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THIOTUNGSTATE ANALOGUES AND USES THEREOF

1. Field

The present invention relates generally to tetrathiotungstate derivatives,
5 methods of making novel tetrathiotungstate derivatives, pharmaceutical compositions
of novel tetrathiotungstate derivatives, methods of using novel tetrathiotungstate
derivatives and pharmaceutical compositions of tetrathiotungstate derivatives to treat
or prevent diseases associated with aberrant vascularization, copper metabolism
disorders, neurodegenerative disorders and obesity.

10

2. Background

Most forms of cancer are derived from solid tumors (Shockley *et al.*, *Ann. N.Y.*
Acad. Sci. **1991**, 617: 367-382, which have proven resistant in the clinic to therapies
such as the use of monoclonal antibodies and immunotoxins. Anti-angiogenic therapy
15 for the treatment of cancer was developed from the recognition that solid tumors
require angiogenesis (*i.e.*, new blood vessel formation) for sustained growth
(Folkman, *Ann. Surg.* **1972**, 175: 409-416; Folkman, *Mol. Med.* **1995**, 1(2): 120-122;
Folkman, *Breast Cancer Res. Treat.* **1995**, 36(2): 109-118; Hanahan *et al.*, *Cell* **1996**,
86(3): 353-364). Efficacy of anti-angiogenic therapy in animal models has been
20 demonstrated (Millauer *et al.*, *Cancer Res.* **1996**, 56:1615-1620; Borgstrom *et al.*,
Prostate **1998**, 35:1-10; Benjamin *et al.*, *J. Clin. Invest.* **1999**, 103: 159-165;
Merajver *et al.*, *Proceedings of Special AACR Conference on Angiogenesis and*
Cancer **1998**, Abstract #B-11, January 22-24). In the absence of angiogenesis,
internal cell layers of solid tumors are inadequately nourished. Further, angiogenesis
25 (*i.e.*, aberrant vascularization) has been implicated in numerous other diseases (*e.g.*,
ocular neovascular disease, macular degeneration, rheumatoid arthritis, *etc.*). More
recently, angiogenesis inhibition has been directly correlated with adipose tissue loss
and weight loss in animal models, which suggests anti-angiogenic therapy may be
useful in prevention of obesity (Rupnick *et al.*, *Proc. Natl. Acad. Sci.* **2002**,
30 99:10730-10735).

Contrastingly, normal tissue does not require angiogenesis except under
specialized circumstances (*e.g.*, wound repair, proliferation of the internal lining of
the uterus during the menstrual cycle, *etc.*). Accordingly, a requirement for
angiogenesis is a significant difference between tumor cells and normal tissue.



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Importantly, the dependency of tumor cells on angiogenesis, when compared to normal cells, is quantitatively greater than differences in cell replication and cell death, between normal tissue and tumor tissue, which are often exploited in cancer therapy.

5 Angiogenesis requires copper, as has been shown by numerous studies (Parke *et al.*, *Am. J. Pathol.* **1988**, 137:173-178; Raju *et al.*, *Natl. Cancer Inst.* **1982**, 69: 1183-1188; Ziche *et al.*, *Natl. Cancer Inst.* **1982**, 69: 475-482; Gullino, *Anticancer Res.* **1986**, 6(2): 153-158). Attempts at preventing angiogenesis and hence tumor growth in animal models by reducing *in vivo* amounts of copper have been reported in
10 the art (Brem *et al.*, *Neurosurgery* **1990**, 26:391-396; Brem *et al.*, *Am. J. Pathol.* **1990**, 137(5): 1121-1142; Yoshida *et al.*, *Neurosurgery* **1995** 37(2): 287-295). These approaches incorporated both copper chelators and low copper diets. More recently, Brewer *et al.*, International Application No. PCT/US99/20374 have shown that the copper chelators, (*e.g.*, tetrathiomolybdate) may be effective in treating diseases (*e.g.*,
15 solid tumor growth), which require angiogenesis.

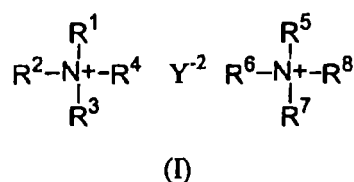
 In addition to the induction of angiogenesis, copper may also have a direct role in tumor cell growth and survival. High copper levels exist in both the plasma and in tumor tissue from patients with many different solid cancers (Chakravarty *et al.*, *J Cancer Res. Clin. Oncol.* **1984**, 108: 312-315). Recently, tetrathiomolybdate has been
20 shown to down-regulate the expression of NF- κ B as well as inhibit its translocation to the nucleus in the inflammatory breast cancer cell line SUM 149 (Pan *et al.*, *Cancer Res.* **2002**, 62: 4854-4859). The NF- κ B system may be involved in mediating tumor cell survival and thus its down-regulation in tumor cells by tetrathiomolybdate suggests a direct effect of copper chelation on tumor survival.

25 Accordingly, novel compounds such as tetrathiotungstate compounds, which are copper chelators, are required to fully explore the potential of copper chelators in treating and/or preventing angiogenesis and in tumor cell viability. Such novel tetrathiotungstate compounds may be effective in treating various diseases associated with angiogenesis such as cancer and obesity along with copper metabolism disorders
30 neurodegenerative disorders, obesity as well as treating diseases where the NF- κ B pathway is dysregulated such as inflammatory disorders.

3. Summary

The present invention satisfies this and other needs by providing novel tetrathiotungstate derivatives, methods of making novel tetrathiotungstate derivatives, pharmaceutical compositions of novel tetrathiotungstate derivatives, methods of using novel tetrathiotungstate derivatives to treat diseases associated with aberrant vascularization, copper metabolism disorders, neurodegenerative disorders and obesity and methods of using pharmaceutical compositions of tetrathiotungstate derivatives to treat or prevent diseases associated with aberrant vascularization, copper metabolism disorders, neurodegenerative disorders, obesity or NF-κB dysregulation.

In a first aspect, the present invention provides a compound of structural formula (I):



or a solvate, hydrate or N-oxide thereof wherein:

R¹, R², R³, R⁵, R⁶ and R⁷ are independently hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, heteroaryl, substituted heteroaryl, heteroarylalkyl, substituted heteroarylalkyl, heteroalkyl or substituted heteroalkyl;

R⁴ and R⁸ are independently hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, heteroaryl, substituted heteroaryl, heteroarylalkyl, substituted heteroarylalkyl, heteroalkyl or substituted heteroalkyl or are absent when N is part of an aromatic ring;

optionally, R¹ and R² taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl;

optionally, R^5 and R^6 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl;

optionally, R^1 and R^2 taken together, R^2 and R^3 taken together and R^2 and R^4 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl;

optionally, R^5 and R^6 taken together, R^6 and R^7 taken together and R^6 and R^8 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl;

optionally, R^3 and R^7 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl; and

Y^{-2} is $(WS_4)^{-2}$, $(W_2S_{12})^{-2}$, $(W_2S_9)^{-2}$, $(W_2S_7)^{-2}$, $(W_2S_8)^{-2}$, $(W_2S_{11})^{-2}$, $(W_2S_6)^{-2}$ or $(W_2S_{13})^{-2}$.

In a second aspect, the present invention provides pharmaceutical compositions of compounds of the invention. The pharmaceutical compositions generally comprise one or more compounds of the invention, hydrates or solvates thereof and a pharmaceutically acceptable diluent, carrier, excipient and adjuvant. The choice of diluent, carrier, excipient and adjuvant will depend upon, among other factors, the desired mode of administration.

In a third aspect, the present invention provides methods for treating or preventing diseases or disorders characterized by aberrant vascularization, copper metabolism disorders, neurodegenerative disorders, obesity or NF- κ B dysregulation. The methods generally involve administering to a patient in need of such treatment or prevention a therapeutically effective amount of a compound and/or pharmaceutical composition of the invention.

In a fourth aspect, the current invention provides pharmaceutical compositions for treating or preventing diseases or disorders characterized by aberrant vascularization, copper metabolism disorders neurodegenerative disorders, obesity or NF- κ B dysregulation in a patient in need of such treatment or prevention.

4. Brief Description of the Drawings

Figure 1 illustrates inhibition of angiogenesis by ammonium tetrathiotungstate in Matrigel® plug assay.

5. Detailed Description

5.1 Definitions

“Compounds” refers to compounds encompassed by structural formula (I) disclosed herein and includes any specific compounds within that generic formula whose structure is disclosed herein. Compounds may be identified either by chemical structure and/or chemical name. When the chemical structure and chemical name conflict, the chemical structure is determinative of the identity of the compound. Compounds may contain one or more chiral centers and/or double bonds and therefore, may exist as stereoisomers, such as double-bond isomers (*i.e.*, geometric isomers), enantiomers or diastereomers. Accordingly, the chemical structures depicted herein encompass all possible enantiomers and stereoisomers of the illustrated compounds including the stereoisomerically pure form (*e.g.*, geometrically pure, enantiomerically pure or diastereomerically pure) and enantiomeric and stereoisomeric mixtures. Enantiomeric and stereoisomeric mixtures can be resolved into their component enantiomers or stereoisomers using separation techniques or chiral synthesis techniques well known to the skilled artisan. Compounds may also exist in several tautomeric forms including the enol form, the keto form and mixtures thereof. Accordingly, the chemical structures depicted herein encompass all possible tautomeric forms of the illustrated compounds. Compounds also include isotopically labeled compounds where one or more atoms have an atomic mass different from the atomic mass conventionally found in nature. Examples of isotopes that may be incorporated into the compounds of the invention include, but are not limited to, ^2H , ^3H , ^{13}C , ^{14}C , ^{15}N , ^{18}O , ^{17}O and ^{35}S . Compounds may exist in unsolvated forms as well as solvated forms, including hydrated forms and as N-oxides. In general, the hydrated, solvated forms and N-oxides are within the scope of the present invention. Certain compounds may exist in multiple crystalline or amorphous forms. In general, all physical forms are equivalent for the uses contemplated herein. Further, it should

be understood, when partial structures of the compounds are illustrated, that brackets indicate the point of attachment of the partial structure to the rest of the molecule.

“Alkyl” by itself or as part of another substituent, refers to a saturated or
5 unsaturated, branched, straight-chain or cyclic monovalent hydrocarbon radical derived by the removal of one hydrogen atom from a single carbon atom of a parent alkane, alkene or alkyne. Typical alkyl groups include, but are not limited to, methyl; ethyls such as ethanyl, ethenyl, ethynyl; propyls such as propan-1-yl, propan-2-yl, cyclopropan-1-yl, prop-1-en-1-yl, prop-1-en-2-yl, prop-2-en-1-yl (allyl),
10 cycloprop-1-en-1-yl; cycloprop-2-en-1-yl, prop-1-yn-1-yl, prop-2-yn-1-yl, *etc.*; butyls such as butan-1-yl, butan-2-yl, 2-methyl-propan-1-yl, 2-methyl-propan-2-yl, cyclobutan-1-yl, but-1-en-1-yl, but-1-en-2-yl, 2-methyl-prop-1-en-1-yl, but-2-en-1-yl, but-2-en-2-yl, buta-1,3-dien-1-yl, buta-1,3-dien-2-yl, cyclobut-1-en-1-yl, cyclobut-1-en-3-yl, cyclobuta-1,3-dien-1-yl, but-1-yn-1-yl, but-1-yn-3-yl,
15 but-3-yn-1-yl, *etc.*; and the like.

The term “alkyl” is specifically intended to include groups having any degree or level of saturation, *i.e.*, groups having exclusively single carbon-carbon bonds, groups having one or more double carbon-carbon bonds, groups having one or more triple carbon-carbon bonds and groups having mixtures of single, double and triple
20 carbon-carbon bonds. Where a specific level of saturation is intended, the expressions “alkanyl,” “alkenyl,” and “alkynyl” are used. Preferably, an alkyl group comprises from 1 to 20 carbon atoms, more preferably, from 1 to 10 carbon atoms, most preferably, from 1 to 6 carbon atoms.

25 “Alkanyl” by itself or as part of another substituent, refers to a saturated branched, straight-chain or cyclic alkyl radical derived by the removal of one hydrogen atom from a single carbon atom of a parent alkane. Typical alkanyl groups include, but are not limited to, methanyl; ethanyl; propanyls such as propan-1-yl, propan-2-yl (isopropyl), cyclopropan-1-yl, *etc.*; butanyls such as butan-1-yl,
30 butan-2-yl (*sec*-butyl), 2-methyl-propan-1-yl (isobutyl), 2-methyl-propan-2-yl (*t*-butyl), cyclobutan-1-yl, *etc.*; and the like.

“Alkenyl” by itself or as part of another substituent, refers to an unsaturated branched, straight-chain or cyclic alkyl radical having at least one carbon-carbon

double bond derived by the removal of one hydrogen atom from a single carbon atom of a parent alkene. The group may be in either the *cis* or *trans* conformation about the double bond(s). Typical alkenyl groups include, but are not limited to, ethenyl; propenyls such as prop-1-en-1-yl, prop-1-en-2-yl, prop-2-en-1-yl (allyl),
 5 prop-2-en-2-yl, cycloprop-1-en-1-yl; cycloprop-2-en-1-yl; butenyls such as but-1-en-1-yl, but-1-en-2-yl, 2-methyl-prop-1-en-1-yl, but-2-en-1-yl, but-2-en-1-yl, but-2-en-2-yl, buta-1,3-dien-1-yl, buta-1,3-dien-2-yl, cyclobut-1-en-1-yl, cyclobut-1-en-3-yl, cyclobuta-1,3-dien-1-yl, *etc.*; and the like.

- 10 “Alkynyl” by itself or as part of another substituent, refers to an unsaturated branched, straight-chain or cyclic alkyl radical having at least one carbon-carbon triple bond derived by the removal of one hydrogen atom from a single carbon atom of a parent alkyne. Typical alkynyl groups include, but are not limited to, ethynyl; propynyls such as prop-1-yn-1-yl, prop-2-yn-1-yl, *etc.*; butynyls such as
 15 but-1-yn-1-yl, but-1-yn-3-yl, but-3-yn-1-yl, *etc.*; and the like.

- “Alkyldiyl” by itself or as part of another substituent, refers to a saturated or unsaturated, branched, straight-chain or cyclic divalent hydrocarbon group derived by the removal of one hydrogen atom from each of two different carbon atoms of a
 20 parent alkane, alkene or alkyne, or by the removal of two hydrogen atoms from a single carbon atom of a parent alkane, alkene or alkyne. The two monovalent radical centers or each valency of the divalent radical center can form bonds with the same or different atoms. Typical alkyldiyl groups include, but are not limited to methandiyl; ethyldiyls such as ethan-1,1-diyl, ethan-1,2-diyl, ethen-1,1-diyl, ethen-1,2-diyl;
 25 propyldiyls such as propan-1,1-diyl, propan-1,2-diyl, propan-2,2-diyl, propan-1,3-diyl, cyclopropan-1,1-diyl, cyclopropan-1,2-diyl, prop-1-en-1,1-diyl, prop-1-en-1,2-diyl, prop-2-en-1,2-diyl, prop-1-en-1,3-diyl, cycloprop-1-en-1,2-diyl, cycloprop-2-en-1,2-diyl, cycloprop-2-en-1,1-diyl, prop-1-yn-1,3-diyl, *etc.*; butyldiyls such as, butan-1,1-diyl, butan-1,2-diyl, butan-1,3-diyl, butan-1,4-diyl, butan-2,2-diyl,
 30 2-methyl-propan-1,1-diyl, 2-methyl-propan-1,2-diyl, cyclobutan-1,1-diyl; cyclobutan-1,2-diyl, cyclobutan-1,3-diyl, but-1-en-1,1-diyl, but-1-en-1,2-diyl, but-1-en-1,3-diyl, but-1-en-1,4-diyl, 2-methyl-prop-1-en-1,1-diyl, 2-methanylidene-propan-1,1-diyl, buta-1,3-dien-1,1-diyl, buta-1,3-dien-1,2-diyl, buta-1,3-dien-1,3-diyl, buta-1,3-dien-1,4-diyl, cyclobut-1-en-1,2-diyl,

cyclobut-1-en-1,3-diyl, cyclobut-2-en-1,2-diyl, cyclobuta-1,3-dien-1,2-diyl, cyclobuta-1,3-dien-1,3-diyl, but-1-yn-1,3-diyl, but-1-yn-1,4-diyl, buta-1,3-diyn-1,4-diyl, *etc.*; and the like. Where specific levels of saturation are intended, the nomenclature alkanyldiyl, alkenyldiyl and/or alkynyldiyl is used.

- 5 Preferably, the alkylidiyl group is (C₁-C₂₀) alkylidiyl, more preferably, (C₁-C₁₀) alkylidiyl, most preferably, (C₁-C₆) alkylidiyl. Preferred are saturated acyclic alkanyldiyl groups in which the radical centers are at the terminal carbons, *e.g.*, methandiyl (methano); ethan-1,2-diyl (ethano); propan-1,3-diyl (propano); butan-1,4-diyl (butano); and the like (also referred to as alkyleno, defined *infra*).

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- “Alkyleno” by itself or as part of another substituent, refers to a straight-chain alkylidiyl group having two terminal monovalent radical centers derived by the removal of one hydrogen atom from each of the two terminal carbon atoms of straight-chain parent alkane, alkene or alkyne. Typical alkyleno groups include, but
15 are not limited to, methano; ethylenos such as ethano, etheno, ethyno; propylenos such as propano, prop[1]eno, propa[1,2]dieno, prop[1]yno, *etc.*; butylenos such as butano, but[1]eno, but[2]eno, buta[1,3]dieno, but[1]yno, but[2]yno, but[1,3]diyno, *etc.*; and the like. Where specific levels of saturation are intended, the nomenclature alkano, alkeno and/or alkyno is used. Preferably, the alkyleno group is (C₁-C₂₀)
20 alkyleno, more preferably, (C₁-C₁₀) alkyleno, most preferably, (C₁-C₆) alkyleno. Preferred are straight-chain saturated alkano groups, *e.g.*, methano, ethano, propano, butano, and the like.

- “Acyl” by itself or as part of another substituent, refers to a radical -C(O)R³⁰,
25 where R³⁰ is hydrogen, alkyl, cycloalkyl, cycloheteroalkyl, aryl, arylalkyl, heteroalkyl, heteroaryl, heteroarylalkyl as defined herein. Representative examples include, but are not limited to formyl, acetyl, cyclohexylcarbonyl, cyclohexylmethylcarbonyl, benzoyl, benzylcarbonyl and the like.

- 30 “Acylamino” by itself or as part of another substituent, refers to a radical -NR³¹C(O)R³², where R³¹ and R³² are each independently hydrogen, alkyl, cycloalkyl, cycloheteroalkyl, aryl, arylalkyl, heteroalkyl, heteroaryl, heteroarylalkyl, as defined herein. Representative examples include, but are not limited to, formylamino,

acetylamino, cyclohexylcarbonylamino, cyclohexylmethyl-carbonylamino, benzoylamino, benzylcarbonylamino and the like.

“Alkoxy” by itself or as part of another substituent, refers to a radical -OR³³ where R³³ represents an alkyl or cycloalkyl group as defined herein. Representative examples include, but are not limited to, methoxy, ethoxy, propoxy, butoxy, cyclohexyloxy and the like.

“Alkoxy carbonyl” by itself or as part of another substituent, refers to a radical -C(O)OR³³ where R³³ is as defined above.

“Aryl” by itself or as part of another substituent, refers to a monovalent aromatic hydrocarbon radical derived by the removal of one hydrogen atom from a single carbon atom of a parent aromatic ring system. Typical aryl groups include, but are not limited to, groups derived from aceanthrylene, acenaphthylene, acephenanthrylene, anthracene, azulene, benzene, chrysene, coronene, fluoranthene, fluorene, hexacene, hexaphene, hexalene, *as*-indacene, *s*-indacene, indane, indene, naphthalene, octacene, octaphene, octalene, ovalene, penta-2,4-diene, pentacene, pentalene, pentaphene, perylene, phenalene, phenanthrene, picene, pleiadene, pyrene, pyranthrene, rubicene, triphenylene, trinaphthalene and the like. Preferably, an aryl group comprises from 6 to 20 carbon atoms, more preferably, from 6 to 12 carbon atoms.

“Arylalkyl” by itself or as part of another substituent, refers to an acyclic alkyl radical in which one of the hydrogen atoms bonded to a carbon atom, typically a terminal or *sp*³ carbon atom, is replaced with an aryl group. Typical arylalkyl groups include, but are not limited to, benzyl, 2-phenylethan-1-yl, 2-phenylethen-1-yl, naphthylmethyl, 2-naphthylethan-1-yl, 2-naphthylethen-1-yl, naphthobenzyl, 2-naphthophenylethan-1-yl and the like. Where specific alkyl moieties are intended, the nomenclature arylalkanyl, arylalkenyl and/or arylalkynyl is used. Preferably, an arylalkyl group is (C₆-C₃₀) arylalkyl, *e.g.*, the alkanyl, alkenyl or alkynyl moiety of the arylalkyl group is (C₁-C₁₀) and the aryl moiety is (C₆-C₂₀), more preferably, an arylalkyl group is (C₆-C₂₀) arylalkyl, *e.g.*, the alkanyl, alkenyl or alkynyl moiety of the arylalkyl group is (C₁-C₈) and the aryl moiety is (C₆-C₁₂).

“Cycloalkyl” by itself or as part of another substituent, refers to a saturated or unsaturated cyclic alkyl radical. Where a specific level of saturation is intended, the nomenclature “cycloalkanyl” or “cycloalkenyl” is used. Typical cycloalkyl groups
 5 include, but are not limited to, groups derived from cyclopropane, cyclobutane, cyclopentane, cyclohexane and the like. Preferably, the cycloalkyl group is (C₃–C₁₀) cycloalkyl, more preferably, (C₃–C₇) cycloalkyl.

“Cycloheteroalkyl” by itself or as part of another substituent, refers to a
 10 saturated or unsaturated cyclic alkyl radical in which one or more carbon atoms (and any associated hydrogen atoms) are independently replaced with the same or different heteroatom. Typical heteroatoms to replace the carbon atom(s) include, but are not limited to, N, P, O, S, Si, *etc.* Where a specific level of saturation is intended, the nomenclature “cycloheteroalkanyl” or “cycloheteroalkenyl” is used. Typical
 15 cycloheteroalkyl groups include, but are not limited to, groups derived from epoxides, azirines, thiiranes, imidazolidine, morpholine, piperazine, piperidine, pyrazolidine, pyrrolidine, quinuclidine and the like.

“Heteroalkyl, Heteroalkanyl, Heteroalkenyl, Heteroalkanyl, Heteroalkyldiyl and Heteroalkyleno” by themselves or as part of another substituent, refer to alkyl, alkanyl, alkenyl, alkynyl, alkyldiyl and alkyleno groups, respectively, in which one or more of the carbon atoms (and any associated hydrogen atoms) are each
 20 independently replaced with the same or different heteroatomic groups. Typical heteroatomic groups which can be included in these groups include, but are not limited to, -O-, -S-, -O-O-, -S-S-, -O-S-, -NR³⁵R³⁶-, =N-N=, -N=N-, -N=N-NR³⁷R³⁸-, -PR³⁹-, -P(O)₂-, -POR³⁹-, -O-P(O)₂-, -SO-, -SO₂-, -SnR⁴¹R⁴²- and the like, where R³⁵, R³⁶, R³⁷, R³⁸, R³⁹, R⁴⁰, R⁴¹ and R⁴² are independently hydrogen, alkyl, substituted
 25 alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, heteroalkyl, substituted heteroalkyl, heteroaryl, substituted heteroaryl, heteroarylalkyl or substituted
 30 heteroarylalkyl.

“Heteroaryl” by itself or as part of another substituent, refers to a monovalent heteroaromatic radical derived by the removal of one hydrogen atom from a single

atom of a parent heteroaromatic ring system. Typical heteroaryl groups include, but are not limited to, groups derived from acridine, arsindole, carbazole, β -carboline, chromane, chromene, cinnoline, furan, imidazole, indazole, indole, indoline, indolizine, isobenzofuran, isochromene, isoindole, isoindoline, isoquinoline, isothiazole, isoxazole, naphthyridine, oxadiazole, oxazole, perimidine, phenanthridine, phenanthroline, phenazine, phthalazine, pteridine, purine, pyran, pyrazine, pyrazole, pyridazine, pyridine, pyrimidine, pyrrole, pyrrolizine, quinazoline, quinoline, quinolizine, quinoxaline, tetrazole, thiadiazole, thiazole, thiophene, triazole, xanthene, and the like. Preferably, the heteroaryl group is from 5-20 membered heteroaryl, more preferably from 5-10 membered heteroaryl. Preferred heteroaryl groups are those derived from thiophene, pyrrole, benzothiophene, benzofuran, indole, pyridine, quinoline, imidazole, oxazole and pyrazine.

"Heteroarylalkyl" by itself or as part of another substituent, refers to an acyclic alkyl radical in which one of the hydrogen atoms bonded to a carbon atom, typically a terminal or sp^3 carbon atom, is replaced with a heteroaryl group. Where specific alkyl moieties are intended, the nomenclature heteroarylalkanyl, heteroarylalkenyl and/or heteroarylalkynyl is used. In preferred embodiments, the heteroarylalkyl group is a 6-30 membered heteroarylalkyl, e.g., the alkanyl, alkenyl or alkynyl moiety of the heteroarylalkyl is 1-10 membered and the heteroaryl moiety is a 5-20-membered heteroaryl, more preferably, 6-20 membered heteroarylalkyl, e.g., the alkanyl, alkenyl or alkynyl moiety of the heteroarylalkyl is 1-8 membered and the heteroaryl moiety is a 5-12-membered heteroaryl.

"Parent Aromatic Ring System" by itself or as part of another substituent, refers to an unsaturated cyclic or polycyclic ring system having a conjugated π electron system. Specifically included within the definition of "parent aromatic ring system" are fused ring systems in which one or more of the rings are aromatic and one or more of the rings are saturated or unsaturated, such as, for example, fluorene, indane, indene, phenalene, etc. Typical parent aromatic ring systems include, but are not limited to, aceanthrylene, acenaphthylene, acephenanthrylene, anthracene, azulene, benzene, chrysene, coronene, fluoranthene, fluorene, hexacene, hexaphene, hexalene, *as*-indacene, *s*-indacene, indane, indene, naphthalene, octacene, octaphene,

octalene, ovalene, penta-2,4-diene, pentacene, pentalene, pentaphene, perylene, phenalene, phenanthrene, picene, pleiadene, pyrene, pyranthrene, rubicene, triphenylene, trinaphthalene and the like.

- 5 "Parent Heteroaromatic Ring System" by itself or as part of another substituent, refers to a parent aromatic ring system in which one or more carbon atoms (and any associated hydrogen atoms) are independently replaced with the same or different heteroatom. Typical heteroatoms to replace the carbon atoms include, but are not limited to, N, P, O, S, Si, *etc.* Specifically included within the definition of
- 10 "parent heteroaromatic ring systems" are fused ring systems in which one or more of the rings are aromatic and one or more of the rings are saturated or unsaturated, such as, for example, arsindole, benzodioxan, benzofuran, chromane, chromene, indole, indoline, xanthene, *etc.* Typical parent heteroaromatic ring systems include, but are not limited to, arsindole, carbazole, β -carboline, chromane, chromene, cinnoline,
- 15 furan, imidazole, indazole, indole, indoline, indolizine, isobenzofuran, isochromene, isoindole, isoindoline, isoquinoline, isothiazole, isoxazole, naphthyridine, oxadiazole, oxazole, perimidine, phenanthridine, phenanthroline, phenazine, phthalazine, pteridine, purine, pyran, pyrazine, pyrazole, pyridazine, pyridine, pyrimidine, pyrrole, pyrrolizine, quinazoline, quinoline, quinolizine, quinoxaline, tetrazole, thiadiazole,
- 20 thiazole, thiophene, triazole, xanthene, and the like.

- "Pharmaceutical composition" refers to at least one compound of the invention and a pharmaceutically acceptable vehicle, with which the compound is administered to a patient.

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- "Pharmaceutically acceptable salt" refers to a salt of a compound of the invention, which possesses the desired pharmacological activity of the parent compound. Such salts include: (1) acid addition salts, formed with inorganic acids such as hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric
- 30 acid, and the like; or formed with organic acids such as acetic acid, propionic acid, hexanoic acid, cyclopentanepropionic acid, glycolic acid, pyruvic acid, lactic acid, malonic acid, succinic acid, malic acid, maleic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, 3-(4-hydroxybenzoyl) benzoic acid, cinnamic acid, mandelic acid,

methanesulfonic acid, ethanesulfonic acid, 1,2-ethane-disulfonic acid,
2-hydroxyethanesulfonic acid, benzenesulfonic acid, 4-chlorobenzenesulfonic acid,
2-naphthalenesulfonic acid, 4-toluenesulfonic acid, camphorsulfonic acid,
4-methylbicyclo[2.2.2]-oct-2-ene-1-carboxylic acid, glucoheptonic acid,
5 3-phenylpropionic acid, trimethylacetic acid, tertiary butylacetic acid, lauryl sulfuric
acid, gluconic acid, glutamic acid, hydroxynaphthoic acid, salicylic acid, stearic acid,
muconic acid, and the like; or (2) salts formed when an acidic proton present in the
parent compound is replaced by a metal ion, e.g., an alkali metal ion, an alkaline earth
ion, or an aluminum ion; or coordinates with an organic base such as ethanolamine,
10 diethanolamine, triethanolamine, N-methylglucamine and the like.

"Pharmaceutically acceptable vehicle" refers to a diluent, adjuvant, excipient
or carrier with which a compound of the invention is administered.

15 "Patient" includes humans. The terms "human" and "patient" are used
interchangeably herein.

"Preventing" or "prevention" refers to a reduction in risk of acquiring a
disease or disorder (*i.e.*, causing at least one of the clinical symptoms of the disease
20 not to develop in a patient that may be exposed to or predisposed to the disease but
does not yet experience or display symptoms of the disease).

"Prodrug" refers to a derivative of a drug molecule that requires a
transformation within the body to release the active drug. Prodrugs are frequently,
25 although not necessarily, pharmacologically inactive until converted to the parent
drug. A hydroxyl containing drug may be converted to, for example, to a sulfonate,
ester or carbonate prodrug, which may be hydrolyzed *in vivo* to provide the hydroxyl
compound. An amino containing drug may be converted, for example, to a
carbamate, amide, enamine, imine, N-phosphonyl, N-phosphoryl or N-sulfonyl
30 prodrug, which may be hydrolyzed *in vivo* to provide the amino compound. A
carboxylic acid drug may be converted to an ester (including silyl esters and
thioesters), amide or hydrazide prodrug, which be hydrolyzed *in vivo* to provide the
carboxylic acid compound. Prodrugs for drugs which functional groups different than
those listed above are well known to the skilled artisan.

“Promoiety” refers to a form of protecting group that when used to mask a functional group within a drug molecule converts the drug into a prodrug. Typically, the promoiety will be attached to the drug *via* bond(s) that are cleaved by enzymatic or non-enzymatic means *in vivo*.

“Protecting group” refers to a grouping of atoms that when attached to a reactive functional group in a molecule masks, reduces or prevents reactivity of the functional group. Examples of protecting groups can be found in Green *et al.*, “Protective Groups in Organic Chemistry”, (Wiley, 2nd ed. 1991) and Harrison *et al.*, “Compendium of Synthetic Organic Methods”, Vols. 1-8 (John Wiley and Sons, 1971-1996). Representative amino protecting groups include, but are not limited to, formyl, acetyl, trifluoroacetyl, benzyl, benzyloxycarbonyl (“CBZ”), *tert*-butoxycarbonyl (“Boc”), trimethylsilyl (“TMS”), 2-trimethylsilyl-ethanesulfonyl (“SES”), trityl and substituted trityl groups, allyloxycarbonyl, 9-fluorenylmethyloxycarbonyl (“Fmoc”), nitro-veratryloxycarbonyl (“NVOC”) and the like. Representative hydroxy protecting groups include, but are not limited to, those where the hydroxy group is either acylated or alkylated such as benzyl, and trityl ethers as well as alkyl ethers, tetrahydropyranyl ethers, trialkylsilyl ethers and allyl ethers.

“Substituted” refers to a group in which one or more hydrogen atoms are independently replaced with the same or different substituent(s). Typical substituents include, but are not limited to, -M, -R⁶⁰, -O⁻, =O, -OR⁶⁰, -SR⁶⁰, -S⁻, =S, -NR⁶⁰R⁶¹, =NR⁶⁰, -CF₃, -CN, -OCN, -SCN, -NO, -NO₂, =N₂, -N₃, -S(O)₂O⁻, -S(O)₂OH, -S(O)₂R⁶⁰, -OS(O)₂O⁻, -OS(O)₂R⁶⁰, -P(O)(O⁻)₂, -P(O)(OR⁶⁰)(O⁻), -OP(O)(OR⁶⁰)(OR⁶¹), -C(O)R⁶⁰, -C(S)R⁶⁰, -C(O)OR⁶⁰, -C(O)NR⁶⁰R⁶¹, -C(O)O⁻, -C(S)OR⁶⁰, -NR⁶²C(O)NR⁶⁰R⁶¹, -NR⁶²C(S)NR⁶⁰R⁶¹, -NR⁶²C(NR⁶³)NR⁶⁰R⁶¹ and -C(NR⁶²)NR⁶⁰R⁶¹ where M is independently a halogen; R⁶⁰, R⁶¹, R⁶² and R⁶³ are independently hydrogen, alkyl, substituted alkyl, alkoxy, substituted alkoxy, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, aryl, substituted aryl, heteroaryl or substituted heteroaryl, or optionally R⁶⁰ and R⁶¹ together with the nitrogen atom to which they are bonded form a cycloheteroalkyl or substituted cycloheteroalkyl ring; and R⁶⁴ and R⁶⁵ are independently hydrogen, alkyl,

substituted alkyl, aryl, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, aryl, substituted aryl, heteroaryl or substituted heteroaryl, or optionally R^{64} and R^{65} together with the nitrogen atom to which they are bonded form a cycloheteroalkyl or substituted cycloheteroalkyl ring. Preferably, substituents

5 include $-M$, $-R^{60}$, $=O$, $-OR^{60}$, $-SR^{60}$, $-S^-$, $=S$, $-NR^{60}R^{61}$, $=NR^{60}$, $-CF_3$, $-CN$, $-OCN$, $-SCN$, $-NO$, $-NO_2$, $=N_2$, $-N_3$, $-S(O)_2R^{60}$, $-OS(O)_2O^-$, $-OS(O)_2R^{60}$, $-P(O)(O^-)_2$, $-P(O)(OR^{60})(O^-)$, $-OP(O)(OR^{60})(OR^{61})$, $-C(O)R^{60}$, $-C(S)R^{60}$, $-C(O)OR^{60}$, $-C(O)NR^{60}R^{61}$, $-C(O)O^-$, $-NR^{62}C(O)NR^{60}R^{61}$, more preferably, $-M$, $-R^{60}$, $=O$, $-OR^{60}$, $-SR^{60}$, $-NR^{60}R^{61}$, $-CF_3$, $-CN$, $-NO_2$, $-S(O)_2R^{60}$, $-P(O)(OR^{60})(O^-)$, $-OP(O)(OR^{60})(OR^{61})$,

10 $-C(O)R^{60}$, $-C(O)OR^{60}$, $-C(O)NR^{60}R^{61}$, $-C(O)O^-$, most preferably, $-M$, $-R^{60}$, $=O$, $-OR^{60}$, $-SR^{60}$, $-NR^{60}R^{61}$, $-CF_3$, $-CN$, $-NO_2$, $-S(O)_2R^{60}$, $-OP(O)(OR^{60})(OR^{61})$, $-C(O)R^{60}$, $-C(O)OR^{60}$, $-C(O)O^-$, where R^{60} , R^{61} and R^{62} are as defined above.

"Treating" or "treatment" of any disease or disorder refers, in one

15 embodiment, to ameliorating the disease or disorder (*i.e.*, arresting or reducing the development of the disease or at least one of the clinical symptoms thereof). In another embodiment "treating" or "treatment" refers to ameliorating at least one physical parameter, which may not be discernible by the patient. In yet another embodiment, "treating" or "treatment" refers to inhibiting the disease or disorder,

20 either physically, (*e.g.*, stabilization of a discernible symptom), physiologically, (*e.g.*, stabilization of a physical parameter), or both. In yet another embodiment, "treating" or "treatment" refers to delaying the onset of the disease or disorder.

"Therapeutically effective amount" means the amount of a compound that,

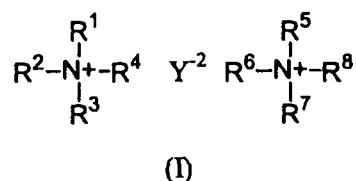
25 when administered to a patient for treating a disease, is sufficient to effect such treatment for the disease. The "therapeutically effective amount" will vary depending on the compound, the disease and its severity and the age, weight, *etc.*, of the patient to be treated.

30 Reference will now be made in detail to preferred embodiments of the invention. While the invention will be described in conjunction with the preferred embodiments, it will be understood that it is not intended to limit the invention to those preferred embodiments. To the contrary, it is intended to cover alternatives,

modifications, and equivalents as may be included within the spirit and scope of the invention as defined by the appended claims.

5.2 Compounds of Structural Formula (I)

In a first embodiment, the compounds of the invention include compound of structural formula (I):



or a solvate or hydrate or N-oxide thereof wherein:

$\text{R}^1, \text{R}^2, \text{R}^3, \text{R}^5, \text{R}^6$ and R^7 are independently hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, heteroaryl, substituted heteroaryl, heteroarylalkyl, substituted heteroarylalkyl, heteroalkyl or substituted heteroalkyl;

R^4 and R^8 are independently hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, heteroaryl, substituted heteroaryl, heteroarylalkyl, substituted heteroarylalkyl, heteroalkyl or substituted heteroalkyl or are absent when N is part of an aromatic ring;

optionally, R^1 and R^2 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl;

optionally, R^5 and R^6 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl;

optionally, R^1 and R^2 taken together, R^2 and R^3 taken together and R^2 and R^4 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl;

optionally, R^5 and R^6 taken together, R^6 and R^7 taken together and R^6 and R^8 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl;

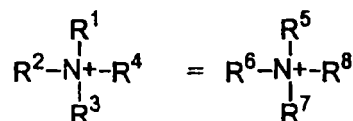
5 optionally, R^3 and R^7 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl; and

Y^{-2} is $(WS_4)^{-2}$, $(W_2S_{12})^{-2}$, $(W_2S_9)^{-2}$, $(W_2S_7)^{-2}$, $(W_2S_8)^{-2}$, $(W_2S_{11})^{-2}$, $(W_2S_6)^{-2}$ or $(W_2S_{13})^{-2}$.

10

In some embodiment, Y is $(WS_4)^{-2}$ and all of R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 and R^8 are not hydrogen. In other embodiments, Y is $(WS_4)^{-2}$ and all of R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 and R^8 are not alkyl.

In still other embodiments,



15

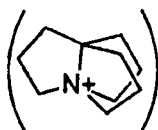
Preferably, Y is $(WS_4)^{-2}$.

In some embodiments, at least one of R^1 , R^2 , R^3 and R^4 is not alkyl. In other embodiments, R^1 , R^2 and R^4 are hydrogen, alkanyl or substituted alkanyl. Preferably, R^1 , R^2 and R^4 are hydrogen, methyl or ethyl.

20 In still other embodiments, R^1 and R^2 are alkanyl. Preferably, R^1 and R^2 are methyl or ethyl.

In still other embodiments, R^1 is alkanyl, substituted alkanyl, alkenyl, substituted alkenyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl or substituted cycloalkyl. Preferably, R^1 and R^2 taken together are alkyleno, substituted alkyleno, heteroalkyleno or substituted heteroalkyleno. More preferably, R^1 and R^2 taken together are alkyleno or heteroalkyleno.

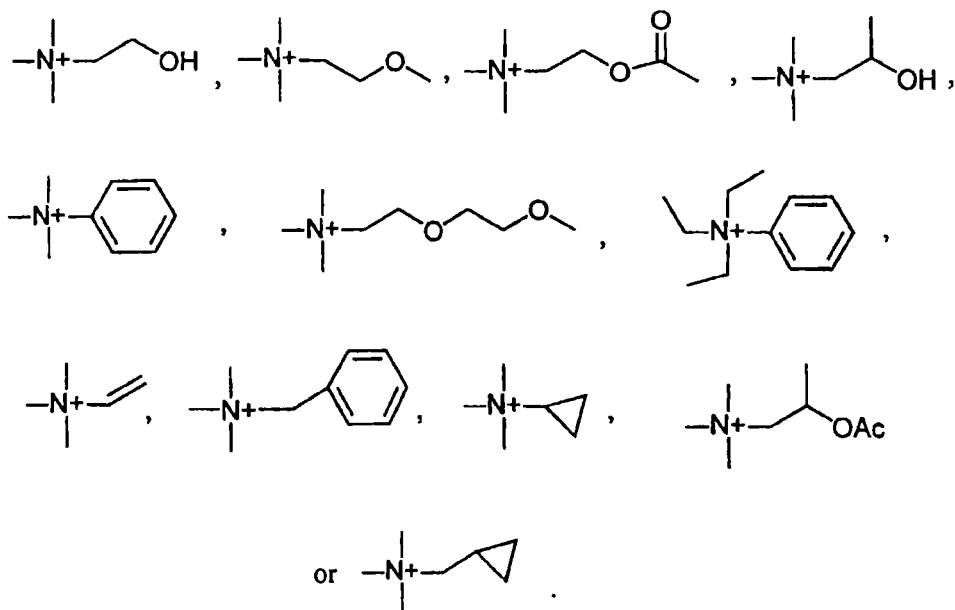
In still other embodiments, R^1 and R^2 taken together, R^2 and R^3 taken together and R^2 and R^4 taken together are alkyleno, substituted alkyleno, heteroalkyleno or substituted heteroalkyleno. Preferably, R^1 and R^2 taken together, R^2 and R^3 taken together and R^2 and R^4 taken together are alkyleno. Preferably, $R^1(R^2)(R^3)(R^4)N$ has the structure:



In still other embodiments, R^3 and R^7 taken together are alkylene, substituted alkylene, heteroalkylene or substituted heteroalkylene. Preferably, R^3 and R^7 taken together are alkylene or heteroalkylene.

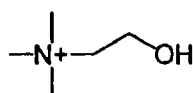
- 5 In still other embodiments, R^1 , R^2 and R^4 are hydrogen, alkanyl or substituted alkanyl and R^3 is alkyl, substituted alkyl, alkenyl, aryl, arylalkyl, cycloalkyl or R^3 and R^7 taken together are alkylene, substituted alkylene, heteroalkylene or substituted heteroalkylene. Preferably, R^1 , R^2 and R^4 are methyl or ethyl and R^3 is alkyl, substituted alkyl, alkenyl, aryl, arylalkyl, cycloalkyl or R^3 and R^7 taken together are alkylene or heteroalkylene. Preferably, R^1 , R^2 and R^4 are methyl or ethyl and R^3 is alkyl, substituted alkyl, alkenyl, aryl, arylalkyl or cycloalkyl.

In still other embodiments, $R^1(R^2)(R^3)(R^4)N$ is

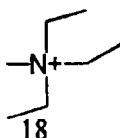


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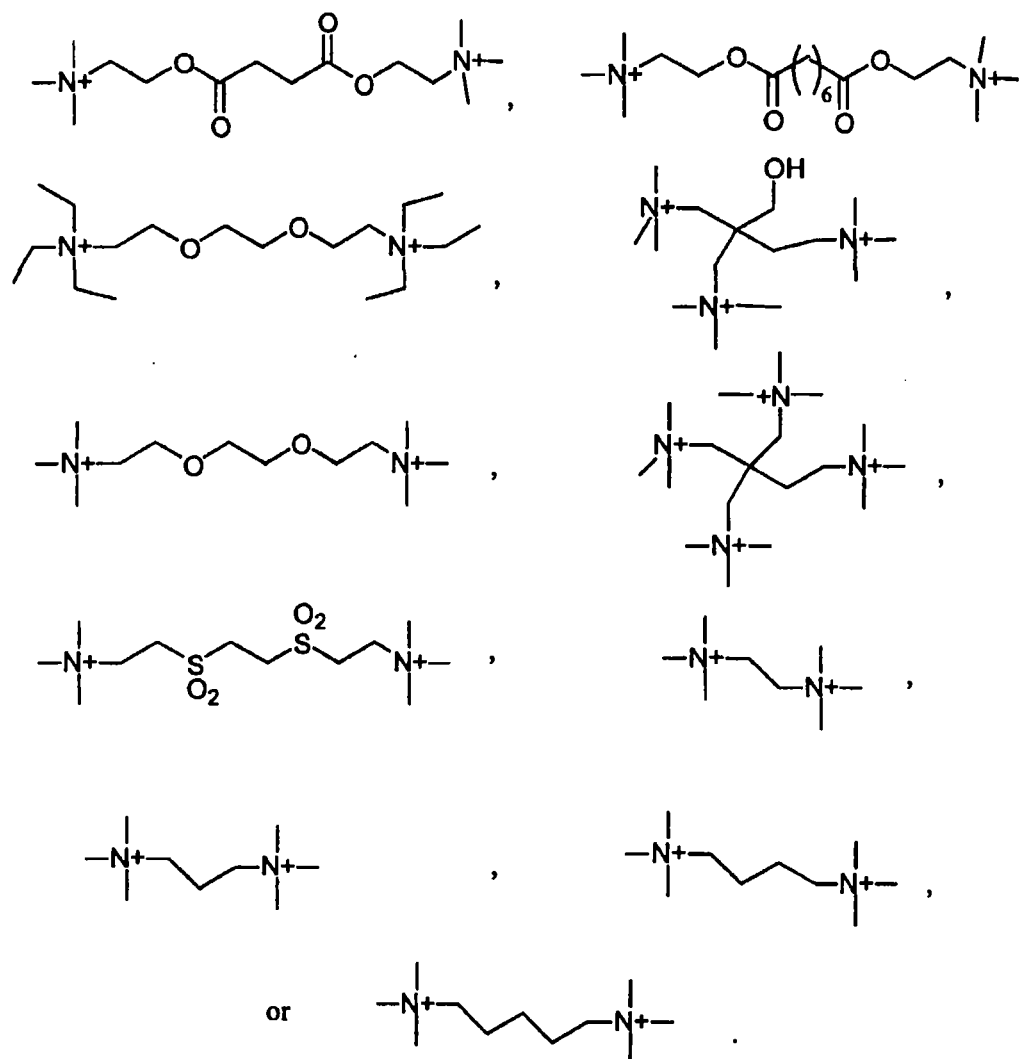
In still other embodiments, $R^1(R^2)(R^3)(R^4)N$ is



In still other embodiments, $R^1(R^2)(R^3)(R^4)N$ is



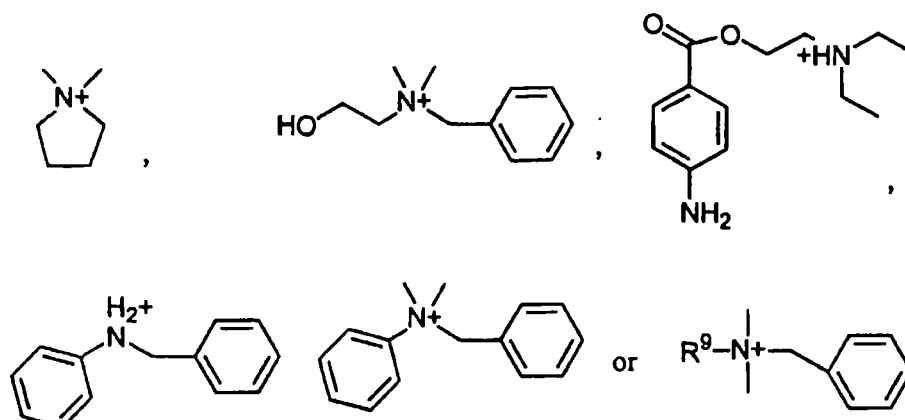
In still other embodiments, R¹, R² and R⁴ are methyl or ethyl and R³ and R⁷ taken together are alkyleno or heteroalkyleno. Preferably, R¹(R²)(R³)(R⁴)N has the structure:



5

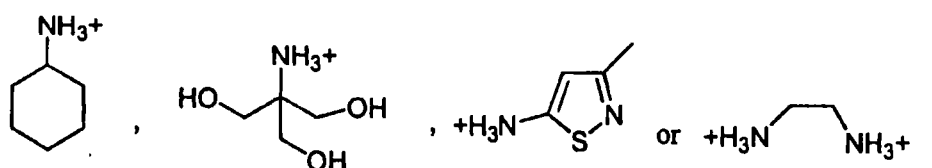
In still other embodiments, R¹, R² and R⁴ are hydrogen and R³ is substituted alkyl, cycloalkyl or substituted heteroaryl or R³ and R⁷ taken together are alkylene. In still another embodiment, R¹ and R² are alkanyl and R³ and R⁴ are alkyl, substituted alkyl, aryl, arylalkyl or alkylene. Preferably, R¹ and R² are methyl or ethyl and R³ and R⁴ are alkyl, substituted alkyl, aryl, arylalkyl or alkylene.

In still other embodiments, $R^1(R^2)(R^3)(R^4)N$ are

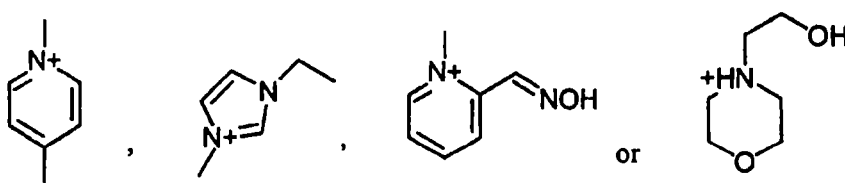


wherein R^9 is a mixture of straight chain alkanyl groups which have at least eight carbon atoms and not more than eighteen carbon atoms.

In still other embodiments, R^1 , R^2 and R^4 are hydrogen and R^3 is substituted alkyl, substituted heteroaryl, cycloalkyl or alkylene. Preferably, $R^1(R^2)(R^3)(R^4)N$ has the structure:



In still other embodiments, R^1 and R^2 taken together are alkylene, substituted alkylene, heteroalkylene or substituted heteroalkylene, R^3 is alkyl or substituted alkyl and R^4 is hydrogen or is absent. Preferably, $R^1(R^2)(R^3)N$ or $R^1(R^2)(R^3)(R^4)N$ has the structure:

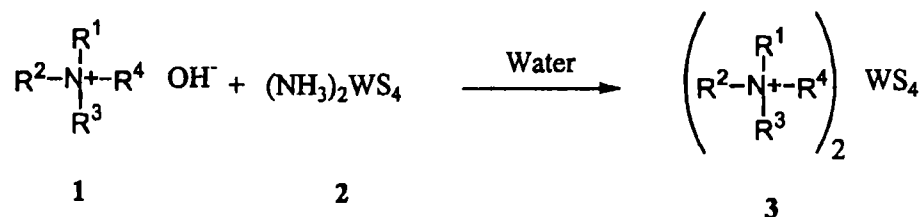


5.3 Synthesis of Compounds

The compounds described herein may be obtained *via* conventional synthetic methods illustrated in Schemes 1 and 2. Starting materials useful for preparing compounds described herein and intermediates thereof are commercially available or can be prepared by well-known synthetic methods. Substituted ammonium salts (*e.g.*, ammonium hydroxide and ammonium halides) may be either purchased from commercial sources or may be readily synthesized using well-known synthetic

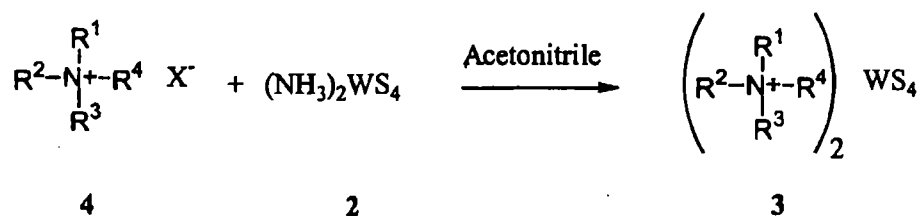
- methods (Harrison *et al.*, "Compendium of Synthetic Organic Methods", Vols. 1-8 (John Wiley and Sons, 1971-1996); "Beilstein Handbook of Organic Chemistry," Beilstein Institute of Organic Chemistry, Frankfurt, Germany; Feiser *et al.*, "Reagents for Organic Synthesis," Volumes 1-17, Wiley Interscience; Trost *et al.*,
- 5 "Comprehensive Organic Synthesis," Pergamon Press, 1991; "Theilheimer's Synthetic Methods of Organic Chemistry," Volumes 1-45, Karger, 1991; March, "Advanced Organic Chemistry," Wiley Interscience, 1991; Larock "Comprehensive Organic Transformations," VCH Publishers, 1989; Paquette, "Encyclopedia of Reagents for Organic Synthesis," John Wiley & Sons, 1995). Other methods for
- 10 synthesis of the compounds described herein and/or starting materials are either described in the art or will be readily apparent to the skilled artisan. Accordingly, the methods presented in Schemes 1 and 2 herein are illustrative rather than comprehensive.

Scheme 1



- 15 As shown above, in Scheme 1, addition of a quaternary ammonium hydroxide to thiotungstate in the presence of water leads to cation exchange (equilibrium to product is driven by removal of volatile ammonia) to provide the desired thiotungstate derivative.

Scheme 2



- 20 As shown above, in Scheme 2, addition of a quaternary ammonium halide to thiotungstate in the presence of acetonitrile leads to cation exchange (equilibrium to product is driven by formation of ammonium halide) to provide the desired thiotungstate derivative.

It should be noted that thiotungstate derivatives where the ammonium counterions are different may be prepared from compounds 3 by treating with one equivalent of a different ammonium counterion. Such a reaction would be expected to produce a statistical mixture of products.

- 5 Further, those of skill in the art will appreciate that conventional methods comprising treating tungstate with an ammonium salt and hydrogen sulfide may be used to synthesize many of the compounds described herein.

5.4 Assays

- 10 Those of skill in the art will appreciate that the *in vitro* and *in vivo* assays useful for measuring the activity of the compounds described herein are illustrative rather than comprehensive.

5.4.1 Assay for Endothelial Cell Migration

- 15 For endothelial cell migration, transwells are coated with type I collagen (50 µg/mL) by adding 200 µL of the collagen solution per transwell, then incubating overnight at 37 °C. The transwells are assembled in a 24-well plate and a chemoattractant (*e.g.*, FGF-2) is added to the bottom chamber in a total volume of 0.8 mL media. Endothelial cells such as human umbilical vein endothelial cells
20 ("HUVEC"), which have been detached from monolayer culture using trypsin, are diluted to a final concentration of about 10⁶ cells/mL with serum-free media and 0.2 mL of this cell suspension is added to the upper chamber of each transwell. Inhibitors are added to both the upper and lower chambers, and the migration is allowed to proceed for 5 hrs in a humidified atmosphere at 37°C. The transwells are removed
25 from the plate stained using DiffQuik®. Cells which did not migrate are removed from the upper chamber by scraping with a cotton swab and the membranes are detached, mounted on slides, and counted under a high-power field (400x) to determine the number of cells migrated.

30 5.4.2 Biological Assay of Anti-Invasive Activity

Compounds and/or compositions are tested for their anti-invasive capacity. The ability of cells such as endothelial cells or tumor cells (*e.g.*, PC-3 human prostatic carcinoma) cells to invade through a reconstituted basement membrane (Matrigel®)

in an assay known as a Matrigel® invasion assay (Kleinman *et al.*, *Biochemistry* 1986, 25: 312-318; Parish *et al.*, *Int. J. Cancer* 1992, 52:378-383). Matrigel® is a reconstituted basement membrane containing type IV collagen, laminin, heparan sulfate proteoglycans such as perlecan, which bind to and localize bFGF, vitronectin
5 as well as transforming growth factor- β (TGF β , urokinase-type plasminogen activator (uPA), tissue plasminogen activator (tPA), and the serpin known as plasminogen activator inhibitor type 1 (PAI-1)) (Chambers *et al.*, *Canc. Res.* 1995, 55:1578-1585). Results obtained in this assay for compounds which target extracellular receptors or enzymes are typically predictive of the efficacy of these compounds *in vivo* (Rabbani
10 *et al.*, 1995, *Int. J. Cancer* 63: 840-845).

Such assays employ transwell tissue culture inserts. Invasive cells are defined as cells which are able to traverse through the Matrigel® and upper aspect of a polycarbonate membrane and adhere to the bottom of the membrane. Transwells (Costar) containing polycarbonate membranes (8.0 μ m pore size) are coated with
15 Matrigel® (Collaborative Research), which has been diluted in sterile PBS to a final concentration of 75 μ g/mL (60 μ L of diluted Matrigel® per insert), and placed in the wells of a 24-well plate. The membranes are dried overnight in a biological safety cabinet, then rehydrated by adding 100 μ L of DMEM containing antibiotics for 1 hour on a shaker table. The DMEM is removed from each insert by aspiration and 0.8
20 mL of DMEM/10 % FBS/antibiotics is added to each well of the 24-well plate such that it surrounds the outside of the transwell ("lower chamber"). Fresh DMEM/antibiotics (100 μ L), human Glu-plasminogen (5 μ g/mL), and any compounds to be tested are added to the top, inside of the transwell ("upper chamber"). The cells which are to be tested are trypsinized and resuspended in DMEM/antibiotics, then
25 added to the top chamber of the transwell at a final concentration of 800,000 cells/mL. The final volume of the upper chamber is adjusted to 200 μ L. The assembled plate is then incubated in a humid 5% CO₂ atmosphere for 72 hours. After incubation, the cells are fixed and stained using DiffQuik® (Giemsa stain) and the upper chamber is then scraped using a cotton swab to remove the Matrigel® and any cells which did not
30 invade through the membrane. The membranes are detached from the transwell using an X-acto® blade, mounted on slides using Permount® and cover-slips, then counted under a high-powered (400x) field. An average of the cells invaded is determined from 5-10 fields counted and plotted as a function of inhibitor concentration.

5.4.3 Tube-Formation Assays of Anti-Angiogenic Activity

Compounds may be tested for anti-angiogenic activity in one of two different assay systems *in vitro*.

5 Endothelial cells, for example, human umbilical vein endothelial cells ("HUVEC") or human microvascular endothelial cells ("HMVEC") which can be prepared or obtained commercially, are mixed at a concentration of 2×10^5 cells/mL with fibrinogen (5mg/mL in phosphate buffered saline ("PBS") in a 1:1 (v/v) ratio. Thrombin is added (5 units/mL final concentration) and the mixture is immediately
10 transferred to a 24-well plate (0.5 mL per well). The fibrin gel is allowed to form and then VEGF and bFGF are added to the wells (each at 5 ng/mL final concentration) along with the test compound. The cells are incubated at 37°C in 5% CO₂ for 4 days at which time the cells in each well are counted and classified as either rounded,
15 elongated with no branches, elongated with one branch, or elongated with 2 or more branches. Results are expressed as the average of 5 different wells for each concentration of compound. Typically, in the presence of angiogenic inhibitors, cells remain either rounded or form undifferentiated tubes (*e.g.*, 0 or 1 branch). This assay is recognized in the art to be predictive of angiogenic (or anti-angiogenic) efficacy *in vivo* (Min *et al.*, *Cancer Res.* 1996, 56: 2428-2433).

20 In an alternate assay, endothelial cell tube formation is observed when endothelial cells are cultured on Matrigel® (Schnaper *et al.*, *J. Cell. Physiol.* 1995, 165:107-118). Endothelial cells (1×10^4 cells/well) are transferred onto Matrigel®-coated 24-well plates and tube formation is quantitated after 48 hours. Inhibitors are tested by addition at either the same time as the endothelial cells or at various time
25 points thereafter. Tube formation can also be stimulated by adding angiogenic growth factors such as bFGF or VEGF, differentiation stimulating agents (*e.g.*, PMA) or combinations thereof.

 This assay models angiogenesis by presenting a particular type of basement membrane to the endothelial cells, namely the layer of matrix, which migrating and
30 differentiating endothelial cells might be expected to first encounter. In addition, the matrix components found in Matrigel® (and in basement membranes *in situ*) or proteolytic products thereof may also be stimulatory for endothelial cell tube formation which makes this model complementary to the fibrin gel angiogenesis model previously described (Blood *et al.*, *Biochim. Biophys. Acta* 1990, 1032:89-118;

Odedra *et al.*, *Pharmac. Ther.* 1991, 49:111-124). The compounds inhibit endothelial cell tube formation in both assays, which suggests that the compounds will also have anti-angiogenic activity.

5 **5.4.4 Assays for Inhibition of Proliferation**

 The ability of the compounds to inhibit the proliferation of endothelial cells may be determined in a 96-well format. Type I collagen (gelatin) is used to coat the wells of the plate (0.1-1 mg/mL in PBS, 0.1 mL per well for 30 minutes at room temperature). After washing the plate (3x w/PBS), 3-6,000 cells are plated per well and allowed to attach for 4 hours (37 °C/5% CO₂) in Endothelial Growth Medium (EGM; Clonetics) or M199 media containing 0.1-2% FBS. The media and any unattached cells are removed at the end of 4 hours and fresh media containing bFGF (1-10 ng/mL) or VEGF (1-10 ng/mL) is added to each well. Compounds to be tested are added last and the plate is allowed to incubate (37 °C/5% CO₂) for 24-48 hrs.

15 MTS (Promega) is added to each well and allowed to incubate from 1-4 hrs. The absorbance at 490nm, which is proportional to the cell number, is then measured to determine the differences in proliferation between control wells and those containing test compounds. A similar assay system can be set up with cultured adherent tumor cells. However, collagen may be omitted in this format. Tumor cells (e.g., 3,000-

20 10,000/well) are plated and allowed to attach overnight. Serum free medium is then added to the wells and the cells are synchronized for 24 hrs. Medium containing 10% FBS is then added to each well to stimulate proliferation. Compounds to be tested are included in some of the wells. After 24 hrs, MTS is added to the plate and the assay developed and read as described above. A similar methodology may also be

25 employed to evaluate the effects of the compounds of the invention on the proliferation of other cell types including tumor cells except that VEGF and bFGF would not be used to stimulate growth of the cells. If there is evidence of anti-proliferative activity, induction of apoptosis may be measured using TumorTACS (Genzyme).

30

5.4.5 Assays of Cytotoxicity

 The cytotoxic effects of compounds described herein may be determined for various cell types including tumor cells, endothelial cells, fibroblasts and macrophages.

A typical assay involves plating cells at a density of 5-10,000 cells per well in a 96-well plate. Compounds are then added at a variety of concentrations and allowed to incubate with the cells for 24 hours. The cells are washed 3X with media. For cytotoxicity assays (measuring cell lysis), a Promega 96-well cytotoxicity kit is used.

5

5.4.6 Corneal Angiogenesis Model

The protocol used is essentially identical to that described by Volpert *et al.*, *J. Clin. Invest.* **1996**, *98*:671-679. Briefly, female Fischer rats (120-140 gms) are anesthetized and pellets (5 μ l) comprised of Hydron[®], bFGF (150 nM) and the compounds to be tested are implanted into tiny incisions made in the cornea 1.0-1.5 mm from the limbus. Neovascularization is assessed at 5 and 7 days after implantation. On day 7, animals are anesthetized and infused with a dye such as colloidal carbon to stain the vessels. The animals are then euthanized, the corneas fixed with formalin, and the corneas flattened and photographed to assess the degree of neovascularization. Neovessels may be quantitated by imaging the total vessel area or length or simply by counting vessels.

5.4.7 Matrigel[®] Plug Assay

This assay is performed essentially as described by Passaniti *et al.*, *Lab Invest.* **1992**, *67*:519-528. Ice-cold Matrigel[®] (e.g., 500 μ L) (Collaborative Biomedical Products, Inc., Bedford, MA) is mixed with heparin (e.g., 50 μ g/ml), FGF-2 (e.g., 400 ng/ml) and the compound to be tested. In some assays, bFGF may be substituted with tumor cells as the angiogenic stimulus. The Matrigel[®] mixture is injected subcutaneously into 4-8 week-old athymic nude mice at sites near the abdominal midline, preferably 3 injections per mouse. The injected Matrigel[®] forms a palpable solid gel. Injection sites are chosen such that each animal receives a positive control plug (such as FGF-2 + heparin), a negative control plug (e.g., buffer + heparin) and a plug that includes the compound being tested for its effect on angiogenesis (e.g., FGF-2 + heparin + compound). All treatments are preferably run in triplicate. Animals are sacrificed by cervical dislocation at about 7 days post injection or another time that may be optimal for observing angiogenesis. The mouse skin is detached along the abdominal midline, and the Matrigel[®] plugs are recovered and scanned immediately

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at high resolution. Plugs are then dispersed in water and incubated at 37°C overnight. Hemoglobin (Hb) levels are determined using Drabkin's solution (*e.g.*, obtained from Sigma) according to the manufacturers' instructions. The amount of Hb in the plug is an indirect measure of angiogenesis as it reflects the amount of blood in the sample.

5 In addition, or alternatively, animals may be injected prior to sacrifice with a 0.1 ml buffer (preferably PBS) containing a high molecular weight dextran to which is conjugated a fluorophore. The amount of fluorescence in the dispersed plug which is determined fluorimetrically serves as a measure of angiogenesis in the plug. Staining with mAb anti-CD31 (CD31 is platelet-endothelial cell adhesion molecule or

10 "PECAM") may also be used to confirm neovessel formation and microvessel density in the plugs.

5.4.8 Chick chorioallantoic membrane (CAM) angiogenesis assay

This assay is performed essentially as described by Nguyen *et al.*,

15 *Microvascular Res.* 1994, 47:31-40. A mesh containing either angiogenic factors (bFGF) or tumor cells plus inhibitors is placed onto the CAM of an 8-day old chick embryo and the CAM observed for 3-9 days after implantation of the sample. Angiogenesis is quantitated by determining the percentage of squares in the mesh which contain blood vessels.

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5.4.9 In Vivo Assessment Angiogenesis Inhibition and Anti-Tumor Effects Using the Matrigel® Plug Assay with Tumor Cells

In this assay, tumor cells, for example, $1-5 \times 10^6$ cells of the 3LL Lewis lung carcinoma or the rat prostate cell line MatLyLu, are mixed with Matrigel® and then

25 injected into the flank of a mouse following the protocol described in section 4.4.7 above. A mass of tumor cells and a powerful angiogenic response can be observed in the plugs after about 5 to 7 days. The anti-tumor and anti-angiogenic action of a compound in an actual tumor environment can be evaluated by including it in the plug. Measurement is then made of tumor weight, Hb levels or fluorescence levels

30 (of a dextran-fluorophore conjugate injected prior to sacrifice). To measure Hb or fluorescence, the plugs are first homogenized with a tissue homogenizer.

5.4.10 Xenograft Model of Subcutaneous Tumor Growth

Nude mice are inoculated with MDA-MB-231 cells (human breast carcinoma) and Matrigel® (1 x 10⁶ cells in 0.2mL) subcutaneously in the right flank of the animals. The tumors are staged to 200 mm³ and then treatment with a test compound is initiated. Tumor volumes are obtained every other day and the animals are sacrificed after 2 weeks of treatment. The tumors are excised, weighed and paraffin embedded. Histological sections of the tumors are analyzed by H and E, anti-CD31, Ki-67, TUNEL, and CD68 staining.

Other human tumor cell lines including but not limited to PC-3, CWR22R, SK-OV-3, A2780, A549, HCT116, HT29 may also be used to test the anti-tumor activity of the compounds described herein in a similar manner.

5.4.11 Xenograft Model of Metastasis

The compounds may also be tested for inhibition of late metastasis using an experimental metastasis model (Crowley *et al.*, *Proc. Natl. Acad. Sci. USA* 1993, 90 5021-5025). Late metastasis involves the steps of attachment and extravasation of tumor cells, local invasion, seeding, proliferation and angiogenesis. Human prostatic carcinoma cells (PC-3) transfected with a reporter gene, preferably the green fluorescent protein (GFP) gene, but as an alternative with a gene encoding the enzymes chloramphenicol acetyl-transferase (CAT), luciferase or LacZ, are inoculated into nude mice. This approach permits utilization of either of these markers (fluorescence detection of GFP or histochemical colorimetric detection of enzymatic activity) to follow the fate of these cells. Cells are injected, preferably, i.v., and metastases identified after about 14 days, particularly in the lungs but also in regional lymph nodes, femurs and brain. This mimics the organ tropism of naturally occurring metastases in human prostate cancer. For example, GFP-expressing PC-3 cells (1 x 10⁶ cells per mouse) are injected i.v. into the tail veins of nude (*nu/nu*) mice. Animals are treated with a test composition at 100µg/animal/day given q.d. IP. Single metastatic cells and foci are visualized and quantitated by fluorescence microscopy or light microscopic histochemistry or by grinding the tissue and quantitative colorimetric assay of the detectable label.

5.4.12 Inhibition of Spontaneous Metastasis In Vivo by HPRG and Functional Derivatives

The rat syngeneic breast cancer system (Xing *et al.*, *Int. J. Cancer* 1996, 67:423-429) employs Mat BIII rat breast cancer cells. Tumor cells, for example about
5 10^6 suspended in 0.1 mL PBS, are inoculated into the mammary fat pads of female Fisher rats. At the time of inoculation, a 14-day Alza osmotic mini-pump is implanted intraperitoneally to dispense the test compound. The compound is dissolved in PBS (*e.g.*, 200 mM stock), sterile filtered and placed in the minipump to achieve a release rate of about 4 mg/kg/day. Control animals receive vehicle (PBS)
10 alone or a vehicle control peptide in the mini-pump. Animals are sacrificed at about day 14.

Other models of experimental metastasis may also be used to evaluate the compounds described herein. These models would utilize the human tumor cell lines described, *supra*, injected through the tail vein of a nude mouse. Typically, these
15 mice are sacrificed 28 days after tumor cell inoculation and their lungs evaluated for the presence of metastases.

5.4.13 3LL Lewis Lung Carcinoma: Primary Tumor Growth

This tumor line arose spontaneously in 1951 as carcinoma of the lung in a
20 C57BL/6 mouse (*Cancer Res.* 1955, 15:39. See, also Malave *et al.*, *J. Nat'l. Canc. Inst.* 1979, 62:83-88). It is propagated by passage in C57BL/6 mice by subcutaneous inoculation and is tested in semiallogeneic C57BL/6 x DBA/2 F₁ mice or in allogeneic C3H mice. Typically six animals per group for subcutaneously implant, or ten for intramuscular implant are used. Tumor may be implanted by subcutaneous
25 inoculation as a 2-4 mm fragment or intramuscularly implanted or subcutaneous implanted as an inoculum of suspended cells of about $0.5-2 \times 10^6$ -cells. Treatment begins 24 hours after implant or is delayed until a tumor of specified size (usually approximately 400 mg) can be palpated. The test compound is administered i.p. daily for 11 days

30 Animals are followed by weighing, palpation, and measurement of tumor size. Typical tumor weight in untreated control recipients on day 12 after intramuscular inoculation is 500-2500 mg. Typical median survival time is 18-28 days. A positive control compound, for example, cyclophosphamide at 20 mg/kg/injection per day on days 1-11 is used. Results computed include mean animal weight, tumor size, tumor

weight and survival time. For confirmed therapeutic activity, the test composition should be tested in two multi-dose assays.

5.4.14 3LL Lewis Lung Carcinoma: Primary Growth and Spontaneous

Metastasis Model

This model has been utilized by a number of investigators (See, for example, Gorelik *et al.*, 1980, *J. Nat'l. Canc. Inst.* 65:1257-1264; Gorelik *et al.*, *Rec. Results Canc. Res.* 1980, 75:20-28; Isakov *et al.*, *Invasion Metas.* 1982, 2:12-32; Talmadge *et al.*, *J. Nat'l. Canc. Inst.* 1982, 69:975-980; Hilgard *et al.*, *Br. J. Cancer* 1977, 35:78-86). Test mice are male C57BL/6 mice, 2-3 months old. Following subcutaneous, intramuscular or intra-footpad implantation, this tumor produces metastases, preferentially in the lungs. With some lines of the tumor, the primary tumor exerts anti-metastatic effects and must first be excised before study of the metastatic phase (see also, O'Reilly *et al.*, United States Patent No. 5,639,725).

Single-cell suspensions are prepared from solid tumors by treating minced tumor tissue with a solution of 0.3% trypsin. Cells are washed 3 times with PBS (pH 7.4) and suspended in PBS. Viability of the 3LL cells prepared in this way is generally about 95-99% (by trypan blue dye exclusion). Viable tumor cells (3×10^4 - 5×10^6) suspended in 0.05 ml PBS are injected subcutaneously, either in the dorsal region or into one hind foot pad of C57BL/6 mice. Visible tumors appear after 3-4 days after dorsal sc injection of 10^6 cells. The day of tumor appearance and the diameters of established tumors are measured by caliper every two days.

The treatment is given as one or two doses of compound, per week. In another embodiment, the compound is delivered by osmotic minipump.

In experiments involving tumor excision of dorsal tumors, when tumors reach about 1500 mm³ in size, mice are randomized into two groups: (1) primary tumor is completely excised; or (2) sham surgery is performed and the tumor is left intact. Although tumors from 500-3000 mm³ inhibit growth of metastases, 1500 mm³ is the largest size primary tumor that can be safely resected with high survival and without local regrowth. After 21 days, all mice are sacrificed and autopsied.

Lungs are removed and weighed. Lungs are fixed in Bouin's solution and the number of visible metastases is recorded. The diameters of the metastases are also measured using a binocular stereoscope equipped with a micrometer-containing ocular under 8X magnification. On the basis of the recorded diameters, it is possible

to calculate the volume of each metastasis. To determine the total volume of metastases per lung, the mean number of visible metastases is multiplied by the mean volume of metastases. To further determine metastatic growth, it is possible to measure incorporation of $^{125}\text{IdUrd}$ into lung cells (Thakur *et al.*, *J. Lab. Clin. Med.*

5 1977, 89:217-228). Ten days following tumor amputation, 25 μg of fluorodeoxyuridine is inoculated into the peritoneums of tumor-bearing (and, if used, tumor-resected mice). After 30 min, mice are given 1 μCi of $^{125}\text{IdUrd}$ (iododeoxyuridine). One day later, lungs and spleens are removed and weighed and a degree of $^{125}\text{IdUrd}$ incorporation is measured using a gamma counter.

10 In mice with footpad tumors, when tumors reach about 8-10 mm in diameter, mice are randomized into two groups: (1) legs with tumors are amputated after ligation above the knee joints; or (2) mice are left intact as nonamputated tumor-bearing controls. (Amputation of a tumor-free leg in a tumor-bearing mouse has no known effect on subsequent metastasis, ruling out possible effects of anesthesia, stress
15 or surgery). Mice are killed 10-14 days after amputation. Metastases are evaluated as described above.

5.5 Therapeutic Uses

A compound of structural formula (I) and/or a pharmaceutical composition
20 thereof is administered to a patient, preferably a human, suffering from a disease characterized by aberrant vascularization. Aberrant vascularization includes abnormal neovascularization such as the formation of new blood vessels, larger blood vessels, more branched blood vessels and any other mechanism, which leads to an increased blood carrying capacity to a diseased tissue or site. The compounds and
25 pharmaceutical compositions thereof may be used to treat and/or prevent aberrant vascularization.

Preferably, diseases characterized by aberrant vascularization include, but are not limited to, cancer (*e.g.*, any vascularized tumor, preferably, a solid tumor, including but not limited to, carcinomas of the lung, breast, ovary, stomach, pancreas,
30 larynx, esophagus, testes, liver, parotid, biliary tract, colon, rectum, cervix, uterus, endometrium, kidney, bladder, prostate, thyroid, squamous cell carcinomas, adenocarcinomas, small cell carcinomas, melanomas, gliomas, neuroblastomas, sarcomas (*e.g.*, angiosarcomas, chondrosarcomas)), arthritis, diabetes, arteriosclerosis, arteriovenous, malformations, corneal graft neovascularization, delayed wound

healing, diabetic retinopathy, age related macular degeneration, granulations, burns, hemophilic joints, rheumatoid arthritis, hypertrophic scars, neovascular glaucoma, nonunion fractures, Osier Weber Syndrome, psoriasis, granuloma, retrolental fibroplasia, pterygium, scleroderma, trachoma, vascular adhesions, ocular
5 neovascularization, parasitic diseases, hypertrophy following surgery, inhibition of hair growth, macular degeneration (including both wet and dry type), rheumatoid arthritis and osteoarthritis. Diseases characterized by aberrant vascularization which are preferably treated and/or prevented by administration of a compound of structural formula (I) and/or a pharmaceutical composition thereof include cancer, macular
10 degeneration and rheumatoid arthritis.

Further, a compound of structural formula (I) and/or a pharmaceutical composition thereof may be administered to a patient, preferably a human, suffering from a disease associated with copper metabolism disorders (*e.g.*, Wilson's disease) to treat and/or prevent such a disease.

15 Still further, a compound of structural formula (I) and/or a pharmaceutical composition thereof may be administered to a patient, preferably a human, to treat and/or prevent obesity. The compounds of structural formula (I) may be also used to reduce levels of inflammatory cytokines (*e.g.*, TNF- α , TNF- β , IL-8, *etc.*) and plasminogen activator inhibitor, which may be associated with angiogenesis and
20 obesity (Loskutoff *et al.*, *Ann. N.Y. Acad. Sci.*, 2000, 902:272-281; Pan *et al.*, *Cancer Res.*, 2002, 62:4854-4859; Hanada *et al.*, *Cytokine Growth Factor Rev.* 2002, 13:413-421; Chen *et al.*, *Science* 2002, 296:1634-5; Miyake *et al.*, *J. Neuropathol. Exp. Neurol.* 59:18-28; Koch *et al.*, *Science* 1992, 258:1798-801; Osawa *et al.*, *Infect. Immun.* 2002, 70:6294-6301; Bajou *et al.*, *Nat. Med.* 1998, 4:923-8).

25 Still further, a compound of structural formula (I) and/or a pharmaceutical composition thereof may be administered to a patient, preferably a human, suffering from a neurodegenerative disorder, to treat and/or prevent the neurodegenerative disorder. Elevated levels of copper have been reported in the art to mediate the pathobiology of various neurodegenerative disorders including Alzheimer's disease,
30 amyotrophic lateral sclerosis (ALS) and prion disease (Llanos *et al.*, *DNA Cell Biol.* 2002, 21: 259-270; Carri *et al.*, *Funct. Neurol* 2001, 16:181-188; Perry *et al.*, *CNS Drugs* 2002, 16:339-352; Kowalik-Jankowska *et al.*, *Environ Health Perspect.* 2002, 5: 869-870; Maynard *et al.*, *J. Biol. Chem.* 2002, 277, 44670-44676; Gnjec *et al.*,

Front Biosci. **2002**, 16-23; Strausak *et al.*, *Brain Res. Bull.* **2001**, 55: 175-185; Brown, *Brain Res. Bull.* **2001**, 55:165-173; Brown, *Biochem. Soc. Trans* **2002**, 30:742-745).

Still further, in accordance with the invention, a compound of structural formula (I) and/or a pharmaceutical composition thereof may be administered to a
5 patient, preferably a human, to treat diseases characterized by dysregulated activity of the NF- κ B or dysregulated inflammation of inflammatory response.

Further, in certain embodiments, a compounds and and/or pharmaceutical compositions thereof are administered to a patient, preferably, a human, as a preventative measure against various diseases or disorders characterized by aberrant
10 vascularization, copper metabolism disorders, neurodegenerative disorders, obesity or NF- κ B dysregulation. Accordingly, compounds of structural Formula (I) and/or pharmaceutical compositions thereof may be used for the prevention of one disease or disorder and concurrently treating another (*e.g.*, preventing Wilson's disease or Alzheimer's while treating cancer).

15

5.6 Therapeutic/Prophylactic Administration

The compounds of structural Formula (I) and/or pharmaceutical compositions thereof may be advantageously used in human medicine. As previously described in Section 4.5, *supra*, compounds of structural Formula (I) and/or pharmaceutical
20 compositions thereof are useful for the treatment and/or prevention of various diseases or disorders characterized by aberrant vascularization, copper metabolism disorders, neurodegenerative disorders, obesity or NF- κ B dysregulation.

When used to treat and/or prevent the above disease or disorders, compounds of structural Formula (I) and/or pharmaceutical compositions may be administered or
25 applied singly, or in combination with other agents. The compounds of structural Formula (I) and/or pharmaceutical compositions thereof may also be administered or applied singly, in combination with other pharmaceutically active agents (*e.g.*, other anti-cancer agents, other anti-angiogenic agents, other chelators such as zinc, penicillamine, *etc.* and other anti-obesity agents), including other compounds of
30 structural Formula (I) and/or pharmaceutical compositions thereof.

Methods of treatment and prophylaxis by administration to a patient of a therapeutically effective amount of a compound of structural Formula (I) and/or

pharmaceutical composition thereof are provided herein. The patient may be an animal, more preferably, a mammal and most preferably, a human.

The present compounds of structural Formula (I) and/or pharmaceutical compositions thereof, are preferably administered orally. The compounds of structural Formula (I) and/or pharmaceutical compositions thereof may also be administered by any other convenient route, for example, by infusion or bolus injection, by absorption through epithelial or mucocutaneous linings (*e.g.*, oral mucosa, rectal and intestinal mucosa, *etc.*). Administration can be systemic or local. Various delivery systems are known, (*e.g.*, encapsulation in liposomes, microparticles, microcapsules, capsules, *etc.*) that can be used to administer a compound of structural Formula (I) and/or pharmaceutical composition thereof. Methods of administration include, but are not limited to, intradermal, intramuscular, intraperitoneal, intravenous, subcutaneous, intranasal, epidural, oral, sublingual, intranasal, intracerebral, intravaginal, transdermal, rectally, by inhalation, or topically, particularly to the ears, nose, eyes, or skin. The preferred mode of administration is left to the discretion of the practitioner and will depend in-part upon the site of the medical condition. In most instances, administration will result in the release of the compounds of structural Formula (I) and/or pharmaceutical compositions thereof into the bloodstream.

In specific embodiments, it may be desirable to administer one or more compounds of structural Formula (I) and/or pharmaceutical composition thereof locally to the area in need of treatment. This may be achieved, for example, and not by way of limitation, by local infusion during surgery, topical application, *e.g.*, in conjunction with a wound dressing after surgery, by injection, by means of a catheter, by means of a suppository, or by means of an implant, said implant being of a porous, non-porous, or gelatinous material, including membranes, such as sialastic membranes, or fibers. In one embodiment, administration can be by direct injection at the site (or former site) of aberrant vascularization (*e.g.*, cancer or arthritis).

In certain embodiments, it may be desirable to introduce one or more compounds of structural Formula (I) and/or pharmaceutical compositions thereof into the central nervous system by any suitable route, including intraventricular, intrathecal and epidural injection. Intraventricular injection may be facilitated by an intraventricular catheter, for example, attached to a reservoir, such as an Ommaya reservoir.

A compound of structural Formula (I) and/or pharmaceutical composition thereof may also be administered directly to the lung by inhalation. For administration by inhalation, a compound of structural Formula (I) and/or pharmaceutical composition thereof may be conveniently delivered to the lung by a number of different devices. For example, a Metered Dose Inhaler ("MDI"), which utilizes canisters that contain a suitable low boiling propellant, (*e.g.*, dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon dioxide or any other suitable gas) may be used to deliver compounds of structural Formula (I) and/or pharmaceutical compositions thereof directly to the lung.

Alternatively, a Dry Powder Inhaler ("DPI") device may be used to administer a compound of structural Formula (I) and/or pharmaceutical thereof to the lung. DPI devices typically use a mechanism such as a burst of gas to create a cloud of dry powder inside a container, which may then be inhaled by the patient. DPI devices are also well known in the art. A popular variation is the multiple dose DPI ("MDDPI") system, which allows for the delivery of more than one therapeutic dose. For example, capsules and cartridges of gelatin for use in an inhaler or insufflator may be formulated containing a powder mix of a compound of structural Formula (I) and a suitable powder base such as lactose or starch for these systems.

Another type of device that may be used to deliver a compound of structural Formula (I) and/or pharmaceutical composition thereof to the lung is a liquid spray device supplied, for example, by Aradigm Corporation, Hayward, CA. Liquid spray systems use extremely small nozzle holes to aerosolize liquid drug formulations that may then be directly inhaled into the lung.

In one embodiment, a nebulizer is used to deliver a compound of structural Formula (I) and/or pharmaceutical composition thereof to the lung. Nebulizers create aerosols from liquid drug formulations by using, for example, ultrasonic energy to form fine particles that may be readily inhaled (see *e.g.*, Verschoyle *et al.*, *British J. Cancer*, 1999, 80, Suppl. 2, 96). Examples of nebulizers include devices supplied by Sheffield Pharmaceuticals, Inc (See, Armer *et al.*, United States Patent No. 5,954,047; van der Linden *et al.*, United States Patent No. 5,950,619; van der Linden *et al.*, United States Patent No. 5,970,974), and Batelle Pulmonary Therapeutics, Columbus, OH.

In another embodiment, an electrohydrodynamic ("EHD") aerosol device is used to deliver a compound of structural Formula (I) and/or pharmaceutical

composition thereof to the lung. EHD aerosol devices use electrical energy to aerosolize liquid drug solutions or suspensions (see *e.g.*, Noakes *et al.*, United States Patent No. 4,765,539). The electrochemical properties of the formulation may be important parameters to optimize when delivering a compound of structural Formula (I) and/or pharmaceutical composition thereof to the lung with an EHD aerosol device and such optimization is routinely performed by one of skill in the art. EHD aerosol devices may more efficiently deliver compounds to the lung than other pulmonary delivery technologies.

In some embodiments, the compounds of structural Formula (I) and/or pharmaceutical compositions thereof can be delivered in a vesicle, in particular a liposome (See, Langer, 1990, *Science* 249:1527-1533; Treat *et al.*, in "Liposomes in the Therapy of Infectious Disease and Cancer," Lopez-Berestein and Fidler (eds.), Liss, New York, pp. 353-365 (1989); see generally "Liposomes in the Therapy of Infectious Disease and Cancer," Lopez-Berestein and Fidler (eds.), Liss, New York, pp.353-365 (1989)).

In other embodiments, the compounds of structural Formula (I) and/or pharmaceutical compositions thereof can be delivered via sustained release systems, preferably oral sustained release systems. In one embodiment, a pump may be used (See, Langer, *supra*, Sefton, 1987, *CRC Crit. Ref. Biomed Eng.* 14:201; Saudek *et al.*, 1989, *N. Engl. J Med.* 321:574).

In yet other embodiments, polymeric materials can be used (see "Medical Applications of Controlled Release," Langer and Wise (eds.), CRC Pres., Boca Raton, Florida (1974); "Controlled Drug Bioavailability," Drug Product Design and Performance, Smolen and Ball (eds.), Wiley, New York (1984); Langer *et al.*, 1983, *J Macromol. Sci. Rev. Macromol Chem.* 23:61; see also Levy *et al.*, 1985, *Science* 228: 190; During *et al.*, 1989, *Ann. Neurol.* 25:351; Howard *et al.*, 1989, *J. Neurosurg.* 71:105). In still other embodiments, polymeric materials are used for oral sustained release delivery. Preferred polymers include sodium carboxymethylcellulose, hydroxypropylcellulose, hydroxypropylmethylcellulose and hydroxyethylcellulose (most preferred, hydroxypropyl methylcellulose). Other preferred cellulose ethers have been described (Alderman, *Int. J. Pharm. Tech. & Prod. Mfr.*, 1984, 5(3) 1-9). Factors affecting drug release are well known to the skilled artisan and have been described in the art (Bamba *et al.*, *Int. J. Pharm.*, 1979, 2, 307).

In still other embodiment, enteric-coated preparations can be used for oral sustained release administration. Preferred coating materials include polymers with a pH-dependent solubility (*i.e.*, pH-controlled release), polymers with a slow or pH-dependent rate of swelling, dissolution or erosion (*i.e.*, time-controlled release),
5 polymers that are degraded by enzymes (*i.e.*, enzyme-controlled release) and polymers that form firm layers that are destroyed by an increase in pressure (*i.e.*, pressure-controlled release).

In still other embodiments, osmotic delivery systems are used for oral sustained release administration (Verma *et al.*, *Drug Dev. Ind. Pharm.*, 2000,
10 26:695-708). In yet other embodiments, OROSTM osmotic devices are used for oral sustained release delivery devices (Theeuwes *et al.*, United States Patent No. 3,845,770; Theeuwes *et al.*, United States Patent No. 3,916,899).

In still other embodiments, a controlled-release system can be placed in proximity of the target of the compounds and/or pharmaceutical composition of the invention, thus requiring only a fraction of the systemic dose (See, *e.g.*, Goodson, in
15 "Medical Applications of Controlled Release," *supra*, vol. 2, pp. 115-138 (1984). Other controlled-release systems discussed in Langer, 1990, *Science* 249:1527-1533 may also be used.

20 5.7 Pharmaceutical Compositions

The present pharmaceutical compositions contain a therapeutically effective amount of one or more compounds of structural Formula (I), preferably in purified form, together with a suitable amount of a pharmaceutically acceptable vehicle, so as to provide the form for proper administration to a patient. When administered to a
25 patient, the compounds of structural Formula (I) and pharmaceutically acceptable vehicles are preferably sterile. Water is a preferred vehicle when the compound of structural Formula (I) is administered intravenously. Saline solutions and aqueous dextrose and glycerol solutions can also be employed as liquid vehicles, particularly for injectable solutions. Suitable pharmaceutical vehicles also include excipients such
30 as starch, glucose, lactose, sucrose, gelatin, malt, rice, flour, chalk, silica gel, sodium stearate, glycerol monostearate, talc, sodium chloride, dried skim milk, glycerol, propylene, glycol, water, ethanol and the like. The present pharmaceutical compositions, if desired, can also contain minor amounts of wetting or emulsifying

agents, or pH buffering agents. In addition, auxiliary, stabilizing, thickening, lubricating and coloring agents may be used.

Pharmaceutical compositions comprising a compound of structural Formula (I) may be manufactured by means of conventional mixing, dissolving, granulating, dragee-making, levigating, emulsifying, encapsulating, entrapping or lyophilizing processes. Pharmaceutical compositions may be formulated in conventional manner using one or more physiologically acceptable carriers, diluents, excipients or auxiliaries, which facilitate processing of compounds of structural Formula (I) into preparations which can be used pharmaceutically. Proper formulation is dependent upon the route of administration chosen.

The present pharmaceutical compositions can take the form of solutions, suspensions, emulsion, tablets, pills, pellets, capsules, capsules containing liquids, powders, sustained-release formulations, suppositories, emulsions, aerosols, sprays, suspensions, or any other form suitable for use. In one embodiment, the pharmaceutically acceptable vehicle is a capsule (see *e.g.*, Grosswald *et al.*, United States Patent No. 5,698,155). Other examples of suitable pharmaceutical vehicles have been described in the art (see Remington: The Science and Practice of Pharmacy, Philadelphia College of Pharmacy and Science, 20th Edition, 2000).

For topical administration a compound of structural Formula (I) may be formulated as solutions, gels, ointments, creams, suspensions, *etc.* as is well-known in the art.

Systemic formulations include those designed for administration by injection, *e.g.*, subcutaneous, intravenous, intramuscular, intrathecal or intraperitoneal injection, as well as those designed for transdermal, transmucosal, oral or pulmonary administration. Systemic formulations may be made in combination with a further active agent that improves mucociliary clearance of airway mucus or reduces mucous viscosity. These active agents include, but are not limited to, sodium channel blockers, antibiotics, N-acetyl cysteine, homocysteine and phospholipids.

In some embodiments, the compounds of structural Formula (I) are formulated in accordance with routine procedures as a pharmaceutical composition adapted for intravenous administration to human beings. Typically, compounds of structural Formula (I) for intravenous administration are solutions in sterile isotonic aqueous buffer. For injection, a compound of structural Formula (I) may be formulated in aqueous solutions, preferably in physiologically compatible buffers such as Hanks'

solution, Ringer's solution, or physiological saline buffer. The solution may contain formulatory agents such as suspending, stabilizing and/or dispersing agents. When necessary, the pharmaceutical compositions may also include a solubilizing agent. Pharmaceutical compositions for intravenous administration may optionally include a
5 local anesthetic such as lignocaine to ease pain at the site of the injection. Generally, the ingredients are supplied either separately or mixed together in unit dosage form, for example, as a lyophilized powder or water free concentrate in a hermetically sealed container such as an ampoule or sachette indicating the quantity of active agent. When the compound of structural Formula (I) is administered by infusion, it
10 can be dispensed, for example, with an infusion bottle containing sterile pharmaceutical grade water or saline. When the compound of structural Formula (I) is administered by injection, an ampoule of sterile water for injection or saline can be provided so that the ingredients may be mixed prior to administration.

For transmucosal administration, penetrants appropriate to the barrier to be
15 permeated are used in the formulation. Such penetrants are generally known in the art.

Pharmaceutical compositions for oral delivery may be in the form of tablets, lozenges, aqueous or oily suspensions, granules, powders, emulsions, capsules, syrups, or elixirs, for example. Orally administered pharmaceutical compositions
20 may contain one or more optionally agents, for example, sweetening agents such as fructose, aspartame or saccharin; flavoring agents such as peppermint, oil of wintergreen, or cherry coloring agents and preserving agents, to provide a pharmaceutically palatable preparation. Moreover, where in tablet or pill form, the pharmaceutical compositions may be coated to delay disintegration and absorption in
25 the gastrointestinal tract, thereby providing a sustained action over an extended period of time. Selectively permeable membranes surrounding an osmotically active driving compound are also suitable for orally administered compounds of structural Formula (I). In these later platforms, fluid from the environment surrounding the capsule is imbibed by the driving compound, which swells to displace the agent or agent
30 composition through an aperture. These delivery platforms can provide an essentially zero order delivery profile as opposed to the spiked profiles of immediate release formulations. A time delay material such as glycerol monostearate or glycerol stearate may also be used. Oral compositions can include standard vehicles such as

mannitol, lactose, starch, magnesium stearate, sodium saccharine, cellulose, magnesium carbonate, *etc.* Such vehicles are preferably of pharmaceutical grade.

For oral liquid preparations such as, for example, suspensions, elixirs and solutions, suitable carriers, excipients or diluents include water, saline,
5 alkylene glycols (*e.g.*, propylene glycol), polyalkylene glycols (*e.g.*, polyethylene glycol) oils, alcohols, slightly acidic buffers between pH 4 and pH 6 (*e.g.*, acetate, citrate, ascorbate at between about 5.0 mM to about 50.0 mM) *etc.* Additionally, flavoring agents, preservatives, coloring agents, bile salts, acylcarnitines and the like may be added.

10 For buccal administration, the pharmaceutical compositions may take the form of tablets, lozenges, *etc.* formulated in conventional manner.

Liquid drug formulations suitable for use with nebulizers and liquid spray devices and EHD aerosol devices will typically include a compound of structural Formula (I) with a pharmaceutically acceptable vehicle. Preferably, the
15 pharmaceutically acceptable vehicle is a liquid such as alcohol, water, polyethylene glycol or a perfluorocarbon. Optionally, another material may be added to alter the aerosol properties of the solution or suspension of compounds of the invention. Preferably, this material is liquid such as an alcohol, glycol, polyglycol or a fatty acid. Other methods of formulating liquid drug solutions or suspension suitable for use in
20 aerosol devices are known to those of skill in the art (see, *e.g.*, Biesalski, United States Patent No. 5,112,598; Biesalski, United States Patent No. 5,556,611).

A compound of structural Formula (I) may also be formulated in rectal or vaginal pharmaceutical compositions such as suppositories or retention enemas, *e.g.*, containing conventional suppository bases such as cocoa butter or other glycerides.

25 In addition to the formulations described previously, a compound of structural Formula (I) may also be formulated as a depot preparation. Such long acting formulations may be administered by implantation (for example subcutaneously or intramuscularly) or by intramuscular injection. Thus, for example, a compound of structural Formula (I) may be formulated with suitable polymeric or hydrophobic
30 materials (for example, as an emulsion in an acceptable oil) or ion exchange resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt.

When a compound of structural Formula (I) is acidic, it may be included in any of the above-described formulations as the free acid, a pharmaceutically acceptable salt, a solvate or hydrate. Pharmaceutically acceptable salts substantially

retain the activity of the free acid, may be prepared by reaction with bases and tend to be more soluble in aqueous and other protic solvents than the corresponding free acid form.

5 5.8 Therapeutic Doses

A compound of structural Formula (I), and/or pharmaceutical composition thereof, will generally be used in an amount effective to achieve the intended purpose. For use to treat or prevent diseases or disorders characterized by aberrant vascularization, copper metabolism disorders, neurodegenerative disorders and obesity the compounds of structural Formula (I) and/or pharmaceutical compositions thereof, are administered or applied in a therapeutically effective amount.

The amount of a compound of structural Formula (I) that will be effective in the treatment of a particular disorder or condition disclosed herein will depend on the nature of the disorder or condition and can be determined by standard clinical techniques known in the art, as previously described. In addition, *in vitro* or *in vivo* assays may optionally be employed to help identify optimal dosage ranges. The amount of a compound of structural Formula (I) administered will, of course, be dependent on, among other factors, the subject being treated, the weight of the subject, the severity of the affliction, the manner of administration and the judgment of the prescribing physician.

For example, the dosage may be delivered in a pharmaceutical composition by a single administration, by multiple applications or controlled release. In one embodiment, the compounds of structural Formula (I) are delivered by oral sustained release administration. Preferably, in this embodiment, the compounds of structural Formula (I) are administered twice per day (more preferably, once per day). Dosing may be repeated intermittently, may be provided alone or in combination with other drugs and may continue as long as required for effective treatment of the disease state or disorder.

Suitable dosage ranges for oral administration depend on potency, but are generally between about 0.001 mg to about 200 mg of a compound of structural Formula (I) per kilogram body weight. Dosage ranges may be readily determined by methods known to the artisan of ordinary skill the art.

Suitable dosage ranges for intravenous (i.v.) administration are about 0.01 mg to about 100 mg per kilogram body weight. Suitable dosage ranges for intranasal

administration are generally about 0.01 mg/kg body weight to about 1 mg/kg body weight. Suppositories generally contain about 0.01 milligram to about 50 milligrams of a compound of structural Formula (I) per kilogram body weight and comprise active ingredient in the range of about 0.5% to about 10% by weight. Recommended dosages for intradermal, intramuscular, intraperitoneal, subcutaneous, epidural, sublingual or intracerebral administration are in the range of about 0.001 mg to about 200 mg per kilogram of body weight. Effective doses may be extrapolated from dose-response curves derived from *in vitro* or animal model test systems. Such animal models and systems are well-known in the art.

The compounds of structural Formula (I) are preferably assayed *in vitro* and *in vivo*, for the desired therapeutic or prophylactic activity, prior to use in humans. For example, *in vitro* assays can be used to determine whether administration of a specific compound of structural Formula (I) or a combination of compounds of structural Formula (I) is preferred for treating or preventing diseases or disorders characterized by aberrant vascularization, copper metabolism disorders, neurodegenerative disorders and obesity. The compounds of structural Formula (I) may also be demonstrated to be effective and safe using animal model systems.

Preferably, a therapeutically effective dose of a compound of structural Formula (I) described herein will provide therapeutic benefit without causing substantial toxicity. Toxicity of compounds of structural Formula (I) may be determined using standard pharmaceutical procedures and may be readily ascertained by the skilled artisan. The dose ratio between toxic and therapeutic effect is the therapeutic index. A compound of structural Formula (I) will preferably exhibit particularly high therapeutic indices in treating disease and disorders. The dosage of a compound of structural Formula (I) described herein will preferably be within a range of circulating concentrations that include an effective dose with little or no toxicity.

5.9 Combination Therapy

In certain embodiments of the present invention, the compounds of structural Formula (I) and/or pharmaceutical compositions thereof can be used in combination therapy with at least one other therapeutic agent or with radiation therapy. The compound of structural Formula (I) and/or pharmaceutical composition thereof and the other therapeutic agent can act additively or, more preferably, synergistically. In some embodiments, a compound of structural Formula (I) and/or pharmaceutical

composition thereof is administered concurrently with the administration of another therapeutic agent, which may be part of the same pharmaceutical composition as the compound of structural Formula (I) or a different pharmaceutical composition. In other embodiments, a pharmaceutical composition of structural Formula (I) is
5 administered prior or subsequent to administration of another therapeutic agent.

In some embodiments, the compounds of structural Formula (I) and/or pharmaceutical compositions thereof can be used in combination therapy with other chemotherapeutic agents (*e.g.*, alkylating agents (*e.g.*, nitrogen mustards (*e.g.*, cyclophosphamide, ifosfamide, mechlorethamine, melphalen, chlorambucil,
10 hexamethylmelamine, thiotepa), alkyl sulfonates (*e.g.*, busulfan), nitrosoureas, triazines) antimetabolites (*e.g.*, folic acid analogs, pyrimidine analogs (*e.g.*, fluorouracil, floxuridine, cytosine arabinoside, *etc.*), purine analogs (*e.g.*, mercaptopurine, thioguanine, pentostatin, *etc.*), natural products (*e.g.*, vinblastine, vincristine, etoposide, tertiposide, dactinomycin, daunorubicin, doxorubicin,
15 bleomycin, mithramycin, mitomycin C, L-asparaginase, interferon alpha), platinum coordination complexes (*e.g.*, *cis*-platinum, carboplatin, *etc.*), mitoxantrone, hydroxyurea, procarbazine, hormones and antagonists (*e.g.*, prednisone, hydroxyprogesterone caproate, medroxyprogesterone acetate, megestrol acetate, diethylstilbestrol, ethinyl estradiol, tamoxifen, testosterone propionate,
20 fluoxymesterone, flutamide, leuprolide, *etc.*), anti-angiogenesis agents or inhibitors (*e.g.*, angiostatin, retinoic acids and paclitaxel, estradiol derivatives, thiazolopyrimidine derivatives, *etc.*), apoptosis-inducing agents (*e.g.*, antisense nucleotides that block oncogenes which inhibit apoptosis, tumor suppressors, TRAIL, TRAIL polypeptide, Fas-associated factor 1, interleukin-1 β -converting enzyme,
25 phosphotyrosine inhibitors, RXR retinoid receptor agonists, carbostyryl derivatives, *etc.*), chelators (penicillamine, zinc, trientine, *etc.*) and other anti-obesity agents.

5.10 Therapeutic Kits

Therapeutic kits comprising the compounds of structural Formula (I) and/or
30 pharmaceutical compositions thereof are also provided herein. The therapeutic kits may also contain other compounds (*e.g.*, chemotherapeutic agents, natural products, hormones or antagonists, anti-angiogenesis agents or inhibitors, apoptosis-inducing agents or chelators) and/or pharmaceutical compositions thereof.

Therapeutic kits may have a single container which contains the compound of structural Formula (I) and/or pharmaceutical compositions thereof with or without other components (e.g., other compounds and/or pharmaceutical compositions thereof) or may have distinct container for each component. Preferably, therapeutic kits include a compound of structural Formula (I) and/or a pharmaceutical composition thereof packaged for use in combination with the co-administration of a second compound (preferably, a chemotherapeutic agent, a natural product, a hormone or antagonist, an anti-angiogenesis agent or inhibitor, an apoptosis-inducing agent or a chelator) or a pharmaceutical composition thereof. The components of the kit may be pre-complexed or each component may be in a separate distinct container prior to administration to a patient.

The components of the kit may be provided in one or more liquid solutions, preferably, an aqueous solution, more preferably, a sterile aqueous solution. The components of the kit may also be provided as solids, which may be converted into liquids by addition of suitable solvents, which are preferably provided in another distinct container.

The container of a therapeutic kit may be a vial, test tube, flask, bottle, syringe, or any other means of enclosing a solid or liquid. Usually, when there is more than one component, the kit will contain a second vial or other container, which allows for separate dosing. The kit may also contain another container for a pharmaceutically acceptable liquid.

Preferably, a therapeutic kit will contain apparatus (e.g., one or more needles, syringes, eye droppers, pipette, etc.), which enables administration of the components of the kit.

6. Examples

The invention is further defined by reference to the following examples, which describe in detail, preparation of compounds of the invention and methods for assaying for biological activity. It will be apparent to those skilled in the art that many modifications, both to materials and methods, may be practiced without departing from the scope of the invention.

6.1 Example 1: General Procedure for Synthesis of Tetrathiotungstate Derivatives

The quaternary ammonium hydroxide (2 eq.), as a commercially-available aqueous solution, was added to ammonium tetrathiotungstate (1 eq.) and deionized water was added until all the solid material was dissolved. The solution was placed on a rotary evaporator under vacuum (*ca.* 5-10 torr) with the bath at 20 °C for one hour and the water was replaced as needed to maintain a constant volume. The reaction mixture was then allowed to evaporate to dryness and the resulting yellow solid was recrystallized from deionized water and isopropanol. The solid was collected by filtration, washed with isopropanol and diethyl ether, and then dried under high vacuum for 24 hours in a vacuum desiccator in the presence of P₂O₅.

6.2 Example 2: General Procedure for Synthesis of Tetrathiotungstate Derivatives

The quaternary ammonium halide (2 eq.) as a solid was added to a suspension of ammonium tetrathiotungstate (1 eq.) in dry acetonitrile (5 mL per mmol of tetrathiotungstate) and the resulting mixture was stirred at room temperature under nitrogen for 18 hours. If this procedure resulted in a precipitate, the solid was collected by filtration, washed with isopropanol and diethyl ether and was recrystallized from deionized water and isopropanol. The yellow crystals were collected by filtration, washed with isopropanol and diethyl ether and dried under high vacuum for 24 hours in a vacuum desiccator in the presence of P₂O₅. If the solution remained clear, the solvent was removed *in vacuo*, the residue was taken up in dichloromethane, washed three times with water, once with brine, dried (Na₂SO₄), and the solution was concentrated. The resulting oil or solid was dried under high vacuum for 24 hours in a vacuum desiccator in the presence of P₂O₅.

6.3 Example 3: General Procedure for Synthesis of Tetrathiotungstate Derivatives

The quaternary ammonium halide (2 eq.) as a solution in deionized water (10 mL per mmol of tetrathiotungstate) was added to a suspension of ammonium tetrathiotungstate (1 eq.) in dry acetonitrile (20 mL per mmol of tetrathiotungstate) and the resulting mixture was stirred at room temperature for 18 hours. If this

procedure resulted in a precipitate, the solid was collected by filtration, washed with water, isopropanol and diethyl ether, and then dried under high vacuum for 24 hours in a vacuum desiccator in the presence of P_2O_5 . If the solution remained clear, the reaction mixture was first filtered, and the filtrate was concentrated *in vacuo*. The resulting solid was recrystallized from deionized water and isopropanol, the yellow crystals collected by filtration, washed with isopropanol and diethyl ether and then dried under high vacuum for 24 hours in a vacuum desiccator in the presence of P_2O_5 .

6.4 Example 4: Tetrathiotungstate, bis(choline)

This compound was prepared from ammonium tetrathiotungstate (158 mg, 0.454 mmol) and a 50% by weight aqueous solution of choline hydroxide (222 mg, 0.916 mmol) according to the procedure of Example 1, which afforded 151 mg (64%) of the title compound as bright yellow crystals: IR (KBr, cm^{-1}) 3402, 459; 1H NMR (300 MHz, DMSO- d_6) δ 5.21 (t, $J = 4.8$ Hz, 2H), 3.88-3.81 (m, 4H), 3.46-3.43 (m, 4H), 3.14 (s, 18H); ^{13}C NMR (75 MHz, DMSO- d_6) δ 666.8 (2C), 55.2 (2C), 53.1 (6C); ES MS m/z (choline) $^+$ 104.3; UV (H_2O) 393.5 nm ($\epsilon = 16730$). Anal. Calcd for $C_{10}H_{28}N_2O_2S_4W$: C, 23.08; H, 5.42; N, 5.38; S, 24.65. Found: C, 23.17; H, 5.28; N, 5.43; S, 24.87.

6.5 Example 5: Tetrathiotungstate, bis(triethylmethyl ammonium)

This compound was prepared from ammonium tetrathiotungstate (164 mg, 0.471 mmol) and a 20% by weight aqueous solution of triethylmethylammonium hydroxide (627 mg, 0.941 mmol) according to the procedure of Example 1, which provided 147 mg (61%) of the title compound as bright yellow crystals: IR (KBr, cm^{-1}) 460; 1H NMR (300 MHz, DMSO- d_6) δ 3.29 (q, $J = 6.9$ Hz, 12H), 2.91 (s, 6H), 1.21 (t, $J = 6.9$ Hz, 18H); ^{13}C NMR (75 MHz, DMSO- d_6) δ 55.0 (6C), 46.0 (2C), 7.5 (6C); ES MS m/z (triethylmethyl ammonium) $^+$ 116.4; UV (H_2O) 393.5 nm ($\epsilon = 16730$). Anal. Calcd for $C_{14}H_{36}N_2S_4W$: C, 30.88; H, 6.66; N, 5.14; S, 23.55. Found: C, 30.87; H, 6.33; N, 5.18; S, 23.77.

30

6.6 Example 6: Tetrathiotungstate, bis(triethylphenyl ammonium)

This compound was prepared from ammonium tetrathiotungstate (155 mg, 0.444 mmol) and a 10% by weight aqueous solution of triethylphenylammonium

hydroxide (1.74 g, 0.889 mmol) according to the procedure of Example 1, which provided 198 mg (69%) of the title compound as bright yellow crystals: IR (KBr, cm⁻¹) 455; ¹H NMR (300 MHz, DMSO-d₆) δ 7.92 (d, *J* = 8.4 Hz, 4H), 7.71-7.57 (m, 6H), 3.91 (q, *J* = 7.1 Hz, 12H), 1.06 (t, *J* = 7.1 Hz, 18H); ¹³C NMR (75 MHz, DMSO-d₆) δ 141.7 (2C), 130.4 (4C), 130.0 (2C), 122.6 (4C), 55.3 (6C), 7.8 (6C); ES MS *m/z* (triethylphenyl ammonium)⁺ 178.4; UV (H₂O) 393.5 nm (ε = 15600). Anal. Calcd for C₂₄H₄₀N₂S₄W: C, 43.11; H, 6.03; N, 4.19; S, 19.18. Found: C, 42.99; H, 5.73; N, 4.25; S, 19.31.

10 6.7 Example 7: Tetrathiotungstate, bis(1,4-dimethylpyridinium)

This compound was prepared from ammonium tetrathiotungstate (163 mg, 0.467 mmol) and 1,4-dimethylpyridinium iodide (221 mg, 0.940 mmol) according to the procedure of Example 2, which provided 143 mg (58%) of the title compound as bright yellow crystals: IR (KBr, cm⁻¹) 458; ¹H NMR (300 MHz, DMSO-d₆) δ 8.88 (d, *J* = 6.4 Hz, 4H), 7.96 (d, *J* = 6.4 Hz, 4H), 4.32 (s, 6H), 2.60 (s, 6H); ¹³C NMR (75 MHz, DMSO-d₆) δ 158.1 (2C), 144.8 (4C), 128.0 (4C), 47.1 (2C), 21.4 (2C); ES MS *m/z* (1,4-dimethylpyridinium)⁺ 108.3; UV (H₂O) 393.5 nm (ε = 16030). Anal. Calcd for C₁₄H₂₀N₂S₄W: C, 31.82; H, 3.81; N, 5.30; S, 24.27. Found: C, 31.67; H, 3.77; N, 5.32; S, 24.13.

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6.8 Example 8: Tetrathiotungstate, bis(1,1-dimethylpyrrolidinium)

This compound was prepared from ammonium tetrathiotungstate (300 mg, 0.861 mmol) and 1,1-dimethylpyrrolidinium iodide (400 mg, 1.76 mmol) according to the procedure of Example 3, which provided 223 mg (51%) of the title compound as bright yellow crystals: IR (KBr, cm⁻¹) 455; ¹H NMR (300 MHz, DMSO-d₆) δ 3.53-3.47 (m, 8H), 3.13 (s, 12H), 2.14-2.08 (m, 8H); ¹³C NMR (75 MHz, DMSO-d₆) δ 64.8 (4C), 51.0 (4C), 21.5 (4C); ES MS *m/z* (1,1-dimethylpyrrolidinium)⁺ 100.3; UV (H₂O) 393.5 nm (ε = 16950). Anal. Calcd for C₁₂H₂₈N₂S₄W: C, 28.12; H, 5.51; N, 5.47; S, 25.03. Found: C, 27.90; H, 5.47; N, 5.56; S, 25.01.

30

6.9 Example 9: Tetrathiotungstate, bis(trimethylphenylammonium)

This compound was prepared from ammonium tetrathiotungstate (167 mg, 0.479 mmol) and phenyltrimethyl-ammonium chloride (166 mg, 0.968 mmol) according to the procedure of Example 2, which provided 139 mg (50%) of the title compound as bright yellow crystals: IR (KBr, cm⁻¹) 459; ¹H NMR (300 MHz, DMSO-d₆) δ 7.99 (d, *J* = 8.2 Hz, 4H), 7.68-7.55 (m, 6H), 3.64 (s, 18H); ¹³C NMR (75 MHz, DMSO-d₆) δ 147.3 (2C), 130.1 (4C), 130.0 (2C), 120.5 (4C), 56.4 (6C); ES MS *m/z* (trimethylphenylammonium)⁺ 136.2; UV (H₂O) 394.0 nm (ε = 15630). Anal. Calcd for C₁₈H₂₈N₂S₄W: C, 36.99; H, 4.83; N, 4.79; S, 21.94. Found: C, 36.88; H, 4.72; N, 4.90; S, 21.92.

6.10 Example 10: Tetrathiotungstate, bis(acetylcholine)

This compound was prepared from ammonium tetrathiotungstate (171 mg, 0.491 mmol) and acetylcholine chloride (179 mg, 0.987 mmol) according to the procedure of Example 2, which provided 163 mg (55%) of the title compound as bright yellow crystals: IR (KBr, cm⁻¹) 1749, 1729, 473, 456; ¹H NMR (300 MHz, DMSO-d₆) δ 4.47-4.41 (m, 4H), 3.72-3.69 (m, 4H), 3.16 (s, 18H), 2.07 (s, 6H); ¹³C NMR (75 MHz, DMSO-d₆) δ 169.9 (2C), 63.8 (2C), 57.9 (2C), 53.0 (6C), 20.7 (2C); ES MS *m/z* (acetylcholine)⁺ 146.4; UV (H₂O) 393.5 nm (ε = 15400). Anal. Calcd for C₁₄H₃₂N₂O₄S₄W: C, 27.82; H, 5.34; N, 4.63; S, 21.22. Found: C, 27.62; H, 5.12; N, 4.68; S, 20.71.

6.11 Example 11: Tetrathiotungstate, bis[alkyldimethyl(phenylmethyl) ammonium]

This compound was prepared from ammonium tetrathiotungstate (320 mg, 0.920 mmol) and benzalkonium chloride (664 mg, 1.84 mmol) according to the procedure of Example 2, which afforded 651 mg (74%) of the title compound as a thick, red oil: IR (film, cm⁻¹) 466; ¹H NMR (300 MHz, DMSO-d₆) δ 7.59-7.48 (m, 10H), 4.56 (s, 4H), 3.31-3.23 (m, 4H), 2.97 (s, 12H), 1.84-1.72 (m, 4H), 1.32-1.22 (m, 40H), 0.88-0.82 (m, 6H); ES MS *m/z* [dodecyldimethyl(phenylmethyl) ammonium]⁺ 304.7, [tetradecyldimethyl(phenylmethyl) ammonium]⁺ 332.7; UV (DMSO) 399.0 nm (ε = 10400).

6.12 Example 12: Tetrathiotungstate, suberyldicholine

This compound was prepared from ammonium tetrathiotungstate (299 mg, 0.860 mmol) and suberyldicholine diiodide (516 mg, 0.860 mmol) according to the procedure of Example 2, which afforded 115 mg (20%) of the title compound as
5 bright yellow crystals: IR (KBr, cm⁻¹) 1733, 1719, 455; ¹H NMR (300 MHz, DMSO-d₆) δ 4.48-4.42 (m, 4H), 3.73-3.69 (m, 4H), 3.17 (s, 18H), 2.35 (t, *J* = 7.4 Hz, 4H), 1.59-1.48 (m, 4H), 1.32-1.26 (m, 4H); ¹³C NMR (75 MHz, DMSO-d₆) δ 172.3 (2C), 63.7 (2C), 57.7 (2C), 52.8 (6C), 33.2 (2C), 28.0 (2C), 23.9 (2C); UV (H₂O) 394.0 nm (ε = 15570)

10

6.13 Example 13: Tetrathiotungstate, pentane-1,5-bis(trimethylammonium)

This compound was prepared from ammonium tetrathiotungstate (140 mg, 0.402 mmol) and *N,N*-pentamethylenebis(trimethylammonium iodide) (195 mg, 0.442 mmol) according to the procedure of Example 3, which afforded 109 mg (54%) of the
15 title compound as a bright yellow powder: IR (KBr, cm⁻¹) 456; ¹H NMR (300 MHz, D₂O) δ 3.34-3.26 (m, 4H), 3.07 (s, 18H), 1.89-1.77 (m, 4H), 1.45-1.34 (m, 2H); UV (H₂O) 394.0 nm (ε = 15950). Anal. Calcd for C₁₁H₂₈N₂S₄W: C, 26.40; H, 5.64; N, 5.60; S, 25.63. Found: C, 26.60; H, 5.26; N, 5.75; S, 24.64.

20 6.14 Example 14: Tetrathiotungstate, butane-1,4-bis(trimethylammonium)

This compound was prepared from ammonium tetrathiotungstate (200 mg, 0.574 mmol) and *N,N*-tetramethylenebis(trimethylammonium iodide) (271 mg, 0.632 mmol) according to the procedure of Example 3 and afforded 185 mg (66%) of the
25 title compound as a bright yellow powder: IR (KBr, cm⁻¹) 456; ¹H NMR (300 MHz, D₂O) δ 3.45-3.35 (m, 4H), 3.11 (s, 18H), 1.92-1.82 (m, 4H); UV (H₂O) 394.0 nm (ε = 15990). Anal. Calcd for C₁₀H₂₆N₂S₄W: C, 24.69; H, 5.39; N, 5.76; S, 26.37. Found: C, 24.77; H, 5.35; N, 5.85; S, 25.80.

30 6.15 Example 15: Tetrathiotungstate, propane-1,3-bis(trimethylammonium)

This compound was prepared from ammonium tetrathiotungstate (201 mg, 0.578 mmol) and *N,N*-trimethylenebis(trimethylammonium iodide) (263 mg, 0.635 mmol) according to the procedure of Example 3, and afforded 192 mg (70%) of the
title compound as a bright yellow powder: IR (KBr, cm⁻¹) 456; UV (H₂O) 393.5 nm

($\epsilon = 16190$). Anal. Calcd for $C_9H_{24}N_2S_4W$: C, 22.88; H, 5.12; N, 5.93; S, 27.15.

Found: C, 22.94; H, 5.01; N, 6.01; S, 26.79.

6.16 Example 15: Tetrathiotungstate, ethylenebis(trimethylammonium)

5 This compound was prepared from ammonium tetrathiotungstate (200 mg, 0.573 mmol) and ethylenebis(trimethylammonium iodide) (249 mg, 0.623 mmol) according to the procedure of Example 3, and afforded 171 mg (65%) of the title compound as a bright yellow powder: IR (KBr, cm^{-1}) 459; UV (H_2O) 393.5 nm ($\epsilon = 15720$). Anal. Calcd for $C_8H_{22}N_2S_4W$: C, 20.96; H, 4.84; N, 6.11; S, 27.98. Found: C, 20.88; H, 4.71; N, 6.21; S, 27.39.

6.17 Example 17: Tetrathiotungstate, bis(*N*-benzyl-2-phenylethyl ammonium)

This compound was prepared from ammonium tetrathiotungstate (295 mg, 0.848 mmol) and *N*-benzyl-2-phenylethylammonium chloride (422 mg, 1.70 mmol) according to the procedure of Example 3, but with the addition of 6 mL of deionized water, and afforded 317 mg (51%) of the title compound as an orange solid: IR (KBr, cm^{-1}) 455; 1H NMR (300 MHz, DMSO- d_6) δ 8.83 (br s, 4H), 7.57-7.52 (m, 4H), 7.48-7.40 (m, 6H), 7.36-7.23 (m, 10H), 4.22 (s, 4H), 3.21-3.15 (m, 4H), 3.03-2.95 (m, 4H); ^{13}C NMR (75 MHz, DMSO- d_6) δ 137.4 (2C), 132.5 (2C), 130.0 (4C), 128.9 (2C), 128.72 (8C), 128.68 (4C), 126.8 (2C), 50.6 (2C), 48.1 (2C), 31.8 (2C); ES MS m/z [*N*-benzyl-2-phenylethyl ammonium] $^+$ 212.4; UV (DMSO) 399.5 nm ($\epsilon = 16270$).

6.18 Example 18: Tetrathiotungstate, bis(1-ethyl-3-methyl-1*H*-imidazolium)

This compound was prepared from tetrathiotungstate, bis(ammonium) (0.400 g, 1.15 mmol) and 1-ethyl-3-methyl-1*H*-imidazolium chloride (0.354 g, 2.41 mmol) according to the procedure of Example 3 giving the title compound (0.217 g, 35%) as a bright yellow solid: IR (KBr pellet, cm^{-1}) 3438, 3068, 1569, 1560, 1169, 450; 1H NMR (300 MHz, DMSO- d_6) δ 9.22 (s, 1 H), 7.78 (s, 1 H), 7.70 (s, 1 H), 4.21 (q, 2 H, $J = 7.3$ Hz), 3.31 (s, 3 H), 1.42 (t, 3 H, $J = 7.3$ Hz); ^{13}C NMR (75 MHz, DMSO- d_6) δ 136.4, 123.4, 121.8, 44.0, 35.6, 15.1; MS m/z ($C_6H_{11}N_2$) $^+$ 111.3; UV (H_2O) 394 nm ($\epsilon = 15,891$); Anal. calcd for $C_{12}H_{22}N_4WS_4$: C, 26.97; H, 4.15; N, 10.48; S, 24.00. Found: C, 26.91, H, 3.92, N, 10.55; S, 23.67.

6.19 Example 19: Tetrathiotungstate, bis(benzyltrimethylammonium)

This compound was prepared from tetrathiotungstate, bis(ammonium) (0.200 g, 0.574 mmol) and benzyltrimethylammonium hydroxide (0.48 g of a 40% aqueous solution, 1.15 mmol) according to the procedure of Example 1 to provide giving the
 5 title compound (0.246 g, 70 %) as a bright yellow solid: IR (KBr pellet, cm^{-1}) 3446, 2999, 1456, 458; ^1H NMR (300 MHz, DMSO-d_6) δ 7.53-7.55 (m, 10 H), 4.56 (s, 4 H), 3.05 (s, 18 H); ^{13}C NMR (75 MHz, DMSO-d_6) δ 132.7, 130.2, 128.8, 128.3, 67.7, 51.7 (t); MS m/z ($\text{C}_{10}\text{H}_{16}\text{N}^+$) 150.3; UV (H_2O) 394 nm ($\epsilon = 15,027$); Anal. calcd for $\text{C}_{20}\text{H}_{32}\text{N}_2\text{WS}_4$: C, 39.21; H, 5.27; N, 4.57; S, 20.94. Found: C, 39.28, H, 4.88, N,
 10 4.65; S, 20.89.

6.20 Example 20: Tetrathiotungstate, bis(2-hydroxymethyl-1-methylpyridinium)

This compound was prepared from tetrathiotungstate, bis(ammonium) (0.200 g, 0.574 mmol) and 2-pyridinealdoxime methochloride (0.198 g, 1.15 mmol) according to the procedure of Example 3 to give the title compound (0.198 g, 59%) as a bright yellow solid: IR (KBr pellet, cm^{-1}) 3077, 1508, 1005, 455; ^1H NMR (300 MHz, DMSO-d_6) δ 9.04 (d, 1 H, $J = 5.9$ Hz), 8.68 (s, 1 H), 8.55 (app t, 1 H), 8.37 (d, 1 H, $J = 8.0$ Hz), 8.07 (app t, 1 H), 4.39 (s, 3 H); ^{13}C NMR (75 MHz, DMSO-d_6) δ
 20 147.3, 146.7, 144.8, 141.7, 127.1, 124.7, 46.1; MS m/z ($\text{C}_7\text{H}_9\text{N}_2\text{O}^+$) 137.2; UV (H_2O) 394 nm ($\epsilon = 15380$); Anal. calcd for $\text{C}_{14}\text{H}_{18}\text{N}_4\text{O}_2\text{WS}_4$: C, 28.67; H, 3.09; N, 9.55; S, 21.87. Found: C, 28.51, H, 2.87, N, 9.63; S, 21.55.

6.21 Example 21: Tetrathiotungstate, bis(acetyl- β -methylcholine)

This compound was prepared from tetrathiotungstate, bis(ammonium) (0.200 g, 0.574 mmol) and acetyl- β -methylcholine chloride (0.235 g, 1.20 mmol) according to the procedure of Example 3 giving the title compound (0.115 g, 32%) as a bright yellow solid: IR (KBr pellet, cm^{-1}) 3452, 3008, 1735, 1252, 454; ^1H NMR (300 MHz, DMSO-d_6) δ 5.27 (m, 1 H), 3.58-3.74 (m, 2 H), 3.13 (s, 9 H), 2.06 (s, 3 H), 1.24 (d, 3
 30 H, $J = 6.3$ Hz); ^{13}C NMR (75 MHz, DMSO-d_6) δ 169.5, 67.6, 65.2, 53.2, 21.1, 18.5; MS m/z ($\text{C}_8\text{H}_{18}\text{NO}_2^+$) 160.3; UV (H_2O) 394 nm ($\epsilon = 15831$); Anal. calcd for $\text{C}_{16}\text{H}_{36}\text{N}_2\text{O}_4\text{S}_4\text{W}$: C, 30.38; H, 5.74; N, 4.43; S, 20.28. Found: C, 30.10, H, 5.62, N, 4.47; S, 20.47.

6.22 Example 22: Tetrathiotungstate, (succinylcholine)

This compound was prepared from tetrathiotungstate, bis(ammonium) (0.400 g, 1.15 mmol) and succinylcholine chloride dihydrate (0.456 g, 1.15 mmol) according to the procedure of Example 3 to give the title compound (0.414 g, 60%) as a bright yellow solid: IR (KBr pellet, cm^{-1}) 3005, 1732, 1208, 1150, 455; ^1H NMR (300 MHz, DMSO- d_6) δ 4.47 (m, 4 H), 3.68-3.72 (m, 4 H), 3.30 (d, 18 H, $J = 4.2$ Hz), 2.66 (m, 4 H); ^{13}C NMR (75 MHz, DMSO- d_6) δ 171.3, 58.0, 52.9, 28.4; UV (H_2O) 394 nm ($\epsilon = 15513$); Anal. calcd for $\text{C}_{14}\text{H}_{30}\text{N}_2\text{O}_4\text{S}_4\text{W}$: C, 27.91; H, 5.02; N, 4.65; S, 21.29. Found: C, 27.84, H, 4.80, N, 4.66; S, 21.06.

6.23 Example 23: Tetrathiotungstate, (ethylene-1,2-bisammonium)

This compound was prepared from tetrathiotungstate, bis(ammonium) (0.300 g, 0.862 mmol), ammonium chloride (0.092 g, 1.72 mmol) and ethylenediamine (57.6 μl , 0.862 mmol) according to the procedure of Example 3 to give the title compound (0.257 g, 80 %) as a bright yellow solid: IR (KBr pellet, cm^{-1}) 3002, 1435, 1025, 451; ^1H NMR (300 MHz, DMSO- d_6) δ 7.89 (bs, 6 H), 3.09 (s, 4 H); ^{13}C NMR (75 MHz, DMSO- d_6) δ 36.8; UV (H_2O) 394 nm ($\epsilon = 13291$); Anal. calcd for $\text{C}_2\text{H}_{10}\text{N}_2\text{S}_4\text{W}$: C, 6.42; H, 2.69; N, 7.49; S, 34.27. Found: C, 6.58, H, 2.43, N, 7.49; S, 32.98.

6.24 Example 24: Moisture Stability of Tetrathiotungstate Salts

Salts of tetrathiotungstate were placed in an acrylic chamber at room temperature with 95% relative humidity for two weeks. The samples were analyzed for purity according to the previously reported method with the exception that the absorbance was monitored at 493nm and the molar absorptivity was $15710 \text{ M}^{-1}\text{cm}^{-1}$ (McDonald *et. al.*, *Inorg. Chim. Acta* 1983, 72, 205-210). The results are reported below in Table 1

Table 1

Name	% degradation
Tetrathiotungstate, bis(triethylphenyl ammonium)	-0.12
Tetrathiotungstate, bis(ammonium)	-0.46
Tetrathiotungstate, bis(trimethylphenylammonium)	-0.46

Tetrathiotungstate, bis(benzyltrimethylammonium)	0.41
Tetrathiotungstate, bis(acetylcholine)	2.4
Tetrathiotungstate, bis(choline)	3.0
Tetrathiotungstate, bis(1-ethyl-3-methyl-1 <i>H</i> -imidazolium)	1.3
Tetrathiotungstate, bis(1,4-dimethylpyridinium)	2.0
Tetrathiotungstate, bis(acetyl- β -methylcholine)	4.2
Tetrathiotungstate, bis(1,1-dimethylpyrrolidinium)	0.04
Tetrathiotungstate, bis(2-hydroxyiminomethyl-1-methylpyridinium)	5.6
Tetrathiotungstate, pentane-1,5-bis(trimethylammonium)	19.8*
Tetrathiotungstate, ethylenebis(trimethylammonium)	8.6*
Tetrathiotungstate, propane-1,3-bis(trimethylammonium)	23.8*
Tetrathiotungstate, butane-1,4-bis(trimethylammonium)	5.7*
Tetrathiotungstate, (succinylcholine)	23.3*
Tetrathiotungstate, suberyldicholine	18.4*
Tetrathiomolybdate, bis(ammonium)	56 [†] $\pm$ 5
Tetrathiomolybdate, bis(choline)	36 [†] $\pm$ 2

*partially soluble

[†]average of 2 data points**6.25 Example 25: Copper-binding Ability of Tetrathiotungstate salts**

- 5 The copper-binding ability of tetrathiotungstate salts was determined according to the ability of tetrathiotungstate salts to inhibit cysteine autooxidation as reported in Table 2 below.

Table 2

- 10 Inhibition of Cysteine Autooxidation (100 μ M Cys, 100 μ M coumarin-3-carboxylic acid, 100 μ M CuSO₄)

inhibitor	inhibitor concentration	% inhibition
trientine	50 μ M	39.71%
Tetrathiotungstate,	50 μ M	95.94%

bis(ammonium)		
Tetrathiotungstate, bis(triethylphenyl ammonium)	50µM	91.32%
Tetrathiotungstate, bis(trimethylphenylammonium)	50µM	91.53%
Tetrathiotungstate, bis(benzyltrimethylammonium)	50µM	94.62%
Tetrathiotungstate, bis(acetylcholine)	50µM	90.36%
Tetrathiotungstate, bis(choline)	50µM	92.54%
Tetrathiotungstate, bis(1-ethyl-3-methyl-1 <i>H</i> -imidazolium)	50µM	91.16%
Tetrathiotungstate, bis(1,4-dimethylpyridinium)	50µM	89.98%
Tetrathiotungstate, bis(acetyl-β-methylcholine)	50µM	90.40%
Tetrathiotungstate, bis(1,4-dimethylpyridinium)	50µM	89.66%
Tetrathiotungstate, bis(2-hydroxyiminomethyl-1-methylpyridinium)	50µM	91.15%
Tetrathiotungstate, pentane-1,5-bis(trimethylammonium)	50µM	90.69%
Tetrathiotungstate, bis(choline)	50µM	91.99%
Tetrathiomolybdate, bis(ammonium)	50µM	94.24%
Tetrathiotungstate, ethylenebis(trimethylammonium)	50µM	89.60%
Tetrathiotungstate, propane-1,3-bis(trimethylammonium)	50µM	92.95%
Tetrathiotungstate, butane-1,4-bis(trimethylammonium)	50µM	91.75%
trientine	10µM	4.27%
Tetrathiomolybdate, bis(ammonium)	10µM	87.71%
Tetrathiotungstate, ethylenebis(trimethylammonium)	10µM	84.66%
Tetrathiotungstate, propane-1,3-bis(trimethylammonium)	10µM	85.53%
Tetrathiotungstate, butane-1,4-bis(trimethylammonium)	10µM	87.32%
trientine	10µM	17.72%
Tetrathiomolybdate, bis(ammonium)	10µM	76.69%
Tetrathiotungstate, (succinylcholine)	10µM	81.38%
Tetrathiotungstate, suberyldicholine	10µM	77.83%
trientine	1µM	0.00%

Tetrathiomolybdate, bis(ammonium)	1 μ M	68.48%
Tetrathiotungstate, (succinylcholine)	1 μ M	66.06%
Tetrathiotungstate, suberyldicholine	1 μ M	67.74%

6.25 Example 25: Inhibition of Angiogenesis in Matrigel® Plug Assay by Ammonium Tetrathiotungstate

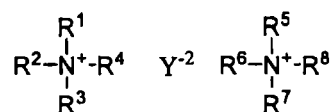
Ammonium tetrathiotungstate was assayed in the matrigel® plug assay as described in Section 5.4.7, *supra*. Two positive controls were used with positive control 1 measured five days after implantation when treatment began and negative control 2 measured five days after implantation when treatment ended. Two negative controls were used with negative control 1 measured five days after implantation when treatment began and negative control 2 measured five days after implantation when treatment ended. As can be seen in Figure 1, treatment with ammonium tetrathiotungstate resulted in about 34% inhibition using this assay.

Finally, it should be noted that there are alternative ways of implementing the present invention. Accordingly, the present embodiments are to be considered as illustrative and not restrictive, and the invention is not to be limited to the details given herein, but may be modified within the scope and equivalents of the appended claims. All publications and patents cited herein are incorporated by reference in their entirety.

CLAIMS

What is Claimed Is:

1. A compound of structural formula (I):



(I)

or a solvate or hydrate or N-oxide thereof wherein:

R^1 , R^2 , R^3 , R^5 , R^6 and R^7 are independently hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, heteroaryl, substituted heteroaryl, heteroarylalkyl, substituted heteroarylalkyl, heteroalkyl or substituted heteroalkyl;

R^4 and R^8 are independently hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, heteroaryl, substituted heteroaryl, heteroarylalkyl, substituted heteroarylalkyl, heteroalkyl or substituted heteroalkyl or are absent when N is part of an aromatic ring;

optionally, R^1 and R^2 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl;

optionally, R^5 and R^6 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl;

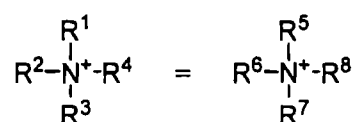
optionally, R^1 and R^2 taken together, R^2 and R^3 taken together and R^2 and R^4 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl;

optionally, R^5 and R^6 taken together, R^6 and R^7 taken together and R^6 and R^8 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl;

optionally, R^3 and R^7 taken together are alkylidiyl, substituted alkylidiyl, heteroalkylidiyl or substituted heteroalkylidiyl; and

Y^{-2} is $(\text{WS}_4)^{-2}$, $(\text{W}_2\text{S}_{12})^{-2}$, $(\text{W}_2\text{S}_9)^{-2}$, $(\text{W}_2\text{S}_7)^{-2}$, $(\text{W}_2\text{S}_8)^{-2}$, $(\text{W}_2\text{S}_{11})^{-2}$, $(\text{W}_2\text{S}_6)^{-2}$ or $(\text{W}_2\text{S}_{13})^{-2}$.

2. The compound of Claim 1, wherein Y^{-2} is $(WS_4)^{-2}$.
3. The compound of Claim 1, wherein



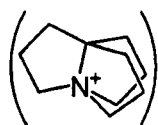
4. The compound of Claim 3, wherein Y^{-2} is $(WS_4)^{-2}$.
5. The compound of any one of Claims 1-4, wherein at least one of R^1 , R^2 , R^3 and R^4 is not alkyl.
6. The compound of any one of Claims 1-4, wherein at least one of R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 and R^8 is not alkyl.
7. The compound of any one of Claims 1-4, wherein R^1 , R^2 and R^4 are hydrogen, alkanyl or substituted alkanyl.
8. The compound of any one of Claims 1-4, wherein R^1 , R^2 and R^4 are hydrogen, methyl or ethyl.
9. The compound of any one of Claims 1-4, wherein R^1 and R^2 are alkanyl.
10. The compound of any one of Claims 1-4, wherein R^1 and R^2 are methyl or ethyl.
11. The compound of any one of Claims 1-4, wherein R^1 is alkanyl, substituted alkanyl, alkenyl, substituted alkenyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl or substituted cycloalkyl.
12. The compound of any one of Claims 1-4, wherein R^1 and R^2 taken together are alkyleno, substituted alkyleno, heteroalkyleno or substituted heteroalkyleno.

13. The compound of any one of Claims 1-4, wherein R^1 and R^2 taken together are alkyleno or heteroalkyleno.

14. The compound of any one of Claims 1-4, wherein R^1 and R^2 taken together, R^2 and R^3 taken together and R^2 and R^4 taken together are alkyleno, substituted alkyleno, heteroalkyleno or substituted heteroalkyleno.

15. The compound of any one of Claims 1-4, wherein R^1 and R^2 taken together, R^2 and R^3 taken together and R^2 and R^4 taken together are alkyleno.

16. The compound of any one of Claims 1-4, wherein $R^1(R^2)(R^3)(R^4)N$ has the structure:



17. The compound of any one of Claims 1-4, wherein R^3 and R^7 taken together are alkyleno, substituted alkyleno, heteroalkyleno or substituted heteroalkyleno.

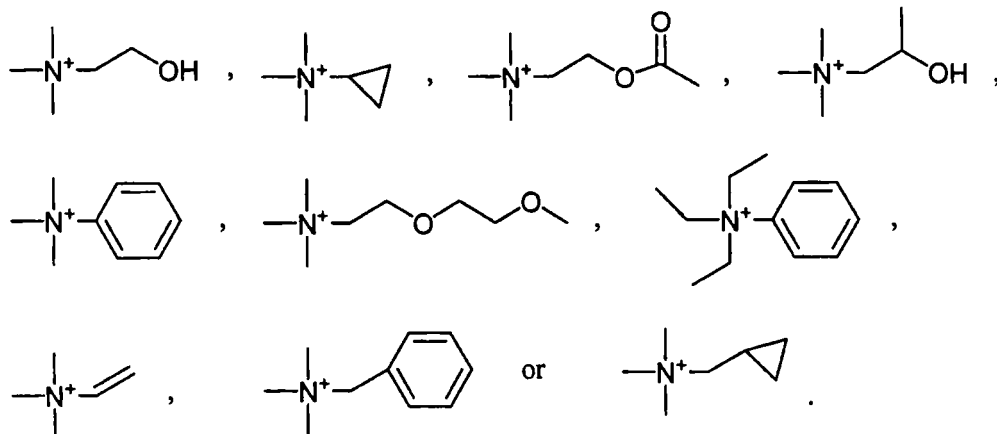
18. The compound of any one of Claims 1-4, wherein R^3 and R^7 taken together are alkyleno or heteroalkyleno.

19. The compound of any one of Claims 1-4, wherein R^1 , R^2 and R^4 are hydrogen, alkanyl or substituted alkanyl and R^3 is substituted alkyl, aryl, arylalkyl, cycloalkyl or R^3 and R^7 taken together are alkyleno, substituted alkyleno, heteroalkyleno or substituted heteroalkyleno.

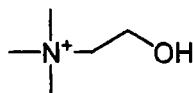
20. The compound of any one of Claims 1-4, wherein R^1 , R^2 and R^4 are methyl or ethyl and R^3 is substituted alkyl, aryl, arylalkyl, cycloalkyl or R^3 and R^7 taken together are alkyleno or heteroalkyleno.

21. The compound of any one of Claims 1-4, wherein R^1 , R^2 and R^4 are methyl or ethyl and R^3 is substituted alkyl, aryl, arylalkyl or cycloalkyl.

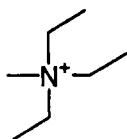
22. The compound of any one of Claims 1-4, wherein $R^1(R^2)(R^3)(R^4)N$ has the structure:



23. The compound of any one of Claims 1-4, wherein $R^1(R^2)(R^3)(R^4)N$ has the structure:

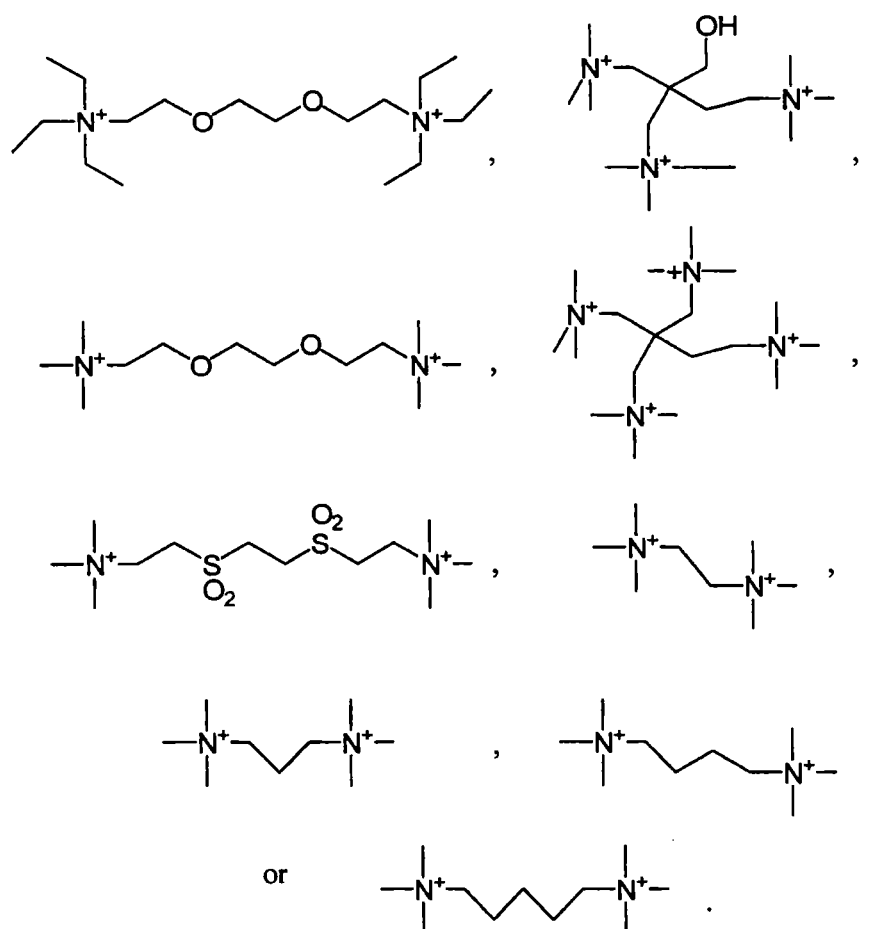


24. The compound of any one of Claims 1-4, wherein $R^1(R^2)(R^3)(R^4)N$ has the structure:



25. The compound of any one of Claims 1-4, wherein R^1 , R^2 and R^4 are methyl or ethyl and R^3 and R^7 taken together are alkylene or heteroalkylene.

26. The compound of any one of Claims 1-4, wherein $R^1(R^2)(R^3)(R^4)N$ has the structure:

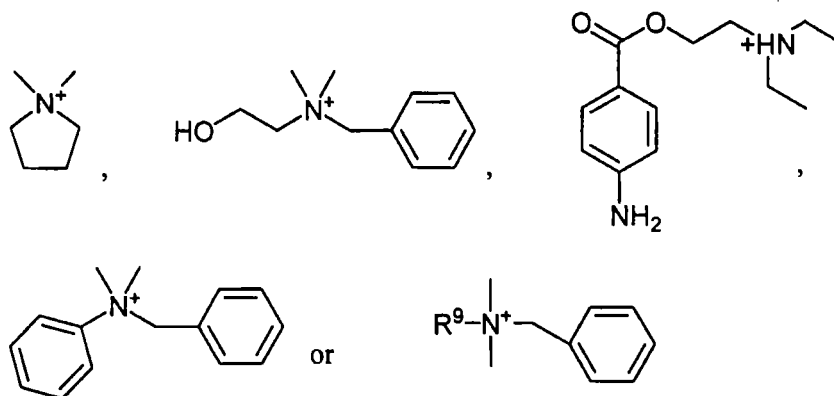


27. The compound of any one of Claims 1-4, wherein R¹, R² and R⁴ are hydrogen and R³ is substituted alkyl, cycloalkyl or substituted heteroaryl or R³ and R⁷ taken together are alkyleno.

28. The compound of any one of Claims 1-4, wherein R¹ and R² are alkanyl and R³ and R⁴ are hydrogen, alkyl, substituted alkyl, aryl, arylalkyl or alkylene.

29. The compound of any one of Claims 1-4, wherein R¹ and R² are methyl or ethyl and R³ and R⁴ are hydrogen, alkyl, substituted alkyl, aryl, arylalkyl or alkyleno.

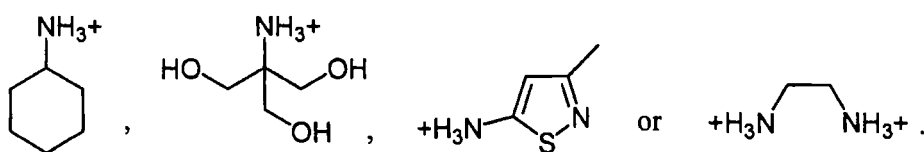
30. The compound of any one of Claims 1-4, wherein $R^1(R^2)(R^3)(R^4)N$ has the structure:



wherein R^9 is $n\text{-C}_8\text{H}_{17}$, $n\text{-C}_9\text{H}_{19}$, $n\text{-C}_{10}\text{H}_{21}$, $n\text{-C}_{11}\text{H}_{23}$, $n\text{-C}_{12}\text{H}_{25}$, $n\text{-C}_{13}\text{H}_{27}$, $n\text{-C}_{14}\text{H}_{29}$, $n\text{-C}_{15}\text{H}_{31}$, $n\text{-C}_{16}\text{H}_{33}$, $n\text{-C}_{17}\text{H}_{35}$, or $n\text{-C}_{18}\text{H}_{37}$.

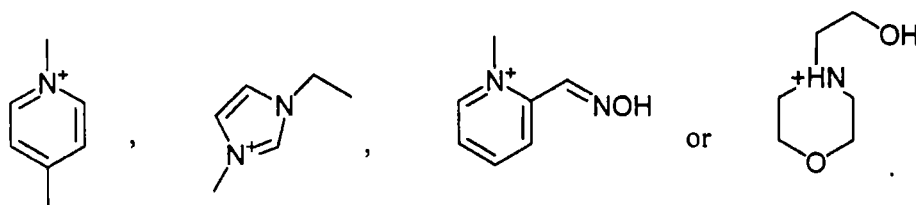
31. The compound of any one of Claims 1-4, wherein R^1 , R^2 and R^4 are hydrogen and R^3 is substituted alkyl, substituted heteroaryl, cycloalkyl or alkyleno.

32. The compound of any one of Claims 1-4, wherein $R^1(R^2)(R^3)(R^4)N$ has the structure:



33. The compound of any one of Claims 1-4, wherein R^1 and R^2 taken together are alkyleno, substituted alkyleno, heteroalkyleno or substituted heteroalkyleno, R^3 is alkyl or substituted alkyl and R^4 is hydrogen or is absent.

34. The compound of any one of Claims 1-4, wherein $R^1(R^2)(R^3)N$ or $R^1(R^2)(R^3)(R^4)N$ has the structure:



35. A pharmaceutical composition comprising the compound of any one of Claims 1-34 and a pharmaceutically acceptable diluent, excipient or adjuvant.

36. A therapeutically effective amount of the compound of any one of Claims 1-34 for use in the treatment of cancer in a patient in need of such treatment.

37. A therapeutically effective amount of the compound of any one of Claims 1-34 and a therapeutically effective amount of another anti-cancer agent for concurrent, separate, or sequential use in the treatment of cancer in a patient in need of such treatment.

38. A therapeutically effective amount of the compound of any one of Claims 1-34 and a therapeutically effective amount of zinc for concurrent, separate, or sequential use in the treatment of cancer in a patient in need of such treatment.

39. The therapeutically effective amount of any one of Claims 36-38, wherein the cancer is breast cancer, renal cancer, brain cancer colon cancer, prostate cancer, chondrosarcoma or angiosarcoma.

40. A therapeutically effective amount of the compound of any one of Claims 1-34 for use in the treatment of wet type macular degeneration or rheumatoid arthritis in a patient in need of such treatment.

41. A therapeutically effective amount of the compound of any one of Claims 1-34 for use in the treatment of aberrant vascularization in a patient in need of such treatment.

42. A therapeutically effective amount of the compound of any one of Claims 1-34 for use in the treatment of excess copper levels in a patient in need of such treatment.

43. A therapeutically effective amount of the compound of any one of Claims 1-34 for use in the treatment of obesity in a patient in need of such treatment.

44. A therapeutically effective amount of the compound of any one of Claims 1-34 and a therapeutically effective amount of another anti-obesity agent for concurrent, separate, or sequential use in the treatment of obesity in a patient in need of such treatment.

45. A therapeutically effective amount of the compound of any one of Claims 1-34 for the use in the treatment of neurodegenerative disease in a patient in need of such treatment.

46. The therapeutically effective amount of Claim 45, wherein the neurodegenerative disease is Alzheimer's disease, amyotrophic lateral sclerosis or prion disease.

47. Use of a therapeutically effective amount of the compound of any one of Claims 1-34 in the manufacture of a medicament for the treatment of cancer in a patient in need of such treatment.

48. The use of Claim 47 wherein the medicament is for use with a therapeutically effective amount of another anti-cancer agent.

49. The use of Claim 47 wherein the medicament is for use with a therapeutically effective amount of zinc.

50. The use of any one of Claims 47-49, wherein the cancer is breast cancer, renal cancer, brain cancer colon cancer, prostate cancer, chondrosarcoma or angiosarcoma.

51. Use of a therapeutically effective amount of the compound of any one of Claims 1-34 in the manufacture of a medicament for the treatment of wet type macular degeneration or rheumatoid arthritis in a patient in need of such treatment.

52. Use of a therapeutically effective amount of the compound of any one of Claims 1-34 in the manufacture of a medicament for the treatment of aberrant vascularization in a patient in need of such treatment.

53. Use of a therapeutically effective amount of the compound of any one of Claims 1-34 in the manufacture of a medicament for the treatment of excess copper levels in a patient in need of such treatment.

54. Use of a therapeutically effective amount of the compound of any one of Claims 1-34 in the manufacture of a medicament for the treatment of obesity in a patient in need of such treatment.

55. The use of Claim 54 wherein the medicament is for use with a therapeutically effective amount of another anti-obesity agent.

56. Use of a therapeutically effective amount of the compound of any one of Claims 1-34 in the manufacture of a medicament for the treatment of neurodegenerative disease in a patient in need of such treatment.

57. The use of claim 56, wherein the neurodegenerative disease is Alzheimer's disease, amyotrophic lateral sclerosis or prion disease.

58. The use of any one of claims 47-57, wherein said patient is a human.

59. A kit for use in the treatment of cancer, comprising in a container a therapeutically effective amount of the compound of any one of Claims 1-34.

60. The kit of claim 59, further comprising in said container or in a separate container a therapeutically effective amount of another anti-cancer agent.

61. The kit of claim 59, further comprising in said container or in a separate container a therapeutically effective amount of zinc.

62. The kit of any one of Claims 59-61, wherein the cancer is breast cancer, renal cancer, brain cancer colon cancer, prostate cancer, chondrosarcoma or angiosarcoma.

63. A kit for use in the treatment of wet type macular degeneration or rheumatoid arthritis, comprising in a container a therapeutically effective amount of the compound of any one of Claims 1-34.

64. A kit for use in the treatment of aberrant vascularization comprising in a container a therapeutically effective amount of the compound of any one of Claims 1-34.

65. A kit for use in the treatment of excess copper levels comprising in a container a therapeutically effective amount of the compound of any one of Claims 1-34.

66. A kit for use in the treatment of obesity comprising in a container a therapeutically effective amount of the compound of any one of Claims 1-34.

67. The kit of claim 66, further comprising in said container or in a separate container a therapeutically effective amount of another anti-obesity agent.

68. A kit for use in the treatment of a neurodegenerative disease comprising in a container a therapeutically effective amount of the compound of any one of Claims 1-34.

69. The kit of claim 68, wherein the neurodegenerative disease is Alzheimer's disease, amyotrophic lateral sclerosis or prion disease.