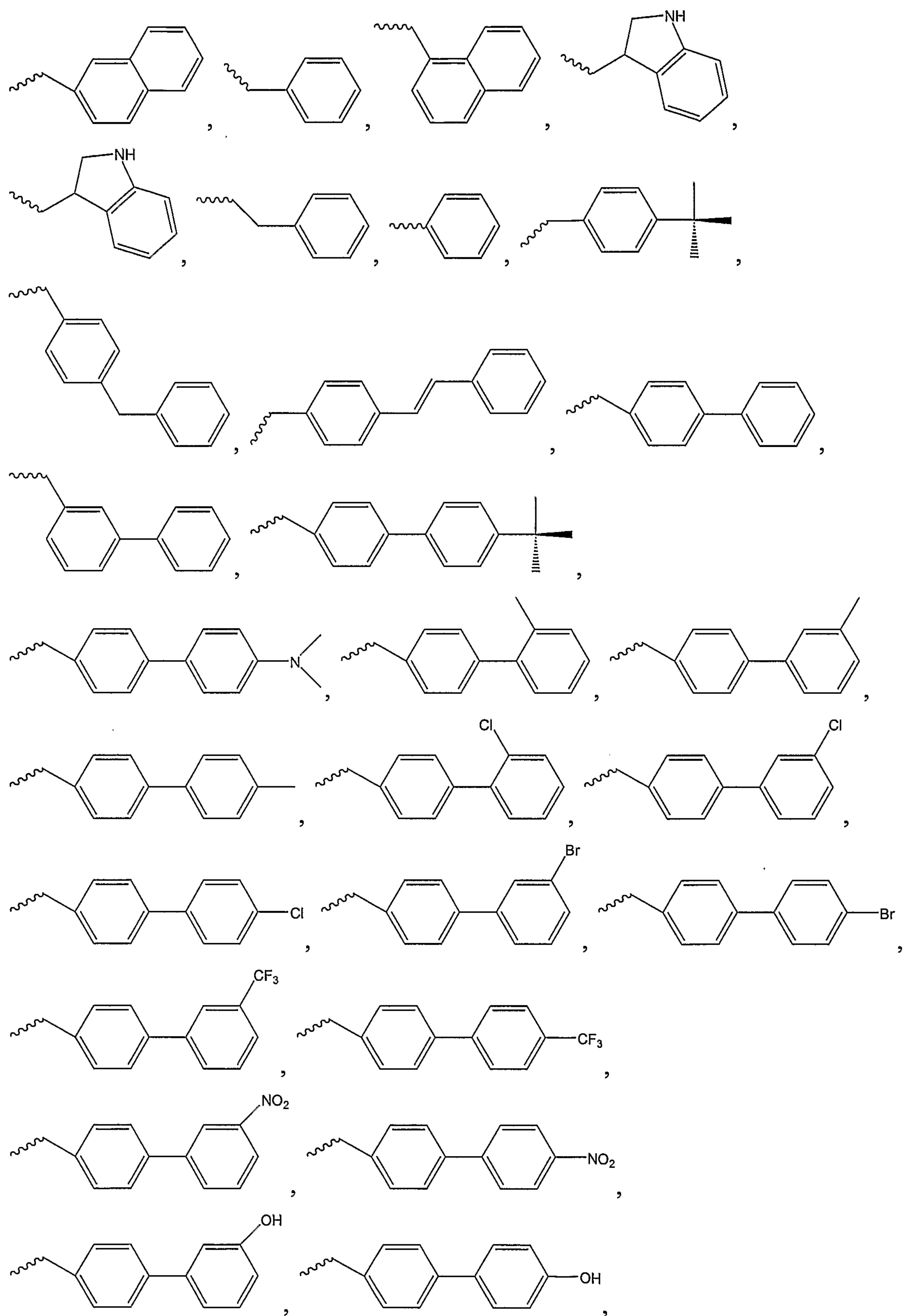


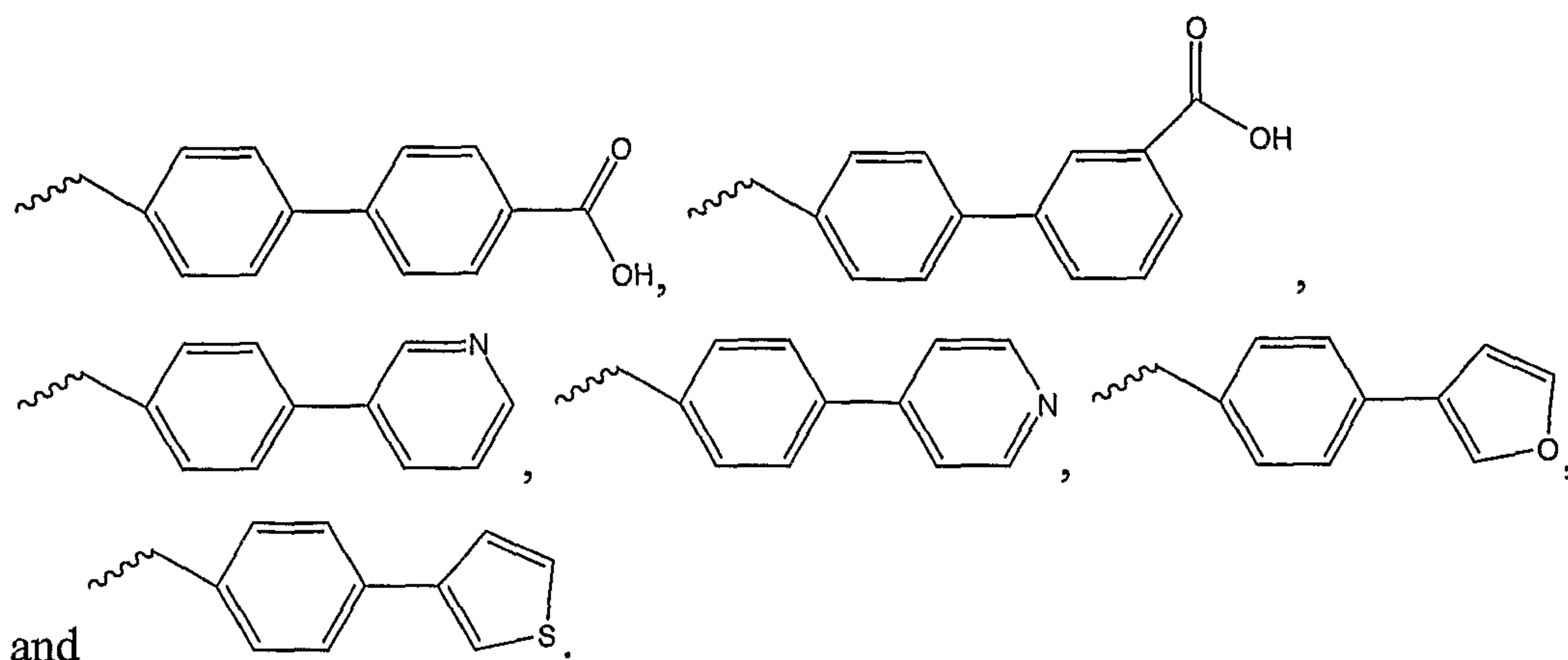
Certain embodiments of the present invention include crystal forms and formulations of exemplary compounds with the following formula:



including both R and S enantiomeric forms and racemic mixtures, wherein R1 is an electron rich heterocycle.

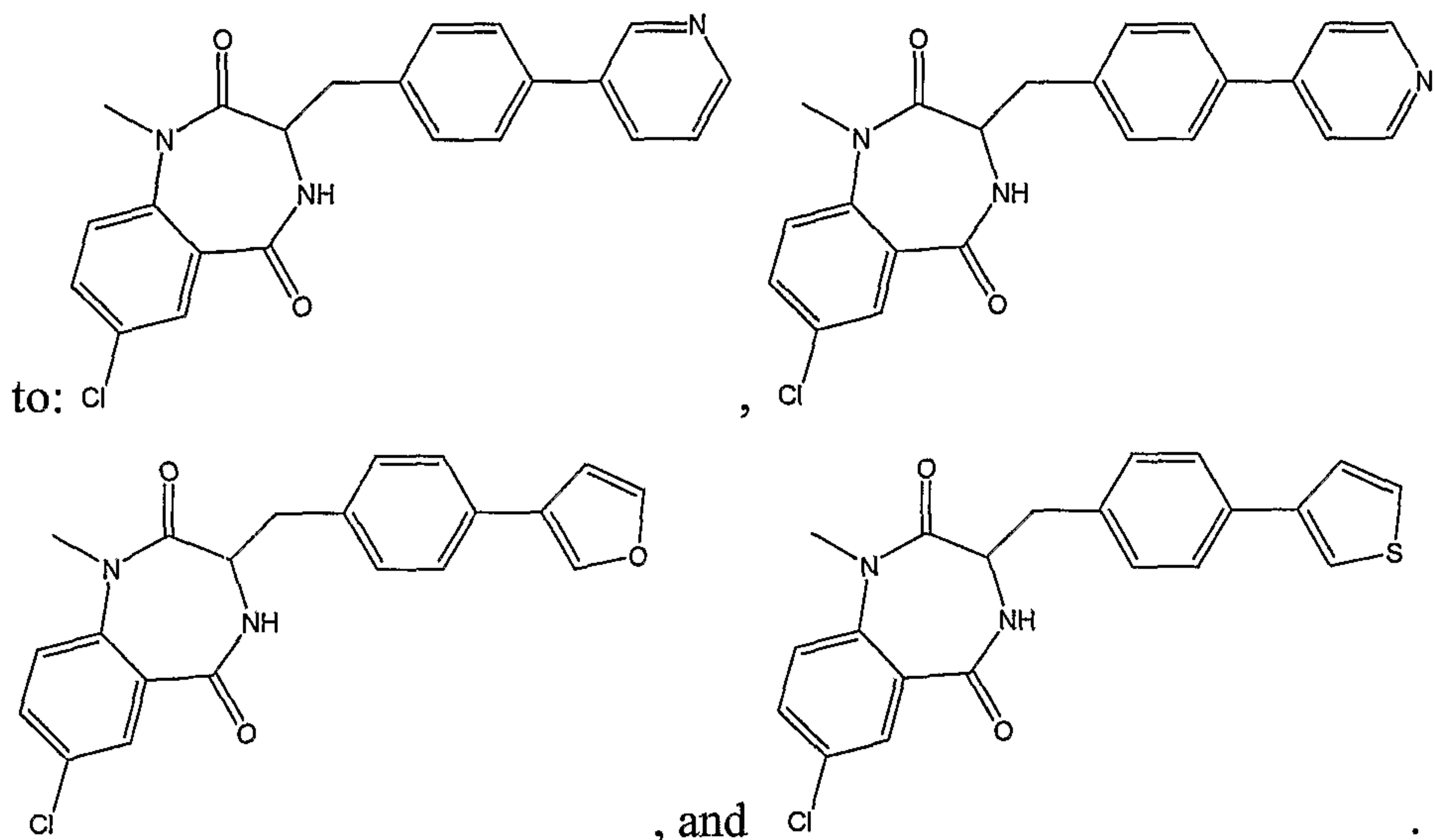
In preferred embodiments, R1 is selected from the group consisting of:



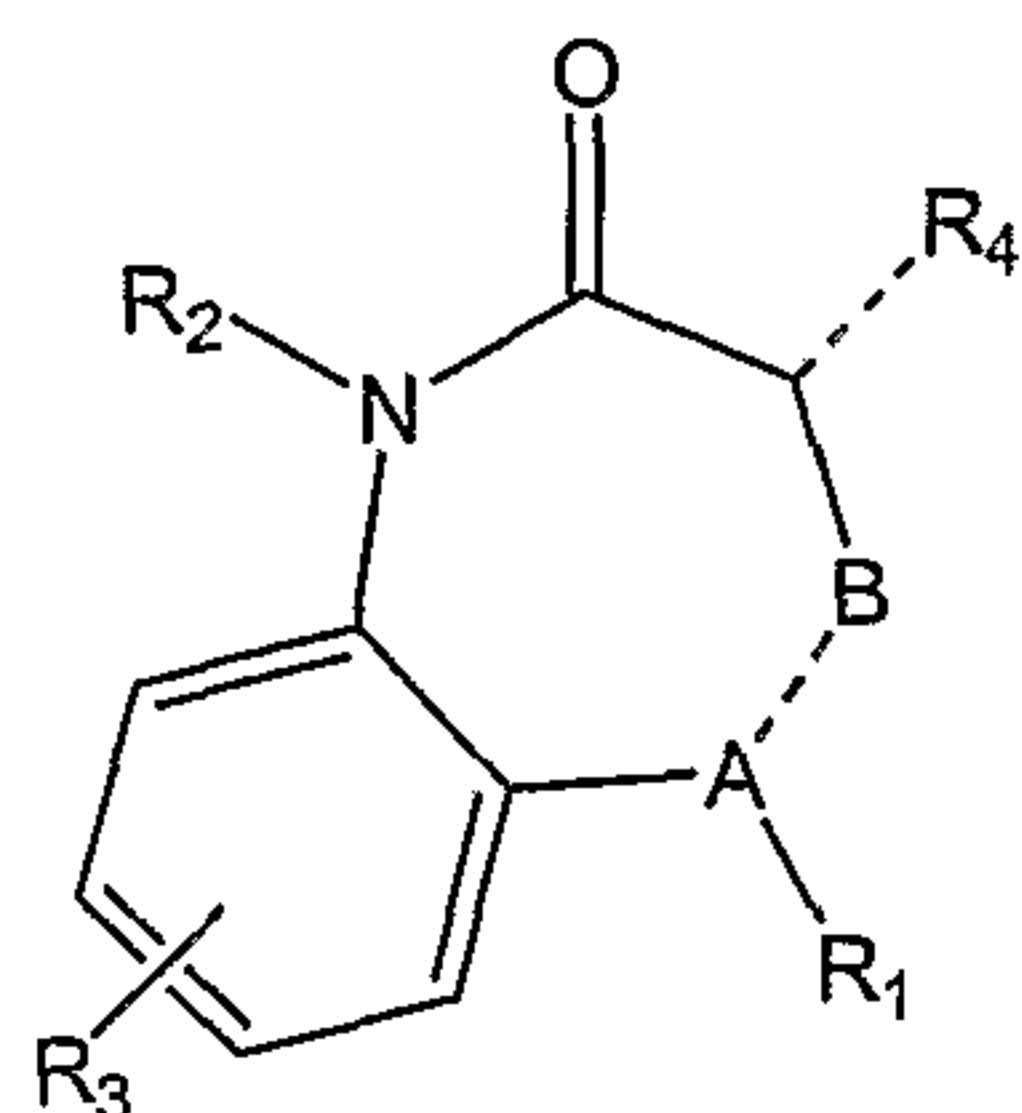


In some embodiments, R₂ is a halogen. In some embodiments, R₂ is Chlorine.

5 In certain embodiments, examples of crystal forms and formulations of the exemplary 1,4-benzodiazepine-2,5-dione compounds include but are not limited



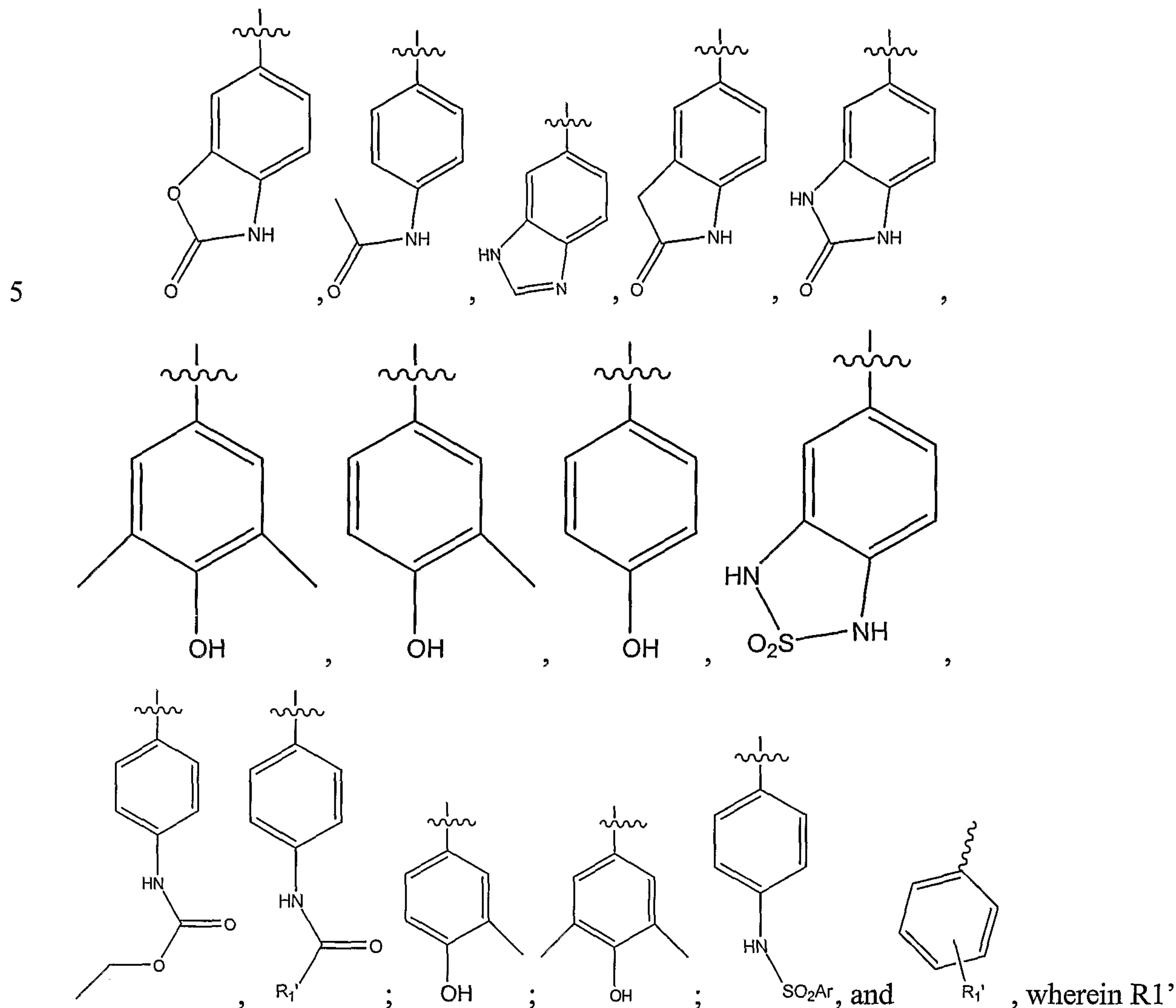
10 Certain embodiments of the present invention include crystal forms and formulations of benzodiazepine (and benzodiazepine related) compounds having a chemical moiety that causes the benzodiazepine to lack a chiral center associated with the third carbon position of the benzodiazepine ring. Certain embodiments of the present invention include crystal forms and formulations of exemplary compounds with the following formula:



15 including both R and S enantiomeric forms and racemic mixtures.

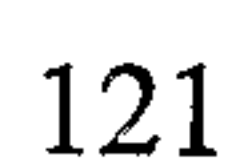
In some embodiments, A---B is selected from the group consisting of N---CH_2 , and C=N .

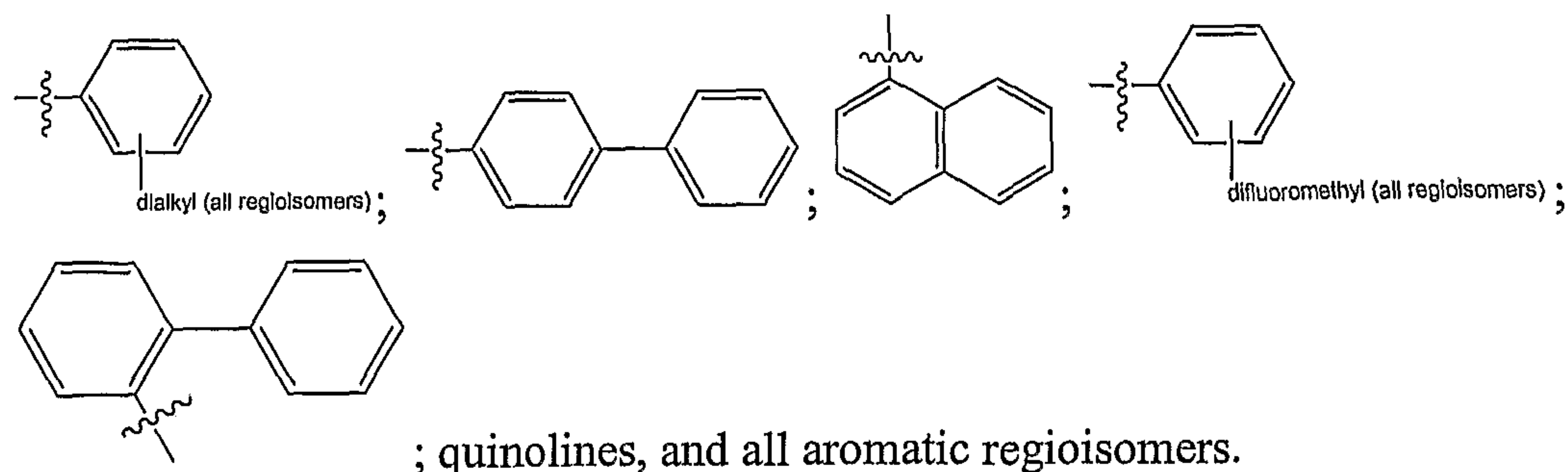
In some embodiments, R1 is selected from the group consisting of



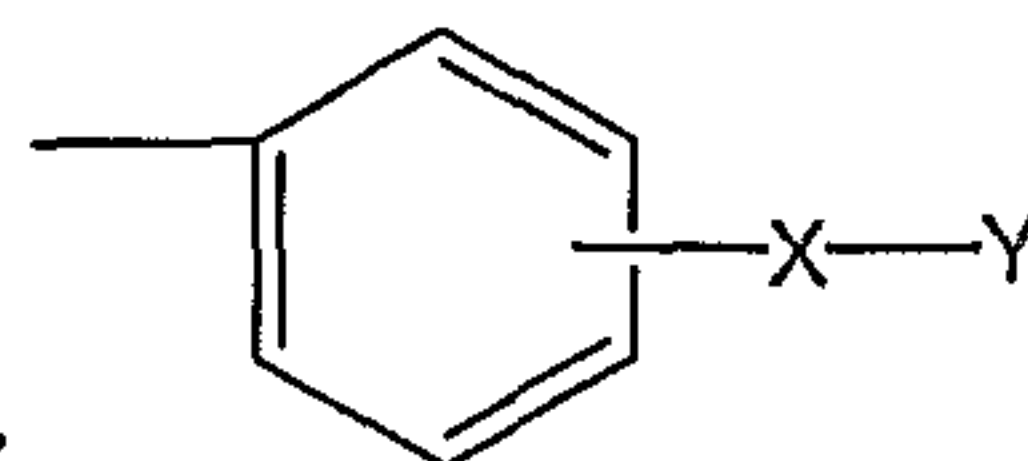
is selected from the group consisting of halogen; alkyl; substituted alkyl; aryl; substituted aryl; amino; carbonyl; sulfone; sulfonamide; ether; OH; a chemical moiety comprising Sulfur; a chemical moiety comprising Nitrogen; CH_3 ; a linear or branched, saturated or unsaturated aliphatic chain having at least 1 carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxy subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein said aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain

In some embodiments, R2 is selected from group consisting of: naphthalalanine;

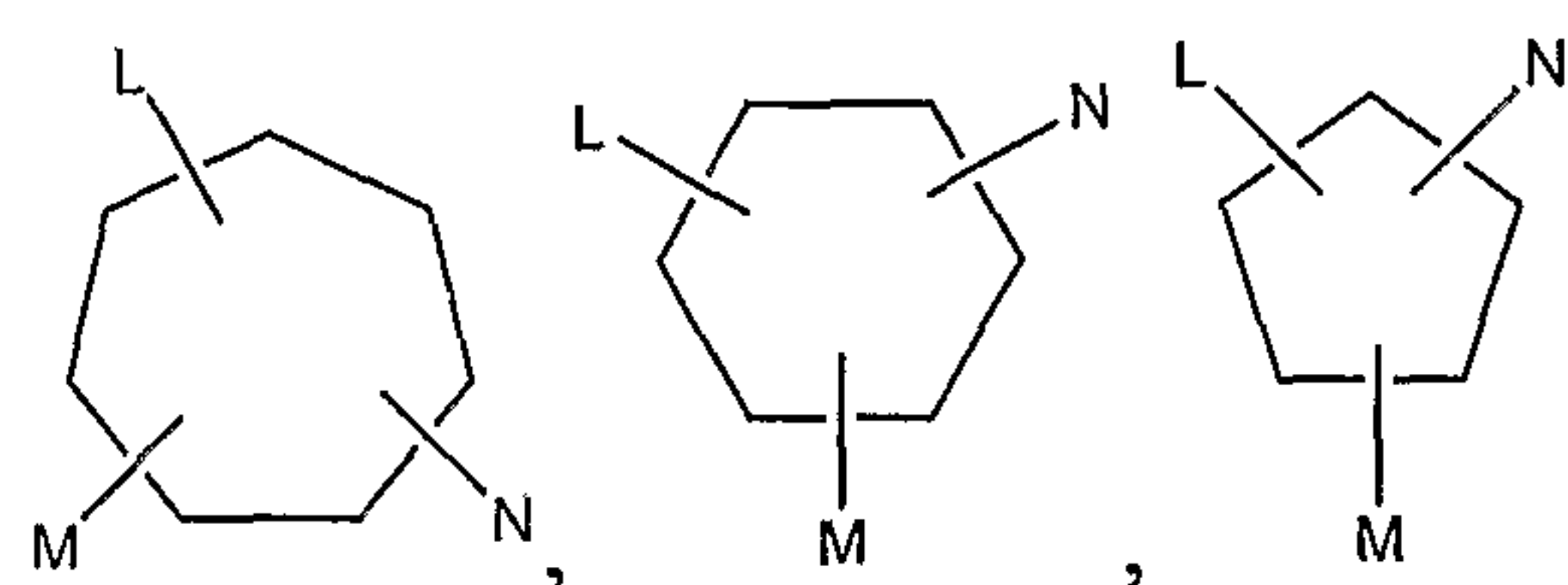
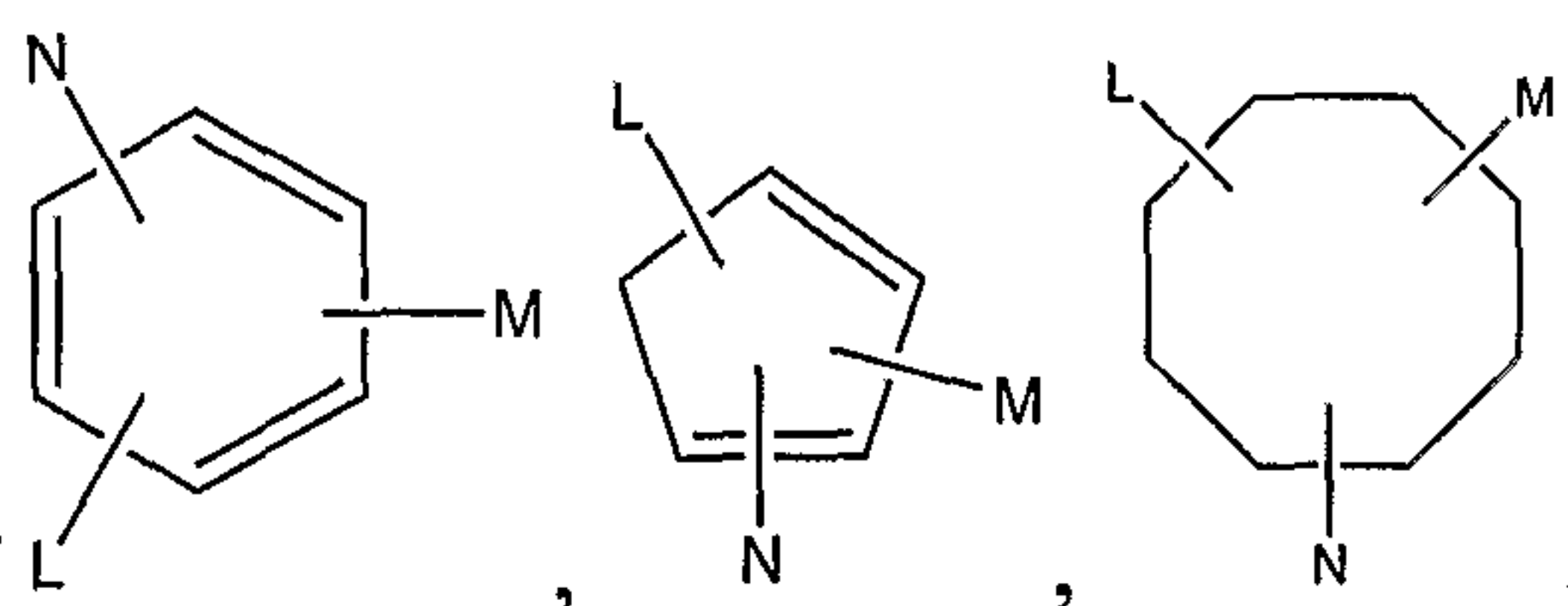




In some embodiments, R2 is:

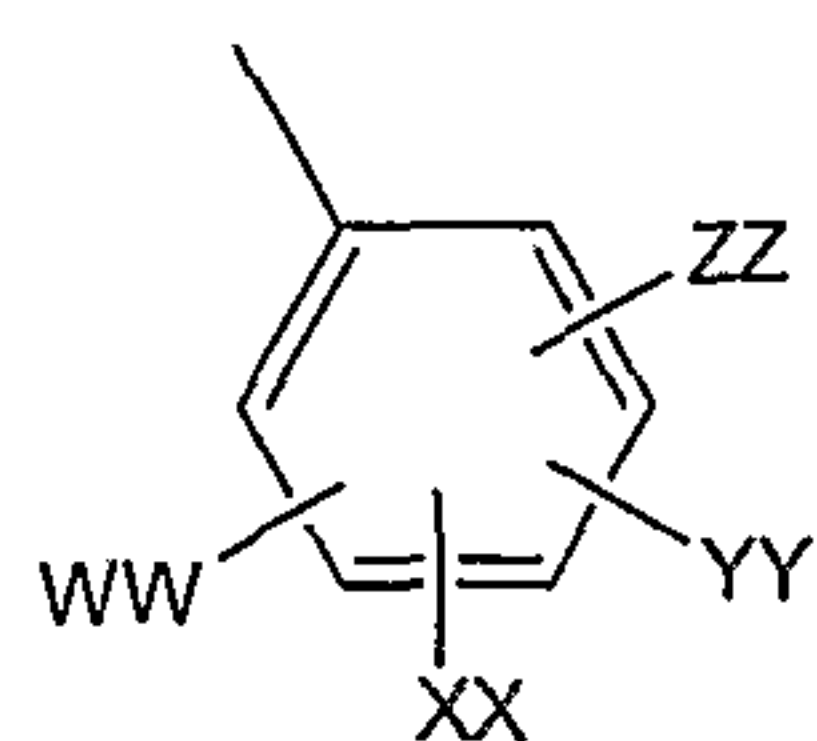


5 wherein X is selected from the group consisting of



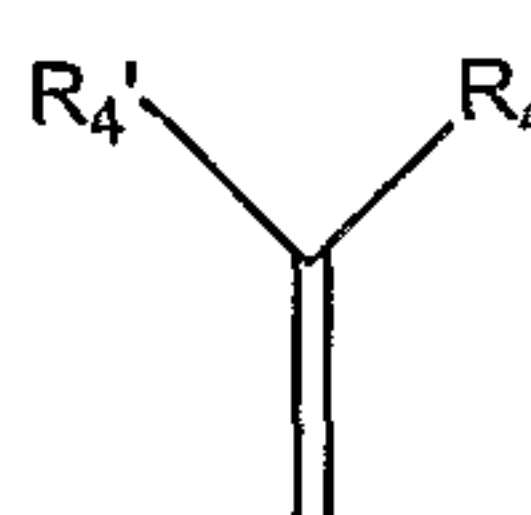
, alkyl, substituted alkyl, sulfonamide, SO₂alkyl, NHSO₂, CH₂, CH₂CH₂, SO₂, CH₂SO₂, SO₂CH₂, OCH₂CH₂O, SO, CH₂CH₂SO, SOCH₂CH₂;

and wherein L, M and N are present or absent, and are selected from the group consisting of alkyl, NO₂, halogen, OH, O-Alkyl, methyl ester, propyl ester, ethyl ester, CO₂H, CF₃,
 10 aniline, nitro, heterocycle, mono-substituted alkyl, di-substituted alkyl, and tri-substituted alkyl, hydrogen, SO₂NH₂, SO₂NH-alkyl, SOalkyl, NHSO₂alkyl; wherein Y is selected from the group consisting of hydrogen, alkyl, substituted alkyl, halogen, OH, O-Alkyl, methyl ester, propyl ester, ethyl ester, CO₂H, nitro, heterocycle, mono-substituted alkyl, di-substituted alkyl, and tri-substituted alkyl, hydrogen, SOalkyl, SO₂NH₂, SO₂NH-alkyl,

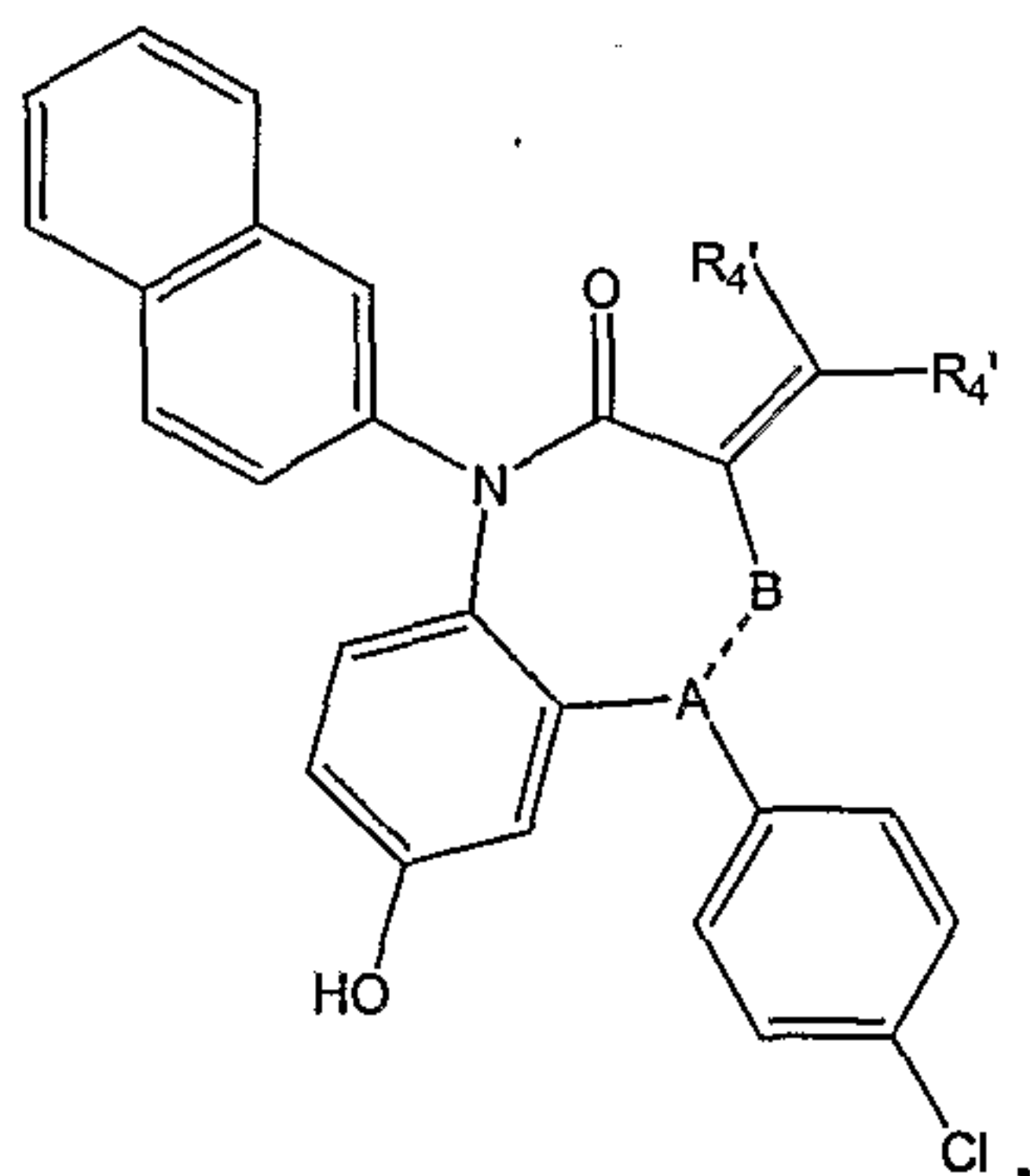


15 NHSO₂alkyl, and ; wherein WW, XX, YY and ZZ are present or absent, and are selected from the group consisting of alkyl, halogen, OH, O-Alkyl, methyl ester, propyl ester, ethyl ester, CO₂H, aniline, nitro, heterocycle, mono-substituted alkyl, di-substituted alkyl, and tri-substituted alkyl, hydrogen, SO₂NH₂, SO₂NH-alkyl, and NHSO₂alkyl.

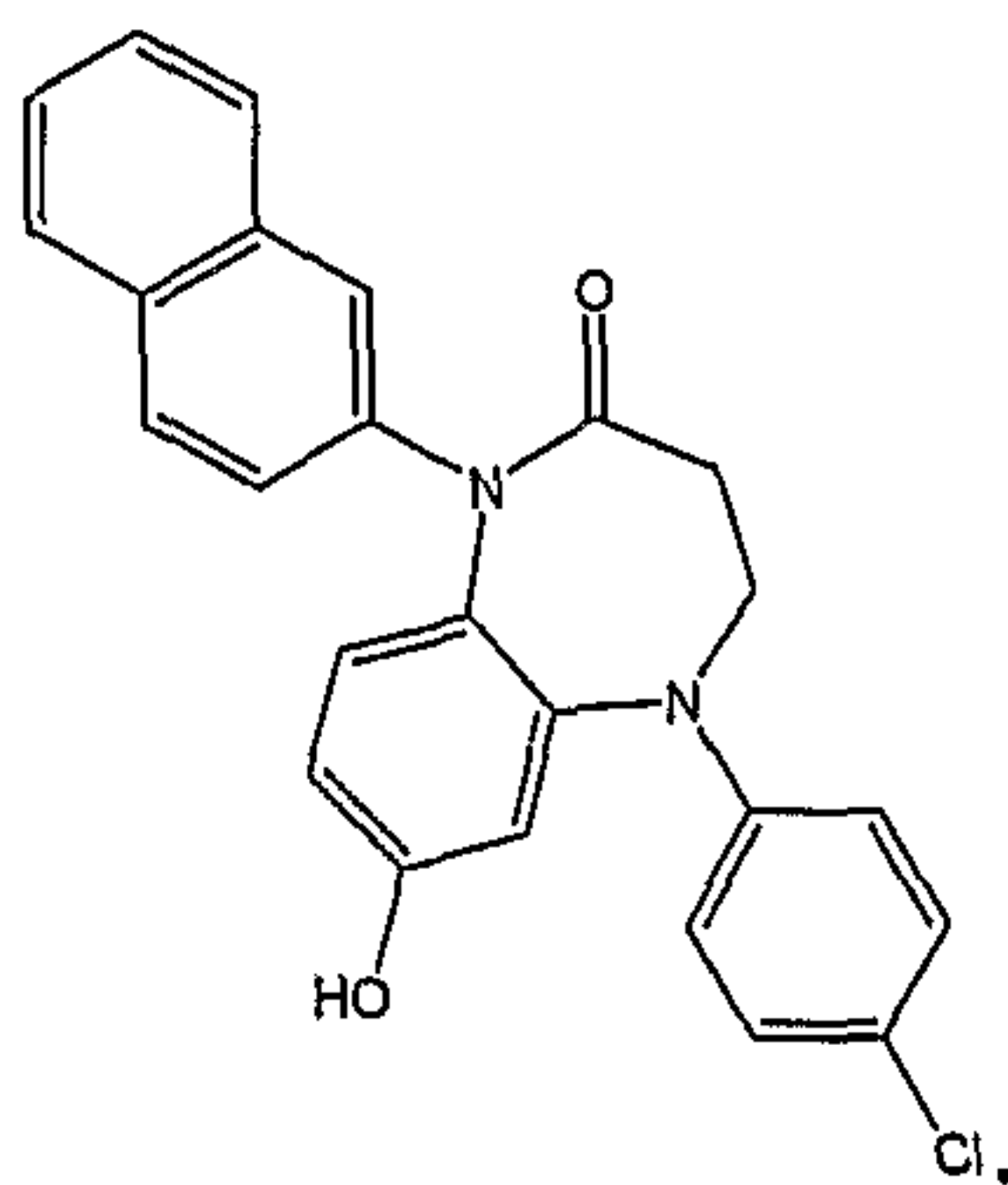
In some embodiments, R3 is an isostere of OH. In some embodiments, R3 is selected from the group consisting of hydrogen; halogen; OH; MnO4; a linear or branched, saturated or unsaturated, substituted or non-substituted, aliphatic chain having at least 2 carbons; a chemical moiety comprising Sulfur; a chemical moiety comprising Nitrogen. In some embodiments, R3 is selected from the group consisting of alkyl; mono-substituted alkyl; di-substituted alkyl; tri-substituted alkyl; (CH₂)_n wherein n=1-6; CN; N₃; CNO; NH₂; SH; CF₃; OCH₃; NCH₂CH(CH₂)N(CH₃)₂; NCH₂CHCH₂N(CH₃)₂; phenyl; 2-pyridyl; 3-pyridyl; 4-pyridyl; NCH₃; NCONHCH₃; CH₂OH; NHCONH₂; NHCOCH₃; NHSO₂CH₃; NHCN; NHCHO; SOCH₃; SO₂CH₃; CHNOH; CHNOCH₃; SCH₃; CH₂CO; CH₂SO₂; CONH; CH₂C(NO₂); CH₂C(NOMe); NHSO₂PH; NHCS; CH₂NHCO; COCH₂; NHCO₂; and NHCOS. In some embodiments, R3 is described by any of the isosteres described in, for example, Patani, G. and LaVoie, E.J., 1996, Chem. Rev. 96:3147-3176; herein incorporated by reference in its entirety.

In some embodiments, R4 is a chemical moiety that causes the benzodiazepine to lack a chiral center. In some embodiments, R4 is hydrogen, , or ; wherein R4' is a linear or branched, saturated or unsaturated, substituted or non-substituted, aliphatic chain having at least 2 carbons.

Certain embodiments of the present invention include crystal forms and formulations of exemplary compounds with the following formula:



In some embodiments, the present invention includes crystal forms and formulations of the following exemplary compound:



Additional exemplary compounds and uses useful in the present invention are described at U.S. Patent Publication 2004/0009972, published January 15, 2004, herein incorporated by reference in its entirety. Additional exemplary compounds and uses useful in the present invention are described in U.S. Provisional Patent Nos. 60/131,761, 60/165,511, 60/191,855, 60/312,560, 60/313,689, 60/396,670, 60/565,788, 60/607,599, 60/641,040, and U.S. Patent Application Nos. 11/324,419, 11/176,719, 11/110,228, 10/935,333, 10/886,450, 10/795,535, 10/634,114, 10/427,211, 10/427,212, 10/217,878, 09/767,283, 09/700,101, and related applications; each herein incorporated by reference in their entireties. Additional exemplary compounds and uses useful in the present invention are described in Atwal, *et al.*, Bioorg. Med. Chem. Lett. 14, 1027-1030 (2004) and Atwal, *et al.*, J. Med. Chem. 47, 1081-1084 (2004); each herein incorporated by reference in their entireties.

Further, it should be understood that the numerical ranges given throughout this disclosure should be construed as a flexible range that contemplates any possible subrange within that range. For example, the description of a group having the range of 1-10 carbons would also contemplate a group possessing a subrange of, for example, 1-3, 1-5, 1-8, or 2-3, 2-5, 2-8, 3-4, 3-5, 3-7, 3-9, 3-10, *etc.*, carbons. Thus, the range 1-10 should be understood to represent the outer boundaries of the range within which many possible subranges are clearly contemplated. Additional examples contemplating ranges in other contexts can be found throughout this disclosure wherein such ranges include analogous subranges within.

In summary, a large number of compounds are presented herein. Any one or more of these compounds can be used to treat a variety of dysregulatory disorders related to cellular death as described elsewhere herein. Additionally, any one or more of these compounds can be used to inhibit ATP Hydrolysis while not affecting cell synthesis or cell viability. Additionally, any one or more of these compounds can be used in combination with at least one other therapeutic agent (e.g., potassium channel openers, calcium channel blockers, sodium hydrogen exchanger inhibitors, antiarrhythmic agents, antiatherosclerotic

agents, anticoagulants, antithrombotic agents, prothrombolytic agents, fibrinogen antagonists, diuretics, antihypertensive agents, ATPase inhibitors, mineralocorticoid receptor antagonists, phosphodiesterase inhibitors, antidiabetic agents, anti-inflammatory agents, antioxidants, angiogenesis modulators, antiosteoporosis agents, hormone replacement therapies, hormone receptor modulators, oral contraceptives, antiobesity agents, antidepressants, antianxiety agents, antipsychotic agents, antiproliferative agents, antitumor agents, antiulcer and gastroesophageal reflux disease agents, growth hormone agents and/or growth hormone secretagogues, thyroid mimetics, anti-infective agents, antiviral agents, antibacterial agents, antifungal agents, cholesterol/lipid lowering agents and lipid profile therapies, and agents that mimic ischemic preconditioning and/or myocardial stunning, antiatherosclerotic agents, anticoagulants, antithrombotic agents, antihypertensive agents, antidiabetic agents, and antihypertensive agents selected from ACE inhibitors, AT-1 receptor antagonists, ET receptor antagonists, dual ET/AII receptor antagonists, and vasopepsidase inhibitors, or an antiplatelet agent selected from GPIIb/IIIa blockers, P2Y₁ and P2Y₁₂ antagonists, thromboxane receptor antagonists, and aspirin) in along with a pharmaceutically-acceptable carrier or diluent in a pharmaceutical composition. Additionally, any one or more of these compounds can be used to treat a mitochondrial F₁F₀ ATP hydrolase associated disorder (e.g., myocardial infarction, ventricular hypertrophy, coronary artery disease, non-Q wave MI, congestive heart failure, cardiac arrhythmias, unstable angina, chronic stable angina, Prinzmetal's angina, high blood pressure, intermittent claudication, peripheral occlusive arterial disease, thrombotic or thromboembolic symptoms of thromboembolic stroke, venous thrombosis, arterial thrombosis, cerebral thrombosis, pulmonary embolism, cerebral embolism, thrombophilia, disseminated intravascular coagulation, restenosis, atrial fibrillation, ventricular enlargement, atherosclerotic vascular disease, atherosclerotic plaque rupture, atherosclerotic plaque formation, transplant atherosclerosis, vascular remodeling atherosclerosis, cancer, surgery, inflammation, systematic infection, artificial surfaces, interventional cardiology, immobility, medication, pregnancy and fetal loss, and diabetic complications comprising retinopathy, nephropathy and neuropathy) in a patient. The above-described compounds can also be used in drug screening assays and other diagnostic methods.

IV. Pharmaceutical compositions, formulations, and exemplary administration routes and dosing considerations

Exemplary embodiments of various contemplated medicaments and pharmaceutical compositions are provided below.

5

A. Preparing Medicaments

The compounds of the present invention are useful in the preparation of medicaments to treat a variety of conditions associated with dysregulation of cell death, aberrant cell growth and hyperproliferation.

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In addition, the compounds are also useful for preparing medicaments for treating other disorders wherein the effectiveness of the compounds are known or predicted. Such disorders include, but are not limited to, neurological (*e.g.*, epilepsy) or neuromuscular disorders. The methods and techniques for preparing medicaments of a compound are well-known in the art. Exemplary pharmaceutical formulations and routes of delivery are

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described below. One of skill in the art will appreciate that any one or more of the compounds described herein, including the many specific embodiments, are prepared by applying standard pharmaceutical manufacturing procedures. Such medicaments can be delivered to the subject by using delivery methods that are well-known in the pharmaceutical arts.

20

B. Exemplary pharmaceutical compositions and formulation

In some embodiments of the present invention, the compositions are administered alone, while in some other embodiments, the compositions are preferably present in a pharmaceutical formulation comprising at least one active ingredient/agent (*e.g.*, benzodiazepine crystal forms and formulations and benzodiazepine related crystal forms and formulations), as defined above, together with a solid support or alternatively, together with one or more pharmaceutically acceptable carriers and optionally other therapeutic agents. Each carrier should be "acceptable" in the sense that it is compatible with the other ingredients of the formulation and not injurious to the subject.

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Contemplated formulations include those suitable oral, rectal, nasal, topical (including transdermal, buccal and sublingual), vaginal, parenteral (including subcutaneous, intramuscular, intravenous and intradermal) and pulmonary administration. In some embodiments, formulations are conveniently presented in unit dosage form and are prepared

by any method known in the art of pharmacy. Such methods include the step of bringing into association the active ingredient with the carrier which constitutes one or more accessory ingredients. In general, the formulations are prepared by uniformly and intimately bringing into association (*e.g.*, mixing) the active ingredient with liquid carriers or finely divided solid carriers or both, and then if necessary shaping the product.

Formulations of the present invention suitable for oral administration may be presented as discrete units such as capsules, cachets or tablets, wherein each preferably contains a predetermined amount of the active ingredient; as a powder or granules; as a solution or suspension in an aqueous or non-aqueous liquid; or as an oil-in-water liquid emulsion or a water-in-oil liquid emulsion. In other embodiments, the active ingredient is presented as a bolus, electuary, or paste, *etc.*

In some embodiments, tablets comprise at least one active ingredient and optionally one or more accessory agents/carriers are made by compressing or molding the respective agents. In preferred embodiments, compressed tablets are prepared by compressing in a suitable machine the active ingredient in a free-flowing form such as a powder or granules, optionally mixed with a binder (*e.g.*, povidone, gelatin, hydroxypropylmethyl cellulose), lubricant, inert diluent, preservative, disintegrant (*e.g.*, sodium starch glycolate, cross-linked povidone, cross-linked sodium carboxymethyl cellulose) surface-active or dispersing agent. Molded tablets are made by molding in a suitable machine a mixture of the powdered compound (*e.g.*, active ingredient) moistened with an inert liquid diluent. Tablets may optionally be coated or scored and may be formulated so as to provide slow or controlled release of the active ingredient therein using, for example, hydroxypropylmethyl cellulose in varying proportions to provide the desired release profile. Tablets may optionally be provided with an enteric coating, to provide release in parts of the gut other than the stomach.

Formulations suitable for topical administration in the mouth include lozenges comprising the active ingredient in a flavored basis, usually sucrose and acacia or tragacanth; pastilles comprising the active ingredient in an inert basis such as gelatin and glycerin, or sucrose and acacia; and mouthwashes comprising the active ingredient in a suitable liquid carrier.

Pharmaceutical compositions for topical administration according to the present invention are optionally formulated as ointments, creams, suspensions, lotions, powders, solutions, pastes, gels, sprays, aerosols or oils. In alternatively embodiments, topical

formulations comprise patches or dressings such as a bandage or adhesive plasters impregnated with active ingredient(s), and optionally one or more excipients or diluents. In preferred embodiments, the topical formulations include a compound(s) that enhances absorption or penetration of the active agent(s) through the skin or other affected areas.

- 5 Examples of such dermal penetration enhancers include dimethylsulfoxide (DMSO) and related analogues.

If desired, the aqueous phase of a cream base includes, for example, at least about 30% w/w of a polyhydric alcohol, *i.e.*, an alcohol having two or more hydroxyl groups such as propylene glycol, butane-1,3-diol, mannitol, sorbitol, glycerol and polyethylene glycol
10 and mixtures thereof.

In some embodiments, oily phase emulsions of this invention are constituted from known ingredients in an known manner. This phase typically comprises an lone emulsifier (otherwise known as an emulgent), it is also desirable in some embodiments for this phase to further comprises a mixture of at least one emulsifier with a fat or an oil or with both a fat
15 and an oil.

Preferably, a hydrophilic emulsifier is included together with a lipophilic emulsifier so as to act as a stabilizer. In some embodiments it is also preferable to include both an oil and a fat. Together, the emulsifier(s) with or without stabilizer(s) make up the so-called emulsifying wax, and the wax together with the oil and/or fat make up the so-called
20 emulsifying ointment base which forms the oily dispersed phase of the cream formulations.

Emulgents and emulsion stabilizers suitable for use in the formulation of the present invention include Tween 60, Span 80, cetostearyl alcohol, myristyl alcohol, glyceryl monostearate and sodium lauryl sulfate.

The choice of suitable oils or fats for the formulation is based on achieving the
25 desired properties (*e.g.*, cosmetic properties), since the solubility of the active compound/agent in most oils likely to be used in pharmaceutical emulsion formulations is very low. Thus creams should preferably be a non-greasy, non-staining and washable products with suitable consistency to avoid leakage from tubes or other containers. Straight or branched chain, mono- or dibasic alkyl esters such as di-isoadipate, isocetyl stearate,
30 propylene glycol diester of coconut fatty acids, isopropyl myristate, decyl oleate, isopropyl palmitate, butyl stearate, 2-ethylhexyl palmitate or a blend of branched chain esters known as Crodamol CAP may be used, the last three being preferred esters. These may be used alone or in combination depending on the properties required. Alternatively, high melting

point lipids such as white soft paraffin and/or liquid paraffin or other mineral oils can be used.

Formulations suitable for topical administration to the eye also include eye drops wherein the active ingredient is dissolved or suspended in a suitable carrier, especially an aqueous solvent for the agent.

Formulations for rectal administration may be presented as a suppository with suitable base comprising, for example, cocoa butter or a salicylate.

Formulations suitable for vaginal administration may be presented as pessaries, creams, gels, pastes, foams or spray formulations containing in addition to the agent, such carriers as are known in the art to be appropriate.

Formulations suitable for nasal administration, wherein the carrier is a solid, include coarse powders having a particle size, for example, in the range of about 20 to about 500 microns which are administered in the manner in which snuff is taken, *i.e.*, by rapid inhalation (*e.g.*, forced) through the nasal passage from a container of the powder held close up to the nose. Other suitable formulations wherein the carrier is a liquid for administration include, but are not limited to, nasal sprays, drops, or aerosols by nebulizer, and include aqueous or oily solutions of the agents.

Formulations suitable for parenteral administration include aqueous and non-aqueous isotonic sterile injection solutions which may contain antioxidants, buffers, bacteriostats and solutes which render the formulation isotonic with the blood of the intended recipient; and aqueous and non-aqueous sterile suspensions which may include suspending agents and thickening agents, and liposomes or other microparticulate systems which are designed to target the compound to blood components or one or more organs. In some embodiments, the formulations are presented/formulated in unit-dose or multi-dose sealed containers, for example, ampoules and vials, and may be stored in a freeze-dried (lyophilized) condition requiring only the addition of the sterile liquid carrier, for example water for injections, immediately prior to use. Extemporaneous injection solutions and suspensions may be prepared from sterile powders, granules and tablets of the kind previously described.

Preferred unit dosage formulations are those containing a daily dose or unit, daily subdose, as herein above-recited, or an appropriate fraction thereof, of an agent.

It should be understood that in addition to the ingredients particularly mentioned above, the formulations of this invention may include other agents conventional in the art

having regard to the type of formulation in question, for example, those suitable for oral administration may include such further agents as sweeteners, thickeners and flavoring agents. It also is intended that the agents, compositions and methods of this invention be combined with other suitable compositions and therapies. Still other formulations
5 optionally include food additives (suitable sweeteners, flavorings, colorings, *etc.*), phytonutrients (*e.g.*, flax seed oil), minerals (*e.g.*, Ca, Fe, K, *etc.*), vitamins, and other acceptable compositions (*e.g.*, conjugated linoelic acid), extenders, and stabilizers, *etc.*

C. Exemplary administration routes and dosing considerations

10 Various delivery systems are known and can be used to administer a therapeutic agents (*e.g.*, benzodiazepine crystal forms and formulations and benzodiazepine related crystal forms and formulations) of the present invention, *e.g.*, encapsulation in liposomes, microparticles, microcapsules, receptor-mediated endocytosis, and the like. Methods of delivery include, but are not limited to, intra-arterial, intra-muscular, intravenous, intranasal,
15 and oral routes. In specific embodiments, it may be desirable to administer the pharmaceutical compositions of the invention locally to the area in need of treatment; this may be achieved by, for example, and not by way of limitation, local infusion during surgery, injection, or by means of a catheter.

The agents identified herein as effective for their intended purpose can be
20 administered to subjects or individuals susceptible to or at risk of developing pathological growth of target cells and condition correlated with this. When the agent is administered to a subject such as a mouse, a rat or a human patient, the agent can be added to a pharmaceutically acceptable carrier and systemically or topically administered to the subject. To determine patients that can be beneficially treated, a tissue sample is removed
25 from the patient and the cells are assayed for sensitivity to the agent.

Therapeutic amounts are empirically determined and vary with the pathology being treated, the subject being treated and the efficacy and toxicity of the agent. When delivered to an animal, the method is useful to further confirm efficacy of the agent. One example of an animal model is MLR/MpJ-*lpr/lpr* ("MLR-*lpr*") (available from Jackson Laboratories,
30 Bal Harbor, Maine). MLR-*lpr* mice develop systemic autoimmune disease. Alternatively, other animal models can be developed by inducing tumor growth, for example, by subcutaneously inoculating nude mice with about 10^5 to about 10^9 hyperproliferative, cancer or target cells as defined herein. When the tumor is established, the compounds

described herein are administered, for example, by subcutaneous injection around the tumor. Tumor measurements to determine reduction of tumor size are made in two dimensions using venier calipers twice a week. Other animal models may also be employed as appropriate. Such animal models for the above-described diseases and conditions are
5 well-known in the art.

In some embodiments, *in vivo* administration is effected in one dose, continuously or intermittently throughout the course of treatment. Methods of determining the most effective means and dosage of administration are well known to those of skill in the art and vary with the composition used for therapy, the purpose of the therapy, the target cell being
10 treated, and the subject being treated. Single or multiple administrations are carried out with the dose level and pattern being selected by the treating physician.

Suitable dosage formulations and methods of administering the agents are readily determined by those of skill in the art. Preferably, the compounds are administered at about 0.01 mg/kg to about 200 mg/kg, more preferably at about 0.1 mg/kg to about 100 mg/kg,
15 even more preferably at about 0.5 mg/kg to about 50 mg/kg. When the compounds described herein are co-administered with another agent (*e.g.*, as sensitizing agents), the effective amount may be less than when the agent is used alone.

The pharmaceutical compositions can be administered orally, intranasally, parenterally or by inhalation therapy, and may take the form of tablets, lozenges, granules,
20 capsules, pills, ampoules, suppositories or aerosol form. They may also take the form of suspensions, solutions and emulsions of the active ingredient in aqueous or nonaqueous diluents, syrups, granulates or powders. In addition to an agent of the present invention, the pharmaceutical compositions can also contain other pharmaceutically active compounds or a plurality of compounds of the invention.

More particularly, an agent of the present invention also referred to herein as the active ingredient, may be administered for therapy by any suitable route including, but not limited to, oral, rectal, nasal, topical (including, but not limited to, transdermal, aerosol, buccal and sublingual), vaginal, parental (including, but not limited to, subcutaneous, intramuscular, intravenous and intradermal) and pulmonary. It is also appreciated that the
30 preferred route varies with the condition and age of the recipient, and the disease being treated.

Ideally, the agent should be administered to achieve peak concentrations of the active compound at sites of disease. This may be achieved, for example, by the intravenous

injection of the agent, optionally in saline, or orally administered, for example, as a tablet, capsule or syrup containing the active ingredient.

Desirable blood levels of the agent may be maintained by a continuous infusion to provide a therapeutic amount of the active ingredient within disease tissue. The use of operative combinations is contemplated to provide therapeutic combinations requiring a lower total dosage of each component antiviral agent than may be required when each individual therapeutic compound or drug is used alone, thereby reducing adverse effects.

D. Exemplary co-administration routes and dosing considerations

The present invention also includes methods involving co-administration of the compounds described herein with one or more additional active agents. Indeed, it is a further aspect of this invention to provide methods for enhancing prior art therapies and/or pharmaceutical compositions by co-administering a compound of this invention. In co-administration procedures, the agents may be administered concurrently or sequentially. In one embodiment, the compounds described herein are administered prior to the other active agent(s). The pharmaceutical formulations and modes of administration may be any of those described above. In addition, the two or more co-administered chemical agents, biological agents or radiation may each be administered using different modes or different formulations.

The agent or agents to be co-administered depends on the type of condition being treated. For example, when the condition being treated is cancer, the additional agent can be a chemotherapeutic agent or radiation. When the condition being treated is an autoimmune disorder, the additional agent can be an immunosuppressant or an anti-inflammatory agent. When the condition being treated is chronic inflammation, the additional agent can be an anti-inflammatory agent. The additional agents to be co-administered, such as anticancer, immunosuppressant, anti-inflammatory, and can be any of the well-known agents in the art, including, but not limited to, those that are currently in clinical use. The determination of appropriate type and dosage of radiation treatment is also within the skill in the art or can be determined with relative ease.

Treatment of the various conditions associated with abnormal apoptosis is generally limited by the following two major factors: (1) the development of drug resistance and (2) the toxicity of known therapeutic agents. In certain cancers, for example, resistance to chemicals and radiation therapy has been shown to be associated with inhibition of

apoptosis. Some therapeutic agents have deleterious side effects, including non-specific lymphotoxicity, renal and bone marrow toxicity.

The methods described herein address both these problems. Drug resistance, where increasing dosages are required to achieve therapeutic benefit, is overcome by co-administering the compounds described herein with the known agent. The compounds described herein appear to sensitize target cells to known agents (and vice versa) and, accordingly, less of these agents are needed to achieve a therapeutic benefit.

The sensitizing function of the claimed compounds also addresses the problems associated with toxic effects of known therapeutics. In instances where the known agent is toxic, it is desirable to limit the dosages administered in all cases, and particularly in those cases where drug resistance has increased the requisite dosage. When the claimed compounds are co-administered with the known agent, they reduce the dosage required which, in turn, reduces the deleterious effects. Further, because the claimed compounds are themselves both effective and non-toxic in large doses, co-administration of proportionally more of these compounds than known toxic therapeutics will achieve the desired effects while minimizing toxic effects.

V. Drug screens

In preferred embodiments of the present invention, the compounds of the present invention, and other potentially useful compounds, are screened for their biological activity (e.g., ability to initiate cell death alone or in combination with other compounds). In preferred embodiments of the present invention, the compounds of the present invention, and other potentially useful compounds, are screened for their binding affinity to the oligomycin sensitivity conferring protein (OSCP) portion of the mitochondrial ATP synthase complex. In particularly preferred embodiments, compounds are selected for use in the methods of the present invention by measuring their binding affinity to recombinant OSCP protein. A number of suitable screens for measuring the binding affinity of drugs and other small molecules to receptors are known in the art. In some embodiments, binding affinity screens are conducted in *in vitro* systems. In other embodiments, these screens are conducted in *in vivo* or *ex vivo* systems. While in some embodiments quantifying the intracellular level of ATP following administration of the compounds of the present invention provides an indication of the efficacy of the methods, preferred embodiments of the present invention do not require intracellular ATP or pH level quantification.

Additional embodiments are directed to measuring levels (*e.g.*, intracellular) of superoxide in cells and/or tissues to measure the effectiveness of particular contemplated methods and compounds of the present invention. In this regard, those skilled in the art will appreciate and be able to provide a number of assays and methods useful for measuring
5 superoxide levels in cells and/or tissues.

In some embodiments, structure-based virtual screening methodologies are contemplated for predicting the binding affinity of compounds of the present invention with OSCP.

In some embodiments, compounds are screened in cell culture or *in vivo* (*e.g.*, non-
10 human or human mammals) for their ability to modulate mitochondrial ATP synthase activity. Any suitable assay may be utilized, including, but not limited to, cell proliferation assays (Commercially available from, *e.g.*, Promega, Madison, WI and Stratagene, La Jolla, CA) and cell based dimerization assays. (*See e.g.*, Fuh *et al.*, Science, 256:1677 [1992]; Colosi *et al.*, J. Biol. Chem., 268:12617 [1993]). Additional assay formats that find use
15 with the present invention include, but are not limited to, assays for measuring cellular ATP levels, and cellular superoxide levels.

Any suitable assay that allows for a measurement of the rate of binding or the affinity of a benzodiazepine or other compound to the OSCP may be utilized. Examples include, but are not limited to, competition binding using Bz-423, surface plasma resonance
20 (SPR) and radio-immunoprecipitation assays (Lowman *et al.*, J. Biol.Chem. 266:10982 [1991]). Surface Plasmon Resonance techniques involve a surface coated with a thin film of a conductive metal, such as gold, silver, chrome or aluminum, in which electromagnetic waves, called Surface Plasmons, can be induced by a beam of light incident on the metal glass interface at a specific angle called the Surface Plasmon Resonance angle. Modulation
25 of the refractive index of the interfacial region between the solution and the metal surface following binding of the captured macromolecules causes a change in the SPR angle which can either be measured directly or which causes the amount of light reflected from the underside of the metal surface to change. Such changes can be directly related to the mass and other optical properties of the molecules binding to the SPR device surface. Several
30 biosensor systems based on such principles have been disclosed (*See e.g.*, WO 90/05305). There are also several commercially available SPR biosensors (*e.g.*, BiaCore, Uppsala, Sweden).

In some embodiments, compounds are screened in cell culture or *in vivo* (e.g., non-human or human mammals) for their ability to modulate mitochondrial ATP synthase activity. Any suitable assay may be utilized, including, but not limited to, cell proliferation assays (Commercially available from, e.g., Promega, Madison, WI and Stratagene, La Jolla, CA) and cell based dimerization assays. (See e.g., Fuh *et al.*, Science, 256:1677 [1992]; Colosi *et al.*, J. Biol. Chem., 268:12617 [1993]). Additional assay formats that find use with the present invention include, but are not limited to, assays for measuring cellular ATP levels, and cellular superoxide levels.

The present invention also provides methods of modifying and derivatizing the compositions of the present invention to increase desirable properties (e.g., binding affinity, activity, and the like), or to minimize undesirable properties (e.g., nonspecific reactivity, toxicity, and the like). The principles of chemical derivatization are well understood. In some embodiments, iterative design and chemical synthesis approaches are used to produce a library of derivatized child compounds from a parent compound. In other embodiments, rational design methods are used to predict and model *in silico* ligand-receptor interactions prior to confirming results by routine experimentation.

VI. Therapeutic Application

A. General Therapeutic Application

In particularly preferred embodiments, the compositions of the present invention provide therapeutic benefits to patients suffering from any one or more of a number of conditions (e.g., diseases characterized by dysregulation of necrosis and/or apoptosis processes in a cell or tissue, disease characterized by aberrant cell growth and/or hyperproliferation, *etc.*) by modulating (e.g., inhibiting or promoting) the activity of the mitochondrial ATP synthase (as referred to as mitochondrial F₀F₁ ATPase) complexes in affected cells or tissues (e.g., myocardial infarction, ventricular hypertrophy, coronary artery disease, non-Q wave MI, congestive heart failure, cardiac arrhythmias, unstable angina, chronic stable angina, Prinzmetal's angina, high blood pressure, intermittent claudication, peripheral occlusive arterial disease, thrombotic or thromboembolic symptoms of thromboembolic stroke, venous thrombosis, arterial thrombosis, cerebral thrombosis, pulmonary embolism, cerebral embolism, thrombophilia, disseminated intravascular coagulation, restenosis, atrial fibrillation, ventricular enlargement, atherosclerotic vascular disease, atherosclerotic plaque rupture, atherosclerotic plaque formation, transplant

atherosclerosis, vascular remodeling atherosclerosis, cancer, surgery, inflammation, systematic infection, artificial surfaces, interventional cardiology, immobility, medication, pregnancy and fetal loss, and diabetic complications comprising retinopathy, nephropathy and neuropathy). In further preferred embodiments, the compositions of the present invention are used to treat autoimmune/chronic inflammatory conditions (*e.g.*, psoriasis). In even further embodiments, the compositions of the present invention are used in conjunction with stenosis therapy to treat compromised (*e.g.*, occluded) vessels. Indeed, any application where benzodiazepines find use is contemplated by the present invention.

In particularly preferred embodiments, the compositions of the present invention inhibit the activity of mitochondrial ATP synthase complex by binding to a specific subunit or subunits of this multi-subunit protein complex. While the present invention is not limited to any particular mechanism, nor to any understanding of the action of the agents being administered, in some embodiments, it is contemplated that the compositions of the present invention bind to the oligomycin sensitivity conferring protein (OSCP) portion of the mitochondrial ATP synthase complex, to the OSCP/F1 junction, or to the F1 subunit. Likewise, it is further contemplated that when the compositions of the present invention bind to the OSCP the initial affect is overall inhibition of the mitochondrial ATP synthase complex, and that the downstream consequence of binding is a change in ATP or pH level and the production of reactive oxygen species (*e.g.*, O₂⁻). In still other preferred embodiments, while the present invention is not limited to any particular mechanism, nor to any understanding of the action of the agents being administered, it is contemplated that the generation of free radicals ultimately results in cell killing. In yet other embodiments, while the present invention is not limited to any particular mechanism, nor to any understanding of the action of the agents being administered, it is contemplated that the inhibiting mitochondrial ATP synthase complex using the compositions and methods of the present invention provides therapeutically useful inhibition of cell proliferation.

Accordingly, preferred methods embodied in the present invention, provide therapeutic benefits to patients by providing compounds of the present invention that modulate (*e.g.*, inhibiting or promoting) the activity of the mitochondrial ATP synthase complexes in affected cells or tissues via binding to the oligomycin sensitivity conferring protein (OSCP) portion of the mitochondrial ATP synthase complex. Importantly, by itself the OSCP, the OSCP/F1 junction, or the F1 subunit has no biological activity.

Thus, in one broad sense, preferred embodiments of the present invention are directed to the discovery that many diseases characterized by dysregulation of necrosis and/or apoptosis processes in a cell or tissue, or diseases characterized by aberrant cell growth and/or hyperproliferation, *etc.*, can be treated by modulating the activity of the mitochondrial ATP synthase complex including, but not limited to, by binding to the oligomycin sensitivity conferring protein (OSCP) / F1 components thereof. The present invention is not intended to be limited, however, to the practice of the compositions and methods explicitly described herein. Indeed, those skilled in the art will appreciate that a number of additional compounds not specifically recited herein (*e.g.*, non-benzodiazepine derivatives) are suitable for use in the methods disclosed herein of modulating the activity of mitochondrial ATP synthase.

The present invention thus specifically contemplates that any number of suitable compounds presently known in the art, or developed later, can optionally find use in the methods of the present invention. For example, compounds including, but not limited to, oligomycin, ossamycin, cytovaricin, apoptolidin, bafilomycin, resveratrol, piceatannol, and dicyclohexylcarbodiimide (DCCD), and the like, find use in the methods of the present invention. The present invention is not intended, however, to be limited to the methods or compounds specified above. In one embodiment, that compounds potentially useful in the methods of the present invention may be selected from those suitable as described in the scientific literature. (*See e.g.*, K.B. Wallace and A.A. Starkov, *Annu. Rev. Pharmacol. Toxicol.*, 40:353-388 [2000]; A.R. Solomon *et al.*, *Proc. Nat. Acad. Sci. U.S.A.*, 97(26):14766-14771 [2000]).

In some embodiments, compounds potentially useful in methods of the present invention are screened against the National Cancer Institute's (NCI-60) cancer cell lines for efficacy. (*See e.g.*, A. Monks *et al.*, *J. Natl. Cancer Inst.*, 83:757-766 [1991]; and K.D. Paull *et al.*, *J. Natl. Cancer Inst.*, 81:1088-1092 [1989]). Additional screens suitable screens (*e.g.*, autoimmunity disease models, *etc.*) are within the skill in the art.

In one aspect, derivatives (*e.g.*, pharmaceutically acceptable salts, analogs, stereoisomers, and the like) of the exemplary compounds or other suitable compounds are also contemplated as being useful in the methods of the present invention.

In other preferred embodiments, the compositions of the present invention are used in conjunction with stenosis therapy to treat compromised (*e.g.*, occluded) vessels. In

further embodiments, the compositions of the present invention are used in conjunction with stenosis therapy to treat compromised cardiac vessels.

Vessel stenosis is a condition that develops when a vessel (*e.g.*, aortic valve) becomes narrowed. For example, aortic valve stenosis is a heart condition that develops
5 when the valve between the lower left chamber (left ventricle) of the heart and the major blood vessel called the aorta becomes narrowed. This narrowing (*e.g.*, stenosis) creates too small a space for the blood to flow to the body. Normally the left ventricle pumps oxygen-rich blood to the body through the aorta, which branches into a system of arteries throughout the body. When the heart pumps, the 3 flaps, or leaflets, of the aortic valve open
10 one way to allow blood to flow from the ventricle into the aorta. Between heartbeats, the flaps close to form a tight seal so that blood does not leak backward through the valve. If the aortic valve is damaged, it may become narrowed (stenosed) and blood flow may be reduced to organs in the body, including the heart itself. The long-term outlook for people with aortic valve stenosis is poor once symptoms develop. People with untreated aortic
15 valve stenosis who develop symptoms of heart failure usually have a life expectancy of 3 years or less.

Several types of treatment exist for treating compromised valves (*e.g.*, balloon dilation, ablation, atherectomy or laser treatment). One type of treatment for compromised cardiac valves is angioplasty. Angioplasty involves inserting a balloon-tipped tube, or
20 catheter, into a narrow or blocked artery in an attempt to open it. By inflating and deflating the balloon several times, physicians usually are able to widen the artery.

A common limitation of angioplasty or valve expansion procedures is restenosis. Restenosis is the reclosure of a peripheral or coronary artery following trauma to that artery caused by efforts to open a stenosed portion of the artery, such as, for example, by balloon
25 dilation, ablation, atherectomy or laser treatment of the artery. For these angioplasty procedures, restenosis occurs at a rate of about 20-50% depending on the definition, vessel location, lesion length and a number of other morphological and clinical variables. Restenosis is believed to be a natural healing reaction to the injury of the arterial wall that is caused by angioplasty procedures. The healing reaction begins with the thrombotic
30 mechanism at the site of the injury. The final result of the complex steps of the healing process can be intimal hyperplasia, the uncontrolled migration and proliferation of medial smooth muscle cells, combined with their extracellular matrix production, until the artery is again stenosed or occluded.

In an attempt to prevent restenosis, metallic intravascular stents have been permanently implanted in coronary or peripheral vessels. The stent is typically inserted by catheter into a vascular lumen and expanded into contact with the diseased portion of the arterial wall, thereby providing mechanical support for the lumen. However, it has been found that restenosis can still occur with such stents in place. Also, the stent itself can cause undesirable local thrombosis. To address the problem of thrombosis, persons receiving stents also receive extensive systemic treatment with anticoagulant and antiplatelet drugs.

To address the restenosis problem, it has been proposed to provide stents which are seeded with endothelial cells (Dichek, D. A. et al Seeding of Intravascular Stents With Genetically Engineered Endothelial Cells; Circulation 1989; 80: 1347-1353). In that experiment, sheep endothelial cells that had undergone retrovirus-mediated gene transfer for either bacterial beta-galactosidase or human tissue-type plasminogen activator were seeded onto stainless steel stents and grown until the stents were covered. The cells were therefore able to be delivered to the vascular wall where they could provide therapeutic proteins. Other methods of providing therapeutic substances to the vascular wall by means of stents have also been proposed such as in international patent application WO 91/12779 "Intraluminal Drug Eluting Prosthesis" and international patent application WO 90/13332 "Stent With Sustained Drug Delivery". In those applications, it is suggested that antiplatelet agents, anticoagulant agents, antimicrobial agents, anti-inflammatory agents, antimetabolic agents and other drugs could be supplied in stents to reduce the incidence of restenosis. Further, other vasoreactive agents such as nitric oxide releasing agents could also be used.

An additional cause of restenosis is the over-proliferation of treated tissue. In preferred embodiments, the anti-proliferative properties of the present invention inhibit restenosis. Drug-eluting stents are well known in the art (*see, e.g.*, U.S. Patent No.: 5,697,967; U.S. Patent No.: 5,599,352; and U.S. Patent No.: 5,591,227; each of which are herein incorporated by reference). In preferred embodiments, the compositions of the present invention are eluted from drug-eluting stents in the treatment of compromised (*e.g.*, occluded) vessels. In further embodiments, the compositions of the present invention are eluted from drug-eluting stents in the treatment of compromised cardiac vessels.

Those skilled in the art of preparing pharmaceutical compounds and formulations will appreciate that when selecting optional compounds for use in the methods disclosed herein, that suitability considerations include, but are not limited to, the toxicity, safety, efficacy, availability, and cost of the particular compounds.

In preferred embodiments, pharmaceutical compositions comprise compounds of the invention and, for example, therapeutic agents (e.g., antiatherosclerotic agents, anticoagulants, antithrombotic agents, antihypertensive agents, and antidiabetic agents). Antihypertensive agents include, but are not limited to, ACE inhibitors, AT-1 receptor
5 antagonists, ET receptor antagonists, dual ET/AII receptor antagonists, and vasopepsidase inhibitors, or an antiplatelet agent selected from GPIIb/IIIa blockers, P2Y₁ and P2Y₁₂ antagonists, thromboxane receptor antagonists, and aspirin.

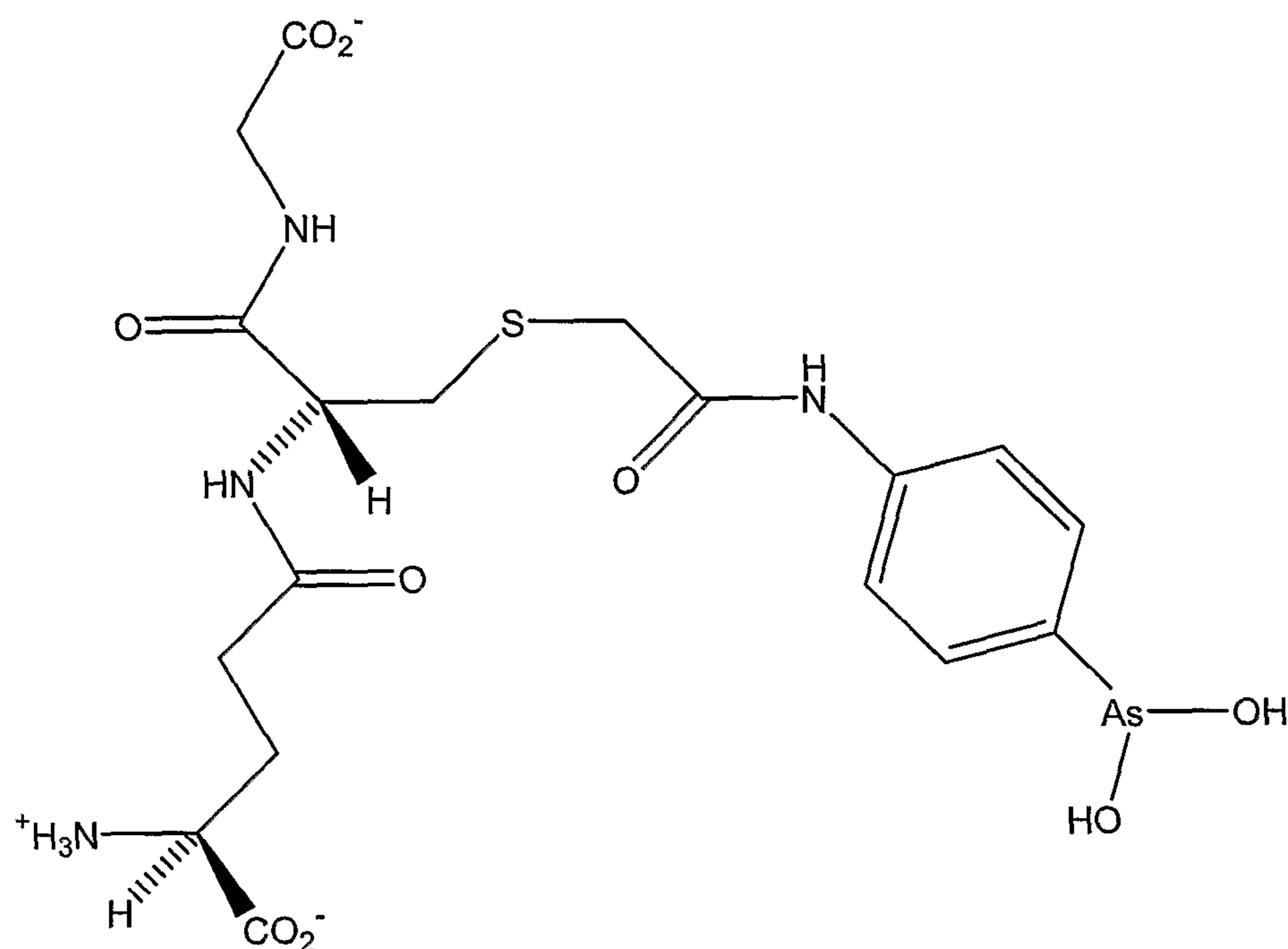
In preferred embodiments, the compounds of the present invention are useful in treating a mitochondrial F₁F₀ ATP hydrolase associated disorder (e.g., myocardial
10 infarction, ventricular hypertrophy, coronary artery disease, non-Q wave MI, congestive heart failure, cardiac arrhythmias, unstable angina, chronic stable angina, Prinzmetal's angina, high blood pressure, intermittent claudication, peripheral occlusive arterial disease, thrombotic or thromboembolic symptoms of thromboembolic stroke, venous thrombosis, arterial thrombosis, cerebral thrombosis, pulmonary embolism, cerebral embolism,
15 thrombophilia, disseminated intravascular coagulation, restenosis, atrial fibrillation, ventricular enlargement, atherosclerotic vascular disease, atherosclerotic plaque rupture, atherosclerotic plaque formation, transplant atherosclerosis, vascular remodeling atherosclerosis, cancer, surgery, inflammation, systematic infection, artificial surfaces, interventional cardiology, immobility, medication, pregnancy and fetal loss, and diabetic
20 complications comprising retinopathy, nephropathy and neuropathy) in a patient.

B. Autoimmune Disorder and Chronic Inflammatory Disorder Therapeutic Application

Autoimmune disorders and chronic inflammatory disorders often result from
25 dysfunctional cellular proliferation regulation and/or cellular apoptosis regulation. Mitochondria perform a key role in the control and execution of cellular apoptosis. The mitochondrial permeability transition pore (MPTP) is a pore that spans the inner and outer mitochondrial membranes and functions in the regulation of proapoptotic particles. Transient MPTP opening results in the release of cytochrome c and the apoptosis inducing
30 factor from the mitochondrial intermembrane space, resulting in cellular apoptosis.

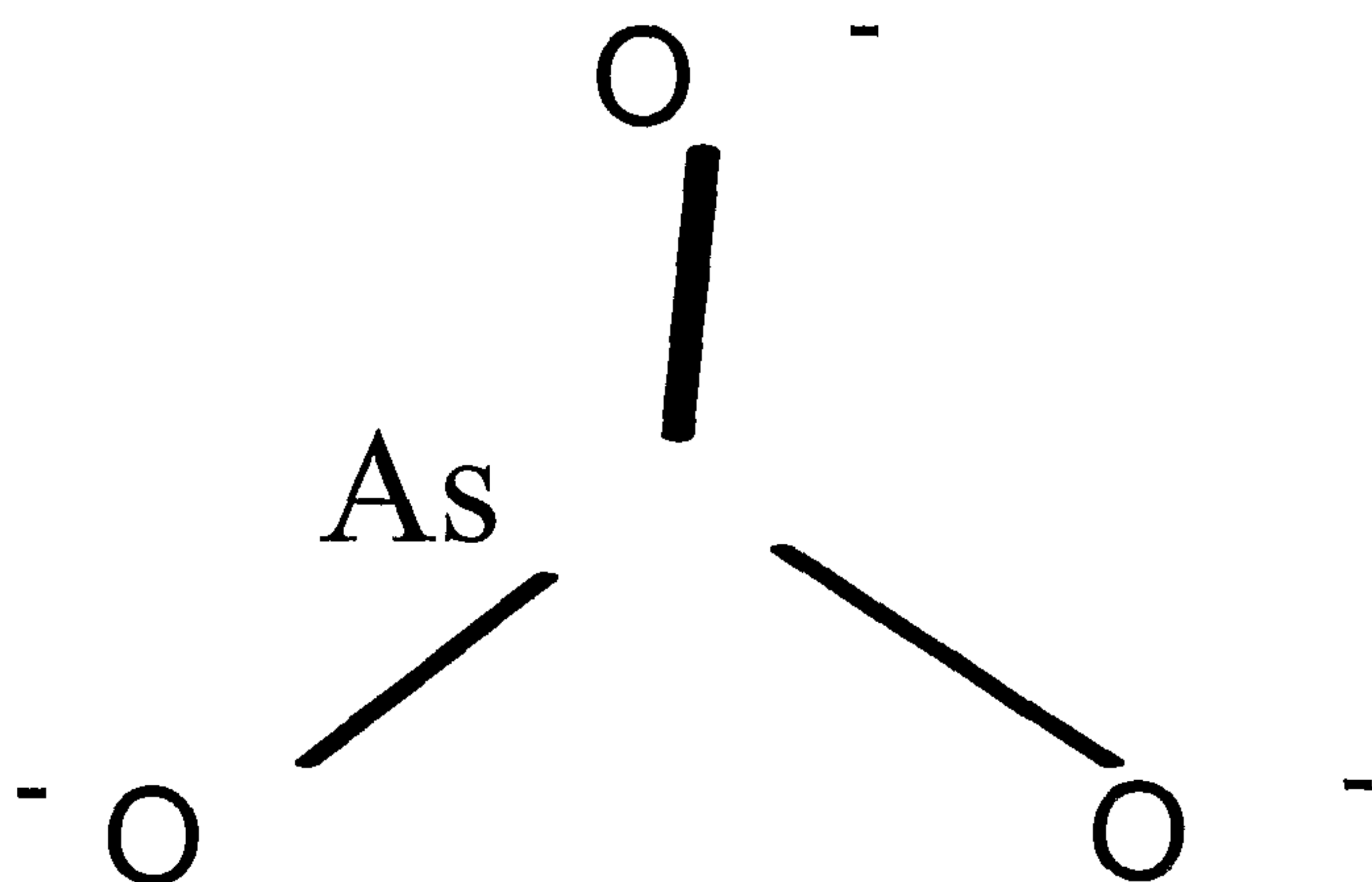
The oligomycin sensitivity conferring protein (OSCP) is a subunit of the F₀F₁ mitochondrial ATP synthase/ATPase and functions in the coupling of a proton gradient across the F₀ sector of the enzyme in the mitochondrial membrane. In preferred

embodiments, compounds of the present invention binds the OSCP, the OSCP / F1 junction, or the F1 subunit increases superoxide and cytochrome c levels, increases cellular apoptosis, and inhibits cellular proliferation. The adenine nucleotide translocator (ANT) is a 30kDa
 5 permeability transition pore (MPTP). Thiol oxidizing or alkylating agents are powerful activators of the MPTP that act by modifying one or more of three unpaired cysteines in the matrix side of the ANT. 4-(N-(S-glutathionylacetyl)amino) phenylarsenoxide,



inhibits the ANT.

10 The compounds and methods of the present invention are useful in the treatment of autoimmune disorders and chronic inflammatory disorders. In such embodiments, the present invention provides a subject suffering from an autoimmune disorder and/or a chronic inflammatory disorder, and a composition comprising, for example,



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entireties). Normal epidermal growth does not appear to be as dependent on EGF receptor function as hyperplastic growth (*see, e.g.,* Varani J, et al., (2001) J. Invest. Dermatol 117:1335-1341; Varani J, et al., (1998) Pathobiology 66:253-259; each herein incorporated by reference in their entireties). Likewise, function of the dermis in intact skin does not
5 depend on EGF receptor function (*see, e.g.,* Varani J, et al., (2001) J. Invest. Dermatol 117:1335-1341; herein incorporated by reference in its entirety).

The central role of the EGF receptor in regulating hyperplastic epithelial growth makes the EGF receptor tyrosine kinase a target for antiproliferative agents. Likewise, the series of signaling molecules engaged downstream of this receptor are additional points at
10 which keratinocyte growth can be interrupted. The mitogen activated protein kinase (MAPK) cascade is activated by the EGF receptor (*see, e.g.,* Marques, S. A., et al., (2002) J Pharmacol Exp Ther 300, 1026-1035; herein incorporated by reference in its entirety). In hyperproliferative epidermis, but not in normal epidermis, extracellular signal-regulated kinases 1/2 (Erk 1/2) are activated in basal and suprabasal keratinocytes and contribute to
15 epidermal hyperproliferation (*see, e.g.,* Haase, I., et al., (2001) J Clin Invest 108, 527-536; Takahashi, H., et al., (2002) J Dermatol Sci 30, 94-99; each herein incorporated by reference in their entireties). In culture models, keratinocyte growth regulation through the EGF receptor results in increased MAPK activity. In keratinocytes, growth factor–stimulated MAPK activity is also dependent on integrin engagement and extracellular
20 matrix molecules that bind integrins are capable of independently activating MAPKs and increasing keratinocyte proliferation (*see, e.g.,* Haase, I., et al., (2001) J Clin Invest 108, 527-536; herein incorporated by reference in its entirety). The proliferation of other skin cells, including fibroblasts, is less dependent on Erk 1/2 activity, making Erk inhibition a potentially useful characteristic to evaluate lead compounds for potential utility against
25 epidermal hyperplasia.

In preferred embodiments, compounds of the present invention are used for treating epidermal hyperplasias.

In preferred embodiments, compounds of the present invention are used in treating psoriasis. Psoriasis is common and chronic epidermal hyperplasia. Plaque psoriasis is the
30 most common type of psoriasis and is characterized by red skin covered with silvery scales and inflammation. Patches of circular to oval shaped red plaques that itch or burn are typical of plaque psoriasis. The patches are usually found on the arms, legs, trunk, or scalp but may be found on any part of the skin. The most typical areas are the knees and elbows.

Psoriasis is not contagious and can be inherited. Environmental factors, such as smoking, sun exposure, alcoholism, and HIV infection, may affect how often the psoriasis occurs and how long the flares up last.

Treatment of psoriasis includes topical steroids, coal tar, keratolytic agents, vitamin
5 D-3 analogs, and topical retinoids. Topical steroids are agents used to reduce plaque formation. Topical steroid agents have anti-inflammatory effects and may cause profound and varied metabolic activities. In addition, topical steroid agents modify the body's immune response to diverse stimuli. Examples of topical steroids include, but are not limited to, triamcinolone acetonide (Artistocort, Kenalog) 0.1% cream, and betamethasone
10 dipropionate (Diprolene, Diprosone) 0.05% cream. Coal tar is an inexpensive treatment available over the counter in shampoos or lotions for use in widespread areas of involvement. Coal tar is particularly useful in hair-bearing areas. An example of coal tar is coal tar 2-10% (DHS Tar, Doctar, Theraplex T) – antipruritic. Keratolytic agents are used to remove scale, smooth the skin, and to treat hyperkeratosis. An example of a keratolytic
15 agent is anthralin 0.1-1% (Drithocrema, Anthra-Derm). Vitamin D-3 analogs are used in patients with lesions resistant to older therapy or with lesions on the face or exposed areas where thinning of the skin would pose cosmetic problems. An example of a vitamin D-3 analog is calcipotriene (Dovonex). Topical retinoids are agents that decrease the cohesiveness of follicular epithelial cells and stimulate mitotic activity, resulting in an
20 increase in turnover of follicular epithelial cells. Examples of topical retinoids include, but are not limited to, tretinoin (Retin-A, Avita), and tazarotene (Tazorac).

Approximately 1-2% of people in the United States, or about 5.5 million, have plaque psoriasis. Up to 30% of people with plaque psoriasis also have psoriatic arthritis. Individuals with psoriatic arthritis have inflammation in their joints and may have other
25 arthritis symptoms. Sometimes plaque psoriasis can evolve into more severe disease, such as pustular psoriasis or erythrodermic psoriasis. In pustular psoriasis, the red areas on the skin contain blisters with pus. In erythrodermic psoriasis, a wide area of red and scaling skin is typical, and it may be itchy and painful. The present invention is useful in treating additional types of psoriasis, including but not limited to, guttate psoriasis, nail psoriasis,
30 inverse psoriasis, and scalp psoriasis.

In some embodiments, the compounds of the present invention are useful in treating pigmentation disorders (e.g., albinism, melasma, and vitiligo). The present invention is not limited to a particular mechanism for treating pigment disorders. In preferred embodiments,

pigment disorders are treated through targeting of the F_1F_o -ATPase by the compounds of the present invention. In further embodiments, pigment disorders are treated through the rerouting of tyrosinase by the compounds of the present invention. In further embodiments, pigment disorders are treated through targeting of prohibitin by the compounds of the present invention.

VII. ATPase Inhibitors And Methods For Identifying Therapeutic Inhibitors

The present invention provides compounds that target the F_1F_o -ATPase. In addition, the present invention provides compounds that target the F_1F_o -ATPase as a treatment for autoimmune disorders, and in particular, compounds with low toxicity. The present invention further provides methods of identifying compounds that target the F_1F_o -ATPase. Additionally, the present invention provides therapeutic applications for compounds targeting the F_1F_o -ATPase.

A majority of ATP within eukaryotic cells is synthesized by the mitochondrial F_1F_o -ATPase (*see, e.g.,* C.T. Gregory *et al.*, J. Immunol., 139:313-318 [1987]; J.P. Portanova *et al.*, Mol. Immunol., 32:117-135 [1987]; M.J. Shlomchik *et al.*, Nat. Rev. Immunol., 1:147-153 [2001]; each herein incorporated by reference in their entireties). Although the F_1F_o -ATPase synthesizes and hydrolyzes ATP, during normal physiologic conditions, the F_1F_o -ATPase only synthesizes ATP (*see, e.g.,* Nagyvary J, *et al.*, Biochem. Educ. 1999; 27:193-99; herein incorporated by reference in its entirety). The mitochondrial F_1F_o -ATPase is composed of three major domains: F_o , F_1 and the peripheral stator. F_1 is the portion of the enzyme that contains the catalytic sites and it is located in the matrix (*see, e.g.,* Boyer, PD, Annu Rev Biochem. 1997; 66:717-49; herein incorporated by reference in its entirety). This domain is highly conserved and has the subunit composition $\alpha_3\beta_3\gamma\delta\epsilon$. The landmark X-ray structure of bovine F_1 revealed that $\alpha_3\beta_3$ forms a hexagonal cylinder with the γ subunit in the center of the cylinder. F_o is located within the inner mitochondrial membrane and contains a proton channel. Translocation of protons from the inner-membrane space into the matrix provides the energy to drive ATP synthesis. The peripheral stator is composed of several proteins that physically and functionally link F_o with F_1 . The stator transmits conformational changes from F_o into the catalytic domain that regulate ATP synthesis (*see, e.g.,* Cross RL, Biochim Biophys Acta 2000; 1458:270-75; herein incorporated by reference in its entirety).

Mitochondrial F_1F_0 -ATPase inhibitors are invaluable tools for mechanistic studies of the F_1F_0 -ATPase (*see, e.g.,* James AM, *et al.*, J Biomed Sci 2002; 9:475-87; herein incorporated by reference in its entirety). Because F_1F_0 -ATPase inhibitors are often cytotoxic, they have been explored as drugs for cancer and other hyperproliferative disorders. Macrolides (*e.g.,* oligomycin and apoptoludin) are non-competitive inhibitors of the F_1F_0 -ATPase (*see, e.g.,* Salomon AR, *et al.*, PNAS 2000; 97:14766-71; Salomon AR, *et al.*, Chem Biol 2001; 8:71-80; herein incorporated by reference in its entirety). Macrolides bind to F_0 which blocks proton flow through the channel resulting in inhibition of the F_1F_0 -ATPase. Macrolides are potent (*e.g.,* the IC_{50} for oligomycin = 10 nM) and lead to large decreases in [ATP]. As such, macrolides have an unacceptably narrow therapeutic index and are highly toxic (*e.g.,* the LD_{50} for oligomycin in rodents is two daily doses at 0.5 mg/kg) (*see, e.g.,* Kramar R, *et al.*, Agents & Actions 1984, 15:660-63; herein incorporated by reference in its entirety). Other inhibitors of F_1F_0 -ATPase include Bz-423, which binds to the OSCP in F_1 (as described elsewhere herein). Bz-423 has an $K_i \sim 9 \mu M$. Bz-423 is described in U.S. Patent Application Nos. 10/634,114, 10/427,211, 10/427,212, 10/217,878, 09/767,283, 09/700,101, and related applications, each herein incorporated by reference in their entireties.

In cells that are actively respiring (known as state 3 respiration), inhibiting F_1F_0 -ATPase blocks respiration and places the mitochondria in a resting state (known as state 4). In state 4, the MRC is reduced relative to state 3, which favors reduction of O_2 to O_2^- at complex III (*see, e.g.,* N. Zamzami *et al.*, J. Exp. Med., 181:1661-1672 [1995]; herein incorporated by reference in its entirety). For example, treating cells with either oligomycin leads to a rise of intracellular O_2^- as a consequence of inhibiting complex V. In the case of oligomycin, supplementing cells with ATP protects against death whereas antioxidants do not, indicating that cell death results from the drop in ATP (*see, e.g.,* Zhang JG, *et al.*, Arch Biochem Biophys 2001; 393:87-96; McConkey DJ, *et al.*, The ATP switch in apoptosis. In: Nieminen La, ed. Mitochondria in pathogenesis. New York: Plenum, 2001:265-77; each herein incorporated by reference in their entireties). Bz-423-induced cell death is blocked by antioxidants and is not affected by supplementing cells with ATP, indicating that Bz-423 engages an ROS-dependent death response (*see, e.g.,* N.B. Blatt, *et al.*, J. Clin. Invest., 2002, 110, 1123; herein incorporated by reference in its entirety). As such, F_1F_0 -ATPase inhibitors are either toxic (*e.g.,* oligomycin) or therapeutic (*e.g.,* Bz-423).

The present invention provides a method of distinguishing toxic F_1F_o -ATPase inhibitors from therapeutic F_1F_o -ATPase inhibitors. F_1F_o -ATPase inhibitors with therapeutic potential present a novel mode of inhibition. Specifically, F_1F_o -ATPase inhibitors with beneficial properties are uncompetitive inhibitors that only bind enzyme-substrate complexes at high substrate concentration and do not alter the k_{cat}/K_m ratio. This knowledge forms the basis to identify and distinguish F_1F_o -ATPase inhibitors with therapeutic potential from toxic compounds.

The present invention provides compounds that target the F_1F_o -ATPase as an autoimmune disorder treatment. In particular, the present invention provides methods of identifying compounds that target the F_1F_o -ATPase while not altering the k_{cat}/K_m ratio. Additionally, the present invention provides therapeutic applications for compounds targeting the F_1F_o -ATPase.

A. ATPase Inhibiting Compounds

The present invention provides compounds that inhibit the F_1F_o -ATPase. In some embodiments, the compounds do not bind free F_1F_o -ATPase, but rather bind to an F_1F_o -ATPase-substrate complex. The compounds show maximum activity at high substrate concentration and minimal activity (*e.g.*, F_1F_o -ATPase inhibiting) at low substrate concentration. In preferred embodiments, the compounds do not alter the k_{cat}/K_m ratio of the F_1F_o -ATPase. The properties of the F_1F_o -ATPase inhibitors of the present invention are in contrast with oligomycin, which is a F_1F_o -ATPase inhibitor that is acutely toxic and lethal. Oligomycin is a noncompetitive inhibitor, which binds to both free F_1F_o -ATPase and F_1F_o -ATPase-substrate complexes and alters the k_{cat}/K_m ratio.

The compounds of the present invention that inhibit F_1F_o -ATPase while not altering the k_{cat}/K_m ratio, in some embodiments, have the structure described elsewhere herein. However, compounds of other structures that are identified as therapeutic inhibitors by the methods of the present invention are also encompassed by the present invention.

B. Identifying ATPase Inhibitors

The present invention provides methods of identifying (*e.g.*, screening) compounds useful in treating autoimmune disorders. The present invention is not limited to a particular type compound. In preferred embodiments, compounds of the present invention include, but are not limited to, pharmaceutical compositions, small molecules, antibodies, large molecules,

synthetic molecules, synthetic polypeptides, synthetic polynucleotides, synthetic nucleic acids, aptamers, polypeptides, nucleic acids, and polynucleotides. The present invention is not limited to a particular method of identifying compounds useful in treating autoimmune disorders. In preferred embodiments, compounds useful in treating autoimmune disorders are
5 identified as possessing an ability to inhibit an F_1F_o -ATPase while not altering the k_{cat}/K_m ratio.

C. Therapeutic Applications With F_1F_o -ATPase Inhibitors

The present invention provides methods for treating disorders (*e.g.*, neurodegenerative diseases, Alzheimers, ischemia reprofusion injury, neuromotor disorders, non-Hodgkin's
10 lymphoma, lymphocytic leukemia, cutaneous T cell leukemia, an autoimmune disorder, cancer, solid tumors, lymphomas, leukemias, and tuberculosis). The present invention is not limited to a particular form of treatment. In preferred embodiments, treatment includes, but is not limited to, symptom amelioration, symptom prevention, disorder prevention, and disorder amelioration. The present invention provides methods of treating autoimmune disorders
15 applicable within *in vivo*, *in vitro*, and/or *ex vivo* settings.

In some embodiments, the present invention treats autoimmune disorders through inhibiting of target cells. The present invention is not limited to a particular form of cell inhibition. In preferred embodiments, cell inhibition includes, but is not limited to, cell growth prevention, cell proliferation prevention, and cell death. In preferred embodiments, inhibition
20 of a target cell is accomplished through contacting a target cell with an F_1F_o -ATPase inhibitor of the present invention. In further embodiments, target cell inhibition is accomplished through targeting of the F_1F_o -ATPase with an F_1F_o -ATPase inhibitor of the present invention. The present invention is not limited to a particular F_1F_o -ATPase inhibitor. In preferred embodiments, the F_1F_o -ATPase inhibitor possesses the ability to inhibit an F_1F_o -ATPase while
25 not altering the k_{cat}/K_m ratio.

The present invention further provides methods for selectively inhibiting the pathology of target cells in a subject in need of therapy. The present invention is not limited to a particular method of inhibition target cell pathology. In preferred embodiments, target cell pathology is inhibited through administration of an effective amount of a compound of
30 the invention. The present invention is not limited to a particular compound. In preferred embodiments, the compound is an F_1F_o -ATPase inhibitor. In further preferred embodiments, the compound inhibits the F_1F_o -ATPase while not altering the k_{cat}/K_m ratio.

EXAMPLES

The following examples are provided to demonstrate and further illustrate certain preferred embodiments of the present invention and are not to be construed as limiting the scope thereof. Examples illustrating the various uses and applications of benzodiazepine compounds and benzodiazepine related compounds are described, for example, in U.S. Provisional Patent Nos. 60/131,761, 60/165,511, 60/191,855, 60/312,560, 60/313,689, 60/396,670, 60/565,788, 60/607,599, 60/641,040, and U.S. Patent Application Nos. 11/324,419, 11/176,719, 11/110,228, 10/935,333, 10/886,450, 10/795,535, 10/634,114, 10/427,211, 10/427,212, 10/217,878, 09/767,283, 09/700,101, and related applications; each herein incorporated by reference in their entireties.

Example 1.

This example describes the formation of benzodiazepine crystal forms and formulations. Bz-423 (15.8 mg) was added to a 1 mL vial and dissolved in polyethylene glycol dimethyl ether 500 (0.01mL) at 88 °C. The vial was kept at 88 °C for 16 hours, during which time crystals formed. The vial was then cooled to room temperature and washed with cold ethyl ether (5 x 2 mL) to yield 4.2 mg of anhydrous Bz-423 crystals, a 27 % yield. Figure 1 shows the structural data of anhydrous Bz-423, Figure 2 shows powder x-ray diffraction data for anhydrous Bz-423, and Figure 3 shows Raman spectroscopy data for anhydrous Bz-423.

Bz-423 ethanol solvate was crystallized by evaporation of ethanol at room temperature. Figure 4 shows the structural data of Bz-423 ethanol solvate, Figure 5 shows powder x-ray diffraction data for Bz-423 ethanol solvate, and Figure 6 shows Raman spectroscopy data for Bz-423 ethanol solvate.

Ball milled Bz-423 succinic acid (2:1) was crystallized from tetraethylene glycol dimethyl ether at 88 °C. Figure 7 shows Raman spectroscopy data for ball milled Bz-423 succinic acid (2:1).

Ball milled Bz-423 citric acid (2:1) was generated through ball milling a 2:1 mixture of Bz-423 and citric acid. Figure 8 shows Raman spectroscopy ball milled Bz-423 citric acid (2:1).

Bz-423 biphenyl derivate was crystallized from methanol at room temperature. Figure 9 shows the structural data of Bz-423 biphenyl derivate.

BZ-423-acetic acid was crystallized by evaporation of acetic acid at room temperature. BZ-423-CH₃CN was crystallized by evaporation of CH₃CN at room temperature. BZ-423-methanol was crystallized by evaporation of methanol at room temperature. BZ-423-ethyl acetate was crystallized by evaporation of ethyl acetate at room temperature. BZ-423-toluene was crystallized by evaporation of toluene at room temperature. BZ-423-oxalic acid was crystallized from tetraethylene glycol dimethyl ether at 88 °C. BZ-423-fumaric acid was crystallized from tetraethylene glycol dimethyl ether at 88 °C. BZ-423-octanol was crystallized from a supersaturated solution of octanol. BZ-423-heptanoic acid was crystallized from a supersaturated solution of heptanoic acid. BZ-423-diphenyl ether was crystallized from a supersaturated solution of diphenyl ether. BZ-423-trichlorobenzene was crystallized from a supersaturated solution of trichlorobenzene.

Example 2.

This example demonstrates that anhydrous Bz-423 is more soluble than solvated Bz-423. Water solubility at 37°C analyses were conducted for solvated Bz-423, anhydrous Bz-423, Bz-423 acetic acid, and Bz-423 citric acid. Table 1 shows the solubility results for solvated Bz-423, anhydrous Bz-423, Bz-423 acetic acid, Bz-423 citric acid, and Bz-423 micronized.

20	Table 1.	
	as supplied BZ-423 ball milled	1.00
	anhydrous BZ-423 ball milled	1.4
	BZ-423 acetic acid	1.4
	BZ-423 citric acid	1.4
25	Bz-423 micronized	0.85

Figure 10 shows solubility (e.g., absorbance) as a function of time for solvated Bz-423, anhydrous Bz-423, Bz-423 acetic acid, and Bz-423 citric acid.

30 Example 3.

This example demonstrates that unsolvated Bz-423 is capable of inhibiting ATP hydrolysis, not inhibiting cell synthesis, not affecting cell viability, and its activity is related to binding of the OSCP, along or with other components of the mitochondrial F₀F₁ ATPase (e.g., F₁ subunit). In particular, Figure 11 shows a comparison of ATP hydrolysis between unsolvated Bz-423 and solvated Bz-423, Figure 12 shows a comparison of ATP synthesis

between unsolvated Bz-423 and solvated Bz-423, and Figure 13 shows a comparison of cell viability between unsolvated Bz-423 and solvated Bz-423.

Example 4.

5 This example demonstrates that unsolvated Bz-423 is more soluble in simulated gastric fluid than solvated Bz-423. Simulated gastric fluid solubility at 37°C analyses were conducted for solvated Bz-423, and anhydrous Bz-423. Anhydrous Bz-423 was more soluble than solvated Bz-423. Figure 14 shows a UV-vis spectrum of Bz-423 in simulated gastric fluid. Figure 15 shows a UV-vis spectrum of Bz-423 in simulated gastric fluid
10 before and after addition of K₂CO₃.

Example 5.

 This example describes the formation of benzodiazepine formulations. Figure 16 shows Raman spectroscopy data for Bz-423 ethanol solvate. BZ-423 (12.3 mg) was
15 dissolved completely in 0.5 mL of ethanol at 70 °C. Upon cooling to room temperature crystallization of the solvate occurred. The Raman spectrum above was obtained from the crystals. The characteristic Raman shifts for the ethanol solvate occur at about 1673 and 1560 cm⁻¹.

 Figure 17 shows Raman spectroscopy data for Bz-423 1-propanol solvate. BZ-423
20 (46.6 mg) was dissolved completely in 0.5 mL of 1-propanol at 70 °C. Upon cooling to room temperature crystallization of the solvate occurred. The Raman spectrum above was obtained from the crystals. The characteristic Raman shifts for the 1-propanol solvate occur at about 1671 and 1561 cm⁻¹.

 Figure 18 shows Raman spectroscopy data for Bz-423 2-propanol solvate. BZ-423
25 (53.2 mg) was dissolved completely in 0.5 mL of 2-propanol at 70 °C. Upon cooling to room temperature crystallization of the solvate occurred. The Raman spectrum above was obtained from the crystals. The characteristic Raman shifts for the 2-propanol solvate occur at about 1667 and 1562 cm⁻¹.

 Figure 19 shows Raman spectroscopy data for Bz-423 1-butanol solvate. BZ-423
30 (105.8 mg) was dissolved completely in 0.5 mL of 1-butanol at 70 °C. Upon cooling to room temperature crystallization of the solvate occurred. The Raman spectrum above was obtained from the crystals. The characteristic Raman shifts for the 1-butanol solvate occur at about 1661 and 1556 cm⁻¹.

Figure 20 shows Raman spectroscopy data for Bz-423 2-butanol solvate. BZ-423 (97.3 mg) was dissolved completely in 0.5 mL of 2-butanol at 70 °C. Upon cooling to room temperature crystallization of the solvate occurred. The Raman spectrum above was obtained from the crystals. The characteristic Raman shifts for the 2-butanol solvate occur at about 1666 and 1562 cm^{-1} .

Figure 21 shows Raman spectroscopy data for Bz-423 1-pentanol solvate. BZ-423 (99.3 mg) was dissolved completely in 0.5 mL of 1-pentanol at 70 °C. Upon cooling to room temperature crystallization of the solvate occurred. The Raman spectrum above was obtained from the crystals. The characteristic Raman shifts for the 1-pentanol solvate occur at about 1646 and 1563 cm^{-1} .

Figure 22 shows Raman spectroscopy data for Bz-423 1-octanol solvate. BZ-423 (5.2 mg) was dissolved completely in 0.5 mL of 1-octanol at 70 °C. Upon cooling to room temperature crystallization of the solvate occurred. The Raman spectrum above was obtained from the crystals. The characteristic Raman shifts for the 1-octanol solvate occur at about 1669 and 1559 cm^{-1} .

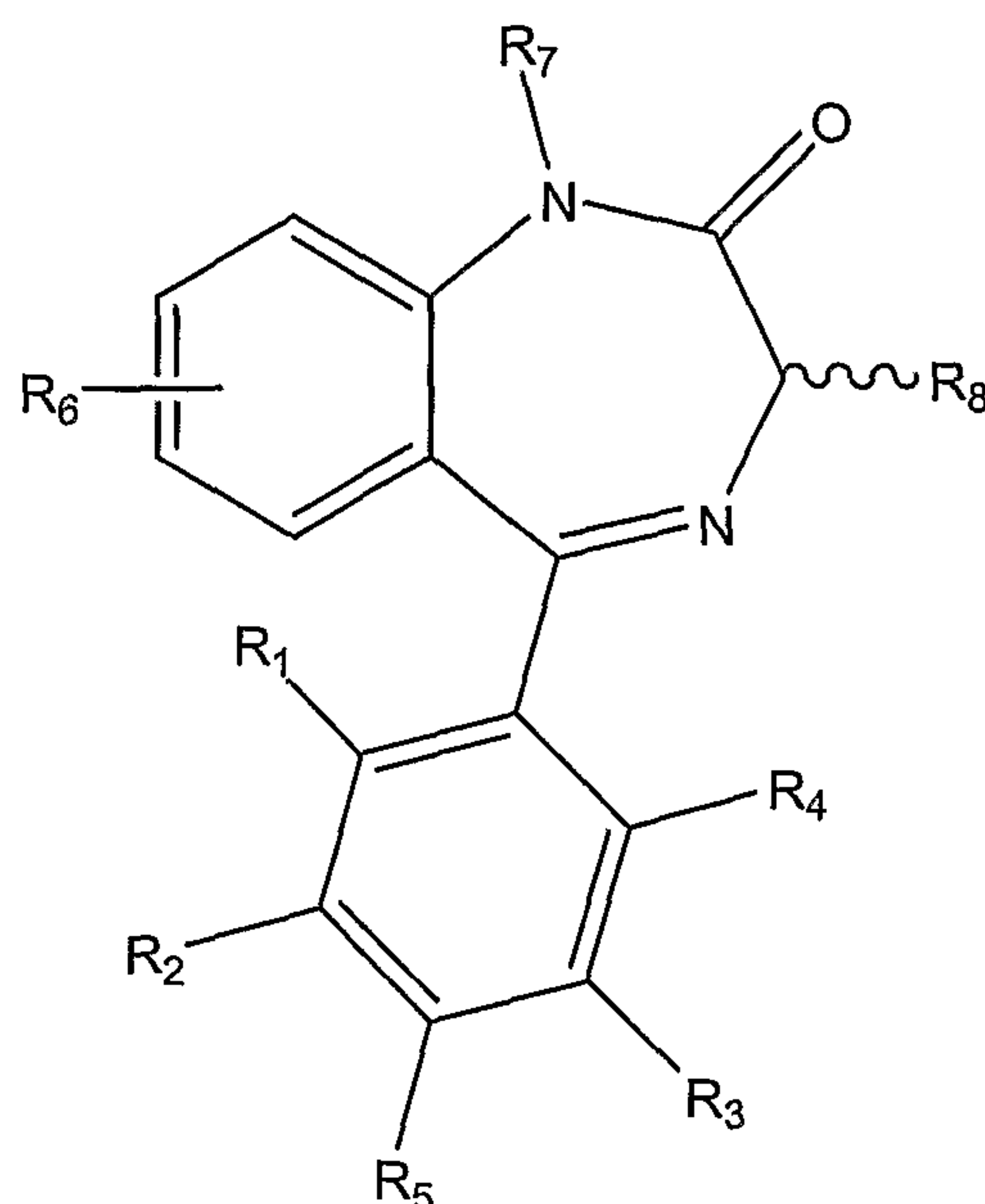
Figure 23 shows Raman spectroscopy data for Bz-423 propylene glycol solvate. BZ-423 (16.8 mg) was dissolved completely in 0.5 mL of propylene glycol at 70 °C. Upon cooling to room temperature crystallization of the solvate occurred. The Raman spectrum above was obtained from the crystals. The characteristic Raman shifts for the propylene glycol solvate occur at about 1660 and 1556 cm^{-1} .

Figure 24 shows Raman spectroscopy data for Bz-423 acetone glass. BZ-423 (14.3 mg) was dissolved completely in 0.5 mL of acetone at room temperature. Upon evaporation a glass was formed. The Raman spectrum above was obtained from the glass. The characteristic Raman shifts for the acetone glass occur at about 1674 and 1559 cm^{-1} .

All publications and patents mentioned in the above specification are herein incorporated by reference. Although the invention has been described in connection with specific preferred embodiments, it should be understood that the invention as claimed should not be unduly limited to such specific embodiments. Indeed, various modifications of the described modes for carrying out the invention that are obvious to those skilled in the relevant fields are intended to be within the scope of the following claims.

We claim:

1. A composition comprising an unsolvated compound having the structure described
 5 by the following formula:



including both R and S enantiomeric forms and racemic mixtures;

wherein R1, R2, R3 and R4 are selected from the group consisting of:

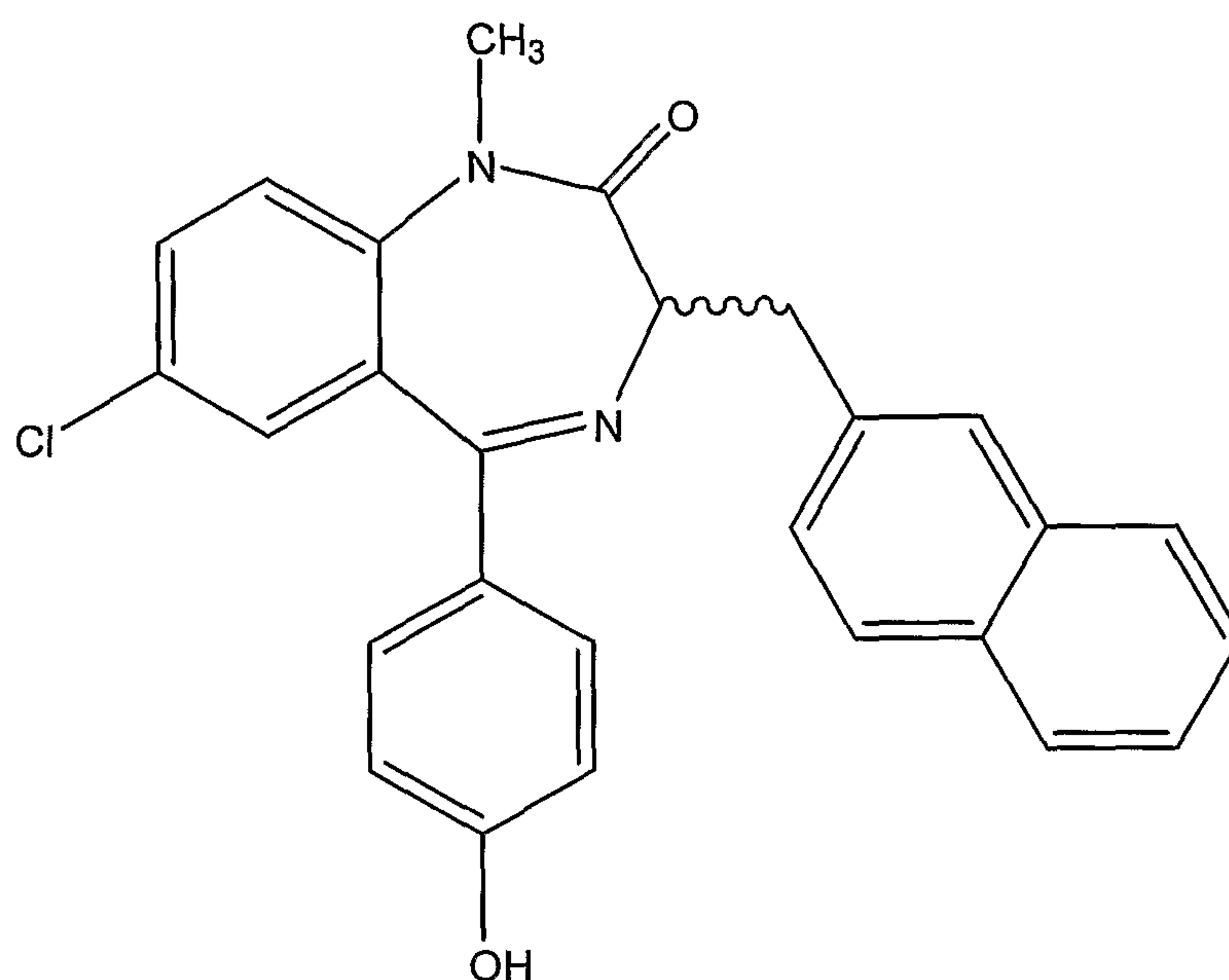
- hydrogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain
 10 having at least 1 carbon; a linear or branched, saturated or unsaturated
 aliphatic chain having at least 2 carbons, and having at least one hydroxy
 subgroup; a linear or branched, saturated or unsaturated aliphatic chain
 having at least 2 carbons, and having at least one thiol subgroup; a linear or
 branched, saturated or unsaturated aliphatic chain having at least 2 carbons,
 15 wherein said aliphatic chain terminates with an aldehyde subgroup; a linear
 or branched, saturated or unsaturated aliphatic chain having at least 2
 carbons, and having at least one ketone subgroup; a linear or branched,
 saturated or unsaturated aliphatic chain having at least 2 carbons; wherein
 said aliphatic chain terminates with a carboxylic acid subgroup; a linear or
 20 branched, saturated or unsaturated aliphatic chain having at least 2 carbons,
 and having at least one amide subgroup; a linear or branched, saturated or
 unsaturated aliphatic chain having at least 2 carbons, and having at least one
 acyl group; a linear or branched, saturated or unsaturated aliphatic chain

having at least 2 carbons, and having at least one nitrogen containing moiety;
a linear or branched, saturated or unsaturated aliphatic chain having at least 2
carbons, and having at least one amine subgroup; a linear or branched,
saturated or unsaturated aliphatic chain having at least 2 carbons, and having
at least one ether subgroup; a linear or branched, saturated or unsaturated
aliphatic chain having at least 2 carbons, and having at least one halogen
subgroup; a linear or branched, saturated or unsaturated aliphatic chain
having at least 2 carbons, and having at least one nitronium subgroup;
wherein R5 is selected from the group consisting of: OH; NO₂; OR';
wherein R' is selected from the group consisting of: a linear or branched,
saturated or unsaturated aliphatic chain having at least one carbon; a linear or
branched, saturated or unsaturated aliphatic chain having at least 2 carbons,
and having at least one hydroxyl subgroup; a linear or branched, saturated or
unsaturated aliphatic chain having at least 2 carbons, and having at least one
thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain
having at least 2 carbons, wherein said aliphatic chain terminates with an
aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic
chain having at least 2 carbons, and having at least one ketone subgroup; a
linear or branched, saturated or unsaturated aliphatic chain having at least 2
carbons; wherein said aliphatic chain terminates with a carboxylic acid
subgroup; a linear or branched, saturated or unsaturated aliphatic chain
having at least 2 carbons, and having at least one amide subgroup; a linear or
branched, saturated or unsaturated aliphatic chain having at least 2 carbons,
and having at least one acyl group; a linear or branched, saturated or
unsaturated aliphatic chain having at least 2 carbons, and having at least one
nitrogen containing moiety; a linear or branched, saturated or unsaturated
aliphatic chain having at least 2 carbons, and having at least one amine
subgroup; a linear or branched, saturated or unsaturated aliphatic chain
having at least 2 carbons, and having at least one halogen subgroup; a linear
or branched, saturated or unsaturated aliphatic chain having at least 2
carbons, and having at least one nitronium subgroup; wherein R6 is selected
from the group consisting of: Hydrogen; NO₂; Cl; F; Br; I; SR'; and NR'₂;
wherein R' is defined as above in R5;

wherein R7 is selected from the group consisting of: Hydrogen; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; and

wherein R8 is an aliphatic cyclic group larger than benzene; wherein said larger than benzene comprises any chemical group containing 7 or more non-hydrogen atoms.

2. The composition of Claim 1, wherein said compound is:

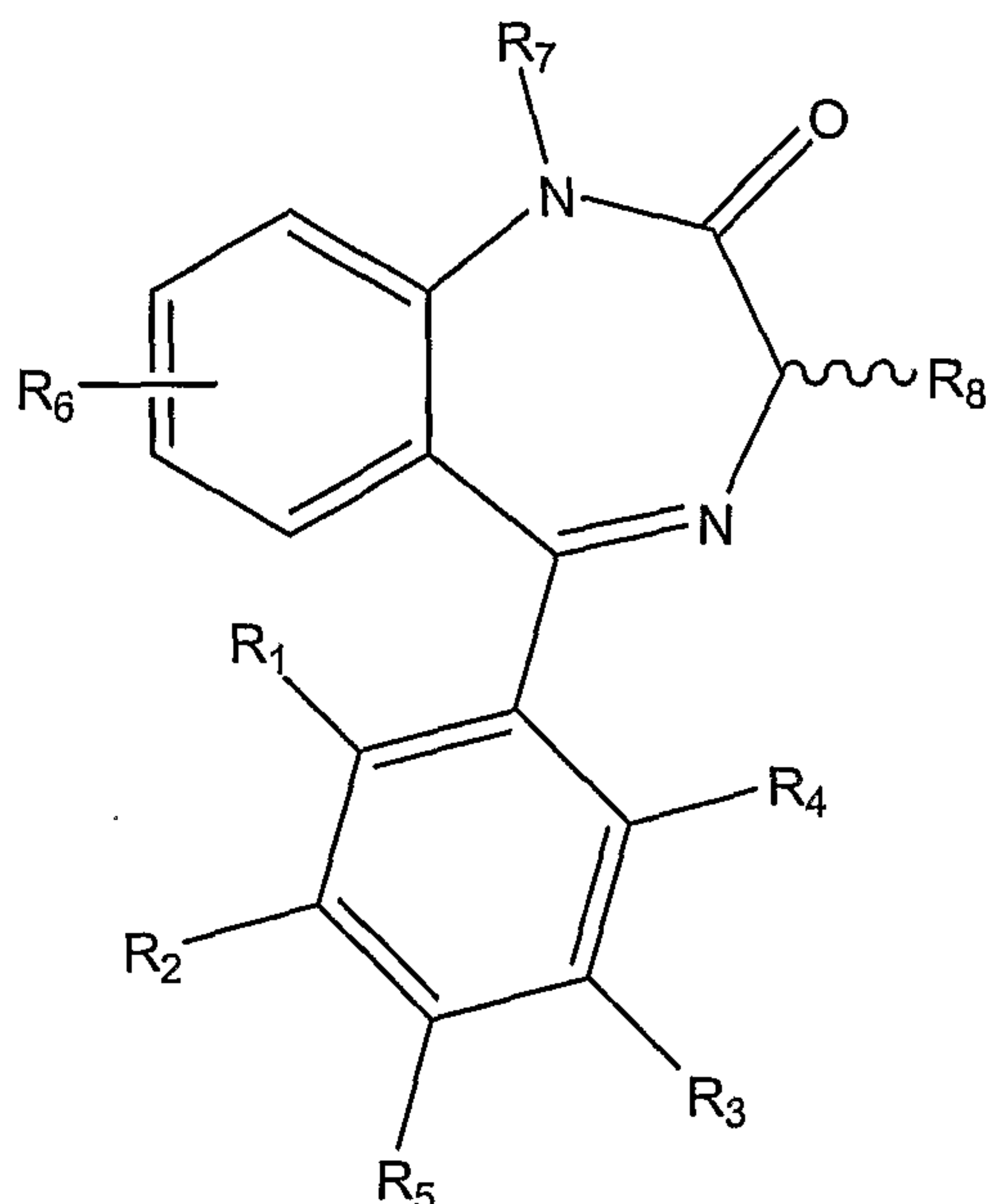


3. The composition of Claim 1, wherein said unsolvated compound is anhydrous.

4. The composition of Claim 1, wherein said unsolvated compound has an orthorhombic crystal structure.

5. A composition comprising a compound selected from the group consisting of Bz-423 ethanol solvate, Bz-423 succinic acid, Bz-423 citric acid, BZ-423-acetic acid, BZ-423-CH₃CN, BZ-423-methanol, BZ-423-ethyl acetate, BZ-423-toluene, BZ-423-oxalic acid, BZ-423-fumaric acid, BZ-423-octanol, BZ-423-heptanoic acid, BZ-423-diphenyl ether, Bz-423 1-propanol solvate, Bz-423 2-propanol solvate, Bz-423 1-butanol solvate, Bz-423 2-butanol solvate, Bz-423 1-pentanol solvate, Bz-423 propylene glycol, Bz-423 1-octanol solvate, Bz-423 acetone glass, and BZ-423-trichlorobenzene.

6. A composition comprising an orthorhombic benzodiazepine crystal, said benzodiazepine having the structure:



including both R and S enantiomeric forms and racemic mixtures;

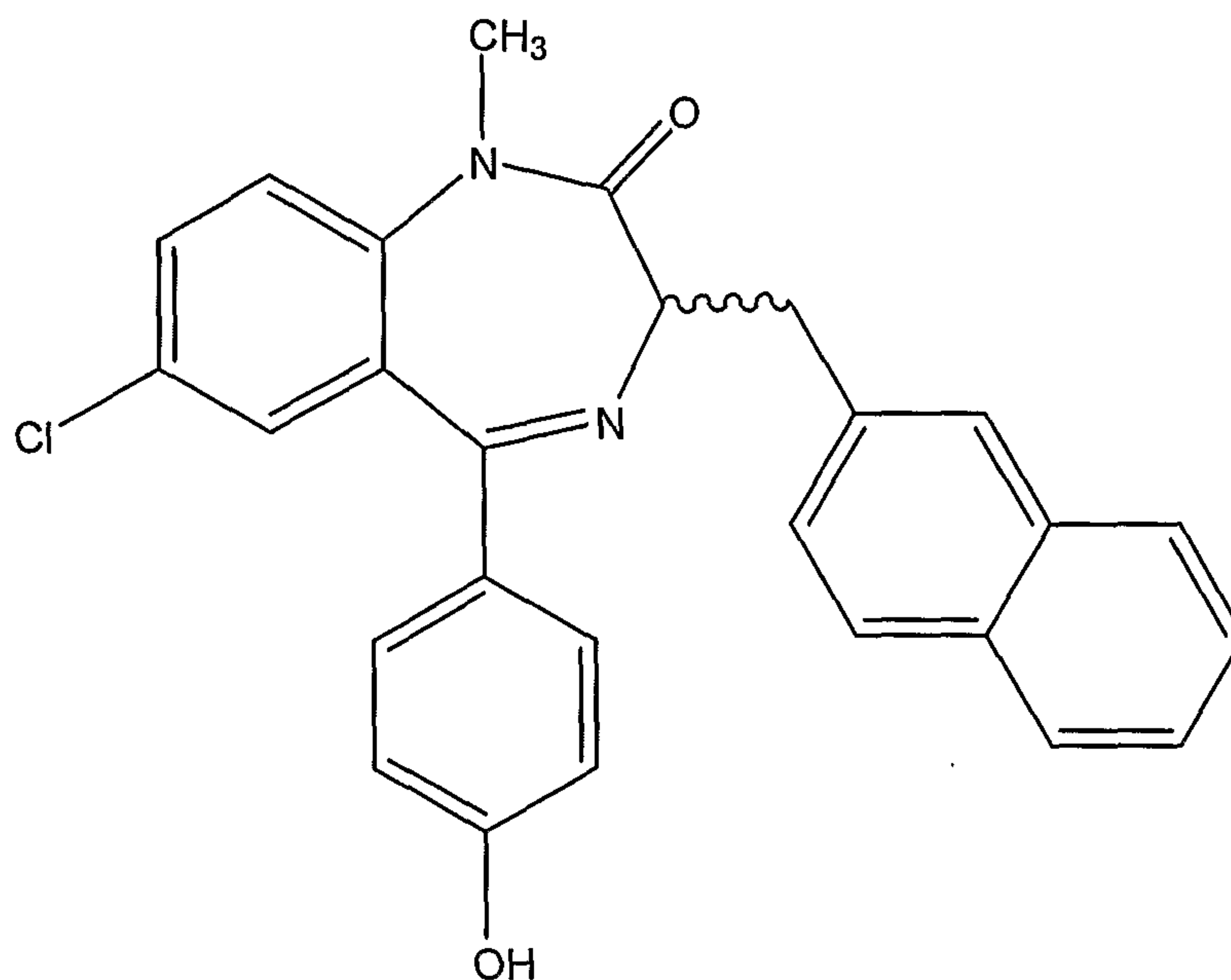
5 wherein R1, R2, R3 and R4 are selected from the group consisting of:

hydrogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 1 carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxy subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein said aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein said aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched,

saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ether subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; wherein R₅ is selected from the group consisting of: OH; NO₂; OR'; wherein R' is selected from the group consisting of: a linear or branched, saturated or unsaturated aliphatic chain having at least one carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxyl subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein said aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein said aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; wherein R₆ is selected from the group consisting of: Hydrogen; NO₂; Cl; F; Br; I; SR'; and NR'₂; wherein R' is defined as above in R₅; wherein R₇ is selected from the group consisting of: Hydrogen; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; and

wherein R8 is an aliphatic cyclic group larger than benzene; wherein
said larger than benzene comprises any chemical group containing 7 or more
non-hydrogen atoms.

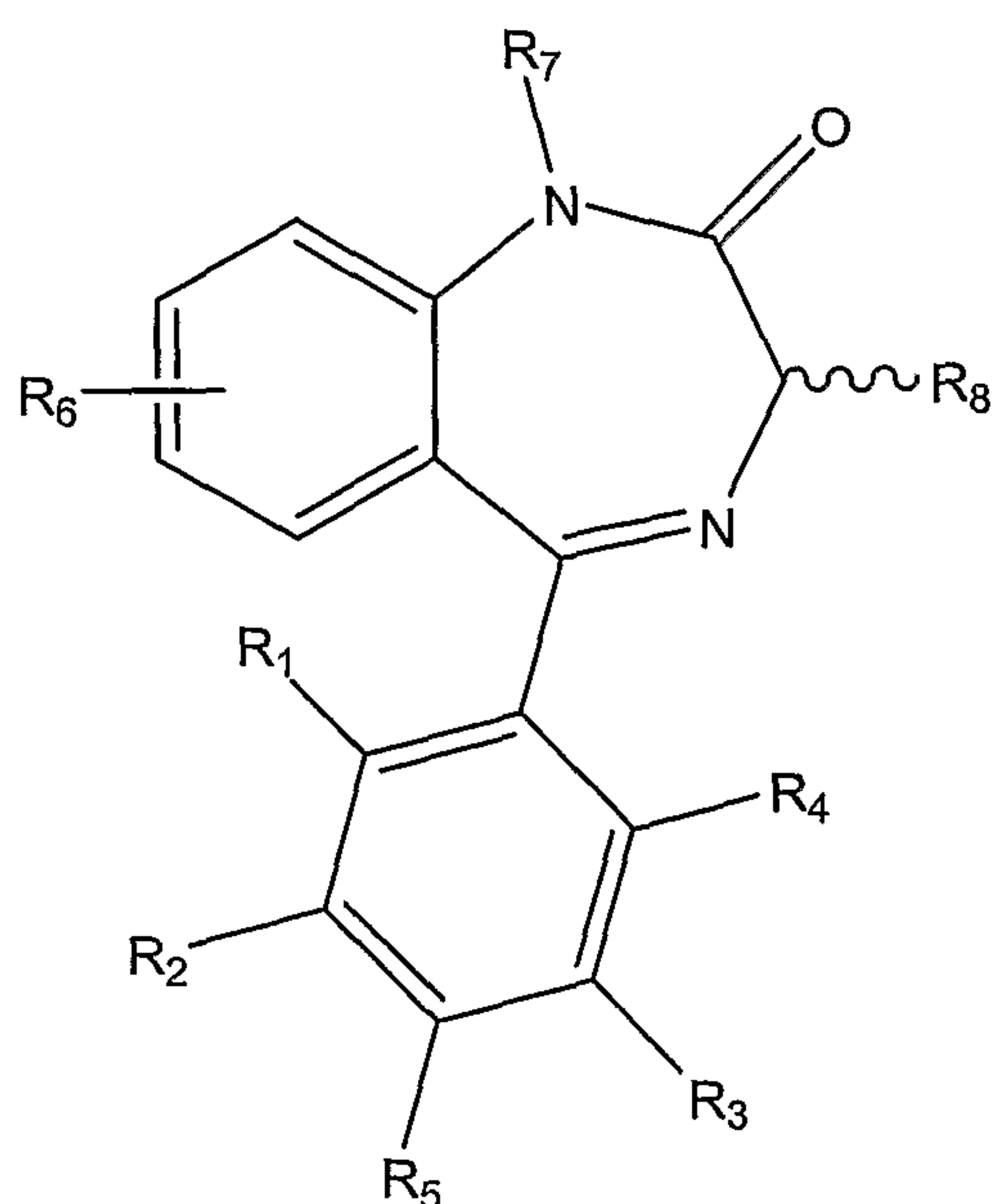
- 5 7. The composition of Claim 6, wherein said compound is:



8. The composition of Claim 6, wherein said orthorhombic benzodiazepine crystal is
anhydrous.

10

9. A composition comprising an oral dose of a benzodiazepine having the structure:



including both R and S enantiomeric forms and racemic mixtures;

wherein R1, R2, R3 and R4 are selected from the group consisting of:

hydrogen; CH₃; a linear or branched, saturated or unsaturated aliphatic chain having at least 1 carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxy subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein said aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein said aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one ether subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup;

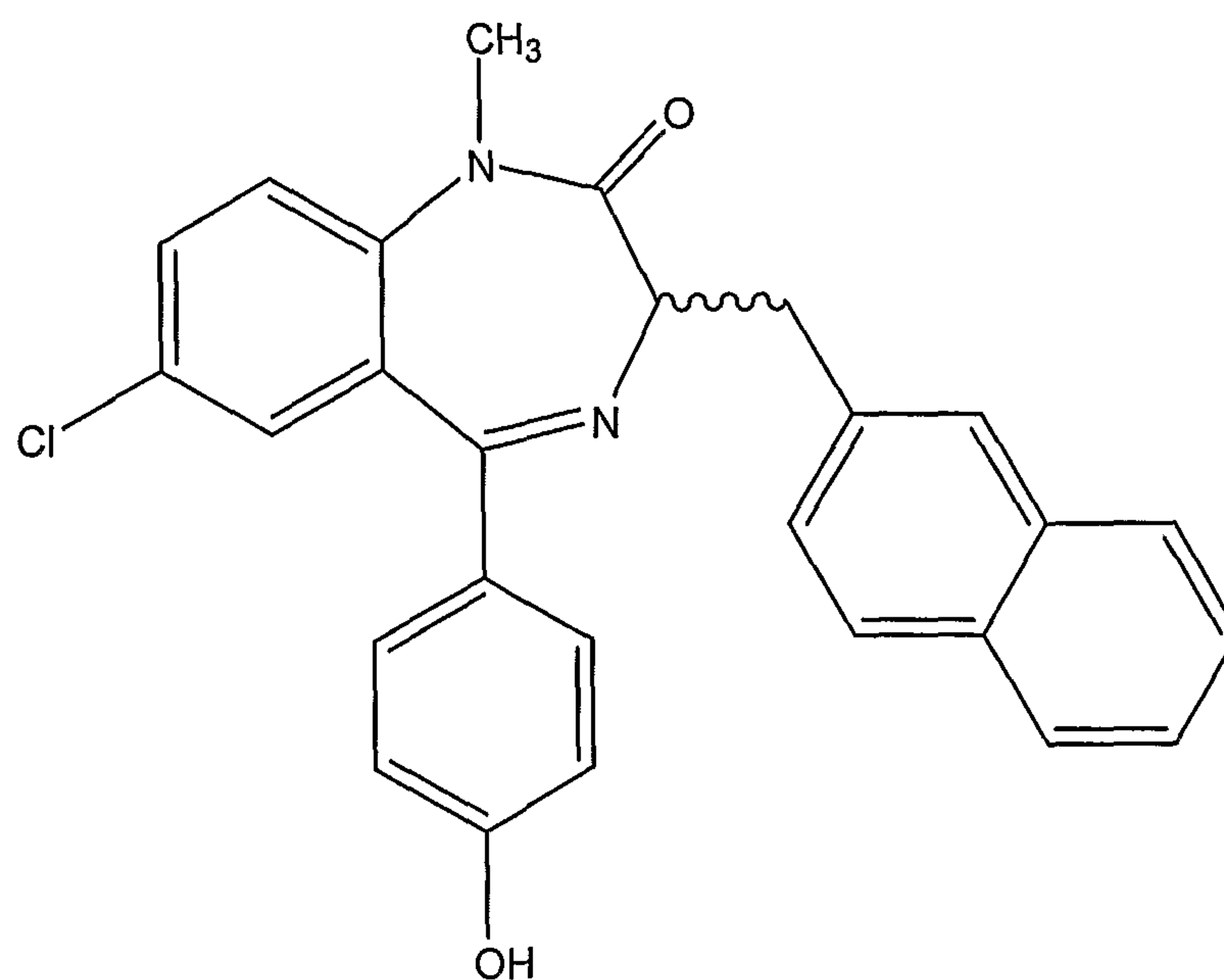
wherein R5 is selected from the group consisting of: OH; NO₂; OR';

wherein R' is selected from the group consisting of: a linear or branched,

saturated or unsaturated aliphatic chain having at least one carbon; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one hydroxyl subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one thiol subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, wherein said aliphatic chain terminates with an aldehyde subgroup; a linear or branched, saturated or unsaturated aliphatic

chain having at least 2 carbons, and having at least one ketone subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; wherein said aliphatic chain terminates with a carboxylic acid subgroup; a linear or branched, saturated or unsaturated aliphatic chain
5 having at least 2 carbons, and having at least one amide subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one acyl group; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitrogen containing moiety; a linear or branched, saturated or unsaturated
10 aliphatic chain having at least 2 carbons, and having at least one amine subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one halogen subgroup; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons, and having at least one nitronium subgroup; wherein R6 is selected from the group consisting of: Hyrdrogen; NO₂; Cl; F; Br; I; SR'; and NR'₂; wherein R' is defined as above in R5;
15 wherein R7 is selected from the group consisting of: Hydrogen; a linear or branched, saturated or unsaturated aliphatic chain having at least 2 carbons; and
20 wherein R8 is an aliphatic cyclic group larger than benzene; wherein said larger than benzene comprises any chemical group containing 7 or more non-hydrogen atoms.

10. The composition of Claim 9, wherein said compound is:



11. The composition of Claim 9, wherein said benzodiazepine compound is anhydrous.
- 5 12. The composition of Claim 9, wherein said benzodiazepine compound has an orthorhombic crystal structure.
13. A method of treating an autoimmune disorder or hyperproliferative disorder comprising administering to a subject an effective amount of a composition comprising the
- 10 composition of Claim 1.

Figure 1. Anhydrous BZ-423

	BZ-423 Anhydrous
crystal system	orthorhombic
space group	<i>Pbca</i>
<i>a</i> (Å)	14.835(3)
<i>b</i> (Å)	14.999(3)
<i>c</i> (Å)	19.901(4)
α (°)	90.00
β (°)	90.00
γ (°)	90.00
<i>Z</i>	8
cell volume	4428.4(14)
Density	1.323
<i>R</i>	0.0378

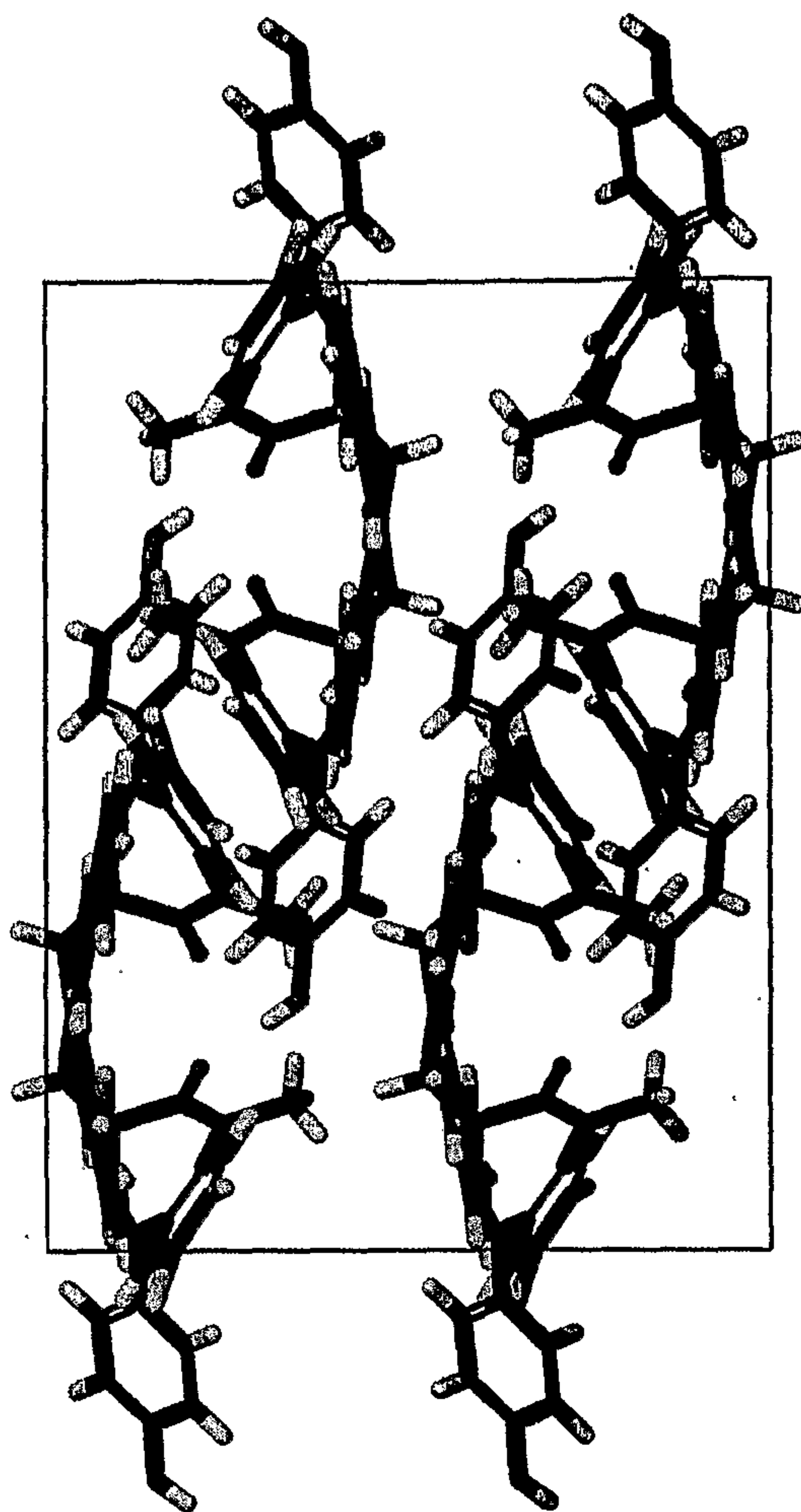


Figure 2. Powder X-ray Diffraction For Anhydrous Bz-423

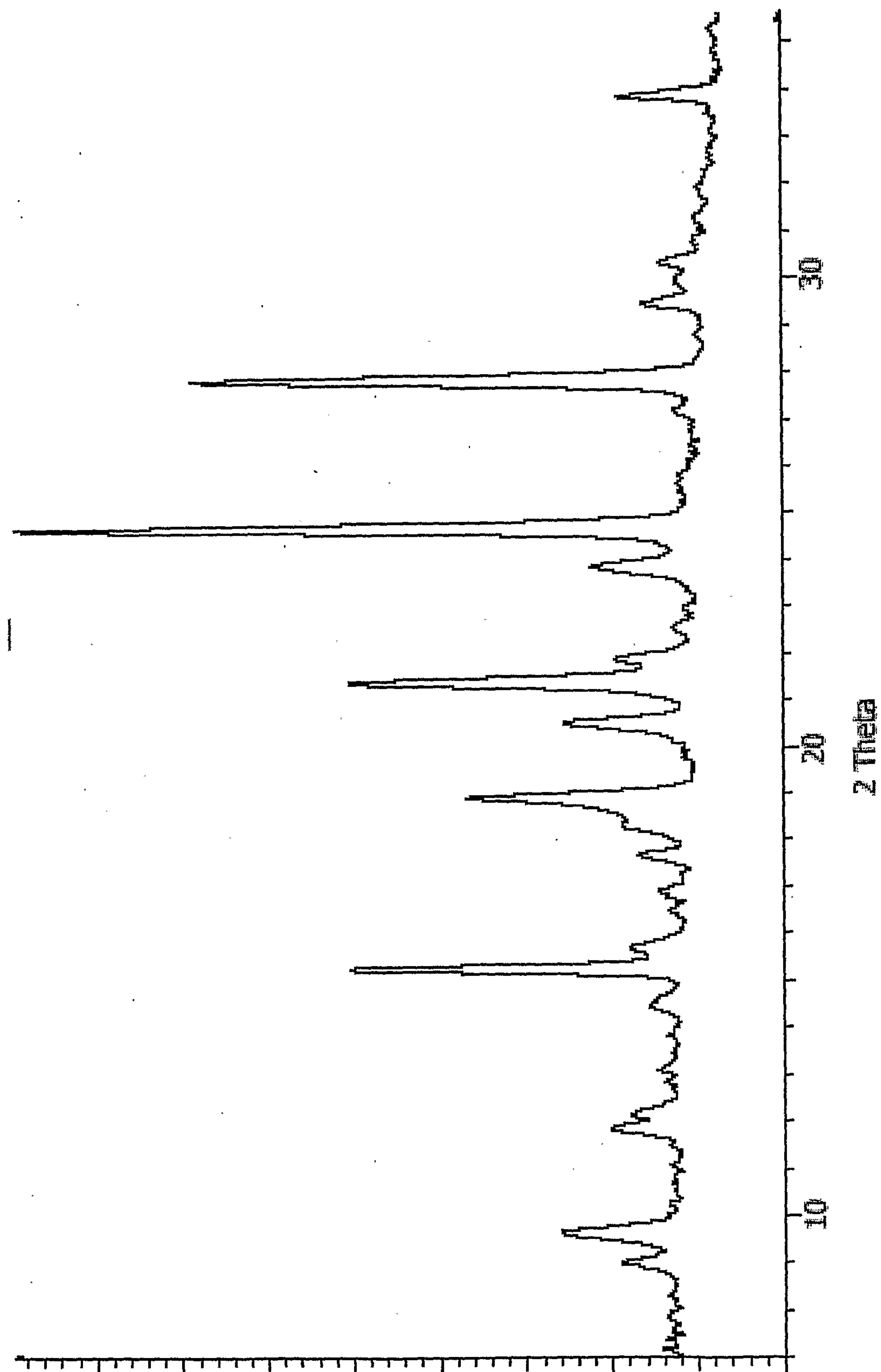


Figure 3. Raman Spectroscopy For Anhydrous Bz-423

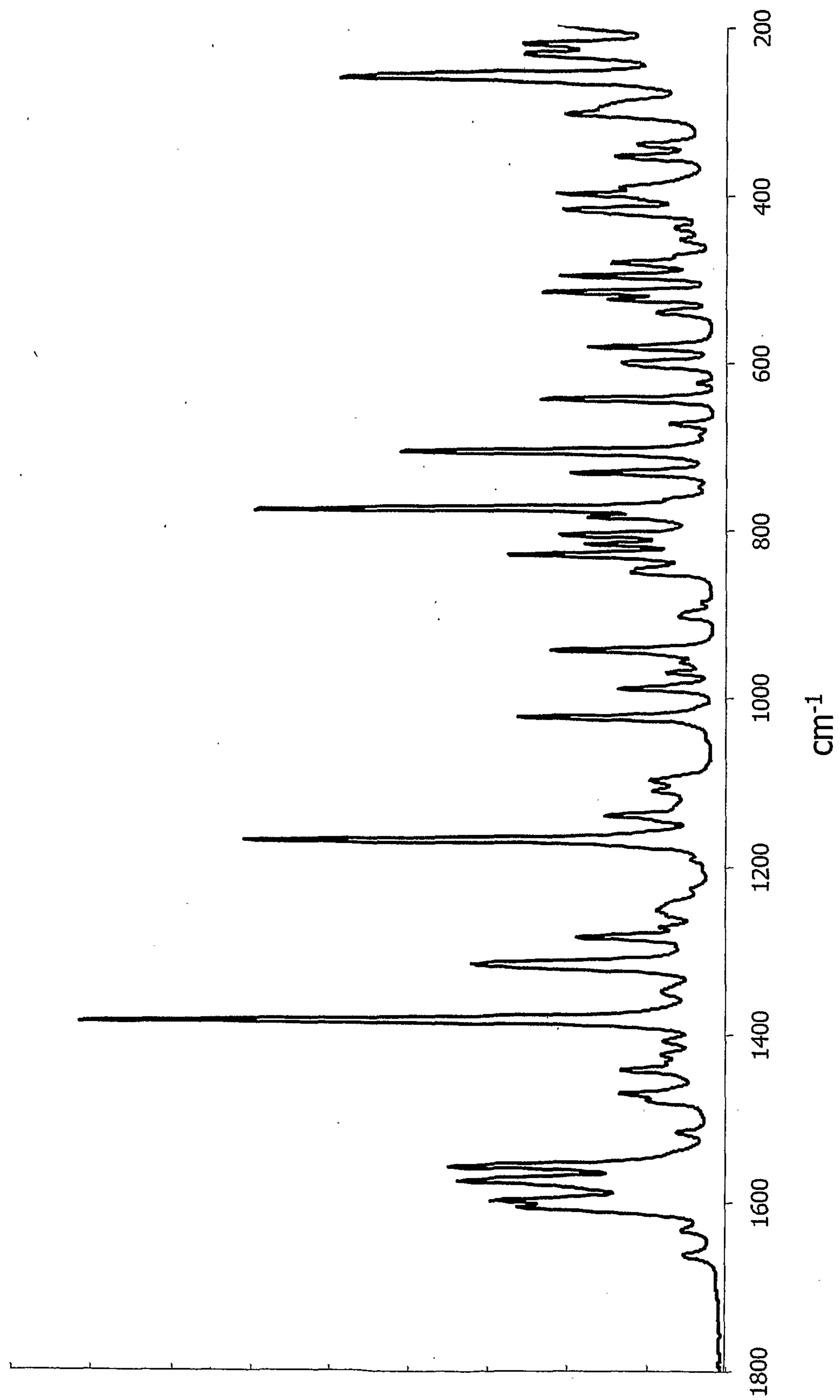


Figure 4. BZ-423 Ethanol Solvate

	BZ-423 ethanol solvate
crystal system	orthorhombic
space group	<i>Pbca</i>
<i>a</i> (Å)	14.7325(11)
<i>b</i> (Å)	15.3206(11)
<i>c</i> (Å)	22.1937(17)
α (°)	90.00
β (°)	90.00
γ (°)	90.00
<i>Z</i>	8
cell volume	5010.3(6)
Density	1.291
<i>R</i>	0.0901

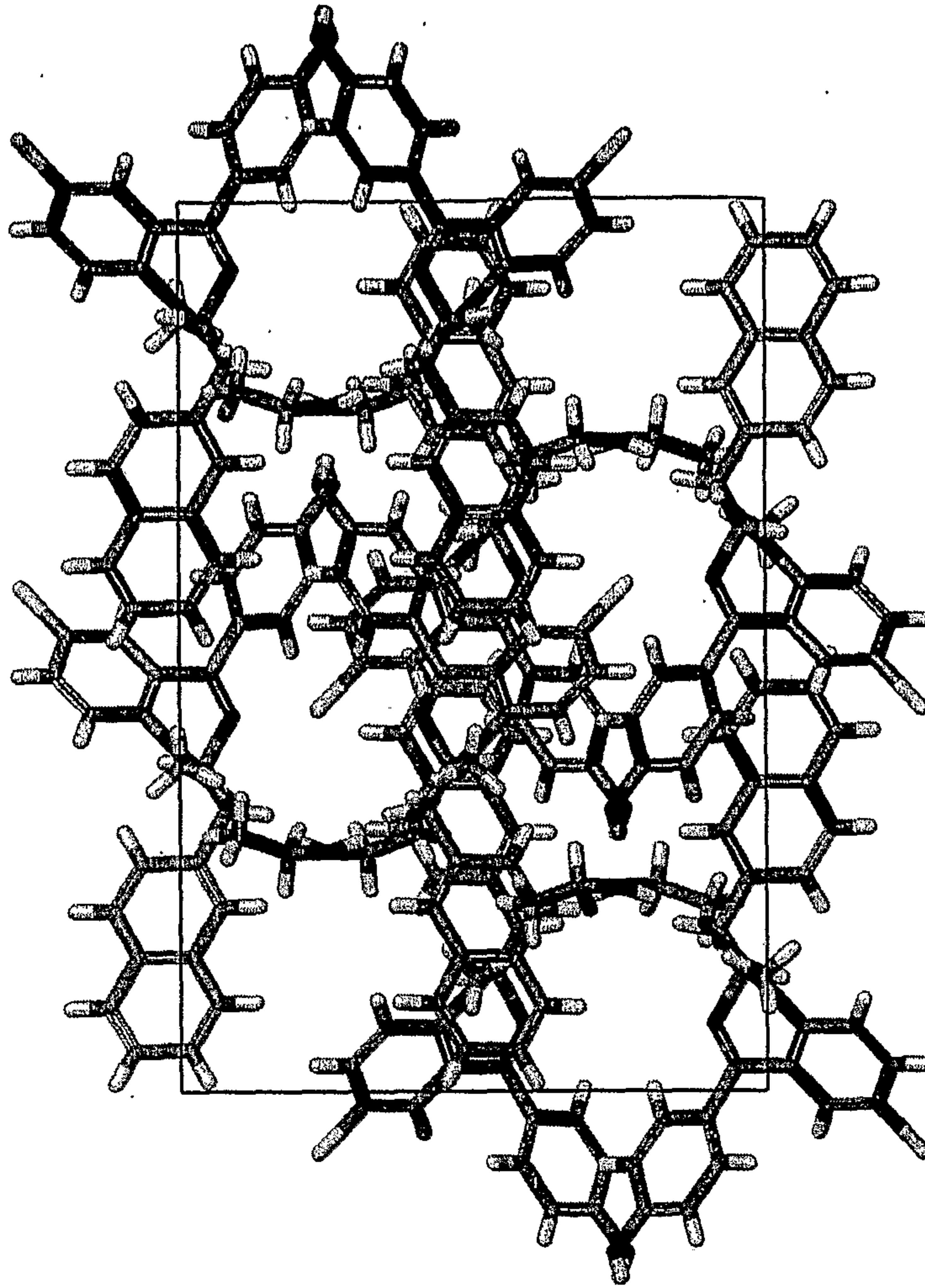
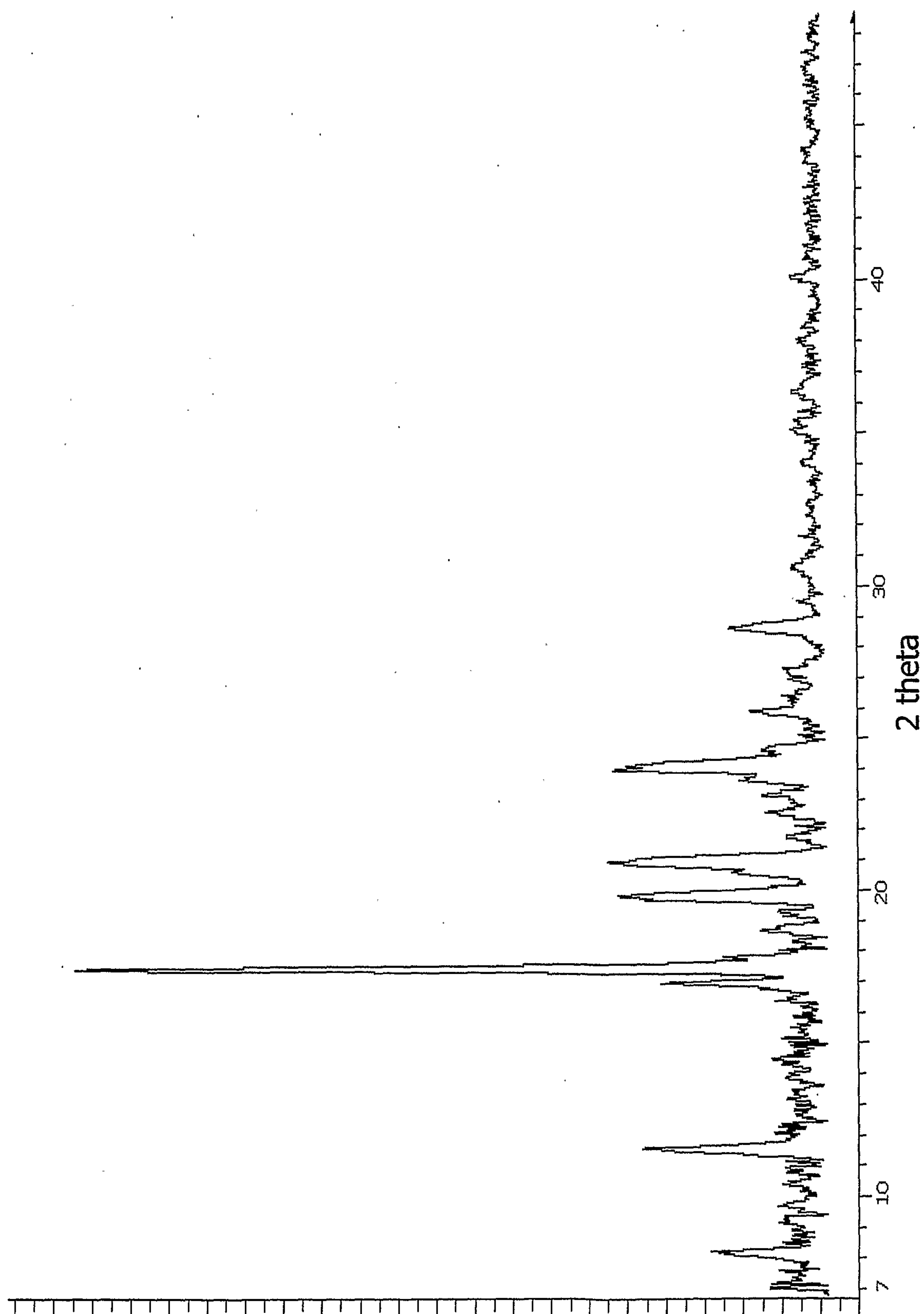


Figure 5. Powder X-ray Diffraction For BZ-423 Ethanol Solvate



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Figure 6. Raman Spectroscopy For BZ-423 Ethanol Solvate

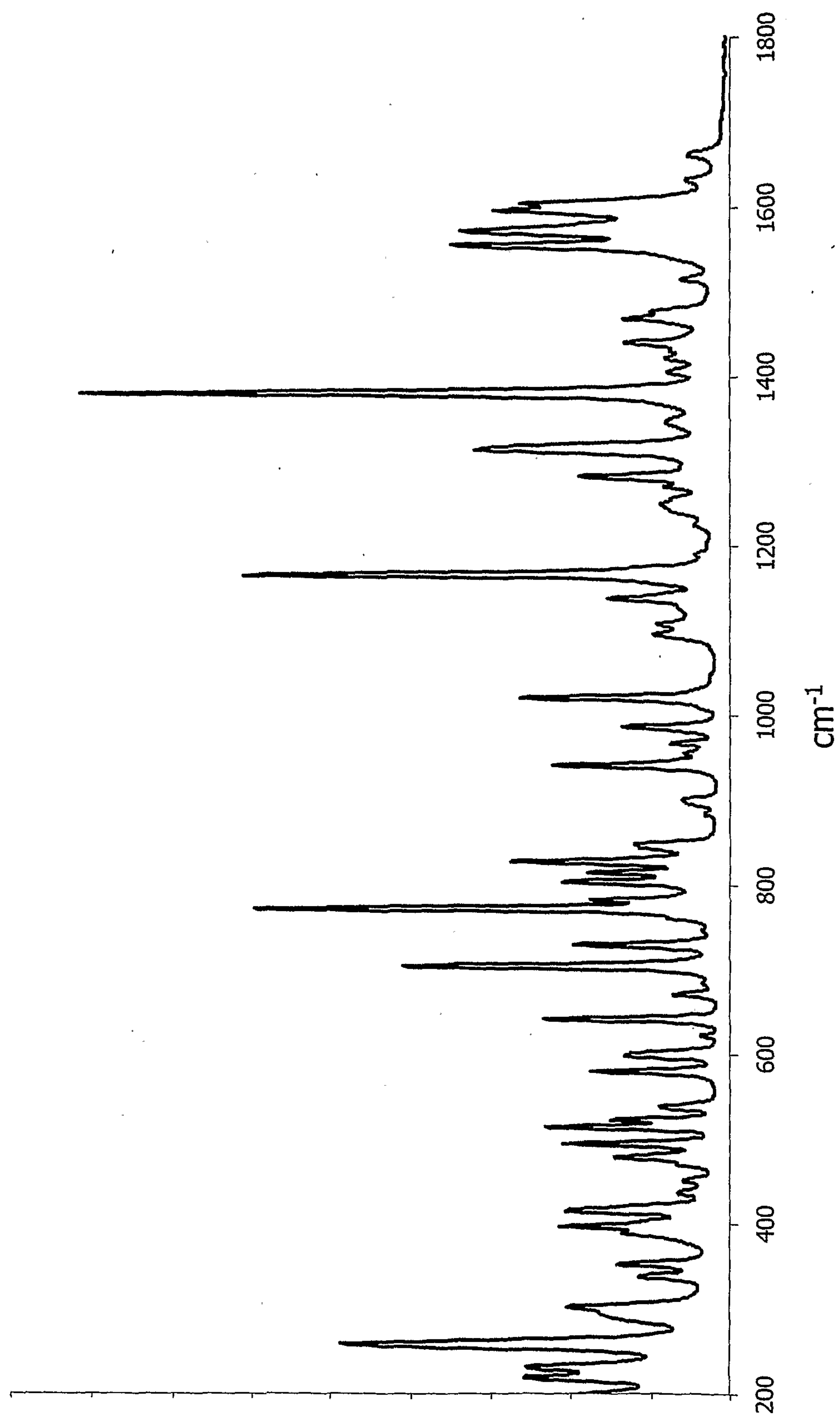


Figure 7. BZ-423 Succinic Acid Raman Spectroscopy

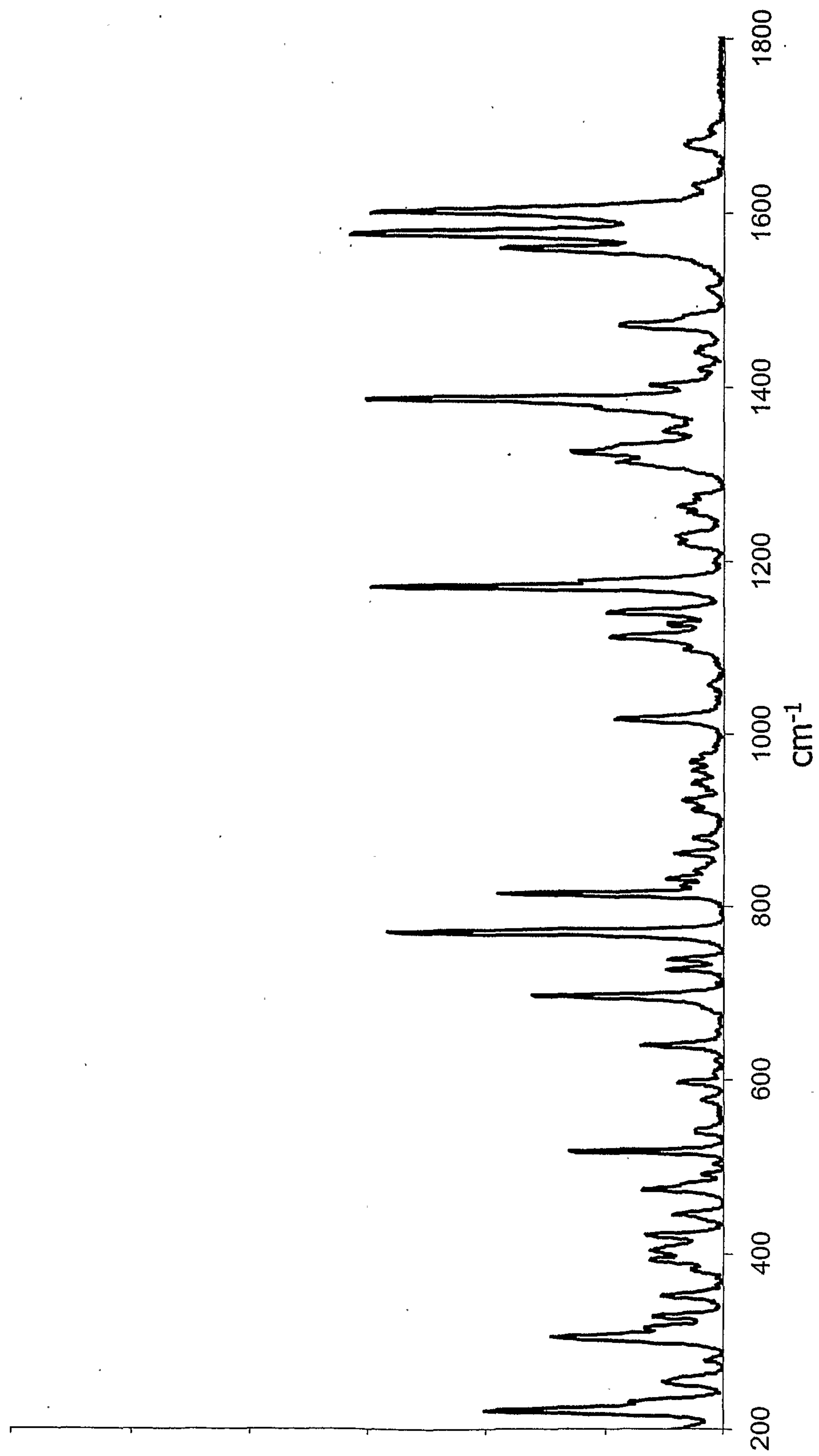


Figure 8. BZ-423 Citric Acid Raman Spectroscopy

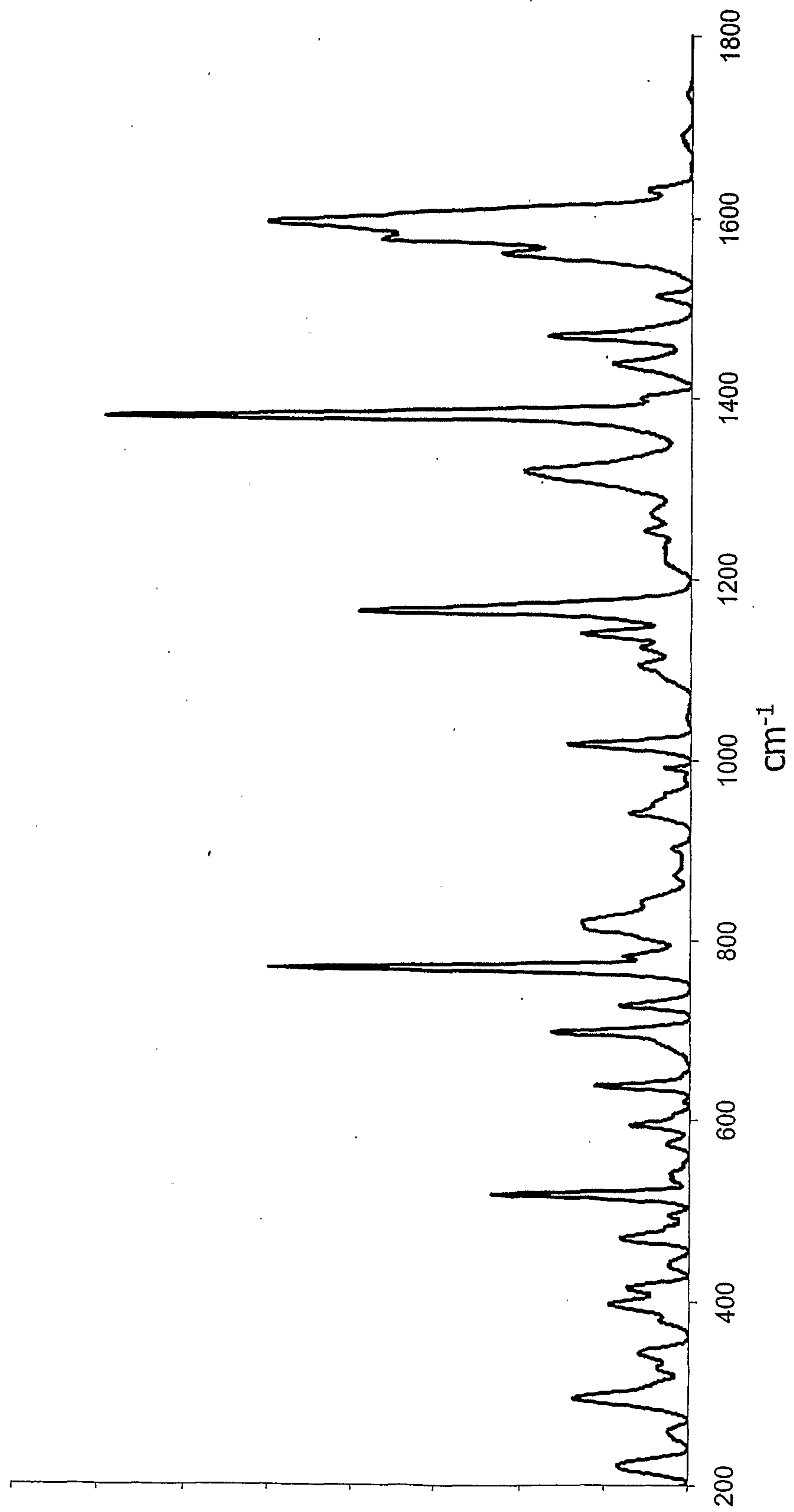
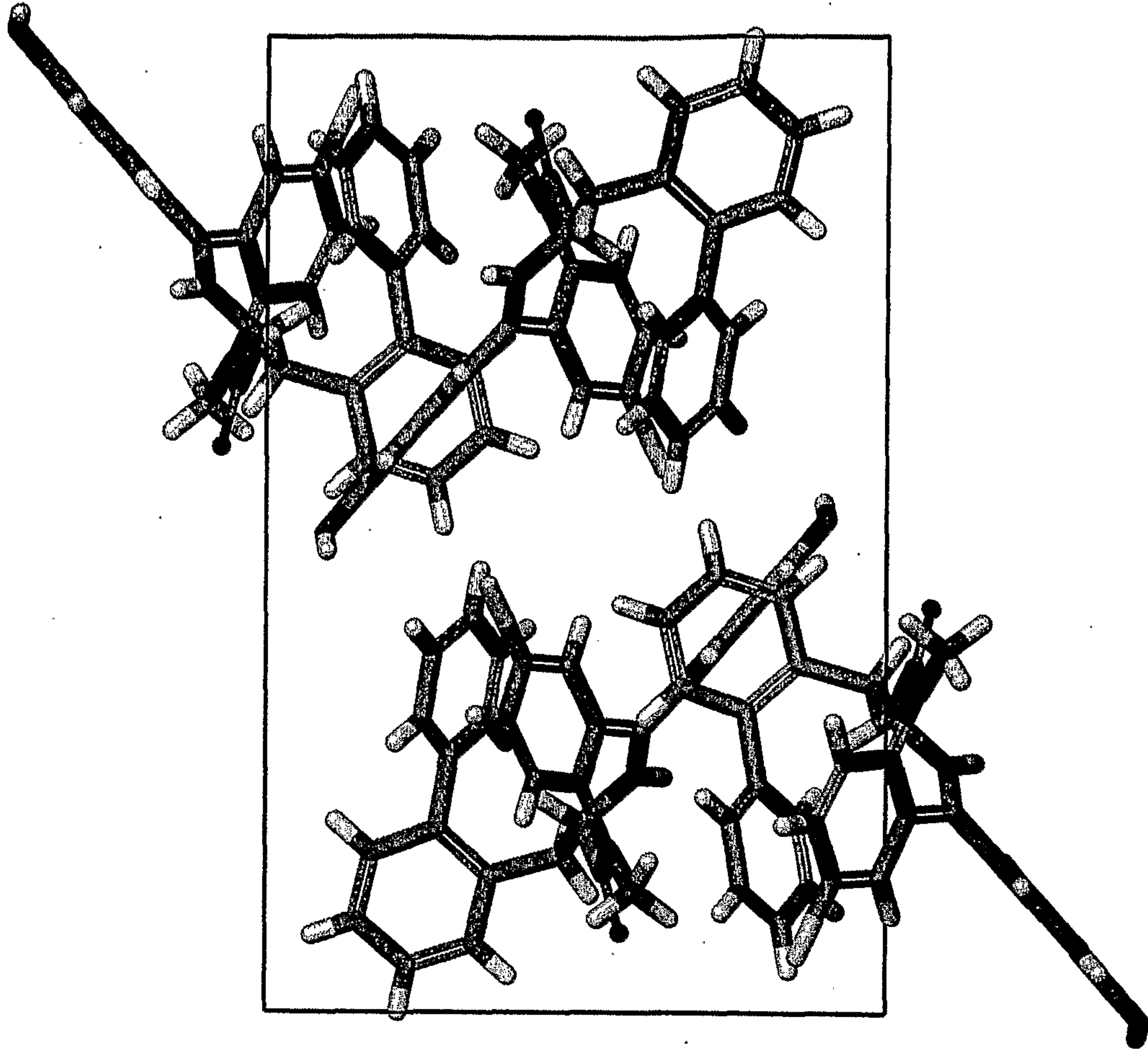


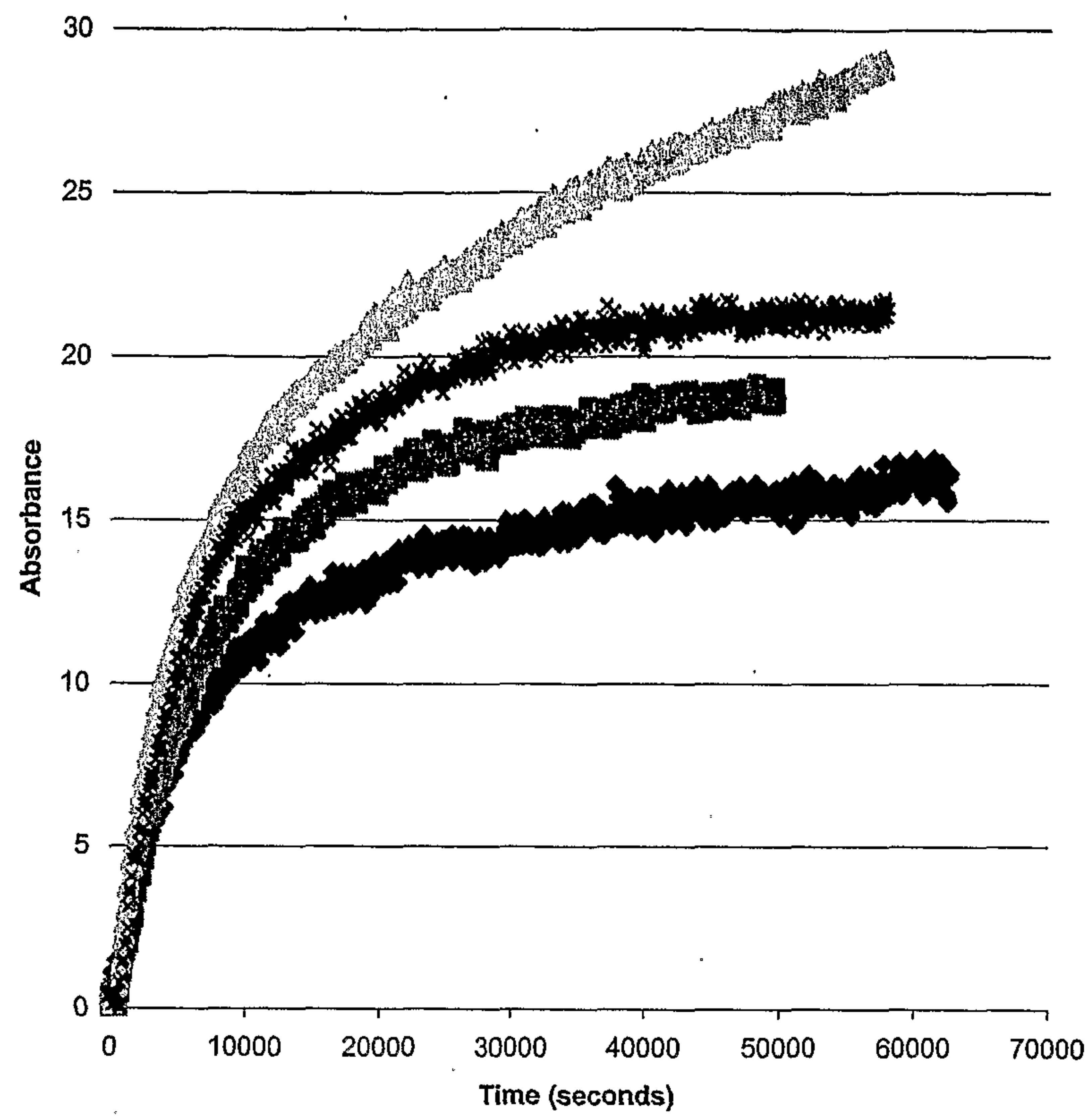
Figure 9. BZ-423 Biphenyl Derivative

	BZ-423 biphenyl derivative
crystal system	monoclinic
space group	$P2_1/n$
a (Å)	9.7205(8)
b (Å)	19.3671(17)
c (Å)	12.8434(10)
α (°)	90.00
β (°)	104.870(3)
γ (°)	90.00
Z	4
cell volume	2336.9(3)
Density	1.327
R	0.0424

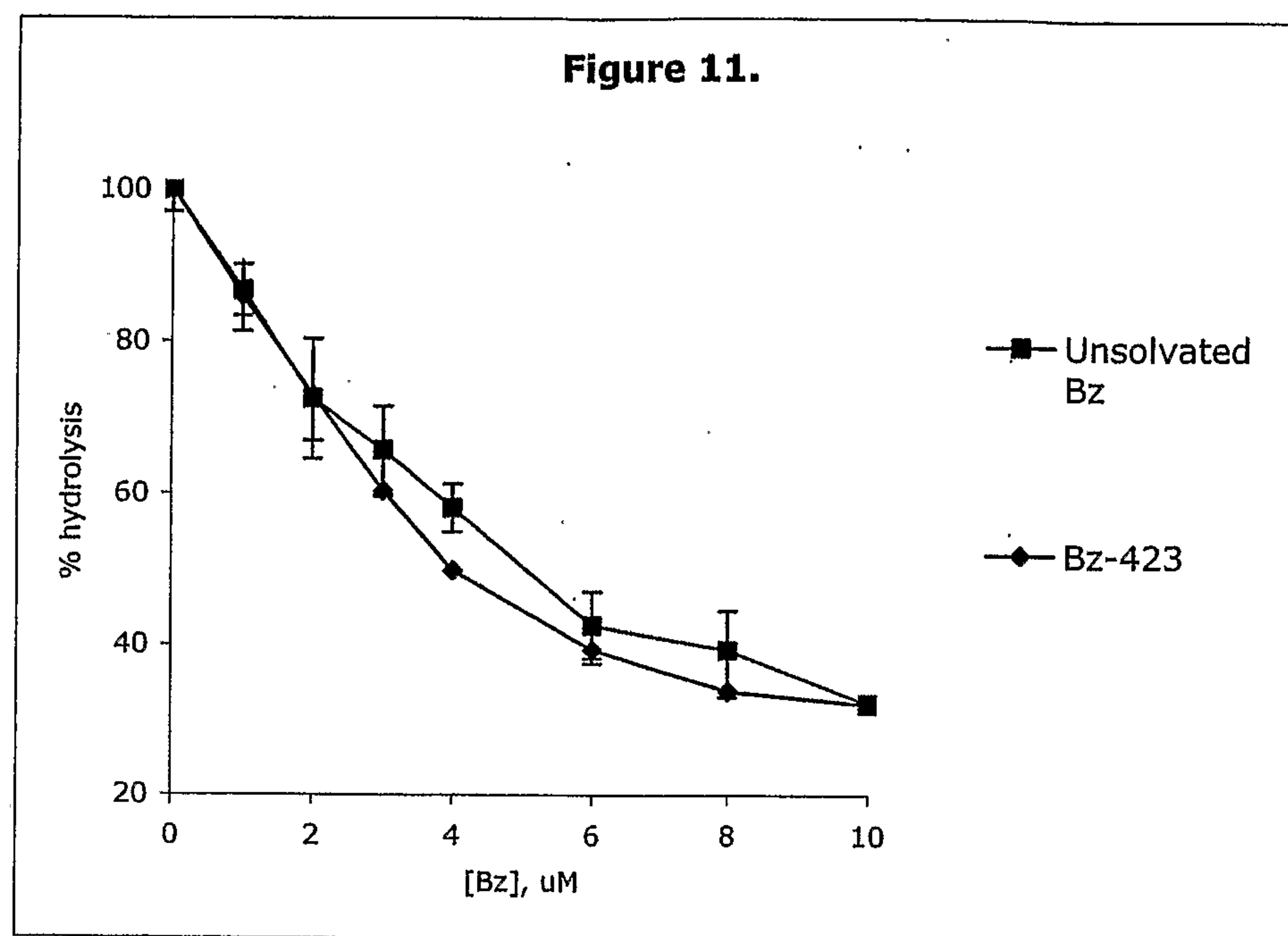


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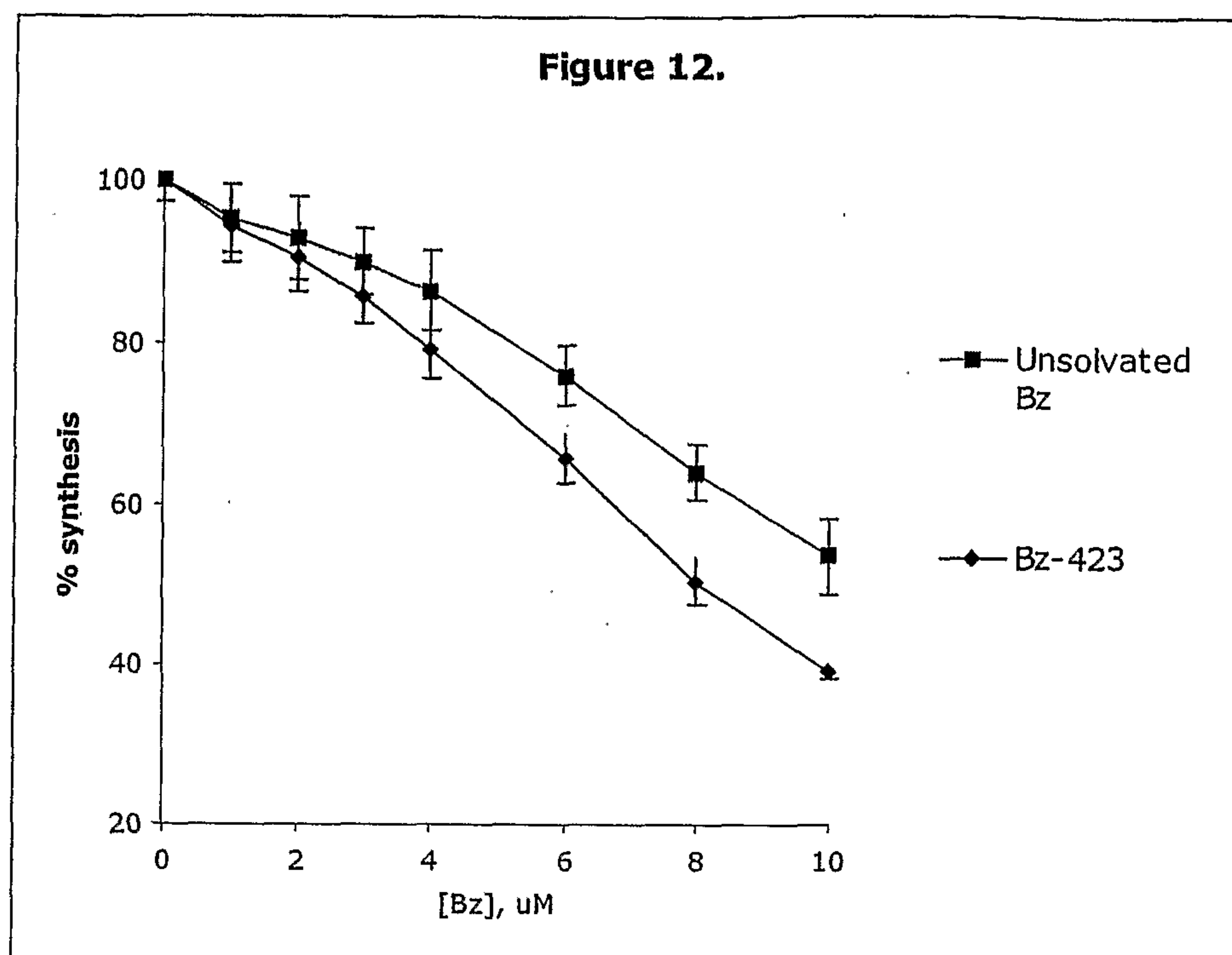
Figure 10.



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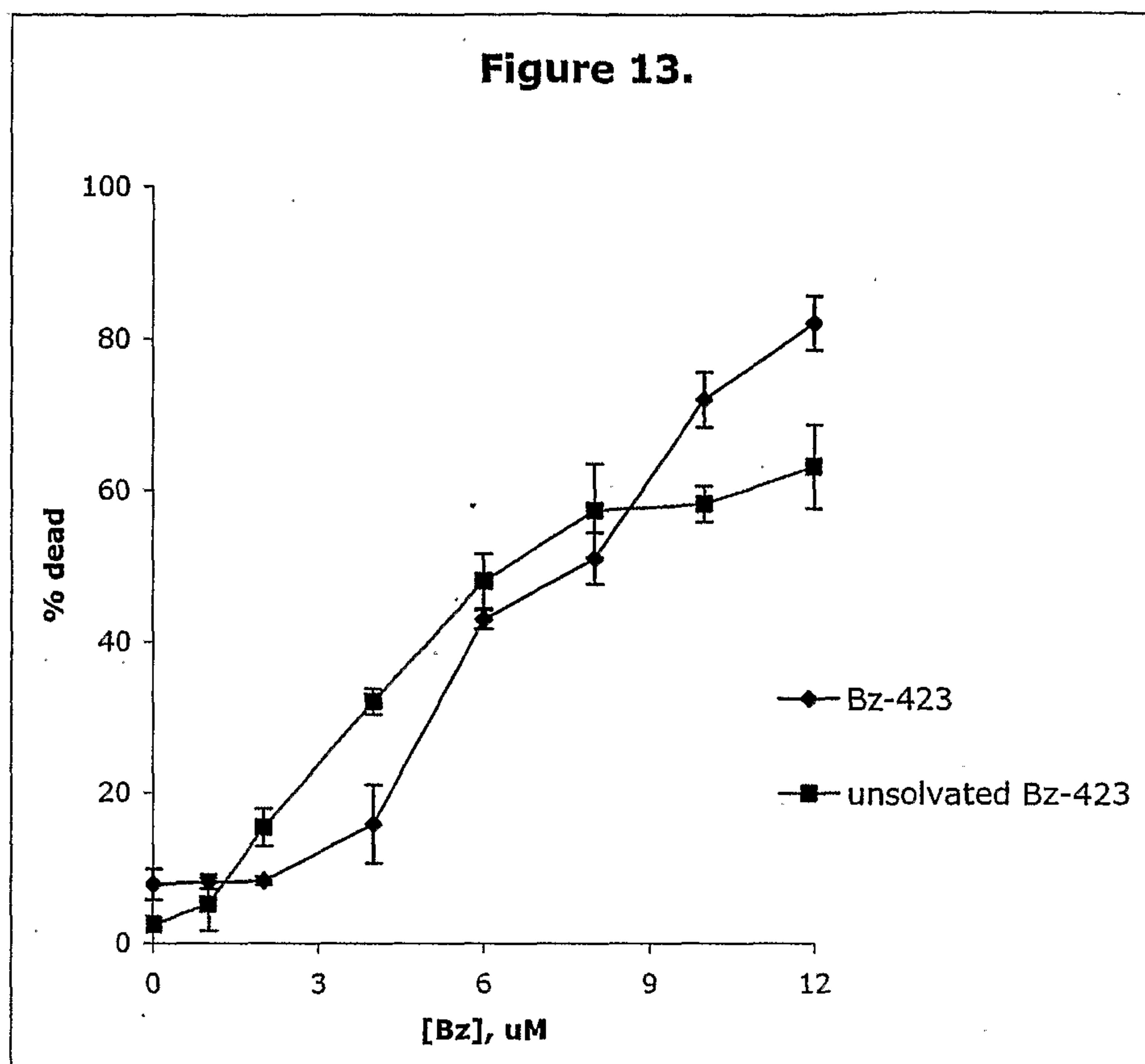


Figure 14. BZ-423 in Simulated Gastric Fluid

- Peak at 370 nm disappears after addition of K_2CO_3 to BZ-423 in gastric fluid.

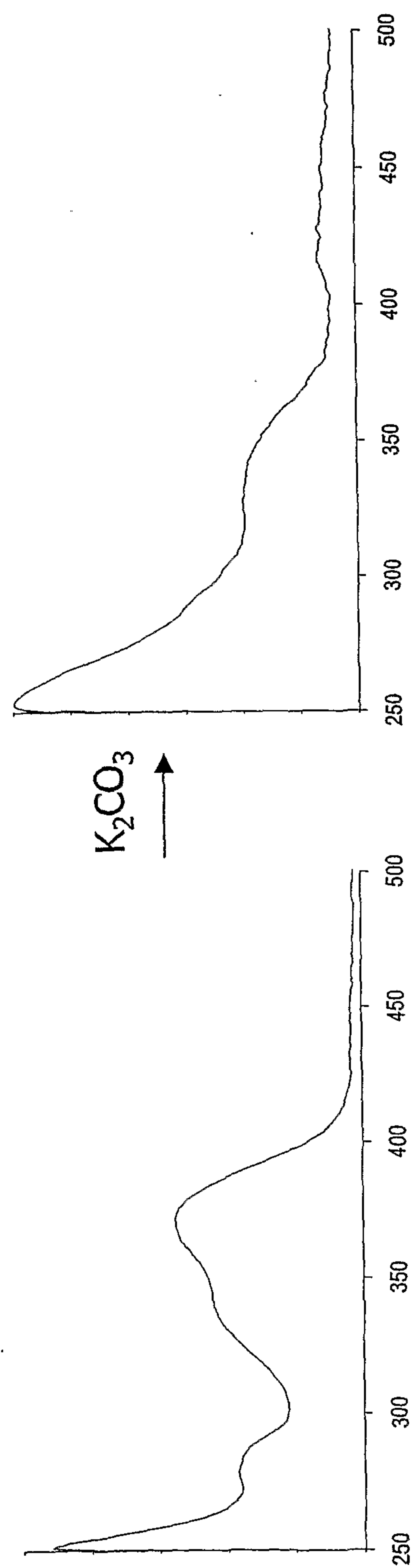
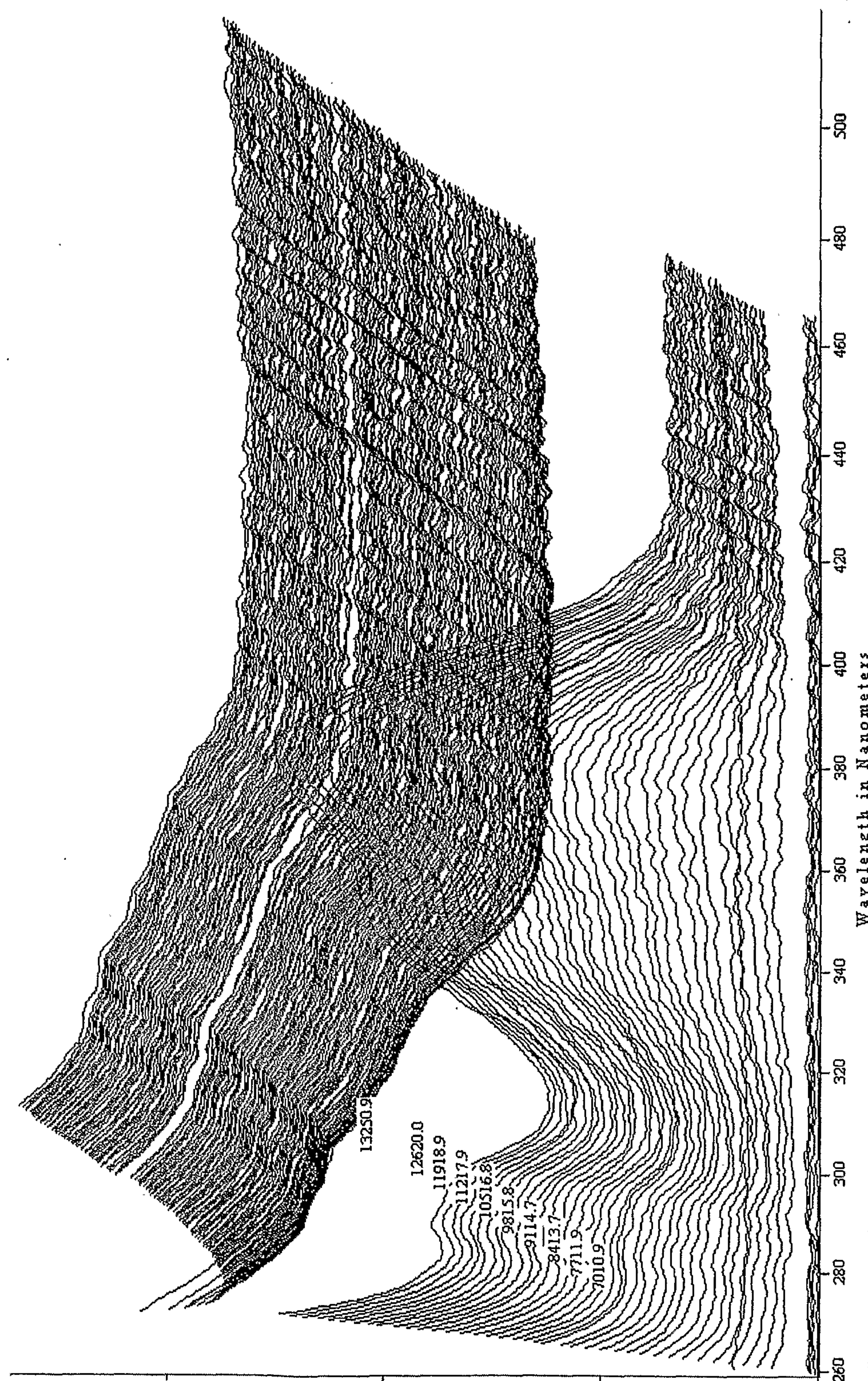
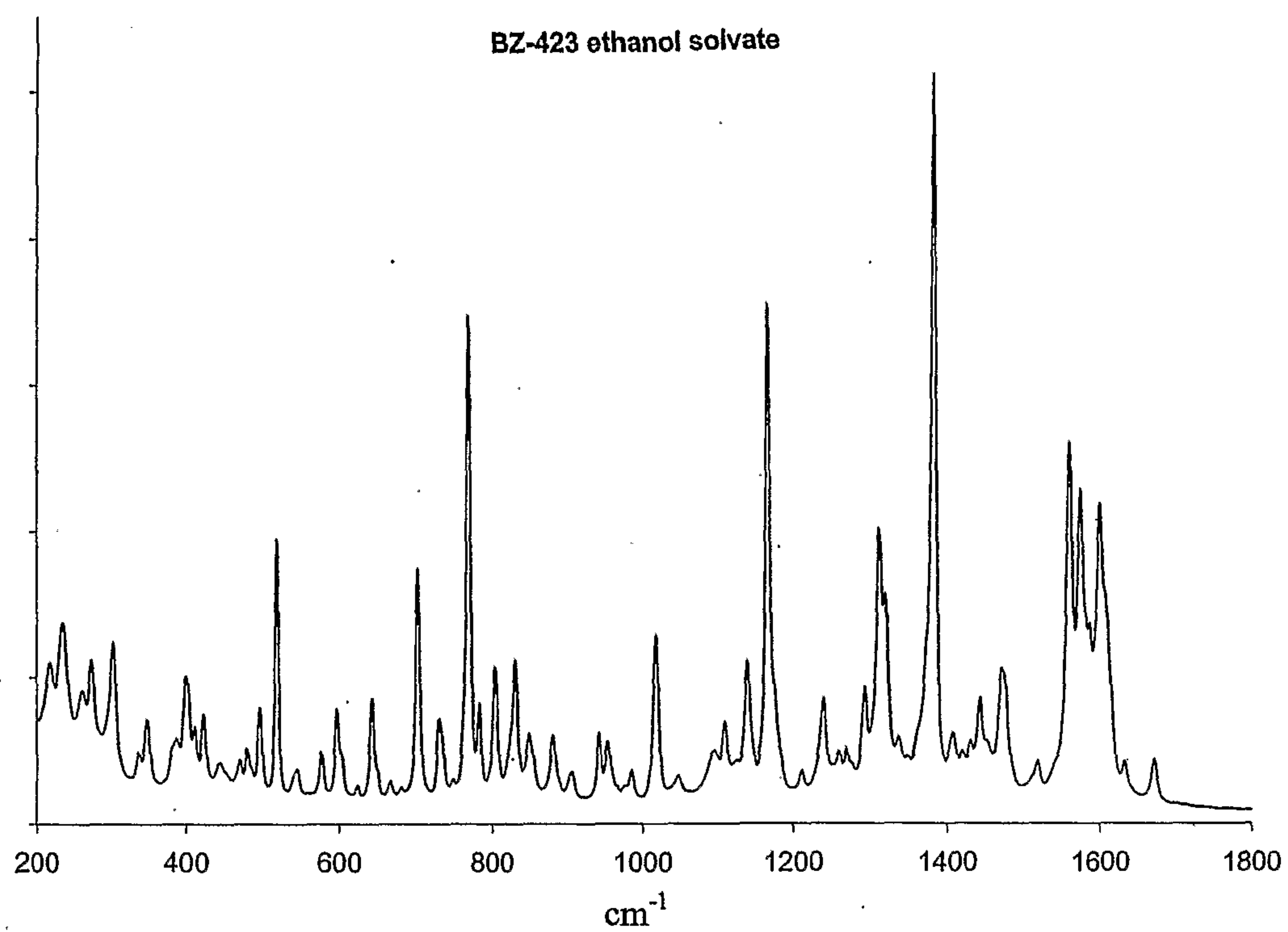
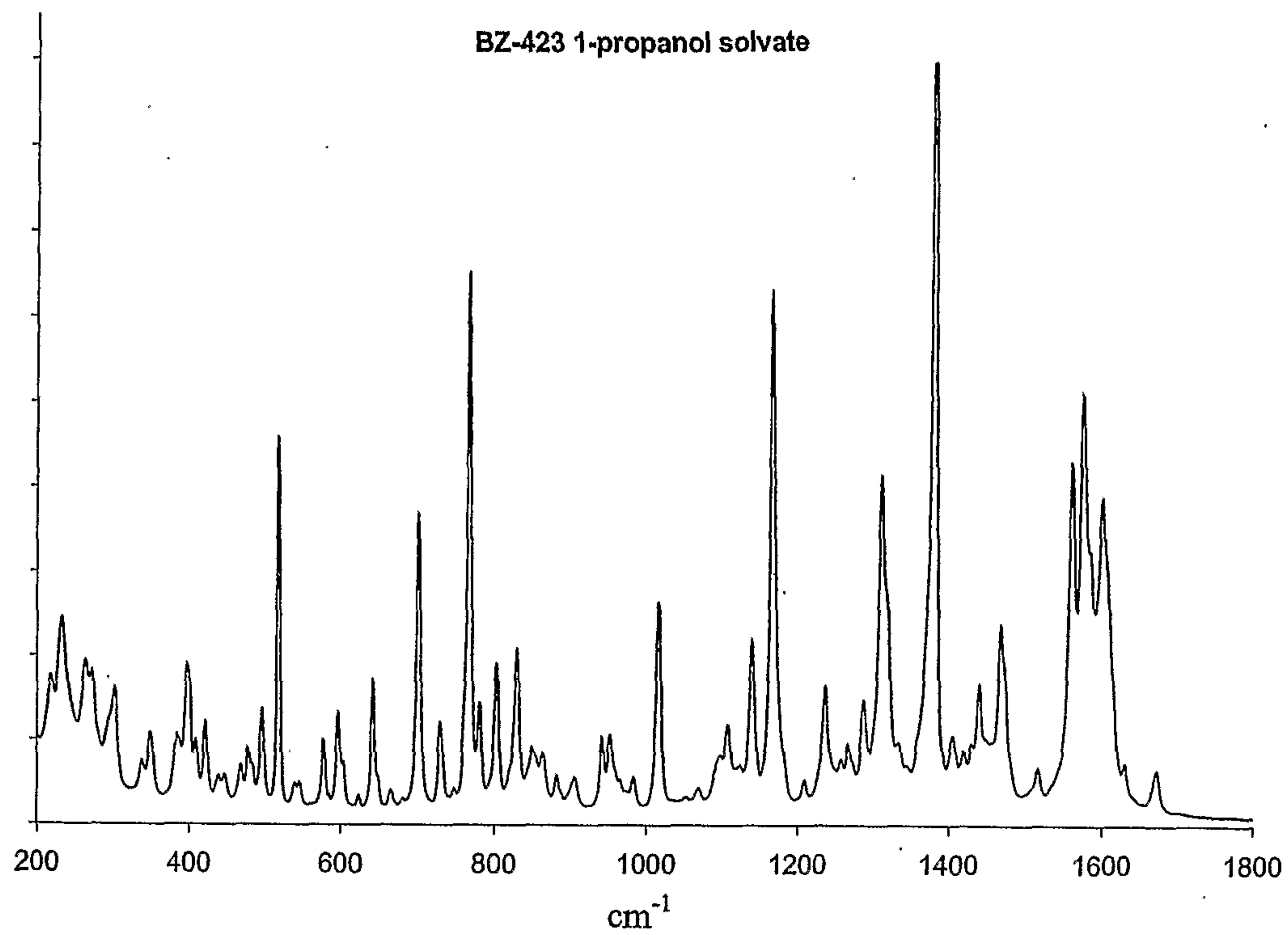


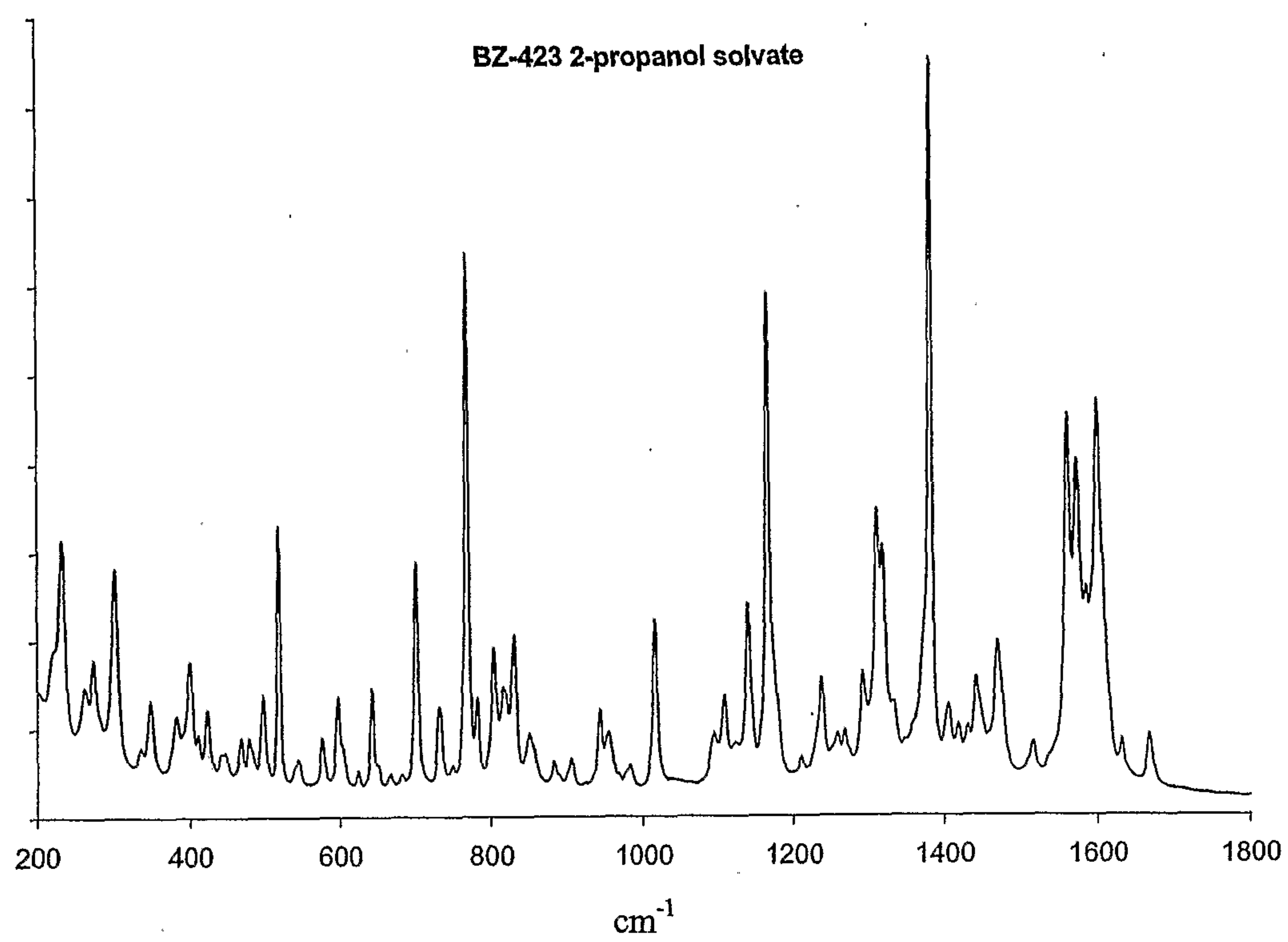
Figure 15. BZ-423 in Simulated Gastric Fluid

- Time lapse UV-vis spectra of BZ-423 in gastric fluid, before and after addition of K_2CO_3 .



**Figure 16.**

**Figure 17.**

**Figure 18.**

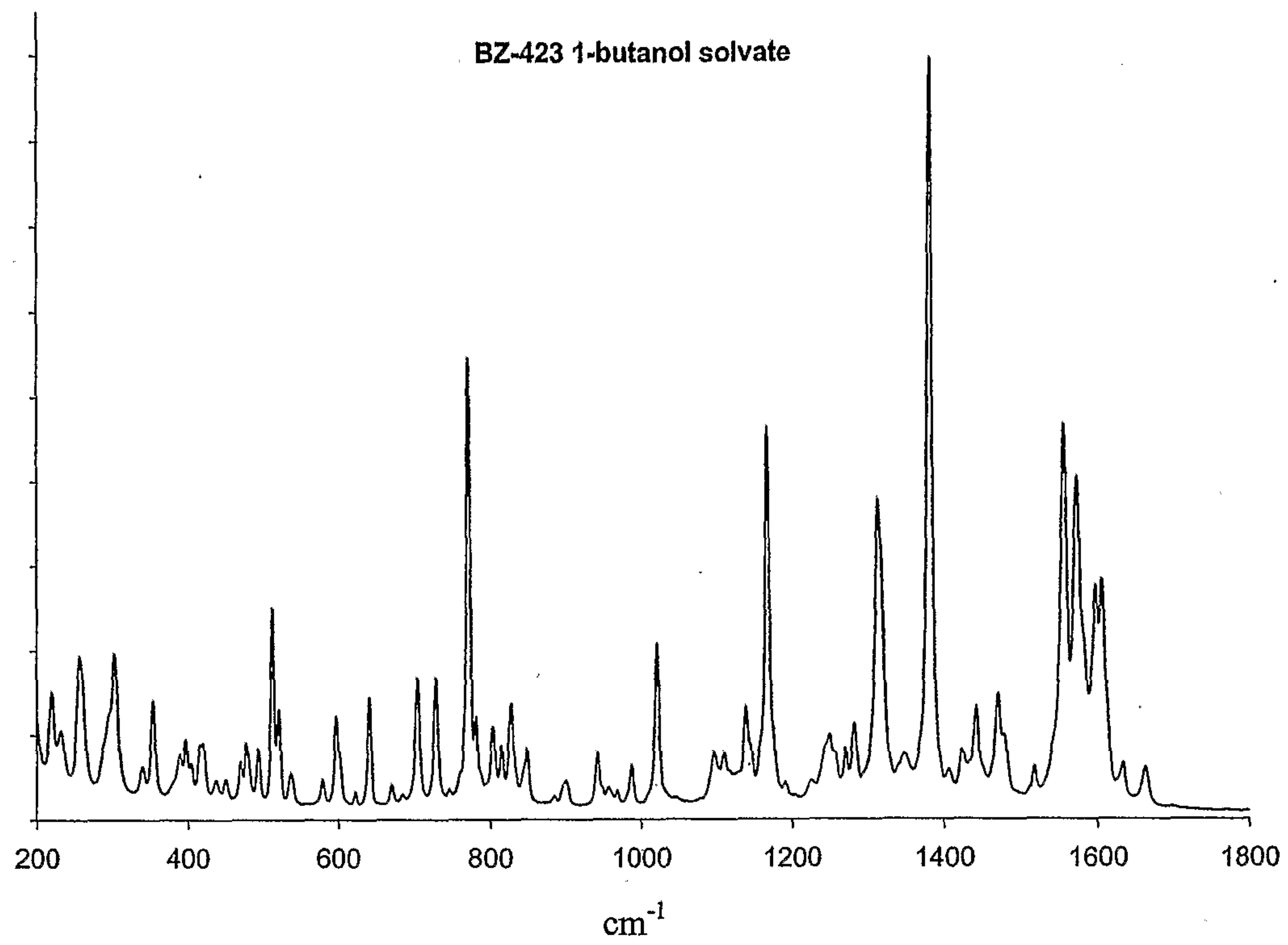
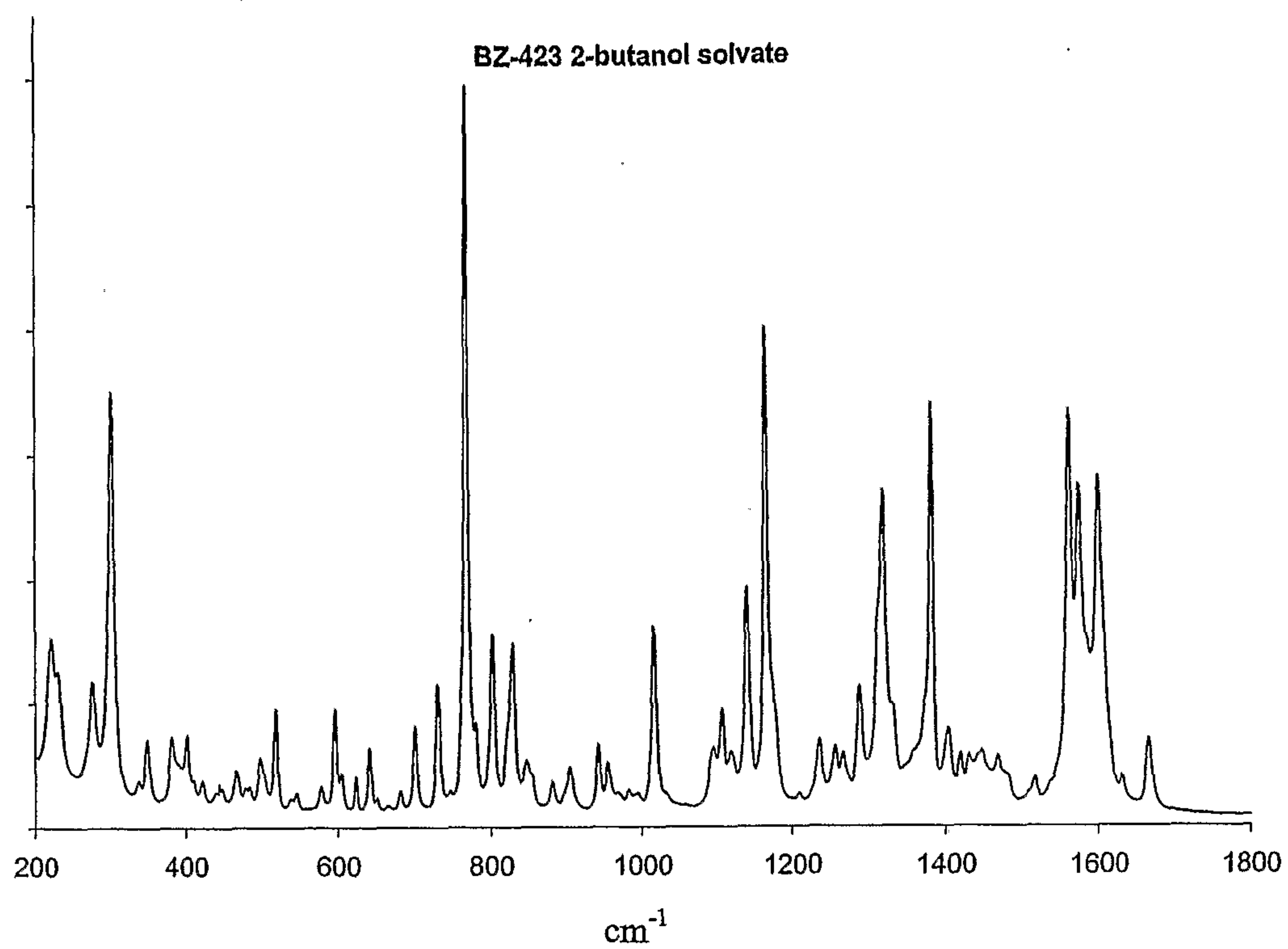
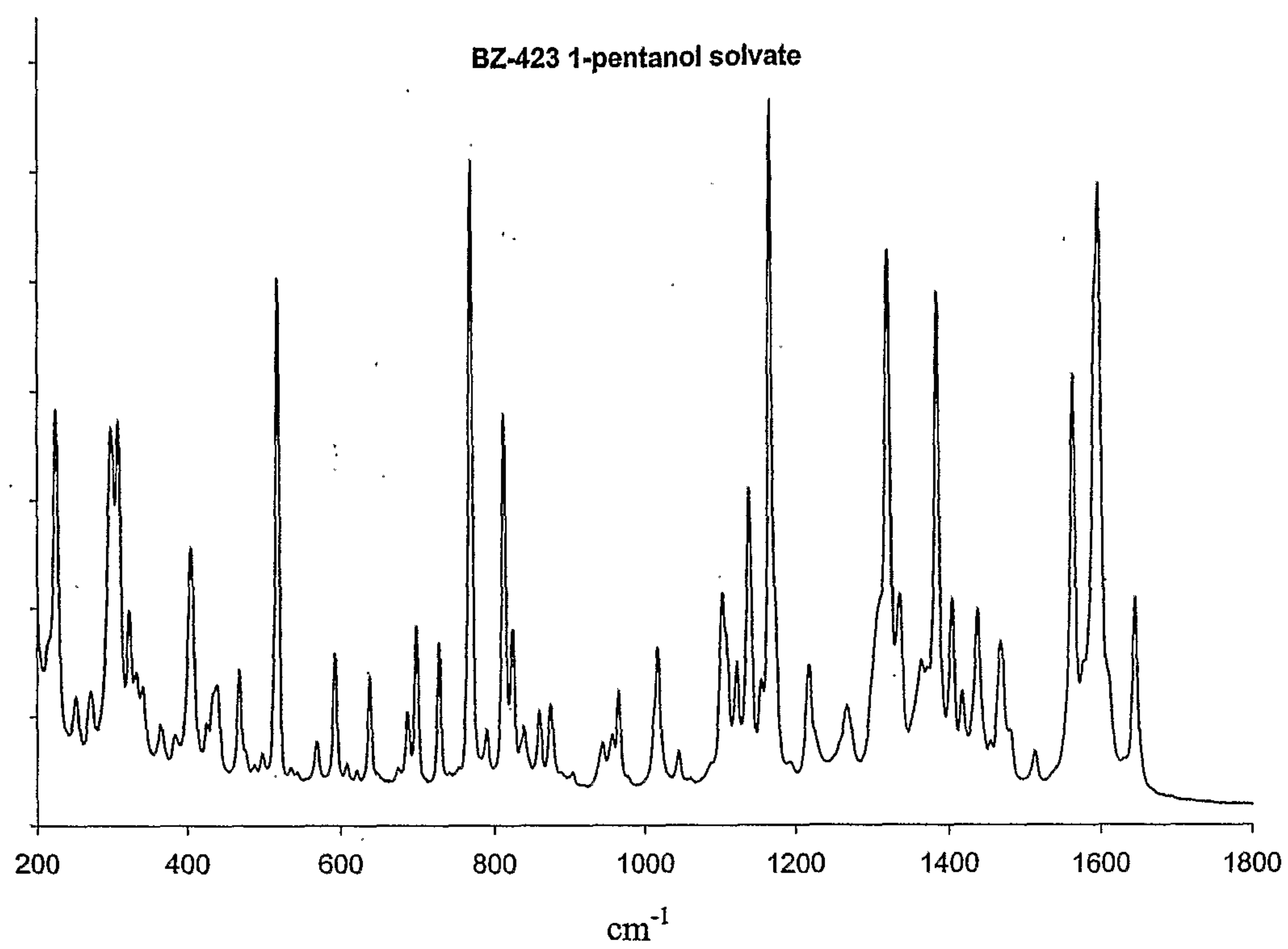
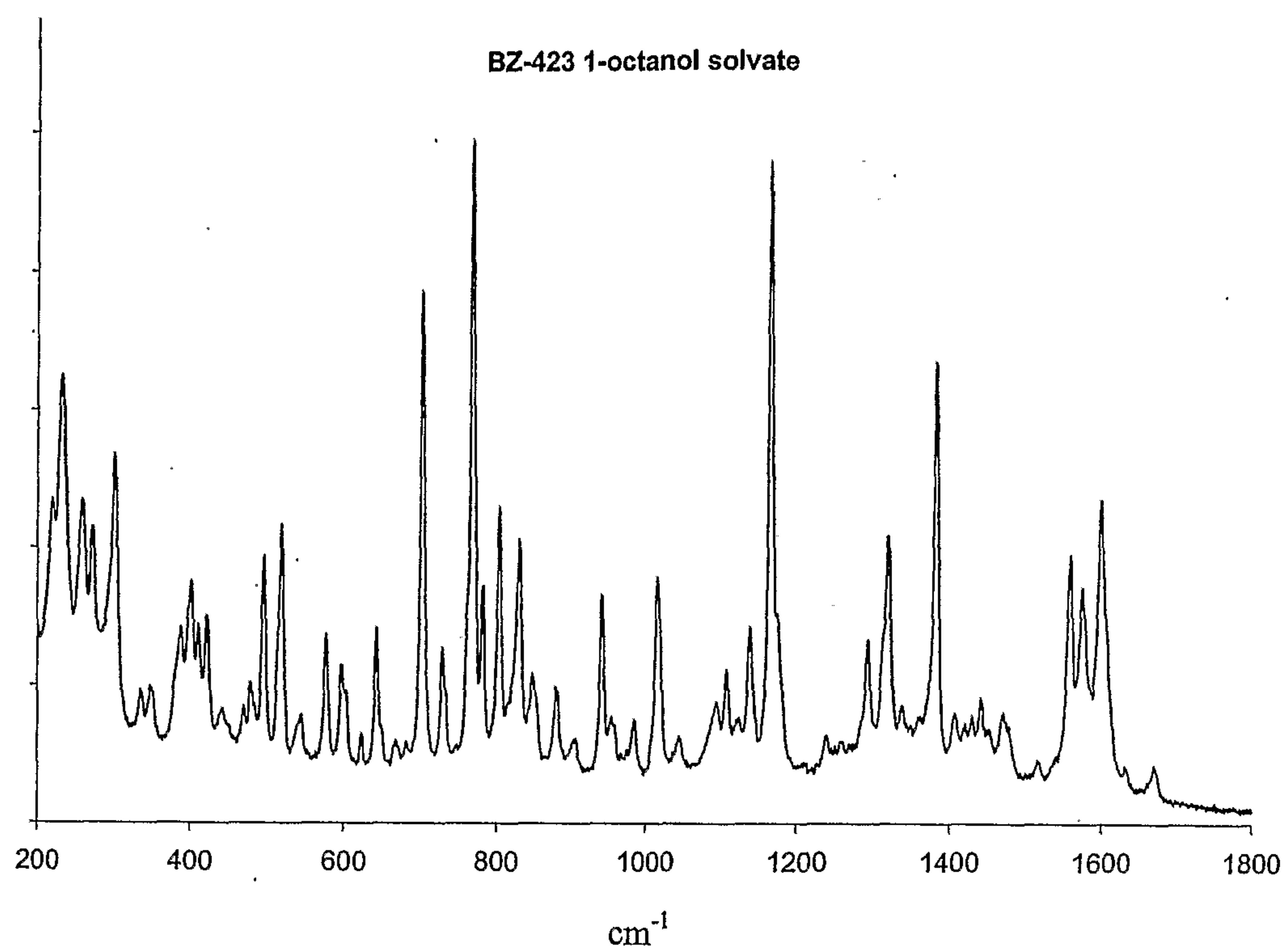
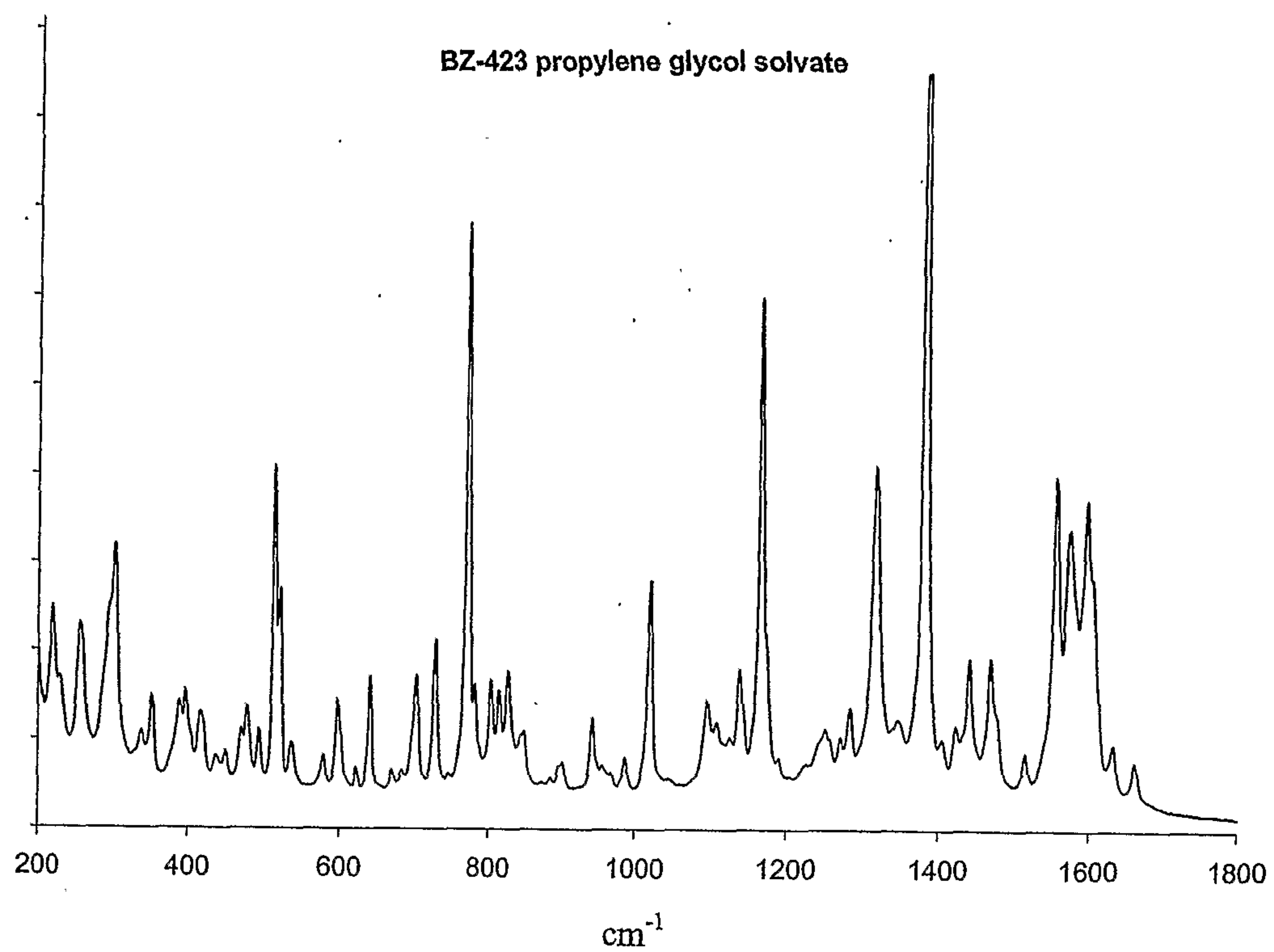


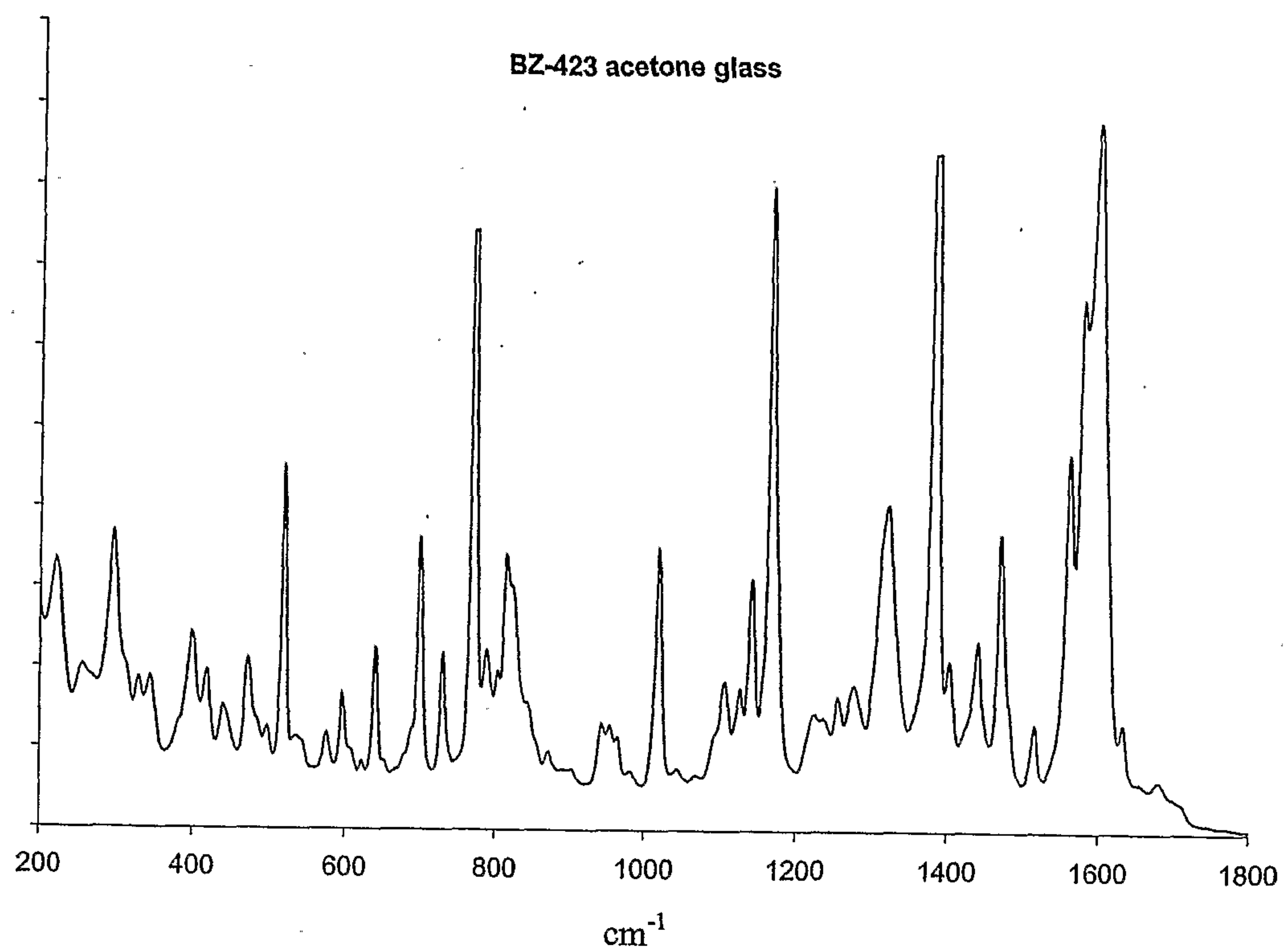
Figure 19.

**Figure 20.**

**Figure 21.**

**Figure 22.**

**Figure 23.**

**Figure 24.**