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DESFERRITHIOCIN POLYETHER ANALOGUES

RELATED APPLICATION

This application claims the benefit of U.S. Provisional Application No. 60/668,045, filed on April 4, 2005. The entire teachings of the above application are
5 incorporated herein by reference.

GOVERNMENT SUPPORT

The invention was supported, in whole or in part, by grant No. R01-DK49108 from the National Diabetes and Digestive and Kidney Diseases Advisory Council (NIDDK) of the National Institute of Health (NIH). The Government has
10 certain rights in the invention.

BACKGROUND OF THE INVENTION

Iron metabolism in primates is characterized by a highly efficient recycling process. Consequently, there is no specific mechanism for eliminating this transition metal. Because of the lack of an iron clearance mechanism, the introduction of
15 "excess iron" into this closed metabolic loop often leads to chronic overload and can ultimately lead to biological damage (e.g., peroxidative tissue damage). There are a number of ways in which excess iron is introduced, including a high-iron diet, acute iron ingestion or malabsorption of the metal. In each of these situations, a subject can typically be treated by phlebotomy to reduce iron levels. However, for iron-
20 overload syndromes resulting from chronic transfusion therapy, e.g., aplastic anemia and thalassemia, phlebotomy is not an option. In these secondary iron overload syndromes, the origin of the excess iron is the transfused red blood cells. Since removing the red blood cells to remedy the iron overload would be counterproductive, an alternative method of removing iron is chelation therapy.

25 Although considerable effort has been invested in the development of new therapeutics for managing iron overload resulting from thalassemia, particularly

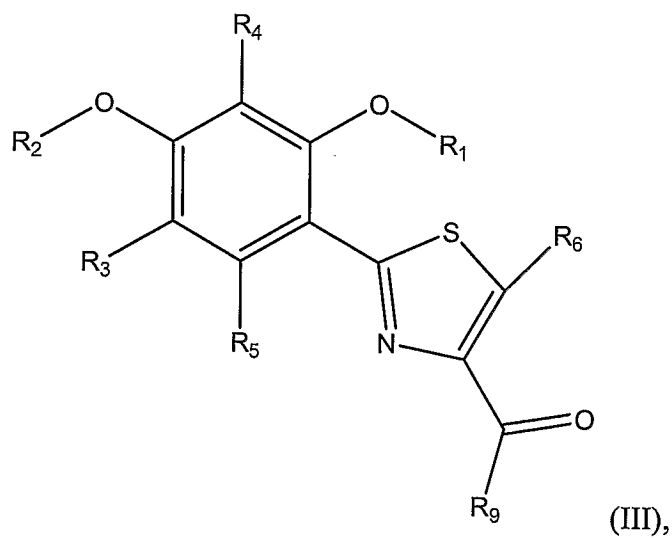
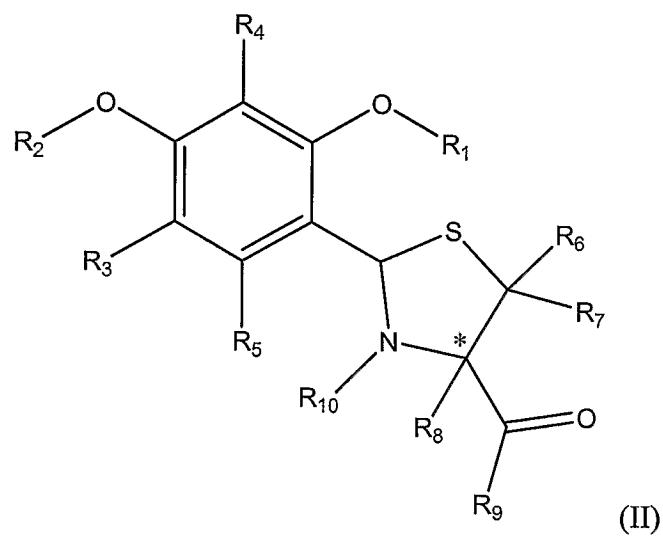
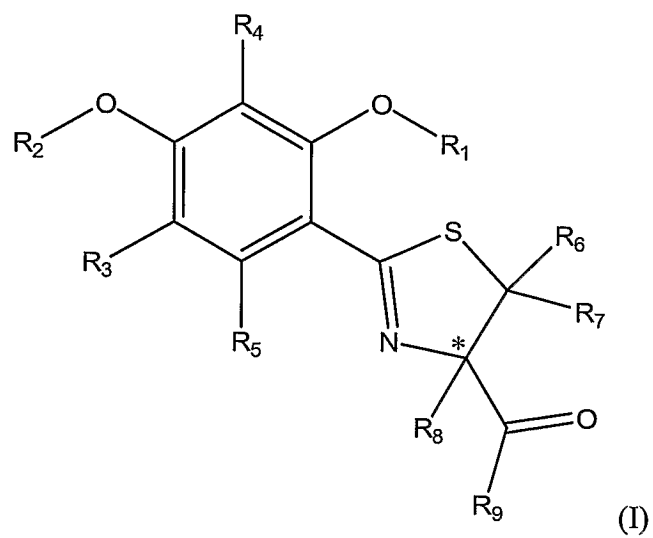
therapeutics that can be administered orally, desferrioxamine B, a hexacoordinate hydroxamate iron chelator produced by *Streptomyces pilosus*, is still the protocol of choice. However, desferrioxamine B is not ideal for chelation therapy, because iron is removed with a low efficiency. In addition, oral activity of desferrioxamine B is
5 marginal, thereby requiring parenteral administration, which can result in poor patient compliance, particularly for patients in need of long-term chelation therapy.

A substantial number of synthetic iron chelators have been studied in recent years as potential orally active therapeutics, e.g., pyridoxal isonicotinoyl hydrazone (PIH), hydroxypyridones and *N, N'*-bis-(2-hydroxybenzylethylenediamine)-*N, N'*-diacetic acid (HBED); however, the synthetic chelators have not yet demonstrated
10 the desired properties (e.g., effective chelation, suitable oral activity, and acceptable toxicity). Siderophores including enterobactin and rhodotorulic acid have also been studied. However, both enterobactin and rhodotorulic acid have exhibited unacceptable toxicity and neither demonstrated measurable oral activity. In general,
15 although a large number of siderophores and synthetic iron chelators have been developed, most have been abandoned because their properties are not suitable for use in treating chronic iron overload.

Therefore, a need still exists for novel iron chelators that can be used in chelation therapy, especially chronic chelation therapy. Suitable chelators can be
20 efficient in chelating and removing iron from an organism, possess suitable oral bioavailability and/or pose minimal toxicity to a subject.

SUMMARY OF THE INVENTION

This application relates to compounds characterized by a structural formula
25 selected from Structural Formulas (I), (II), and (III):



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R_1 is $-H$ or an acyl group;

R_2 is $-[(CH_2)_n-O]_x-[(CH_2)_n-O]_y-R'$;

R_3 , R_4 and R_5 are each independently $-H$, an alkyl group, or $-OR_{11}$;

R_6 , R_7 , and R_8 are each independently $-H$ or an alkyl group;

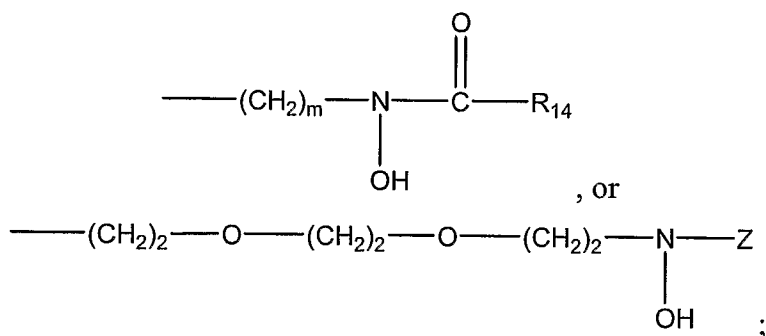
R_9 is $-OR_{12}$ or $-N(OH)R_{13}$;

R_{10} is $-H$ or an alkyl group;

5 R_{11} is $-H$, an alkyl group or an acyl group;

R_{12} is $-H$ or an alkyl group;

R_{13} is an alkyl group,



10 R_{14} is an alkyl group;

R' is an alkyl group;

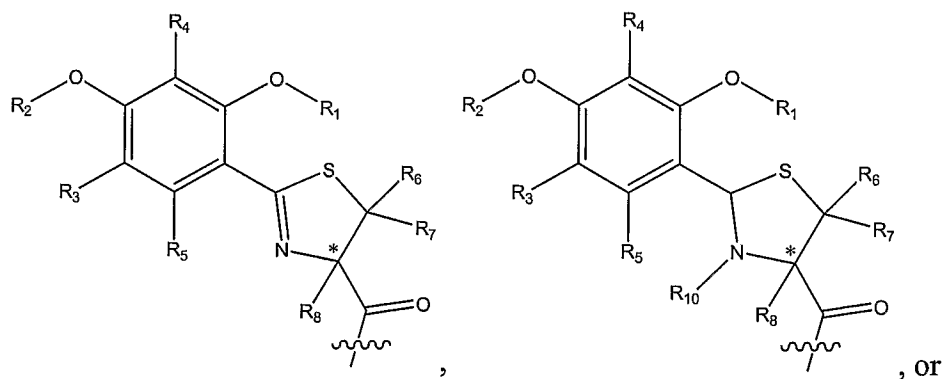
m is an integer from 1 to 8;

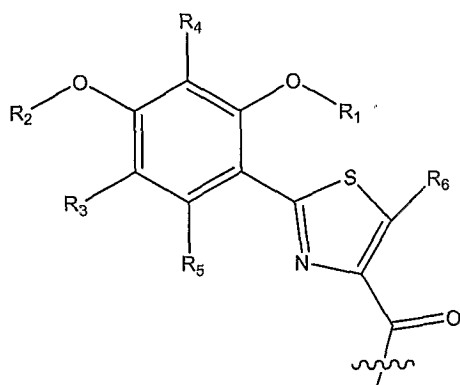
each n is independently an integer from 1 to 8;

x is an integer from 1 to 8;

15 y is an integer from 0 to 8;

Z is $-\text{C}(\text{O})R_{14}$,





or a salt, solvate or hydrate thereof.

The invention includes pharmaceutical compositions comprising a compound of the invention in conjunction with a carrier or diluent. The pharmaceutical
5 compositions can be for use in therapy.

In another embodiment, the present invention is a method of treating a pathological condition responsive to chelation or sequestration of a trivalent metal in a subject, comprising administering to the subject a therapeutically or prophylactically effective amount of a compound represented by a structural formula
10 selected from Structural Formulas (I), (II), and (III) or a pharmaceutical composition including one of these compounds.

The compounds of the invention can also be used in a method of reducing oxidative stress in a subject in need of treatment therefor and a method of treating a subject who is suffering from neoplastic disease or a preneoplastic condition, in
15 which a therapeutically effective amount of one of the compounds or pharmaceutical compositions of the invention is administered to the subject.

The invention also relates to the use of compounds disclosed herein in medical therapy. The invention further relates to the use of the compounds of the invention for the manufacture of a medicament for therapy, for example, for treating
20 pathological conditions responsive to chelation or sequestration of metals, for reducing oxidative stress and for treating neoplastic disease or a preneoplastic condition.

The metal chelators of the invention have the advantage of having a desirable iron clearing efficiency. The metal chelators of the invention can possess a different
25 volume of distribution from present known chelators, resulting in a different distribution among organs. This different distribution can permit penetration into

organs such as the heart, brain and pancreas, as well as result in the majority of clearance of the chelators in the liver, thereby decreasing the risk of toxicity to the kidneys. Advantageously compounds of the invention exhibit a low concentration in the kidneys following administration.

5 BRIEF DESCRIPTION OF THE DRAWINGS

The foregoing and other objects, features and advantages of the invention will be apparent from the following more particular description of preferred embodiments of the invention, as illustrated in the accompanying drawings in which like reference characters refer to the same parts throughout the different views. The
10 drawings are not necessarily to scale, emphasis instead being placed upon illustrating the principles of the invention.

FIG. 1 shows the iron clearing efficiency of compounds of the invention in a non-iron overloaded, bile duct cannulated rat model.

FIG. 2 shows the distribution of compounds of the invention in the kidneys
15 of rodents over time following subcutaneous administration.

FIG. 3 shows the distribution of compounds of the invention in the liver of rodents over time following subcutaneous administration.

FIG. 4 shows the distribution of compounds of the invention in the heart of rodents over time following subcutaneous administration.

FIG. 5 shows the distribution of compounds of the invention in the pancreas
20 of rodents over time following subcutaneous administration.

FIG. 6 shows the distribution of (S)-4'-(HO)-DADFT and (S)-4'-(CH₃O)-DADFT in the liver of rodents over time following subcutaneous administration.

FIG. 7 shows the distribution of (S)-4'-(HO)-DADFT and (S)-4'-(CH₃O)-
25 DADFT in the heart of rodents over time following subcutaneous administration.

FIG. 8 shows the distribution of (S)-4'-(HO)-DADFT and (S)-4'-(CH₃O)-DADFT in the pancreas of rodents over time following subcutaneous administration.

FIG. 9 shows the uranium excretion in rats induced by compounds of the invention following administration.

DETAILED DESCRIPTION OF THE INVENTION

Compounds of the invention are represented by a structural formula selected from Structural Formulas (I), (II), and (III) as described above.

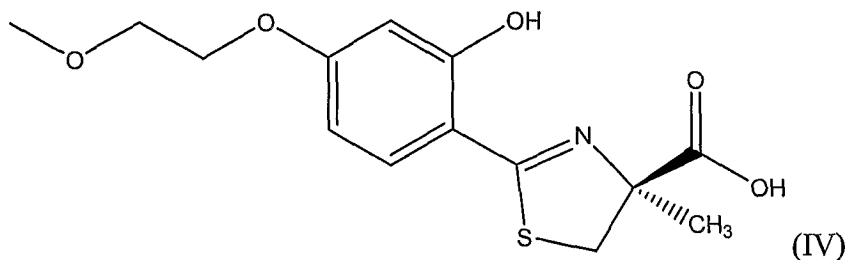
As discussed below, stereoisomers and mixtures of stereoisomers of the
5 compounds disclosed herein are included in the invention.

Typically, compounds of the invention are represented by Structural Formulas (I), where the variables are as defined above.

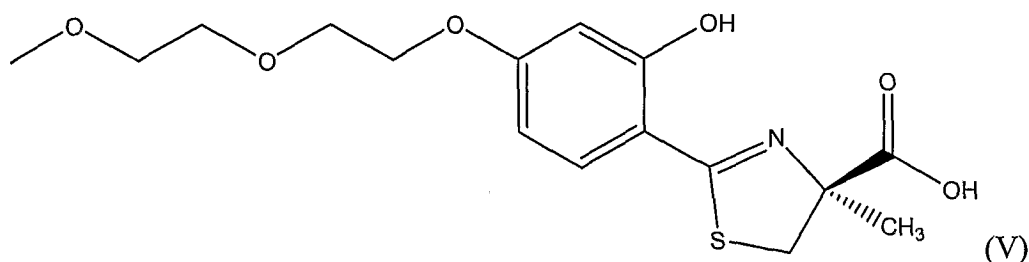
In one embodiment, R_9 in Structural Formulas (I)-(III) is $-OR_{12}$.

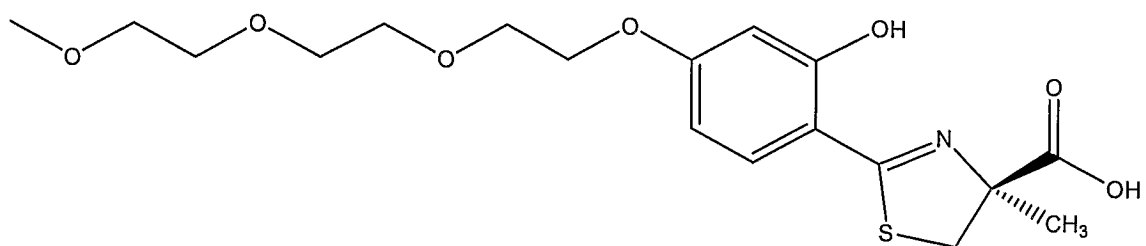
When R_9 is $-OR_{12}$, compounds of the invention can have one or more of the
10 following features: (1) R_8 is $-H$ or $-CH_3$; (2) R_6 and R_7 are each $-H$ or $-CH_3$, preferably $-H$; (3) R_3 , R_4 and R_5 are each $-H$; (4) R_2 is $-[(CH_2)_n-O]_x-R'$; n is independently 1 to 4; and x is 1 to 4; (5) R_1 is $-H$; and (6) R' is $-CH_3$. Preferred compounds of the invention have feature (1), more preferably features (1) and (2), and even more preferably features (1), (2) and (3). Particularly preferred
15 compounds of the invention have features (1), (2), (3) and (4); (1), (2), (3), (4) and (5); or have all six of the above features.

Specific examples of compounds of the invention are represented by Structural Formulas (IV)-(IX):

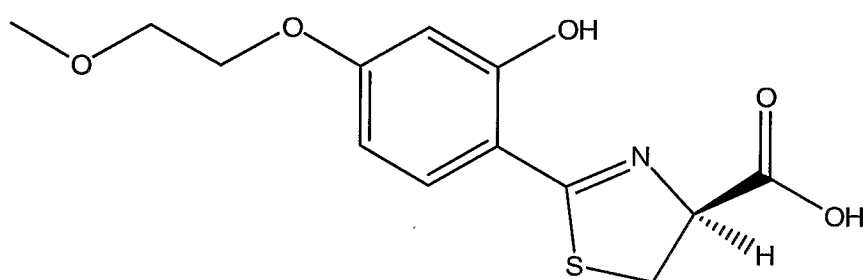


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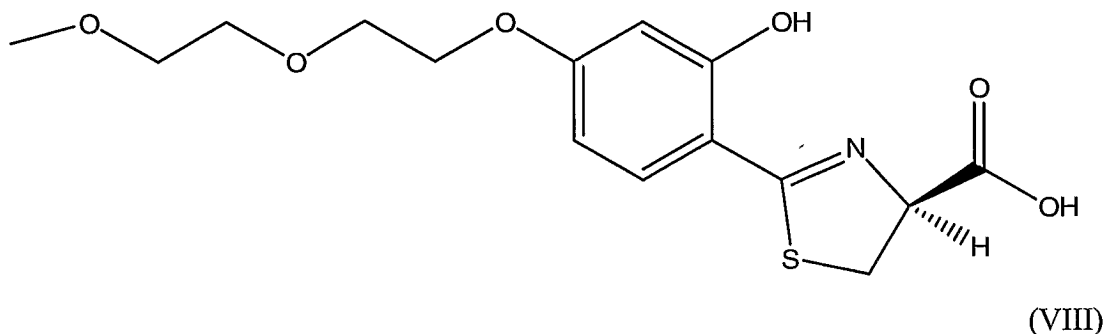




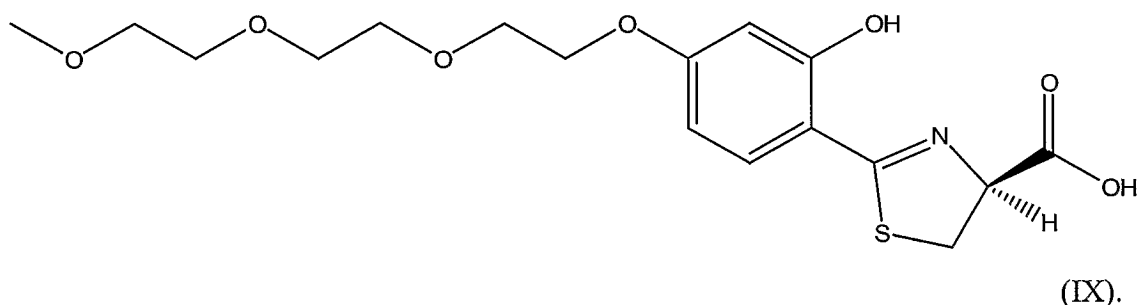
(VI)



(VII)



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The invention also includes enantiomers and mixtures of enantiomers (e.g., racemic mixtures) of the compounds represented by Structural Formulas (I)-(IX), along with their salts (e.g., pharmaceutically acceptable salts), solvates and hydrates.

In addition to compounds represented by Structural Formulas (I)-(IX), compounds of the invention can exist in optically active forms that have the ability to rotate the plane of plane-polarized light. In describing an optically active compound, the prefixes D and L or R and S are used to denote the absolute configuration of the molecule about its chiral center(s). The prefixes d and l or (+) and (-) are employed to designate the sign of rotation of plane-polarized light by the compound, with (-) or l meaning that the compound is levorotatory. A compound prefixed with (+) or d is dextrorotatory. For a given chemical structure, these compounds, called stereoisomers, are identical except that one or more chiral carbons are non-superimposable mirror images of one another. A specific stereoisomer, which is an exact mirror image of another stereoisomer, can also be referred to as an enantiomer, and a mixture of such isomers is often called an enantiomeric mixture. A 50:50 mixture of enantiomers is referred to as a racemic mixture.

Many of the compounds described herein can have one or more chiral centers and therefore can exist in different enantiomeric forms. If desired, a chiral carbon can be designated with an asterisk (*). In the present application, the chiral carbon at the 4-position of the thiazoline or thiazolidine ring has been designated with an asterisk, because the configuration of this carbon is of particular interest. When bonds to chiral carbons are depicted as straight lines in the formulas of the invention, it is understood that both the (R) and (S) configurations of each chiral carbon, and hence both enantiomers and mixtures thereof, are embraced within the formula. As is used in the art, when it is desired to specify the absolute configuration about a chiral carbon, a bond to the chiral carbon can be depicted as a wedge (bonds to atoms above the plane) and another can be depicted as a series or wedge of short parallel lines (bonds to atoms below the plane). The Cahn-Ingold-Prelog system can be used to assign the (R) or (S) configuration to a chiral carbon. A chiral carbon at the 4-position of a thiazoline or thiazolidine ring preferably has an (S) configuration.

When compounds of the present invention contain one chiral center, compounds not prepared by an asymmetric synthesis exist in two enantiomeric forms and the present invention includes either or both enantiomers and mixtures of enantiomers, such as the specific 50:50 mixture referred to as a racemic mixture. The enantiomers can be resolved by methods known to those skilled in the art, for example, by formation of diastereoisomeric salts that may be separated, for example, by crystallization (See, CRC Handbook of Optical Resolutions via Diastereomeric Salt Formation by David Kozma (CRC Press, 2001)); formation of diastereoisomeric derivatives or complexes that may be separated, for example, by crystallization, gas-liquid or liquid chromatography; selective reaction of one enantiomer with an enantiomer-specific reagent, for example, enzymatic esterification; or gas-liquid or liquid chromatography in a chiral environment, for example, on a chiral support (e.g., silica with a bound chiral ligand) or in the presence of a chiral solvent. It will be appreciated that where the desired enantiomer is converted into another chemical entity by one of the separation procedures described above, a further step is required to liberate the desired enantiomeric form. Alternatively, specific enantiomers may be synthesized by asymmetric synthesis

using optically active reagents, substrates, catalysts or solvents, or by converting one enantiomer into the other by asymmetric transformation.

Designation of a specific absolute configuration at a chiral carbon of the compounds of the invention is understood to mean that the designated enantiomeric form of the compounds is in enantiomeric excess (ee) or, in other words, is substantially free from the other enantiomer. For example, the "R" forms of the compounds are substantially free from the "S" forms of the compounds and are, thus, in enantiomeric excess of the "S" forms. Conversely, "S" forms of the compounds are substantially free of "R" forms of the compounds and are, thus, in enantiomeric excess of the "R" forms. Enantiomeric excess, as used herein, is the presence of a particular enantiomer at greater than 50% in an enantiomeric mixture. For example, when a mixture contains 80% of a first enantiomer and 20% of a second enantiomer, the enantiomeric excess of the first enantiomer is 60%. In the present invention, the enantiomeric excess can be about 20% or more, particularly about 40% or more, more particularly about 60% or more, such as about 70% or more, for example about 80% or more, such as about 90% or more. In a particular embodiment when a specific absolute configuration is designated, the enantiomeric excess of depicted compounds is at least about 90%. In a more particular embodiment, the enantiomeric excess of the compounds is at least about 95%, such as at least about 97.5%, for example, at least about 99% enantiomeric excess. When a compound of the present invention has two or more chiral carbons (e.g., compounds of Structural Formula (II) where R₆ and R₇ are not the same), it can have more than two optical isomers and can exist in diastereomeric forms. For example, when there are two chiral carbons, the compound can have up to 4 optical isomers and 2 pairs of enantiomers ((S,S)/(R,R) and (R,S)/(S,R)). The pairs of enantiomers (e.g., (S,S)/(R,R)) are mirror image stereoisomers of one another. The stereoisomers which are not mirror-images (e.g., (S,S) and (R,S)) are diastereomers. The diastereomeric pairs may be separated by methods known to those skilled in the art, for example, chromatography or crystallization and the individual enantiomers within each pair may be separated as described above. The present invention includes each diastereomer of such compounds and mixtures thereof.

An alkyl group is a saturated hydrocarbon in a molecule that is bonded to one other group in the molecule through a single covalent bond from one of its carbon atoms. Alkyl groups can be cyclic or acyclic, branched or unbranched (straight chained) and substituted or unsubstituted when straight chained or
5 branched. An alkyl group typically has from 1 to about 12 carbon atoms, for example, one to about six carbon atoms or one to about four carbon atoms. Lower alkyl groups have one to four carbon atoms and include methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *sec*-butyl and *tert*-butyl. When cyclic, an alkyl group typically contains from about 3 to about 10 carbons, for example, from about 3 to about 8
10 carbon atoms, e.g., a cyclopropyl group, a cyclobutyl group, a cyclopentyl group, a cyclohexyl group, a cycloheptyl group or a cyclooctyl group.

Acyl groups are represented by the formula $-C(O)R$, where R is an alkyl group. Acyl groups can be hydrolyzed or cleaved from a compound by enzymes, acids, or bases. One or more of the hydrogen atoms of an acyl group can be
15 substituted, as described below. Typically, an acyl group is removed before a compound of the present invention binds to a metal ion such as iron(III).

Suitable substituents for alkyl and acyl groups include $-OH$, $-O(R'')$, $-COOH$, $=O$, $-NH_2$, $-NH(R'')$, $-N(R'')_2$, $-COO(R'')$, $-CONH_2$, $-CONH(R'')$, $-CON(R'')_2$, and guanidine. Each R'' is independently an alkyl group or an aryl group. These groups
20 can additionally be substituted by an aryl group (e.g., an alkyl group can be substituted with an aromatic group to form an arylalkyl group). A substituted alkyl or acyl group can have more than one substituent.

Aryl groups include carbocyclic aromatic groups such as phenyl, *p*-tolyl, 1-naphthyl, 2-naphthyl, 1-anthracyl and 2-anthracyl. Aryl groups also include
25 heteroaromatic groups such as *N*-imidazolyl, 2-imidazolyl, 2-thienyl, 3-thienyl, 2-furanyl, 3-furanyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, 2-pyrimidyl, 4-pyrimidyl, 2-pyranyl, 3-pyranyl, 3-pyrazolyl, 4-pyrazolyl, 5-pyrazolyl, 2-pyrazinyl, 2-thiazolyl, 4-thiazolyl, 5-thiazolyl, 2-oxazolyl, 4-oxazolyl and 5-oxazolyl.

Aryl groups also include fused polycyclic aromatic ring systems in which a
30 carbocyclic, alicyclic, or aromatic ring or heteroaryl ring is fused to one or more other heteroaryl or aryl rings. Examples include 2-benzothienyl, 3-benzothienyl, 2-benzofuranyl, 3-benzofuranyl, 2-indolyl, 3-indolyl, 2-quinoliny, 3-quinoliny, 2-

benzothiazolyl, 2-benzoxazolyl, 2-benzimidazolyl, 1-isoquinoliny, 3-isoquinoliny, 1-isoindolyl and 3-isoindolyl.

Also included in the present invention are salts and pharmaceutically acceptable salts of the compounds described herein. Compounds disclosed herein
5 that possess a sufficiently acidic functional group, a sufficiently basic functional group or both, can react with a number of organic or inorganic bases, and inorganic and organic acids, to form salts.

Acidic groups can form salts with one or more of the metals listed above, along with alkali and alkaline earth metals (e.g, sodium, potassium, magnesium,
10 calcium). In addition, acidic groups can form salts with amines. Compounds of the invention can be supplied as a transition, lanthanide, actinide or main group metal salt. For example, the salt can be an iron (iron(II) or iron(III)) salt of a compound. As a transition, lanthanide, actinide or main group metal salt, compounds of the invention tend to form a complex with the metal. For example, if a compound of the
15 invention is tridentate and the metal it forms a salt with has six coordinate sites, then a 2 to 1 compound to metal complex is formed. The ratio of compound to metal will vary according to the denticity of the metal and the number of coordination sites on the metal (preferably each coordination site is filled by a compound of the invention, although a coordination site can be filled with other anions such as hydroxide, halide
20 or a carboxylate). Alternatively, the compound can be a substantially metal-free (e.g. iron-free) salt. Metal-free salts are not typically intended to encompass alkali and alkali earth metal salts. Metal-free salts are advantageously administered to a subject suffering from, for example, a metal overload condition or to an individual suffering from toxic metal exposure or from focal concentrations of metals causing untoward
25 effects

Acids commonly employed to form acid addition salts from compounds with basic groups are inorganic acids such as hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, phosphoric acid, and the like, and organic acids such as *p*-toluenesulfonic acid, methanesulfonic acid, oxalic acid, *p*-bromophenyl-sulfonic
30 acid, carbonic acid, succinic acid, citric acid, benzoic acid, acetic acid, and the like. Examples of such salts include the hydroxide, sulfate, pyrosulfate, bisulfate, sulfite, bisulfite, phosphate, monohydrogenphosphate, dihydrogenphosphate,

metaphosphate, pyrophosphate, chloride, bromide, iodide, acetate, propionate, decanoate, caprylate, acrylate, formate, isobutyrate, caproate, heptanoate, propiolate, oxalate, malonate, succinate, suberate, sebacate, fumarate, maleate, butyne-1,4-dioate, hexyne-1,6-dioate, benzoate, chlorobenzoate, methylbenzoate, 5 dinitrobenzoate, hydroxybenzoate, methoxybenzoate, phthalate, sulfonate, xylenesulfonate, phenylacetate, phenylpropionate, phenylbutyrate, citrate, lactate, gamma-hydroxybutyrate, glycolate, tartrate, methanesulfonate, propanesulfonate, naphthalene-1-sulfonate, naphthalene-2-sulfonate, mandelate, and the like.

The compounds disclosed herein can be prepared in the form of their hydrates, 10 such as hemihydrate, monohydrate, dihydrate, trihydrate, tetrahydrate and the like and as solvates.

Subjects suffering from a pathological condition responsive to chelation or sequestration of a trivalent metal can be treated with a therapeutically or prophylactically effective amount of a compound or pharmaceutical compound of 15 the invention. One particular type of pathological condition that is responsive to chelation of a trivalent metal is a trivalent metal overload condition (e.g., an iron overload condition, an aluminum overload condition, a chromium overload condition). Another type of pathological condition that is responsive to metal chelation or sequestration is when the amount of free trivalent metal is elevated 20 (e.g., in the serum or in a cell), such as when there is insufficient storage capacity for trivalent metals or an abnormality in the metal storage system that leads to metal release.

Iron overload conditions or diseases can be characterized by global iron overload or focal iron overload. Global iron overload conditions generally involve 25 an excess of iron in multiple tissues or excess iron located throughout an organism. Global iron overload conditions can result from excess uptake of iron by a subject, excess storage and/or retention of iron, from, for example, dietary iron or blood transfusions. One global iron overload condition is primary hemochromatosis, which is typically a genetic disorder. A second global iron overload condition is 30 secondary hemochromatosis, which is typically the result of receiving multiple (chronic) blood transfusions. Blood transfusions are often required for subjects suffering from thalassemia or sickle cell anemia. A type of dietary iron overload is

referred to as Bantu siderosis, which is associated with the ingestion of home-brewed beer with high iron content.

In focal iron overload conditions, the excess iron is limited to one or a few cell types or tissues or a particular organ. Alternatively, symptoms associated with the excess iron are limited to a discrete organ, such as the heart, lungs, liver, pancreas, kidneys or brain. It is believed that focal iron overload can lead to neurological or neurodegenerative disorders such as Parkinson's disease, Alzheimer's disease, Huntington's disease, neuroferritinopathy, amyotrophic lateral sclerosis and multiple sclerosis.

Pathological conditions that benefit from metal chelation or sequestration are often associated with deposition of the metal in the tissues of a subject. Deposition can occur globally or focally, as described above.

A subject in need of oxidative stress reduction can have one or more of the following conditions: decreased levels of reducing agents, increased levels of reactive oxygen species, mutations in or decreased levels of antioxidant enzymes (e.g., Cu/Zn superoxide dismutase, Mn superoxide dismutase, glutathione reductase, glutathione peroxidase, thioredoxin, thioredoxin peroxidase, DT-diaphorase), mutations in or decreased levels of metal-binding proteins (e.g., transferrin, ferritin, ceruloplasmin, albumin, metallothionein), mutated or overactive enzymes capable of producing superoxide (e.g., nitric oxide synthase, NADPH oxidases, xanthine oxidase, NADH oxidase, aldehyde oxidase, dihydroorotate dehydrogenase, cytochrome c oxidase), and radiation injury. Increased or decreased levels of reducing agents, reactive oxygen species, and proteins are determined relative to the amount of such substances typically found in healthy persons.

A subject in need of oxidation stress reduction can be suffering from an ischemic episode. Ischemic episodes can occur when there is mechanical obstruction of the blood supply, such as from arterial narrowing or disruption. Myocardial ischemia, which can give rise to angina pectoris and myocardial infarctions, results from inadequate circulation of blood to the myocardium, usually due to coronary artery disease. Ischemic episodes in the brain that resolve within 24 hours are referred to as transient ischemic attacks. A longer-lasting ischemic episode, a stroke, involves irreversible brain damage, where the type and severity of symptoms depend

on the location and extent of brain tissue whose access to blood circulation has been compromised. A subject at risk of suffering from an ischemic episode typically suffers from atherosclerosis, other disorders of the blood vessels, increased tendency of blood to clot, or heart disease. The compounds of this invention can be used to
5 treat these disorders.

A subject in need of oxidation stress reduction can be suffering from inflammation. Inflammation is a fundamental pathologic process consisting of a complex of cytologic and chemical reactions that occur in blood vessels and adjacent tissues in response to an injury or abnormal stimulation caused by a
10 physical, chemical, or biologic agent. Inflammatory disorders are characterized inflammation that lasts for an extended period (i.e., chronic inflammation) or that damages tissue. Such inflammatory disorders can affect a wide variety of tissues, such as respiratory tract, joints, bowels, and soft tissue. The compounds of this invention can be used to treat these disorders.

15 Although not bound by theory, it is believed that the compounds of the invention derive their ability to reduce oxidative stress through various mechanisms. In one mechanism, the compound binds to a metal, particularly a redox-active metal (e.g., iron), and fills all of the coordination sites of the metal. When all of the metal coordination sites are filled, it is believed that oxidation and/or reducing agents have
20 a diminished ability to interact with the metal and cause redox cycling. In another mechanism, the compound stabilizes the metal in a particular oxidation state, such that it is less likely to undergo redox cycling. In yet another mechanism, the compound itself has antioxidant activity (e.g., free radical scavenging, scavenging of reactive oxygen or nitrogen species). Desferrithiocin and its derivatives and
25 analogues are known to have intrinsic antioxidant activity, as described in U.S. Application Publication No. 2004/0044220, published March 4, 2004, and U.S. Application Publication No. 2004/0132789, published July 8, 2004 and PCT Application No. WO2004/017959, published March 4, 2004, US Application Publication No. 2003/0236417, published December 25, 2003, and US Patent Nos:
30 6,083,966, 6,559,315, 6,525,080, 6,521,652 the contents of each of which are incorporated herein by reference.

Imaging or examining one or more organs, tissues, tumors or a combination thereof can be conducted after a metal salt of a compound of the invention is administered to a subject. The methods of imaging and examining are intended to encompass various instrumental techniques used for diagnosis, such as x-ray methods (including CT scans and conventional x-ray images), magnetic imaging (magnetic resonance imaging, electron paramagnetic resonance imaging) and radiochemical methods. Typically, the metal salts used in imaging or examining serve as a contrast agent. Therefore in one embodiment the metal complexes or metal salts of compounds of the present invention can be used as contrast agents for example in imaging or examining one or more organs, for example, the gastrointestinal tract.

Metals that can serve as contrast agents include gadolinium, iron, manganese, chromium, dysprosium, technetium, scandium, barium, aluminum and holmium, preferably as trications. Radioactive metal salts can be made from isotopes including ^{241}Am , ^{51}Cr , ^{60}Co , ^{57}Co , ^{58}Co , ^{64}Cu , ^{153}Gd , ^{67}Ga , ^{198}Au , $^{113\text{m}}\text{In}$, ^{111}In , ^{59}Fe , ^{55}Fe , ^{197}Hg , ^{203}Hg , $^{99\text{m}}\text{Tc}$, ^{201}Tl and ^{169}Yb , again preferably when the metal is present as a trication.

Neoplastic disease is characterized by an abnormal tissue that grows by cellular proliferation more rapidly than normal tissue. The abnormal tissue continues to grow after the stimuli that initiated the new growth cease. Neoplasms show a partial or complete lack of structural organization and functional coordination with the normal tissue, and usually form a distinct mass of tissue that may be either benign or malignant. Neoplasms can occur, for example, in a wide variety of tissues including brain, skin, mouth, nose, esophagus, lungs, stomach, pancreas, liver, bladder, ovary, uterus, testicles, colon, and bone, as well as the immune system (lymph nodes) and endocrine system (thyroid gland, parathyroid glands, adrenal gland, thymus, pituitary gland, pineal gland). The compounds of this invention can be used to treat these disorders.

Examples of tumors or cancers that can be treated by the invention include, but are not limited to, leukemia, Hodgkin's disease, non-Hodgkin's lymphomas, multiple myeloma, macroglobulinemia, polycythemia vera, lung tumors, head and neck tumors, brain tumors (neuroblastoma), endometrial tumors, ovarian tumors, cervical tumors, breast tumors, choriocarcinoma, testicular tumors, prostate tumor,

Wilms' tumor, thyroid tumors, adrenal tumors, stomach tumor, pancreatic tumors, colonic tumors, carcinoids, insulinoma, bone tumors (osteogenic sarcoma), miscellaneous sarcomas and skin cancer (melanoma).

5 A preneoplastic condition precedes the formation of a benign or malignant neoplasm. A precancerous lesion typically forms before a malignant neoplasm. Preneoplasms include photodermatitis, x-ray dermatitis, tar dermatitis, arsenic dermatitis, lupus dermatitis, senile keratosis, Paget disease, condylomata, burn scar, syphilitic scar, fistula scar, ulcer cruris scar, chronic ulcer, varicose ulcer, bone
10 fistula, rectal fistula, Barrett esophagus, gastric ulcer, gastritis, cholelithiasis, kraurosis vulvae, nevus pigmentosus, Bowen dermatosis, xeroderma pigmentosum, erythroplasia, leukoplakia, Paget disease of bone, exostoses, ecchondroma, osteitis fibrosa, leontiasis ossea, neurofibromatosis, polyposis, hydatidiform mole, adenomatous hyperplasia, and struma nodosa. The compounds of this invention can be used to treat these disorders.

15 A "subject" is typically a human, but can also be an animal in need of treatment, e.g., companion animals (e.g., dogs, cats, and the like), farm animals (e.g., cows, pigs, horses, sheep, goats and the like) and laboratory animals (e.g., rats, mice, guinea pigs, non-human primates and the like).

The compounds and pharmaceutical compositions of the present invention can
20 be administered by an appropriate route. Suitable routes of administration include, but are not limited to, orally, intraperitoneally, subcutaneously, intramuscularly, transdermally, rectally, sublingually, intravenously, buccally or via inhalation. Preferably, compounds and pharmaceutical compositions of the invention are administered orally.

25 The pharmaceutical compositions of the invention preferably contain a pharmaceutically acceptable carrier or diluent suitable for rendering the compound or mixture administrable orally, parenterally, intravenously, intradermally, intramuscularly or subcutaneously, rectally, via inhalation or via buccal administration, or transdermally. The active ingredients may be admixed or
30 compounded with a conventional, pharmaceutically acceptable carrier or diluent. It will be understood by those skilled in the art that a mode of administration, vehicle or carrier conventionally employed and which is inert with respect to the active agent

may be utilized for preparing and administering the pharmaceutical compositions of the present invention. Illustrative of such methods, vehicles and carriers are those described, for example, in Remington's Pharmaceutical Sciences, 18th ed. (1990), the disclosure of which is incorporated herein by reference.

5 The formulations of the present invention for use in a subject comprise the agent, together with one or more acceptable carriers or diluents therefor and optionally other therapeutic ingredients. The carriers or diluents must be "acceptable" in the sense of being compatible with the other ingredients of the formulation and not deleterious to the recipient thereof. The formulations can conveniently be presented
10 in unit dosage form and can be prepared by any of the methods well known in the art of pharmacy. All methods include the step of bringing into association the agent with the carrier or diluent which constitutes one or more accessory ingredients. In general, the formulations are prepared by uniformly and intimately bringing into association the agent with the carriers and then, if necessary, dividing the product into unit
15 dosages thereof.

Forms suitable for oral administration include tablets, troches, capsules, elixirs, suspensions, syrups, wafers, chewing gum or the like prepared by art recognized procedures. The amount of active compound in such therapeutically useful compositions or preparations is such that a suitable dosage will be obtained.

20 A syrup formulation will generally consist of a suspension or solution of the compound or salt in a liquid carrier, for example, ethanol, glycerine or water, with a flavoring or coloring agent. Where the composition is in the form of a tablet, one or more pharmaceutical carriers routinely used for preparing solid formulations can be employed. Examples of such carriers include magnesium stearate, starch, lactose and
25 sucrose. Where the composition is in the form of a capsule, the use of routine encapsulation is generally suitable, for example, using the aforementioned carriers in a hard gelatin capsule shell. Where the composition is in the form of a soft gelatin shell capsule, pharmaceutical carriers routinely used for preparing dispersions or suspensions can be considered, for example, aqueous gums, celluloses, silicates or
30 oils, and are incorporated in a soft gelatin capsule shell.

Formulations suitable for parenteral administration conveniently include sterile aqueous preparations of the agents that are preferably isotonic with the blood

of the recipient. Suitable carrier solutions include phosphate buffered saline, saline, water, lactated ringers or dextrose (5% in water). Such formulations can be conveniently prepared by admixing the agent with water to produce a solution or suspension, which is filled into a sterile container and sealed against bacterial
5 contamination. Preferably, sterile materials are used under aseptic manufacturing conditions to avoid the need for terminal sterilization.

Such formulations can optionally contain one or more additional ingredients, which can include preservatives such as methyl hydroxybenzoate, chlorocresol, metacresol, phenol and benzalkonium chloride. Such materials are of special value
10 when the formulations are presented in multidose containers.

Buffers can also be included to provide a suitable pH value for the formulation. Suitable buffer materials include sodium phosphate and acetate. Sodium chloride or glycerin can be used to render a formulation isotonic with the blood.

If desired, a formulation can be filled into containers under an inert
15 atmosphere such as nitrogen and can be conveniently presented in unit dose or multi-dose form, for example, in a sealed ampoule.

Those skilled in the art will be aware that the amounts of the various components of the compositions of the invention to be administered in accordance with the method of the invention to a subject will depend upon those factors noted
20 above.

A typical suppository formulation includes the compound or a pharmaceutically acceptable salt thereof which is active when administered in this way, with a binding and/or lubricating agent, for example, polymeric glycols, gelatins, cocoa-butter or other low melting vegetable waxes or fats.

25 Typical transdermal formulations include a conventional aqueous or non-aqueous vehicle, for example, a cream, ointment, lotion or paste or are in the form of a medicated plastic, patch or membrane.

Typical compositions for inhalation are in the form of a solution, suspension or emulsion that can be administered in the form of an aerosol using a conventional
30 propellant such as dichlorodifluoromethane or trichlorofluoromethane.

The therapeutically effective amount of a compound or pharmaceutical composition of the invention depends, in each case, upon several factors, e.g., the

- health, age, gender, size and condition of the subject to be treated, the intended mode of administration, and the capacity of the subject to incorporate the intended dosage form, among others. A therapeutically effective amount of an active agent is an amount sufficient to have the desired effect for the condition being treated. For
- 5 example, in a method of treating of a neoplastic or a preneoplastic condition, the desired effect is partial or total inhibition, delay or prevention of the progression of cancer or the tumor including cancer metastasis; inhibition, delay or prevention of the recurrence of cancer or the tumor including cancer metastasis; or the prevention of the onset or development of cancer or a tumor (chemoprevention) in a mammal, for
- 10 example a human. In a method of treating a subject with a condition treatable by chelating or sequestering a metal ion, a therapeutically effective amount of an active agent is, for example, an amount sufficient to reduce the burden of the metal in the subject, reduce the symptoms associated with the metal ion or prevent, inhibit or delay the onset and/or severity of symptoms associated with the presence of the metal.
- 15 In a method of reducing oxidative stress in a subject in need of treatment thereof, a therapeutically effective amount of an active agent is, for example, an amount sufficient to reduce symptoms associated with oxidative stress or prevent, inhibit or delay the onset and/or severity of symptoms associated with oxidative stress.

A typical total daily dose of a compound of the invention to be administered

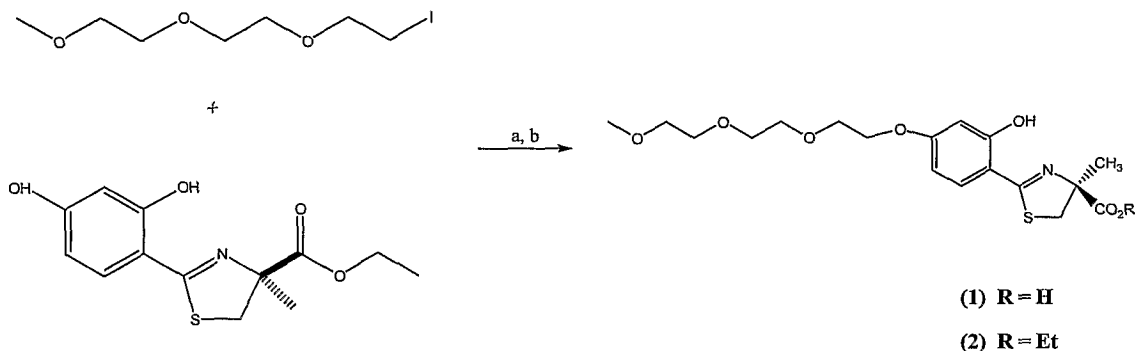
20 to a subject (assuming an average 70 kg subject) is from approximately 5 mg to approximately 10,000 mg, (for example 0.07 mg/kg to 143 mg/kg) and preferably from approximately 50 mg to approximately 5,000 mg approximately 100 mg to approximately 2,000 mg approximately 300 mg to approximately 1,000 mg. For iron overload therapy, a daily dose of a compound of the invention should remove a

25 minimum of from approximately 0.25 to approximately 0.40 mg of iron per kilogram of body mass per day. The dosage can be administered orally in several, for example, one, two, three, four, six, eight, twelve, or more, individual doses.

The present invention will now be illustrated by the following Examples, which are not intended to be limiting in any way.

EXEMPLIFICATION

Scheme 1. (*S*)-4,5-dihydro-2-[2-hydroxy-4-[2-[2-(2-methoxyethoxy)ethoxy]ethoxy]phenyl]-4-methyl-4-thiazolecarboxylic acid (**1**) and Ethyl Ester (**2**). Reagents: (a) 50% molar excess NaOEtH, EtOH, 33%; (b) 50% NaOH, CH₃OH, 91%.

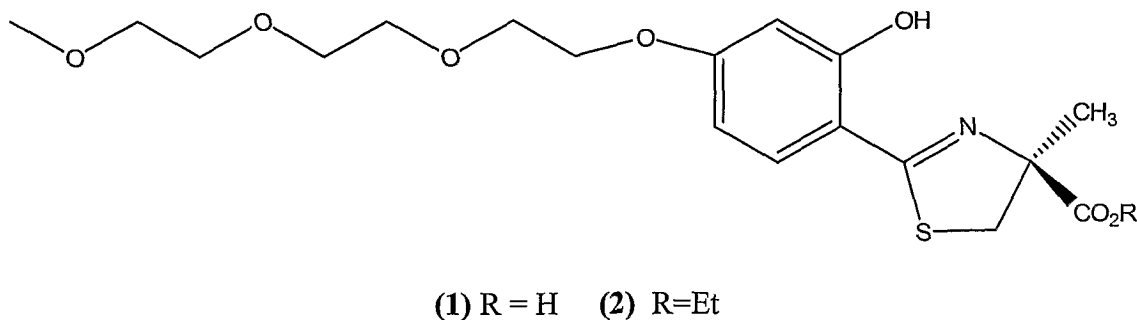


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Example 1

Synthesis of (*S*)-4,5-Dihydro-2-[2-hydroxy-4-[2-[2-(2-methoxyethoxy)ethoxy]ethoxy]ethoxy]phenyl]-4-methyl-4-thiazolecarboxylic Acid (**1**) and Ethyl Ester (**2**)

15



20

The ethyl ester of (*S*)-2-(2,4-dihydroxyphenyl)-4,5-dihydro-4-methyl-4-thiazolecarboxylic acid ((*S*)-4'-(HO)-DADFT) was treated with a 50% excess of

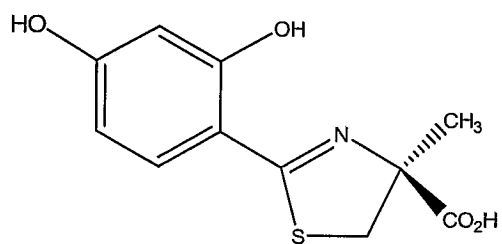
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- NaOEt and heated with 3,6,9-trioxa-1-iododecane (1.3 equiv) in EtOH which effected regiospecific alkylation at the 4'hydroxyl, resulting in adduct **(2)** (Scheme 1). Saponification of ester **(2)** with NaOH in aqueous methanol at room temperature gave (*S*)-DADFT analogue **(1)** (referred to herein as (*S*)-4'-(OH)-DADFT-PE). ¹H
- 5 NMR: (D₂O) δ 1.76 (s, 3 H), 3.35 (s, 3 H), 3.54-3.61 (m, 3H), 3.64-3.72 (m, 4 H), 3.74-3.78 (m, 2 H), 3.90-3.94 (m, 2 H), 3.96 (d, 1 H, *J*=12.0), 4.25-4.29 (m, 2 H), 6.53 (d, 1 H *J*=2.4), 6.4 (dd, 1 H, *J*=9.0, 2.2), 7.61 (d, 1 H, *J*=9.2).

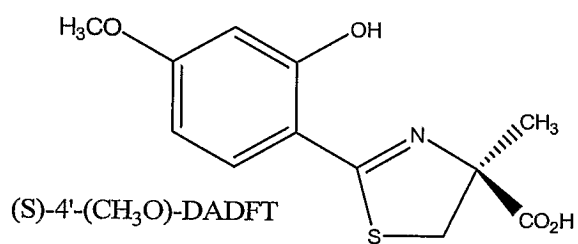
Example 2

- 10 Iron Clearing Efficiency of Iron Chelators in a Non-Iron Overloaded, Bile Duct Cannulated Rat Model

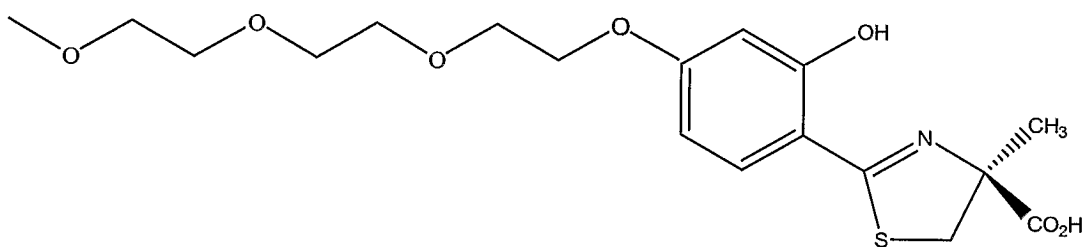
Studies were performed in the non-iron overloaded, bile duct cannulated rodent model with the compounds shown below.



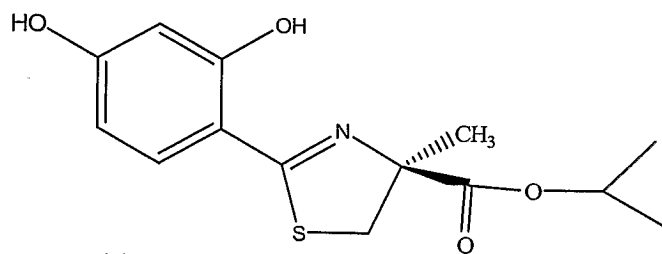
(S)-4'-(HO)-DADFT



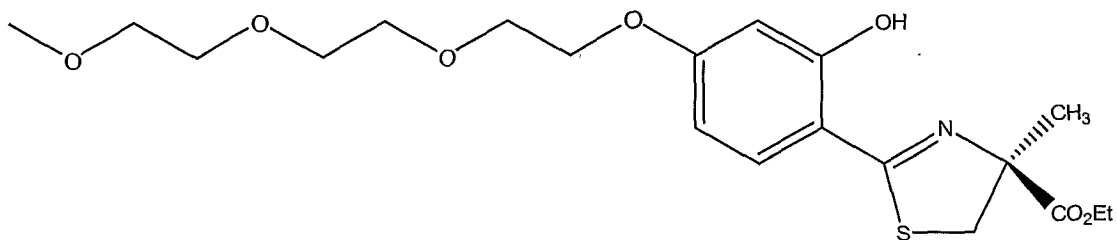
(S)-4'-(CH₃O)-DADFT



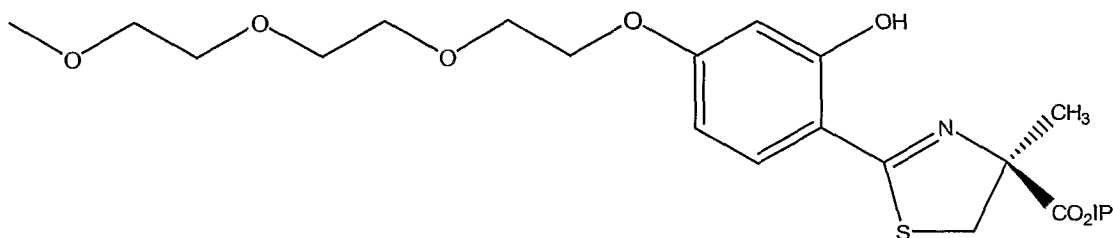
(S)-4'-(HO)-DADFT-PE



(S)-4'-(HO)-DADFT-iPrE



(S)-4'-(HO)-DADFT-PE EE



(S)-4'-(HO)-DADFT-PE iPrE

Briefly, male Sprague-Dawley rats averaging 450 g were housed in Nalgene plastic metabolic cages during the experimental period and given free access to water. The animals were anesthetized using sodium pentobarbital (55 mg/kg) administered intraperitoneally. The bile duct was cannulated using 22-gauge polyethylene tubing. The cannula was inserted into the duct about 1 cm from the duodenum and tied snugly in place. After threading through the shoulder, the cannula was passed from the rat to the swivel inside a metal torque-transmitting tether, which was attached to a rodent jacket around the animal's chest. The cannula was directed from the rat to a Gilson microfraction collector (Middleton, WI) by a fluid swivel mounted above the metabolic cage. Three hour bile samples were continuously collected for a minimum of 24 hours up to 48 hours. However, the efficiency calculations are based on the 24 hour iron excretion. The efficiency of each chelator was calculated on the basis of a 2:1 ligand-iron complex. The efficiencies in the rodent model were

calculated by subtracting the iron excretion of control animals from the iron excretion of treated animals. This number was then divided by the theoretical output; the result is expressed as a percentage (Bergeron, R.J., *et al.*, *J. Med Chem.* 42:95-108 (1999) the entire contents of which are incorporated herein by reference).

- 5 The urine sample was taken at 24 hours and handled as previously described in Bergeron, R.J., *et al.*, *J. Med. Chem.* 34:2072-2078 (1991), the entire contents of which are incorporated herein by reference.

The results of the evaluations are presented in Table 1 and FIG. 1.

10 Table 1. Iron Clearing Efficiencies of Compounds Tested in the Bile Duct Cannulated Rat

Compound	Dose (mg/kg) (μ mol/kg)		Route	Vehicle	N	Efficiency (%)
(S)-4'-(HO)-DADFT	76	300	p.o. ⁺	dH ₂ O*	3	1.0 $\pm$ 0.4
(S)-4'-(HO)-DADFT	76	300	s.c. ⁺	dH ₂ O*	4	1.1 $\pm$ 0.6
(S)-4'-(HO)-DADFT-PE	120	300	p.o.	dH ₂ O	5	6.6 $\pm$ 1.9
(S)-4'-(HO)-DADFT-PE	120	300	s.c.	dH ₂ O	4	8.7 $\pm$ 2.8
(S)-4'-(HO)-DADFT-PE EE	128	300	s.c.	50%EtOH	3	11.4 $\pm$ 0.8
(S)-4'-(HO)-DADFT-PE iPrE	133	300	s.c.	50%EtOH	3	25.9 $\pm$ 9.2
(S)-4'-(HO)-DADFT-PE iPrE	133	300	p.o.	40%cremophor	3	9.2 $\pm$ 4.4
(S)-4'-(HO)-DADFT-iPrE	88	300	s.c.	50%EtOH/ dH ₂ O	3	1.98 $\pm$ 0.91
(S)-4'-(CH ₃ O)-DADFT	80	300	p.o.	40% Cremophor	4	6.6 $\pm$ 2.8

⁺ p.o. is by mouth, s.c is subcutaneously

*brought to neutral pH with NaOH solution

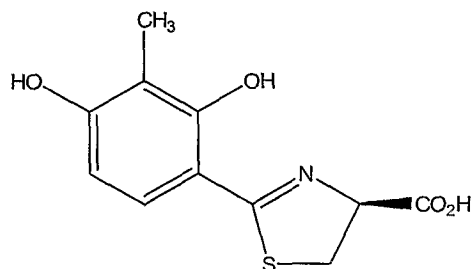
- A comparison of the iron clearing efficiency of the polyethers(S)-4-(OH)-DADFT-PE, (S)-4-(HO)-DADFT-PE EE and (S)-4-(HO)-DADFT-PE iPrE, with that of the parent drug (S)-4-(HO)-DADFT clearly shows the polyethers and their corresponding esters to work better than the parent drug. The most notable feature regarding the esters is the performance of the isopropylester iPrE vs the ethyl ester EE when both are administered subcutaneously. The iPrE is significantly more
- 15

efficient. It is important to note that all drugs are administered at the same micromole per kilogram dose.

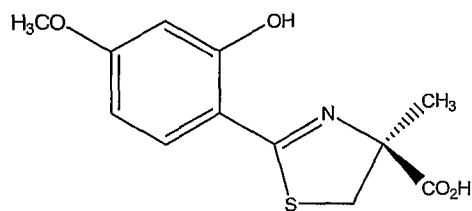
Example 3

5 Iron Chelators in a *Cebus apella* Monkey Model

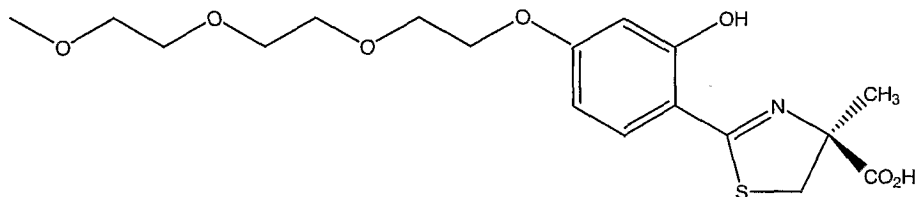
Studies were performed in the iron-overloaded monkey model with the compounds shown below:



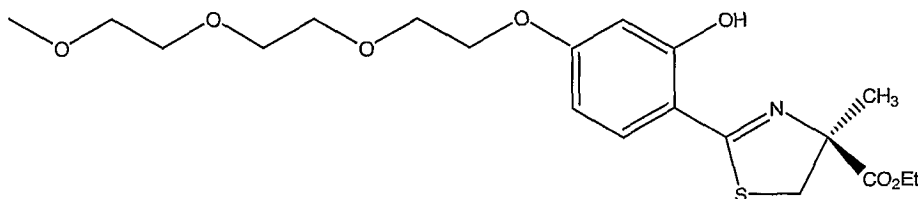
(S)-3'-(CH₃)-4'-(HO)-DADMDFT



(S)-4'-(CH₃O)-DADFT



(S)-4'-(HO)-DADFT-PE



(S)-4'-(HO)-DADFT-PE-EE

10

15

The protocol used can be found in Bergeron, R.J. *et al.*, "Methoxylation of Desazadesferrithiocin Analogues: Enhanced Iron Clearing Efficiency," *J. Med. Chem.* 46:1470-1477 (2003), the contents of which are incorporated herein by reference. Briefly, the monkeys were iron overloaded with iron dextran administered intravenously to result in an iron loading of about 500 mg per kg of

body weight. At least 20 half-lives, 60 days, elapsed before the animals were used in experiments evaluating iron chelators. The iron chelators were suspended in vehicle and administered either p.o. or s.c. as indicated in Table 2. Fecal and urine samples were collected at 24 hour intervals beginning 4 days prior to the

5 administration of an iron chelator and continued for 5 days after the chelator was administered. Iron concentrations in stool and urine were determined by flame atomic absorption spectroscopy. Iron chelator efficiency was calculated by dividing the net iron clearance [total iron excretion (stool plus urine) minus background] by the theoretical iron clearance and multiplying by 100. The

10 theoretical clearance of the iron chelator was generated on the basis of a 2:1 ligand/iron complex.

The results of the evaluations are presented in Table 2.

Table 2. Iron Clearing Efficiencies of Compounds Tested in *Cebus apella* primates

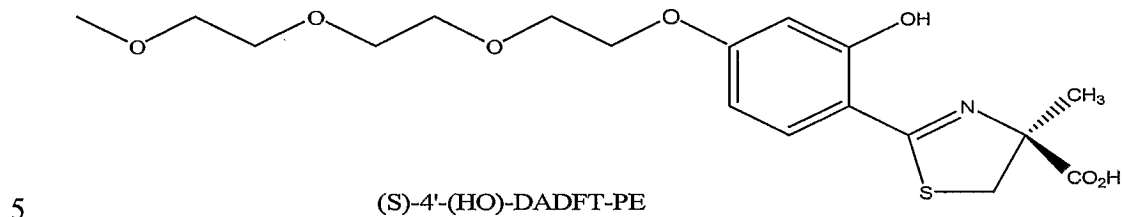
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Compound	Dose (mg/kg)	Route	Vehicle	<i>N</i>	Efficiency (%)
(<i>S</i>)-3'-(HO)-DADMDFT	39.1	p.o.	buffer	1	0.8
(<i>S</i>)-5'-(HO ₂ C)-DADMDFT	40.1	p.o.	buffer	1	0.7
(<i>S</i>)-3'-(CH ₃)-4'-(HO)- DADMDFT	38.0	p.o.	buffer	1	4.0
(<i>S</i>)-4'-(HO)-DADFT	38	p.o.	buffer	4	14± 3.3
(<i>S</i>)-4'-(HO)-DADFT	38	s.c.	water	4	16.6± 2.7
(<i>S</i>)-4'-(CH ₃ O)-DADFT	40.1	s.c.	buffer	4	33.7± 12.1
(<i>S</i>)-4'-(HO)-DADFT-PE	59.9	p.o.	dH ₂ O	4	25.4±7.4
(<i>S</i>)-4'-(HO)-DADFT-PE	59.9	s.c.	dH ₂ O	4	30.4±7.2
(<i>S</i>)-4'-(HO)-DADFT-PE-EE	64.1	s.c.	50%EtOH/ dH ₂ O	2	17.9±1.0

The data shows that the (*S*)-4'-(HO)-DADFT-PE given either p.o. or s.c. is at least as efficient as (*S*)-4'-(HO)-DADFT at clearing iron when given to primates.

Example 4

Tissue Distribution of a Polyether-Substituted (*S*)-4'-Hydroxydesazadesferrithiocin Analogue [(*S*)-4'-(HO)-DADFT-PE] upon Subcutaneous Administration to Rats



10 A measurement was made assessing (*S*)-4'-(HO)-DADFT-PE and (*S*)-4'-(HO)-DADFT tissue and plasma concentrations upon subcutaneous administration at times from 2-8 h post dosing. The rats were given this compound subcutaneously at 300 $\mu\text{mol/kg}$. The tissue and plasma level were obtained as described in Bergeron et al. *J. Med Chem.* 48:821-831 (2005), the entire contents of which are incorporated herein by reference. The results are shown in Table 3 and FIG. 2- FIG. 8.

Table 3. Tissue and Plasma Concentrations of (S)-4'-(HO)-DADFT-PE upon sc Administration vs. Concentrations of (S)-4'-(HO)-DADFT upon sc Administration of (S)-4'-(HO)-DADFT^a

Administered cmpd	Time (h)	(S)-4'-(HO)-DADFT-PE		(S)-4'-(HO)- DADFT	(S)-4'-(CH ₃ O)-DADFT	
		(S)-4'-(HO)- DADFT-PE (metabolite)	(S)-4'-(HO)- DADFT-PE	(S)-4'-(HO)- DADFT	(S)-4'-(CH ₃ O)- DADFT (metabolite)	(S)-4'-(CH ₃ O)- DADFT
Liver	2	2.8±0.1	131.6±32.2	48.3±20.3	43.5±4.5	111.0±14.0
	4	0.0±0.0	66.9±12.4	25.0±3.9	24.6±1.1	73.8±10.8
	6	0.0±0.0	34.3±1.5	19.3±2.0	26.9±5.1	80.9±10.7
	8	0.0±0.0	25.4±4.6	12.3±0.8	25.0±3.6	81.4±15.9
Kidney	2	0.0±0.0	41.1±3.0	97.1±5.0	49.9±21.3	97.5±24.5
	4	0.0±0.0	33.7±13.5	26.6±7.1	23.2±1.7	57.0±15.5
	6	0.0±0.0	13.7±2.1	13.0±8.0	21.7±6.2	67.2±9.9
	8	0.0±0.0	7.4±2.1	8.8±2.9	25.3±2.5	67.4±19.9
Heart	2	0.0±0.0	9.2±2.6	4.6±3	6.4±0.6	66.2±2.2
	4	0.0±0.0	5.3±1.4	0±0	<4.8	32.6±3.6
	6	0.0±0.0	0.0±0.0	0±0	<4.8	28.2±0.7
	8	0.0±0.0	2.0±1.7	0±0	<4.8	26.3±9.7
Pancreas	2	0.0±0.0	29.6±24.8	5.8±1.3	1.3±2.2	37.1±3.8
	4	0.0±0.0	12.6±5.2	0±0	0.0±0.0	19.4±1.4
	6	0.0±0.0	4.6±1.6	0±0	0.0±0.0	16.5±4.5
	8	0.0±0.0	2.7±2.6	0±0	0.0±0.0	9.7±0.7
Brain	2	0.0±0.0	0.5±0.9	0±0	<2.4	13.2±4.8
	4	0.0±0.0	0.0±0.0	0±0	<2.4	7.4±0.7
	6	0.0±0.0	0.0±0.0	0±0	<2.4	4.8±3.1
	8	0.0±0.0	0.0±0.0	0±0	<2.4	<1.6±0.0
Plasma	2	2.7±0.3	16.2±4.2	13.1±2.7	8.4±1.8	478.8±36.8
	4	2.2±0.6	5.3±2.6	2±0.4	4.2±1.0	236.3±42.0
	6	2.3±0.3	1.0±1.0	0±0	1.8±0.6	142.5±29.5
	8	1.2±0.1	0.0±0.0	0±0	0.0±0.0	79.6±13.3

^aReported as nmol/g wet wt (tissue) or μM (plasma), mean ± SD for 3 animals per time point.

5 Table 3 and FIG. 2- FIG. 8 show that there is very little metabolism of (S)-4-(HO)-DADFT-PE back to (S)-4-(HO)DADFT (indicated as (metabolite in Table 3 and FIG. 2- FIG. 5) occurring in any tissue. Most of the PE is found in the liver,

kidney and pancreas. It is important to note that there is much less of the PE in the kidney than there is of (S)-4-(HO)-DADFT at any time point, (FIG. 2).

Example 5

5 Uranium excretion in rats by iron chelators

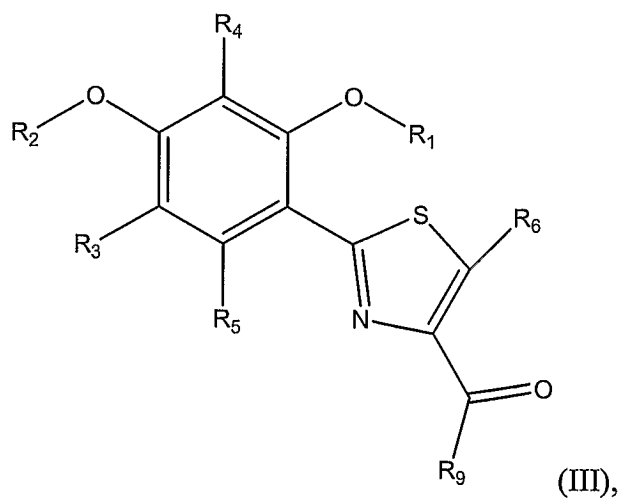
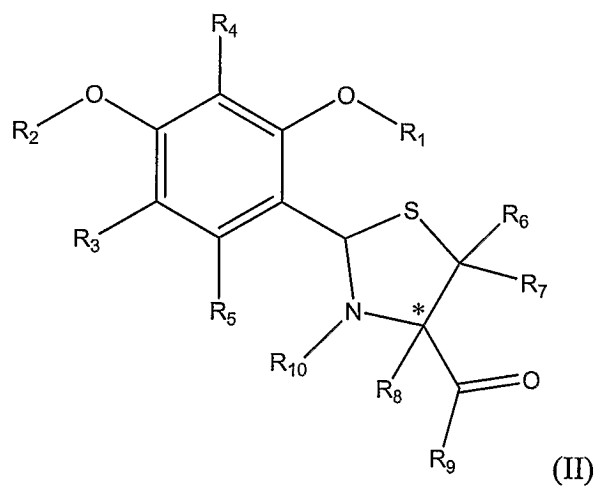
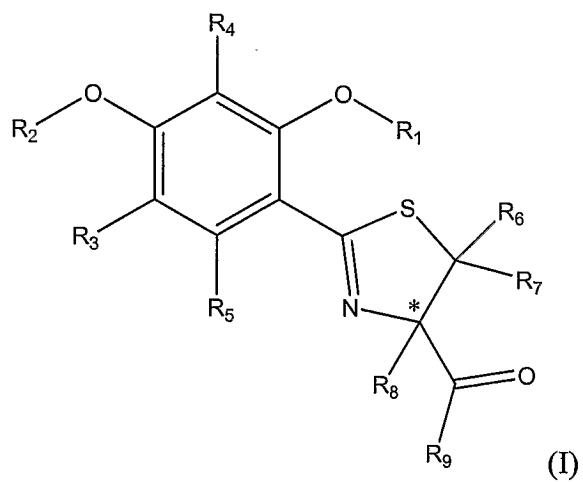
Male Sprague-Dawley rats averaging 450 g were anesthetized using sodium pentobarbital (55 mg/kg) administered intraperitoneally. The bile duct was cannulated using 22-gauge polyethylene tubing. The rats were given uranyl acetate subcutaneously at 5 mg/kg. Immediately thereafter, the rats were given the chelator
10 intraperitoneally at a dose of 300 μ mol/kg. 24-h urine and 24-h bile samples were collected, acidified with 2% concentrated nitric acid and assessed by Inductively Coupled Plasma Mass Spectrometry (ICP-MS) for their uranium content. The results are shown in FIG. 9.

While this invention has been particularly shown and described with
15 references to preferred embodiments thereof, it will be understood by those skilled in the art that various changes in form and details may be made therein without departing from the scope of the invention encompassed by the appended claims.

CLAIMS

What is claimed is:

1. A compound represented by a structural formula selected from Structural Formulas (I), (II), and (III):



wherein:

R₁ is -H or an acyl group;

R₂ is -[(CH₂)_n-O]_x-[(CH₂)_n-O]_y-R';

R₃, R₄ and R₅ are each independently -H, an alkyl group, or -OR₁₁;

5 R₆, R₇, and R₈ are each independently -H or an alkyl group;

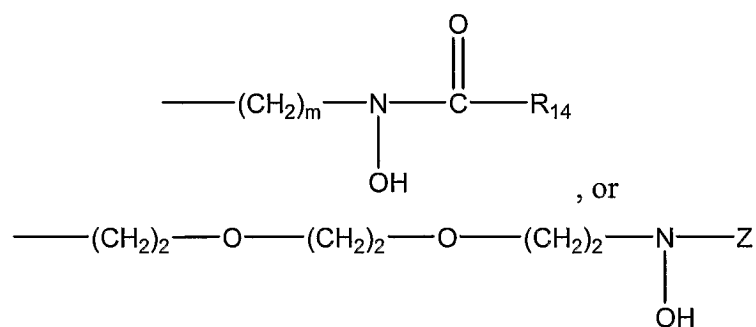
R₉ is -OR₁₂ or -N(OH)R₁₃;

R₁₀ is -H or an alkyl group;

R₁₁ is -H, an alkyl group or an acyl group;

R₁₂ is -H or an alkyl group;

10 R₁₃ is an alkyl group;



R₁₄ is an alkyl group;

R' is an alkyl group;

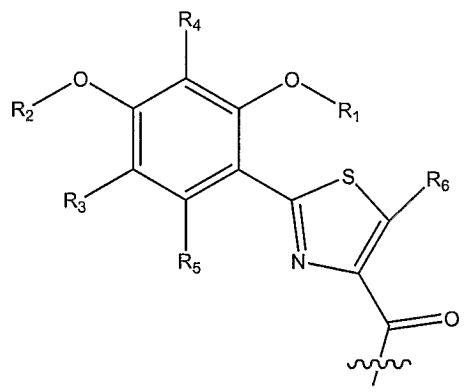
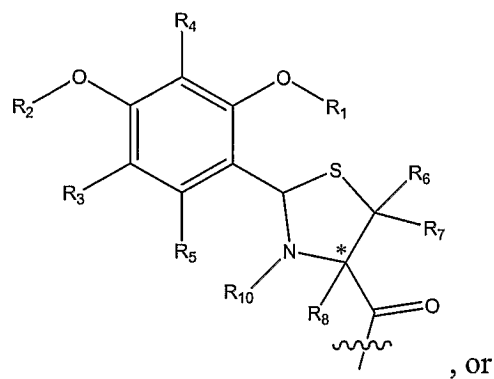
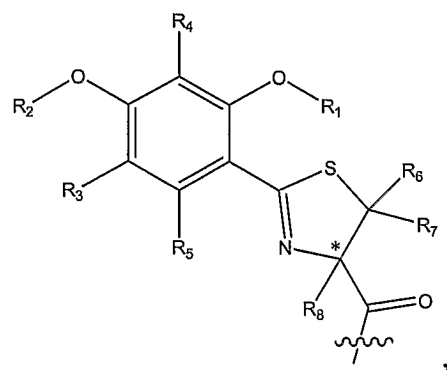
15 m is an integer from 1 to 8;

each n is independently an integer from 1 to 8;

x is an integer from 1 to 8;

y is an integer from 0 to 8;

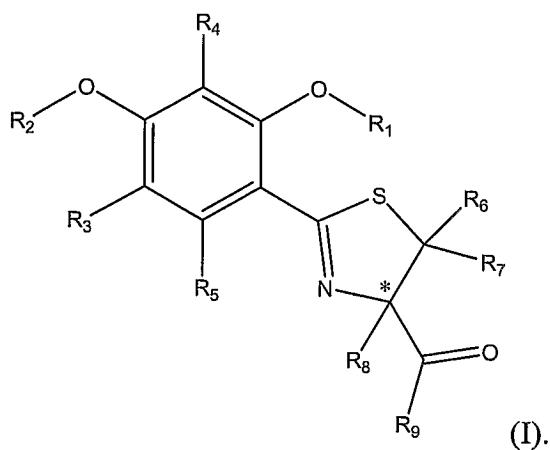
Z is -C(O)R₁₄,



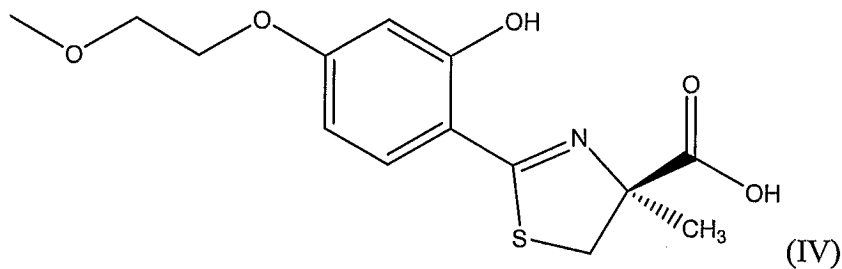
or a salt, solvate or hydrate thereof.

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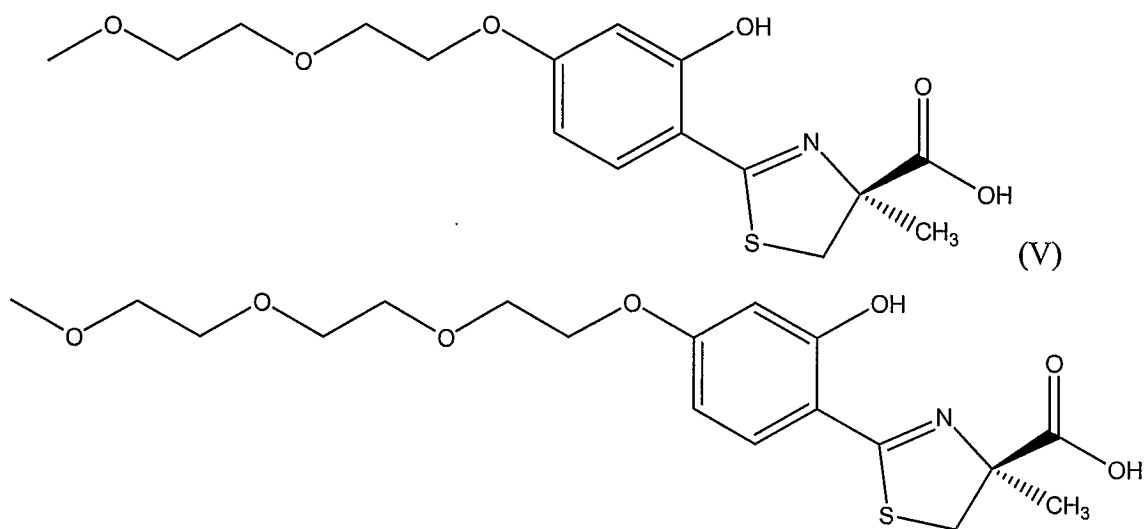
2. The compound of Claim 1, wherein the compound is represented by Structural Formula (I):



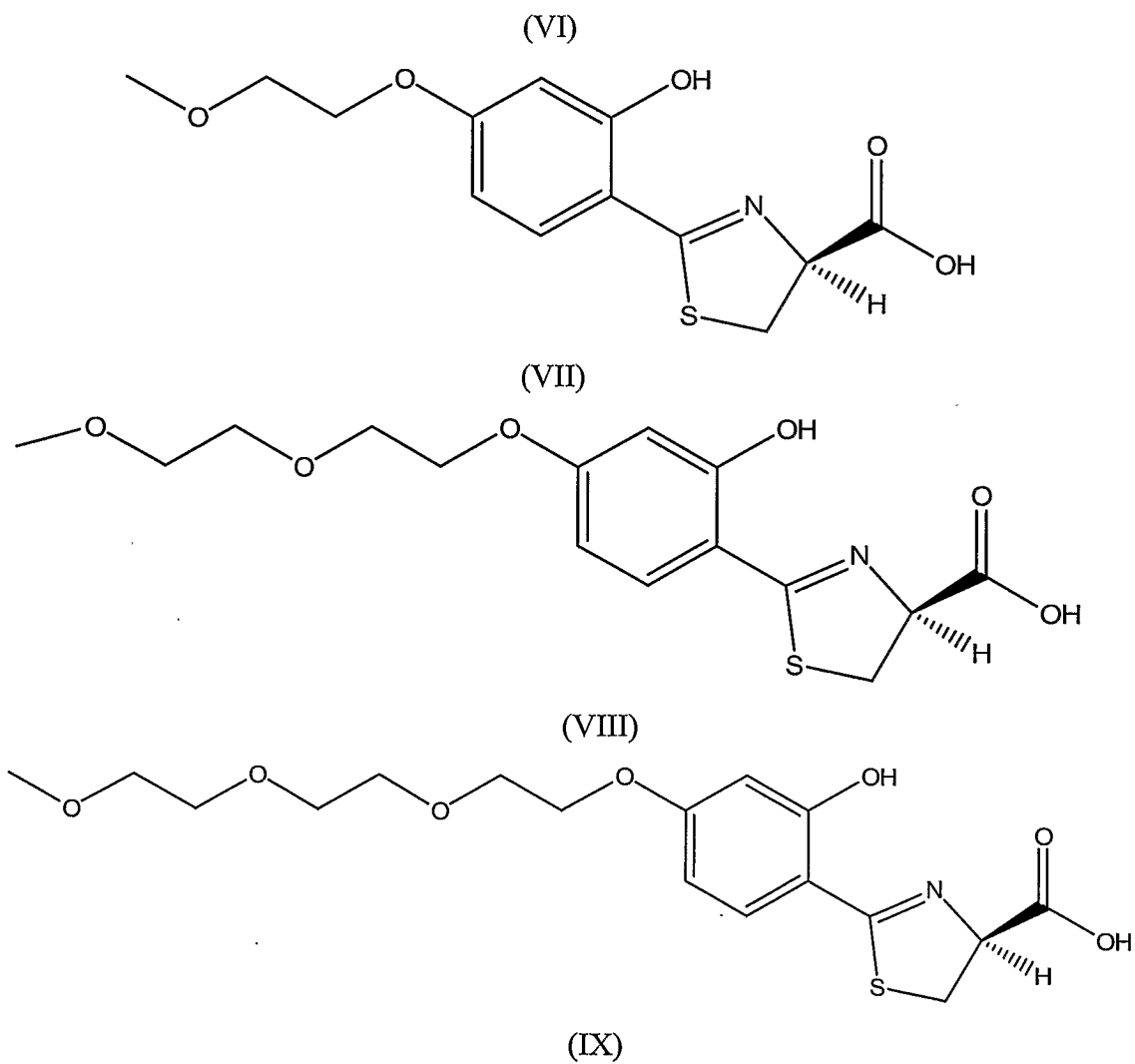
3. The compound of Claim 2, wherein R_9 is $-OR_{12}$.
- 5 4. The compound of Claim 3, wherein R_8 is $-H$ or $-CH_3$.
5. The compound of Claim 4, wherein R_6 and R_7 are independently $-H$ or $-CH_3$.
- 10 6. The compound of Claim 5, wherein R_3 , R_4 and R_5 are each $-H$.
7. The compound of Claim 6, wherein R_2 is $-[(CH_2)_n-O]_x-R'$; n is 1 to 4; and x is 1 to 4.
- 15 8. The compound of Claim 1, wherein the compound is represented by a structural formula selected from one of Structural Formulas (IV)-(IX):



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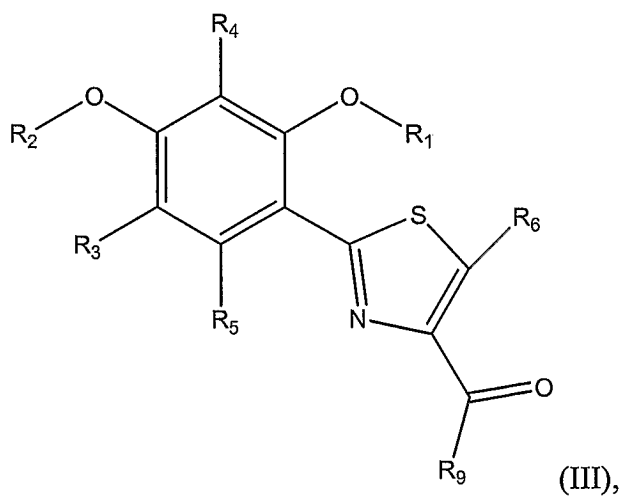
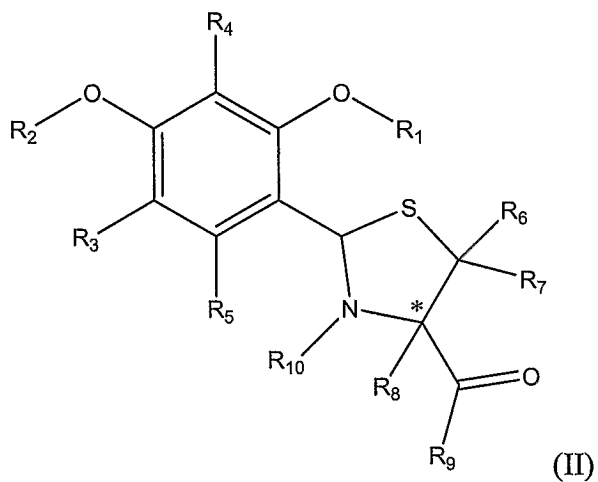
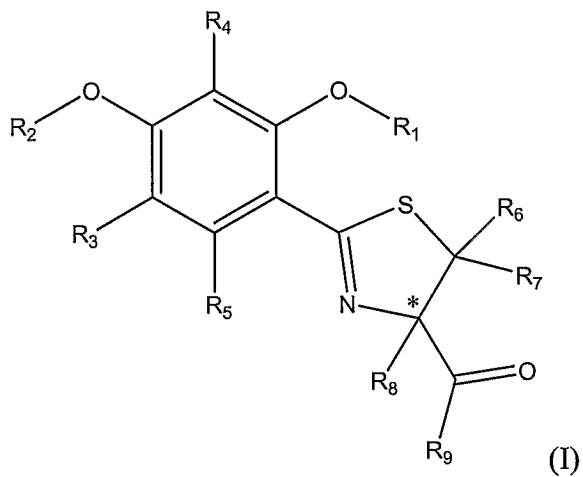


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9. A pharmaceutical composition comprising a pharmaceutically acceptable carrier or diluent and a compound represented by a structural formula selected from Structural Formulas (I), (II), and (III):



wherein:

R_1 is $-H$ or an acyl group;

R_2 is $-[(CH_2)_n-O]_x-[(CH_2)_n-O]_y-R'$;

R_3, R_4 and R_5 are each independently $-H$, an alkyl group, or $-OR_{11}$;

5 R_6, R_7 , and R_8 are each independently $-H$ or an alkyl group;

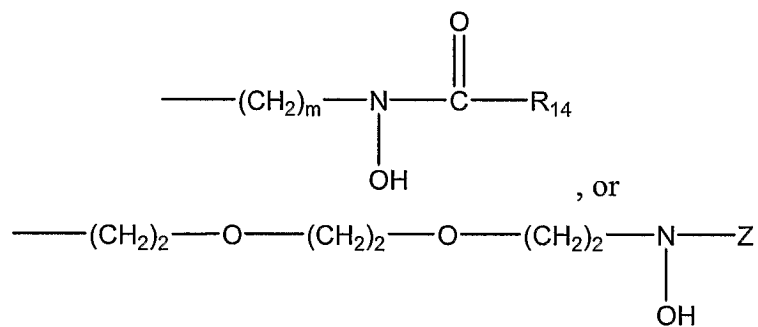
R_9 is $-OR_{12}$ or $-N(OH)R_{13}$;

R_{10} is $-H$ or an alkyl group;

R_{11} is $-H$, an alkyl group or an acyl group;

R_{12} is $-H$ or an alkyl group;

10 R_{13} is an alkyl group;



R_{14} is an alkyl group;

R' is an alkyl group;

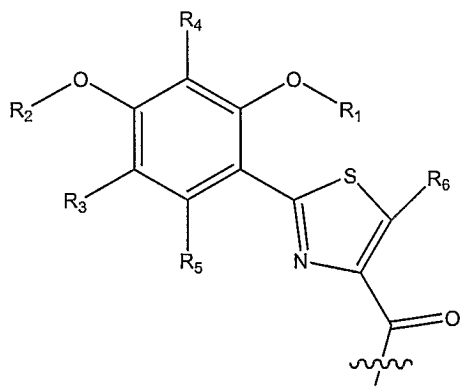
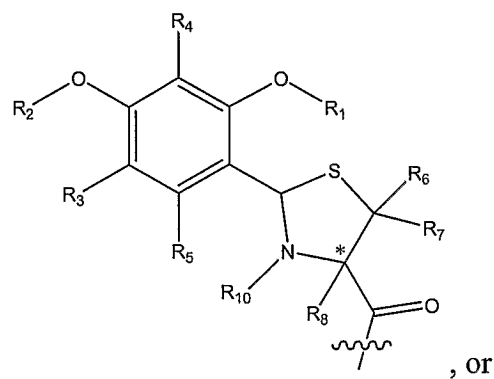
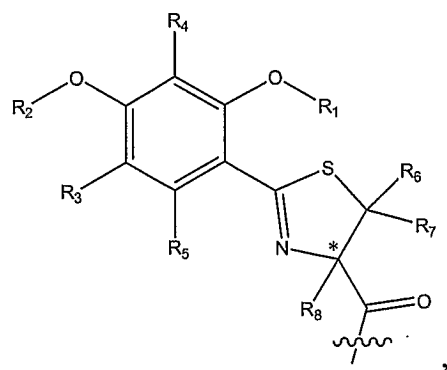
15 m is an integer from 1 to 8;

each n is independently an integer from 1 to 8;

x is an integer from 1 to 8;

y is an integer from 0 to 8;

Z is $-C(O)R_{14}$,

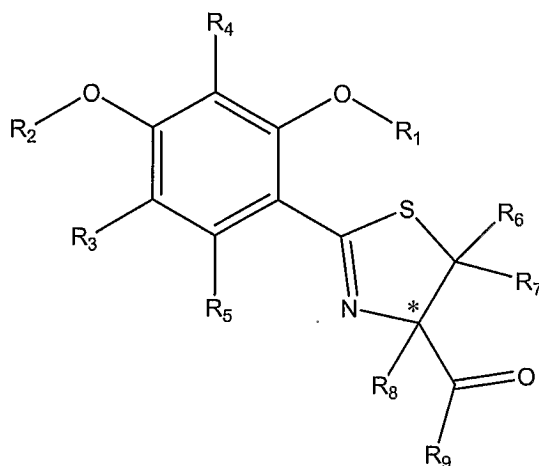


or a pharmaceutically acceptable salt, solvate or hydrate thereof.

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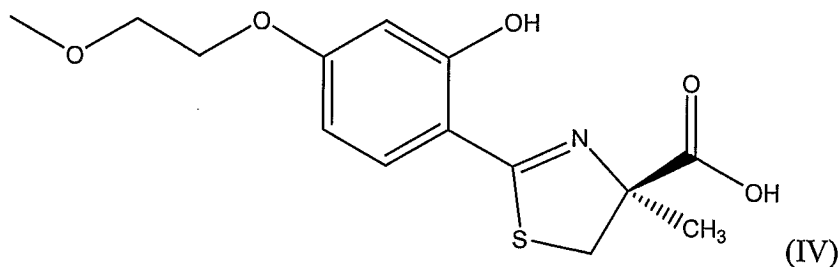
10. The pharmaceutical composition of Claim 9, wherein the compound is represented by Structural Formula (I):

- 40 -

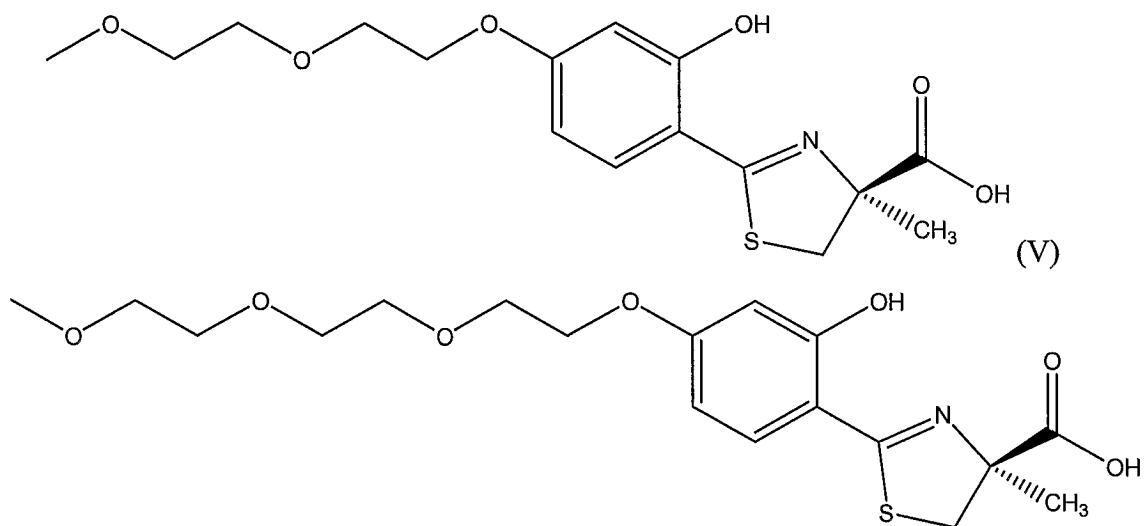


(I).

11. The pharmaceutical composition of Claim 10, wherein R_9 is $-OR_{12}$.
12. The pharmaceutical composition of Claim 11, wherein R_8 is $-H$ or $-CH_3$.
13. The pharmaceutical composition of Claim 12, wherein R_6 and R_7 are independently $-H$ or $-CH_3$.
14. The pharmaceutical composition of Claim 13, wherein R_3 , R_4 and R_5 are each $-H$.
15. The pharmaceutical composition of Claim 14, wherein R_2 is $-[(CH_2)_n-O]_x-R'$; n is 1 to 4; and x is 1 to 4.
16. The pharmaceutical composition of Claim 9, wherein the compound is represented by a structural formula selected from one of Structural Formulas (IV)-(IX):

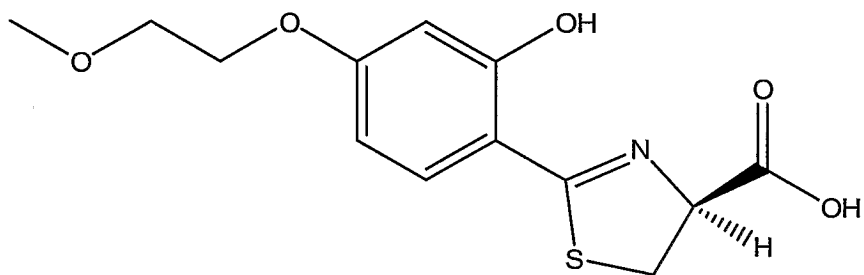


(IV)

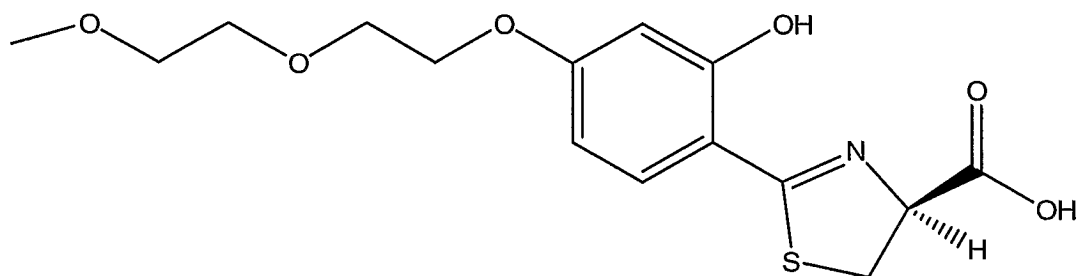


(VI)

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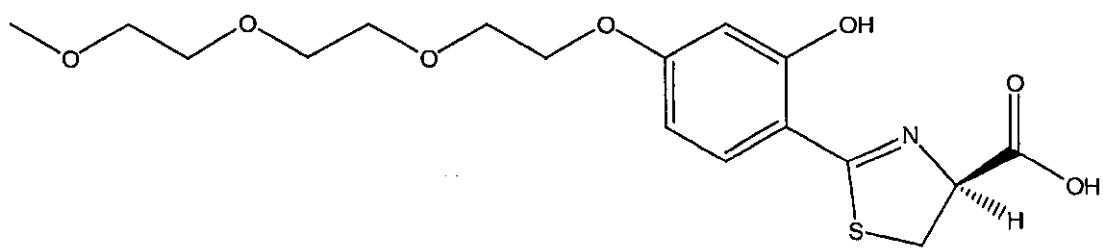
(VII)



(VIII)

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(IX)