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Declarations under Rule 4.17:

- as to the applicant's entitlement to claim the priority of the earlier application (Rule 4.17(iii))
- of inventorship (Rule 4.17(iv))

[Continued on next page]

(54) **Title:** AMINE FUNCTIONAL POLYAMIDES

Figure 1: Susceptibility of *Pseudomonas aeruginosa* Present in Biofilm of Cystic Fibrosis-Derived Human Airway Epithelial Cells of Representative Amine Functional Polyamide

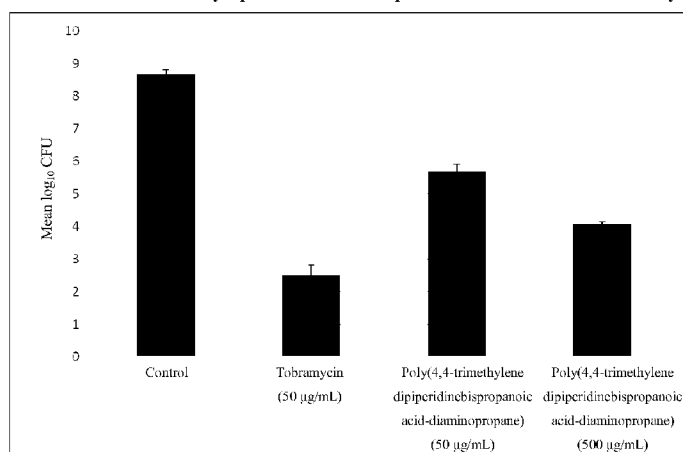


Figure 1 of 1

(57) **Abstract:** Amine functional polyamides comprise amine and ammonium groups along the polymer chain. Amine functional polyamides can be used as pharmaceutical agents and in pharmaceutical compositions. The amine functional polyamides are particularly useful in the treatment or prevention of mucositis and infection, specifically oral mucositis, surgical site infection, and lung infection associated with cystic fibrosis.



Published:

— with international search report (Art. 21(3))

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TITLE OF THE INVENTION

AMINE FUNCTIONAL POLYAMIDES

CROSS-REFERENCE TO RELATED APPLICATIONS

5 This application claims the benefit of U.S. Provisional Application No. 61/790,231,
filed March 15, 2013, and is incorporated by reference in its entirety.

STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH

Not applicable

THE NAMES OF THE PARTIES TO A JOINT RESEARCH AGREEMENT

Not applicable

10 INCORPORATION-BY-REFERENCE OF MATERIAL SUBMITTED ON COMPACT DISC

Not applicable

BACKGROUND OF THE INVENTION

Field of the Invention

15 This invention relates to amine functional polyamides. Amine functional polyamides
comprise amine and ammonium groups along the polymer chain. This invention further
relates to the use of amine functional polyamides as pharmaceutical agents and in
pharmaceutical compositions.

20 Mucositis is defined as inflammation and/or ulceration of a mucous membrane in the
digestive tract. Mucositis can occur in the stomach, intestines and mouth. The disorder is
characterized by breakdown of mucosa, which results in redness, swelling and/or the
formation of ulcerative lesions.

25 Oral mucositis is a common dose-limiting toxicity of drug and radiation therapy for
cancer; it occurs to some degree in more than one third of all patients receiving anti-
neoplastic drug therapy. In granulocytopenic patients, the ulcerations that accompany
mucositis are frequent portals of entry for indigenous oral bacteria leading to sepsis or
bacteremia. There are about one million occurrences of oral mucositis annually in the United
States. Mucositis also includes mucositis that develops spontaneously in a healthy patient not

receiving ant-cancer therapy, as in the case of a canker sore or mouth ulcer. Improved therapies to treat mucositis are needed.

Surgical site infection (SSI) is an infection associated with a surgical procedure. Postoperative SSIs are a major source of illness, and less commonly death, in surgical patients (Nichols RL, 2001). The *Guideline for Prevention of Surgical Site Infection* (1999) sets forth recommendations for preventing SSIs.

- Preoperative measures including proper preparation of the patient, antisepsis for surgical team, management of surgical personnel who exhibit signs of transmissible infectious illness, and antimicrobial prophylaxis.
- Intra-operative measures including proper ventilation in the operating room, cleaning and disinfecting of surfaces in the surgical environment, microbiologic sampling, sterilization of surgical instruments, proper surgical attire and drapes, and proper asepsis and surgical technique.
- Proper incision care post-operation, including sterile dressings and hand washing before and after dressing changes.
- Continued surveillance of the surgical wound during the healing process.

Despite these recommendations, SSIs develop in about 1 to 3 of every 100 patients who have surgery (CDC.gov, 2011). These infections can result in major complications that increase the costs and duration of post-operative hospital stays. Accordingly, novel approaches to mitigating SSIs are needed.

Cystic fibrosis (CF) is a genetic disease caused by a mutation in the cystic fibrosis transmembrane conductor regulator (CFTR) that results in abnormally thick and sticky mucus (Yu Q, et al., 2012). The thick, sticky mucus of a CF patient leads to compromised mucus clearance and lung infection. Chronic airway infections are one of the most common and debilitating manifestations of CF (Tümmeler B and C Kiewitz, 1999). The stagnant mucus becomes a breeding ground for bacteria like *Pseudomonas aeruginosa*, which causes chronic airway infections (Moreau-Marquis S, GA O'Toole and BA Stanton, 2009). Despite the use of traditional antibacterial therapies in CF patients, most CF patients are afflicted with a chronic *P. aeruginosa* infection as teenagers and adults, leading to increased morbidity and mortality (Hoiby N, B Frederiksen B, T Pressler, 2005). In chronic *P. aeruginosa* infection, the *P. aeruginosa* forms biofilms, resulting in a greater tolerance to antibiotics and increasing difficulty in treatment (Yu Q, et al., 2012). Effective, novel treatments to assuage the effects of bacterial infection and biofilm formation in CF patients are needed.

Definitions

As used herein, the term “**amino**” means a functional group having a nitrogen atom and 1 to 2 hydrogen atoms. “Amino” generally may be used herein to describe a primary, secondary, or tertiary amine, and those of skill in the art will readily be able to ascertain the identification of which in view of the context in which this term is used in the present disclosure. The term “**amine**” or “**amine group**” or “**ammonia group**” means a functional group containing a nitrogen atom derived from ammonia (NH₃). The amine groups may be primary amines, meaning the nitrogen is bonded to two hydrogen atoms and one substituent group comprising a substituted or unsubstituted alkyl or aryl group or an aliphatic or aromatic group. The amine groups may be secondary amines meaning, the nitrogen is bonded to one hydrogen atom and two substituent groups comprising a substituted or unsubstituted alkyl or aryl groups or an aliphatic or aromatic group, as defined below. The amine groups may be tertiary amines meaning the nitrogen is bonded to three substituent groups comprising a substituted or unsubstituted alkyl or aryl groups or an aliphatic or aromatic group. The amine groups may also be quaternary amines meaning the designated amine group is bonded to a fourth group, resulting in a positively charged ammonium group.

As used herein, the term “**amide group**” means a functional group comprising a carbonyl group linked to a nitrogen. A “**carbonyl group**” means a functional group comprising a carbon atom double bonded to an oxygen atom, represented by (C=O).

The term “**alkane**” means a saturated hydrocarbon, bonded by single bonds. Alkanes can be linear or branched. “**Cycloalkanes**” are saturated hydrocarbons rings bonded by single bonds.

As used herein, the term “**(C₁-C₁₀)alkyl**” means a saturated straight chained or branched or cyclic hydrocarbon consisting essentially of 1 to 10 carbon atoms and a corresponding number of hydrogen atoms. Typically straight chained or branched groups have from one to ten carbons, or more typically one to five carbons. Exemplary (C₁-C₁₀)alkyl groups include methyl (represented by -CH₃), ethyl (represented by -CH₂-CH₃), n-propyl, isopropyl, n-butyl, isobutyl, etc. Other (C₁-C₁₀)alkyl groups will be readily apparent to those of skill in the art given the benefit of the present disclosure.

As used herein, the term “**(C₂-C₉)heteroalkyl**” means a saturated straight chained or branched or cyclic hydrocarbon consisting essentially of 2 to 10 atoms, wherein 2 to 9 of the atoms are carbon and the remaining atom(s) is selected from the group consisting of nitrogen,

sulfur, and oxygen. Exemplary (C₂-C₉)heteroalkyl groups will be readily apparent to those of skill in the art given the benefit of the present disclosure.

As used herein, the term “(C₃-C₁₀)cycloalkyl” means a nonaromatic saturated hydrocarbon group, forming at least one ring consisting essential of 3 to 10 carbon atoms and a corresponding number of hydrogen atoms. (C₃-C₁₀)cycloalkyl groups can be monocyclic or multicyclic. Individual rings of multicyclic cycloalkyl groups can have different connectivities, for example, fused, bridged, spiro, etc., in addition to covalent bond substitution. Exemplary (C₃-C₁₀)cycloalkyl groups include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, norbornanyl, bicyclo-octanyl, octahydro-pentalenyl, spiro-decanyl, cyclopropyl substituted with cyclobutyl, cyclobutyl substituted with cyclopentyl, cyclohexyl substituted with cyclopropyl, etc. Other (C₃-C₁₀)cycloalkyl groups will be readily apparent to those of skill in the art given the benefit of the present disclosure.

As used herein, the term “(C₂-C₉)heterocycloalkyl” means a nonaromatic group having 3 to 10 atoms that form at least one ring, wherein 2 to 9 of the ring atoms are carbon and the remaining ring atom(s) is selected from the group consisting of nitrogen, sulfur, and oxygen. (C₂-C₉)heterocycloalkyl groups can be monocyclic or multicyclic. Individual rings of such multicyclic heterocycloalkyl groups can have different connectivities, for example, fused, bridged, spiro, etc., in addition to covalent bond substitution. Exemplary (C₂-C₉)heterocycloalkyl groups include pyrrolidinyl, tetrahydrofuranyl, dihydrofuranyl, tetrahydropyranyl, pyranal, thiopyranal, aziridinyl, azetidinal, oxiranyl, methylenedioxy, chromenyl, barbituryl, isoxazolidinyl, 1,3-oxazolidin-3-yl, isothiazolidinyl, 1,3-thiazolidin-3-yl, 1,2-pyrazolidin-2-yl, 1,3-pyrazolidin-1-yl, piperidinyl, thiomorpholinyl, 1,2-tetrahydrothiazin-2-yl, 1,3-tetrahydrothiazin-3-yl, tetrahydrothiadiazinyl, morpholinyl, 1,2-tetrahydrodiazin-2-yl, 1,3-tetrahydrodiazin-1-yl, tetrahydroazepinyl, piperazinyl, piperizin-2-onyl, piperizin-3-onyl, chromanyl, 2-pyrrolinyl, 3-pyrrolinyl, imidazolidinyl, 2-imidazolidinyl, 1,4-dioxanyl, 8-azabicyclo[3.2.1]octanyl, 3-azabicyclo[3.2.1]octanyl, 3,8-diazabicyclo[3.2.1]octanyl, 2,5-diazabicyclo[2.2.1]heptanyl, 2,5-diazabicyclo[2.2.2]octanyl, octahydro-2H-pyrido[1,2-a]pyrazinyl, 3-azabicyclo[4.1.0]heptanyl, 3-azabicyclo[3.1.0]hexanyl, 2-azaspiro[4.4]nonanyl, 7-oxa-1-aza-spiro[4.4]nonanyl, 7-azabicyclo[2.2.2]heptanyl, octahydro-1H-indolyl, etc. The (C₂-C₉)heterocycloalkyl group is typically attached to the main structure via a carbon atom or a nitrogen atom. Other (C₂-C₉)heterocycloalkyl groups will be readily apparent to those of skill in the art given the benefit of the present disclosure.

The term “**aliphatic group**” or “**aliphatic**” means a non-aromatic group consisting of carbon and hydrogen, and may optionally include one or more double and/or triple bonds. An aliphatic group may be straight chained, branched or cyclic and typically contains between about one and about 24 carbon atoms.

5 The term “**aryl group**” may be used interchangeably with “**aryl**,” “**aryl ring**,” “**aromatic**,” “**aromatic group**,” and “**aromatic ring**.” Aryl groups include carbocyclic aromatic groups, typically with six to fourteen ring carbon atoms. Aryl groups also include heteroaryl groups, which typically have five to fourteen ring atoms with one or more heteroatoms selected from nitrogen, oxygen and sulfur.

10 As used herein, the term “**(C₆-C₁₄)aryl**” means an aromatic functional group having 6 to 14 carbon atoms that form at least one ring.

As used herein, the term “**(C₂-C₉)heteroaryl**” means an aromatic functional group having 5 to 10 atoms that form at least one ring, wherein 2 to 9 of the ring atoms are carbon and the remaining ring atom(s) is selected from the group consisting of nitrogen, sulfur, and oxygen. (C₂-C₉)heteroaryl groups can be monocyclic or multicyclic. Individual rings of such multicyclic heteroaryl groups can have different connectivities, for example, fused, etc., in addition to covalent bond substitution. Exemplary (C₂-C₉)heteroaryl groups include furyl, thienyl, thiazolyl, pyrazolyl, isothiazolyl, oxazolyl, isoxazolyl, pyrrolyl, triazolyl, tetrazolyl, imidazolyl, 1,3,5-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,3-oxadiazolyl, 1,3,5-thiadiazolyl, 1,2,3-thiadiazolyl, 1,2,4-thiadiazolyl, pyridyl, pyrimidyl, pyrazinyl, pyridazinyl, 1,2,4-triazinyl, 1,2,3-triazinyl, 1,3,5-triazinyl, pyrazolo[3,4-b]pyridinyl, cinnolinyl, pteridinyl, purinyl, 6,7-dihydro-5H-[1]pyrindinyl, benzo[b]thiophenyl, 5,6,7,8-tetrahydro-quinolin-3-yl, benzoxazolyl, benzothiazolyl, benzisothiazolyl, benzisoxazolyl, benzimidazolyl, thianaphthenyl, isothianaphthenyl, benzofuranyl, isobenzofuranyl, isoindolyl, indolyl, indoliziny, indazolyl, isoquinolyl, quinolyl, phthalazinyl, quinoxalinyl, quinazolinyl and benzoxazinyl, etc. The (C₂-C₉)heteroaryl group is typically attached to the main structure via a carbon atom, however, those of skill in the art will realize when certain other atoms, for example, hetero ring atoms, can be attached to the main structure. Other (C₂-C₉)heteroaryl groups will be readily apparent to those of skill in the art given the benefit of the present disclosure.

As used herein, the term “**alkyl amine**” means an (C₁-C₁₀)alkyl containing a primary, secondary, or tertiary amine group in place of one hydrogen atom, represented by (C₁-C₁₀)alkyl amine and ((C₁-C₁₀)alkyl)₂ amine.

The term “**alkyl ester**” means a (C₁-C₁₀)alkyl containing an ester group in place of one hydrogen atom, represented by -O(O)C-(C₁-C₁₀)alkyl.

The term “**alkyl acid**” means an (C₁-C₁₀)alkyl containing a carboxylic acid group in place of one hydrogen atom, represented by (C₁-C₁₀)alkyl-COOH.

5 The term “**aliphatic acid**” means an acid of nonaromatic hydrocarbons, represented by (C₃-C₁₀)cycloalkyl-COOH.

The term “**halo**” means a fluorine (F), chlorine (Cl), bromine (Br), iodine (I), or astatine (At) ion.

10 The term “**methoxy**” means a (C₁)alkyl containing an oxygen in place of one hydrogen atom, represented by -(O)CH₃.

The term “**polyol**” means an alcohol containing multiple hydroxyl (-OH) groups.

“**Substituted**” means the substitution of a carbon in alkyl, heterocyclic or aryl groups with one or more non-carbon substituent. Non-carbon substituents are selected from nitrogen, oxygen and sulfur.

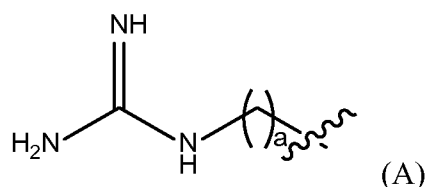
15 “**Unsubstituted**” means the group is comprised of only hydrogen and carbon.

The term “**polymer**” means a molecule comprised of repeating units. The term “**repeat unit**” or “**monomer**” means a group in a polymer that repeats or appears multiple times in a polymer. A polymer may be a copolymer if the repeating units or “**comonomers**” are chemically and structurally different from one another.

20 The term “**pharmaceutically acceptable anion**” means an anion that is suitable for pharmaceutical use. Pharmaceutically acceptable anions include but are not limited to halides, carbonate, bicarbonate, sulfate, bisulfate, hydroxide, nitrate, persulfate, sulfite, acetate, ascorbate, benzoate, citrate, dihydrogen citrate, hydrogen citrate, oxalate, succinate, tartrate, taurocholate, glycocholate, and cholate.

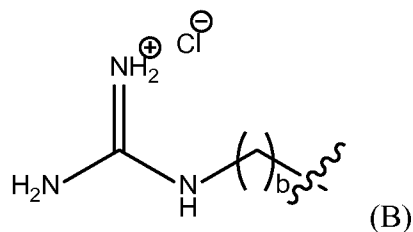
25 The term “**pharmaceutically acceptable end group**” means an end group that is suitable for pharmaceutical use. Examples of pharmaceutically acceptable end groups include but are not limited to H, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl, (C₁-C₁₀)alkylamine, -O(O)C-(C₁-C₁₀)alkyl, (C₁-C₁₀)alkyl-COOH, (C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH,
30 amide, a guanidino group, a guanidinium chloride group, a guanidinobenzene group, a dihydroxy group, and a polyethylene glycol group.

A “**guanidino group**” is represented by Formula (A):



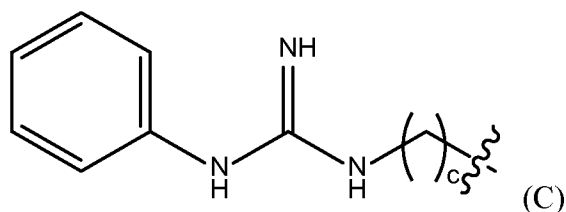
wherein a is an integer from 0 to 25,

A “**guanidinium chloride group**” is represented by Formula (B),



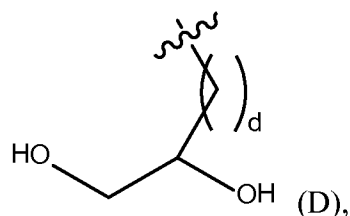
5 wherein b is an integer from 0 to 25,

A “**guanidinobenzene group**” is represented by Formula (C),



wherein c is an integer from 0 to 25,

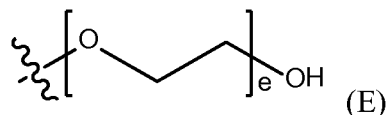
A “**dihydroxy group**” is represented by Formula (D),



10

wherein d is an integer from 0 to 25, or

A “**polyethylene glycol group**” is represented by Formula (E)



wherein e is an integer from 1 to 400.

15 The term “**effective amount**” of a disclosed amine functional polyamides is a quantity sufficient to achieve a therapeutic and/or prophylactic effect on the particular condition being treated, such as an amount which results in the prevention or a decrease in the symptoms associated with mucositis, oral mucositis, infection and surgical site infection, and lung infection associated with cystic fibrosis. The precise amount of the disclosed amine

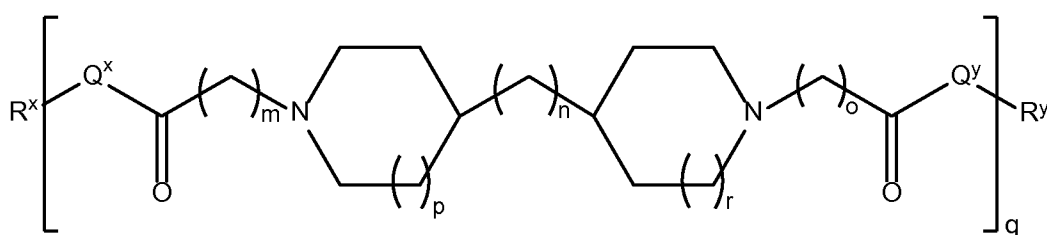
functional polyamides that is administered will depend on the type and severity of mucositis or infection being treated and on the characteristics of the individual, such as general health, age, sex, body weight and tolerance to drugs.

Related Art

5 Not applicable

BRIEF SUMMARY OF THE INVENTION

In one aspect of the invention, the amine functional polyamides are a compound comprising the structure of Formula (I):

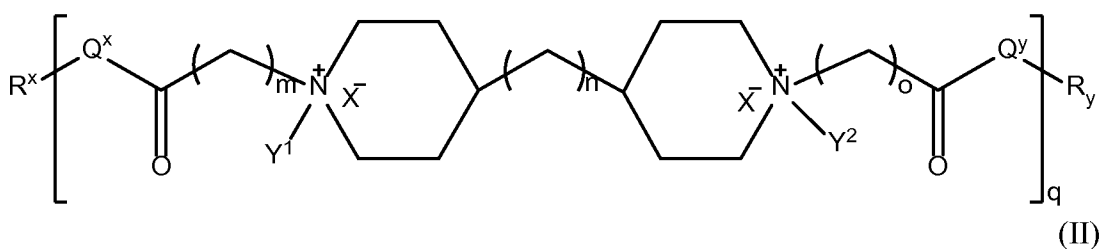


(I)

wherein:

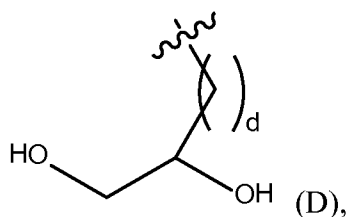
- i) m is 0, 1, 2, or 3;
- ii) n is 0, 1, 2, or 3;
- iii) o is 0, 1, 2, or 3;
- iv) p is 0 or 1;
- v) r is 0 or 1;
- vi) q is an integer from 1 to 400;
- vii) Q^x is NH, (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_3-C_{10}) cycloalkyl, (C_2-C_9) heterocycloalkyl, (C_6-C_{14}) aryl, (C_2-C_9) heteroaryl;
- viii) Q^y is $NH-R^w$, $NH-CH_2-R^w$, (C_1-C_{10}) alkyl, or (C_6-C_{14}) aryl, wherein R^w is absent or a (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_6-C_{14}) aryl, or (C_2-C_9) heteroaryl;
- ix) R^x and R^y are each independently a pharmaceutically acceptable end group.

In another aspect of the invention, the amine functional polyamides are a compound comprising the structure of Formula (II):



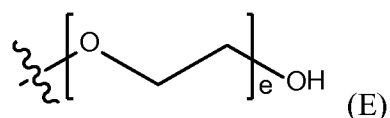
wherein:

- i) m is 0, 1, 2, or 3;
- ii) n is 0, 1, 2, or 3;
- iii) o is 0, 1, 2, or 3;
- iv) p is 0 or 1;
- v) r is 0 or 1;
- vi) q is an integer from 1 to 400;
- vii) Q^x is NH, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl;
- viii) Q^y is NH-R^w, NH-CH₂-R^w, (C₁-C₁₀)alkyl, or (C₆-C₁₄)aryl, wherein R^w is absent or a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl;
- ix) R^x and R^y are each independently a pharmaceutically acceptable end group;
- x) X⁻ is each independently a halo or any pharmaceutically acceptable anion;
- xi) Y¹ and Y₂ are each independently H or (C₁-C₁₀)alkyl optionally substituted by one or more substituents selected from the group consisting of (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl, (C₁-C₁₀)alkylamine, -S-O-(C₁-C₁₀)alkyl, -O(O)C-(C₁-C₁₀)alkyl, -(C₁-C₁₀)alkyl-COOH, (C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH, amide, a dihydroxy group, represented by Formula (D),



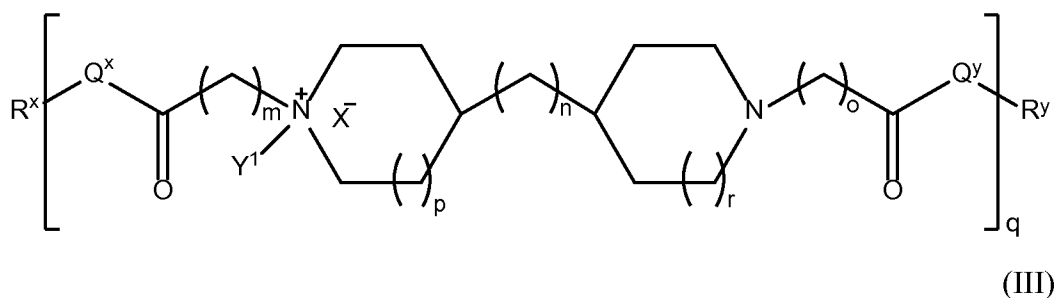
wherein d is an integer from 0 to 25, or

a polyethylene glycol group, represented by Formula (E)



wherein e is an integer from 1 to 25.

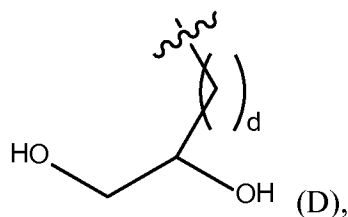
In yet another aspect of the invention, the amine functional polyamides are a compound comprising the structure of Formula (III):



wherein:

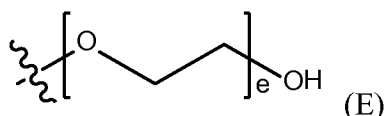
- i) m is 0, 1, 2, or 3;
- ii) n is 0, 1, 2, or 3;
- iii) o is 0, 1, 2, or 3;
- iv) p is 0 or 1;
- v) r is 0 or 1;
- vi) q is an integer from 1 to 400;
- vii) Q^x is NH, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl;
- viii) Q^y is NH-R^w, NH-CH₂-R^w, (C₁-C₁₀)alkyl, or (C₆-C₁₄)aryl, wherein R^w is absent or a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl;
- i) R^x and R^y are each independently a pharmaceutically acceptable end group;
- ix) X⁻ is a halo or any pharmaceutically acceptable anion;
- x) Y¹ is H or (C₁-C₁₀)alkyl optionally substituted by one or more substituents selected from the group consisting of (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl, (C₁-C₁₀)alkylamine, -S-O-(C₁-C₁₀)alkyl, -O(O)C-(C₁-C₁₀)alkyl, -(C₁-C₁₀)alkyl-COOH,

(C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH, amide, a dihydroxy group, represented by Formula (D),



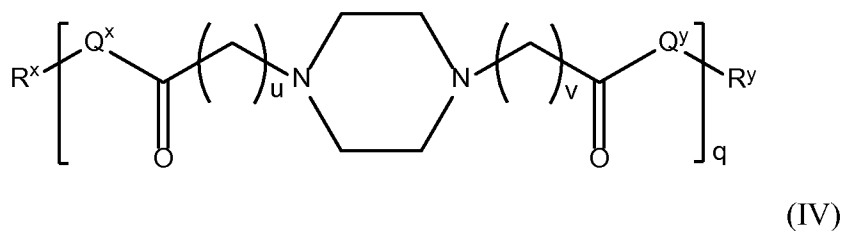
wherein d is an integer from 0 to 25, or

a polyethylene glycol group, represented by Formula (E),



wherein e is an integer from 1 to 400.

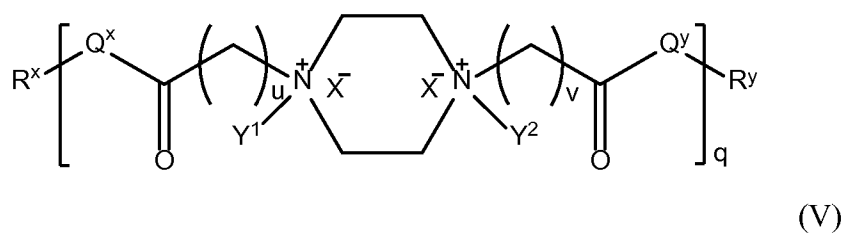
In another aspect of the invention, the amine functional polyamides are a compound comprising the structure of Formula (IV):



wherein:

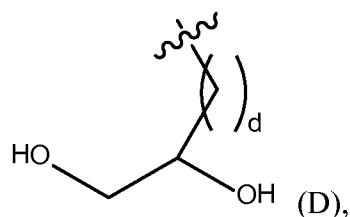
- i) u is 0, 1, 2, or 3;
- ii) v is 0, 1, 2, or 3;
- iii) q is an integer from 1 to 400;
- iv) Q^x is NH, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl;
- v) Q^y is NH-R^w, NH-CH₂-R^w, (C₁-C₁₀)alkyl, or (C₆-C₁₄)aryl, wherein R^w is absent or a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl;
- vi) R^x and R^y are each independently a pharmaceutically acceptable end group.

In yet another aspect of the invention, the amide functional polyamides are a compound comprising the structure of Formula (V):



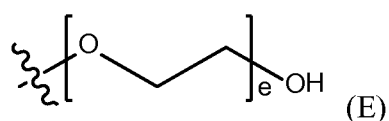
wherein:

- i) u is 0, 1, 2, or 3;
- ii) v is 0, 1, 2, or 3;
- iii) q is an integer from 1 to 400;
- iv) Q^x is NH, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl;
- v) Q^y is NH-R^w, NH-CH₂-R_w, (C₁-C₁₀)alkyl, or (C₆-C₁₄)aryl, wherein R^w is absent or a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl;
- vi) R^x and R^y are each independently a pharmaceutically acceptable end group;
- vii) X is independently a halo or any pharmaceutically acceptable anion,
- viii) Y¹ and Y² are independently H or (C₁-C₁₀)alkyl optionally substituted by one or more substituents selected from the group consisting of (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl, (C₁-C₁₀)alkylamine, -S-O-(C₁-C₁₀)alkyl, -O(O)C-(C₁-C₁₀)alkyl, -(C₁-C₁₀)alkyl-COOH, (C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH, amide, a dihydroxy group, represented by Formula (D),



wherein d is an integer from 0 to 25, or

a polyethylene glycol group, represented by Formula (E)



wherein e is an integer from 1 to 400.

In one aspect of the invention, the amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (I). In another aspect of the invention, amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (II). In yet another aspect of the invention, amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (III). In another aspect of the invention, amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (IV). In another aspect of the invention, amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (V).

In one aspect of the invention, the amine functional polyamides are used for the treatment of mucositis. In another aspect of the invention, the amine functional polyamides are used for the treatment of oral mucositis. In another embodiment of the invention, the amine functional polyamides are used for the treatment of an infection. In yet another embodiment of the invention, the amine functional polyamides are used for the treatment of surgical site infection. In another embodiment of the invention, the amine functional polyamides are used for the treatment of lung infection associated with cystic fibrosis. In another embodiment of the invention, the amine functional polyamides are used for the treatment of *P. aeruginosa* lung infections in CF patients. In yet another embodiment of the invention, the amine functional polyamides are used for the treatment of *P. aeruginosa* lung infections in CF patients where biofilms have formed.

Yet another aspect of the invention is a method of treating a condition selected from mucositis, oral mucositis, and infection comprising administering an amine functional polyamide.

BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWING(S)

Not applicable

DETAILED DESCRIPTION OF THE INVENTION

This invention relates to novel amine functional polyamides. The amine functional polyamides polymers or copolymers and are of varying structures and comprise amine and ammonium groups along the polymer chain.

The amine functional polyamides contain repeat units of amide groups and amine groups; the amine groups can be secondary, tertiary, and quaternary ammonium groups.

Further, the amine functional polyamides of the present invention are of varying molecular weights.

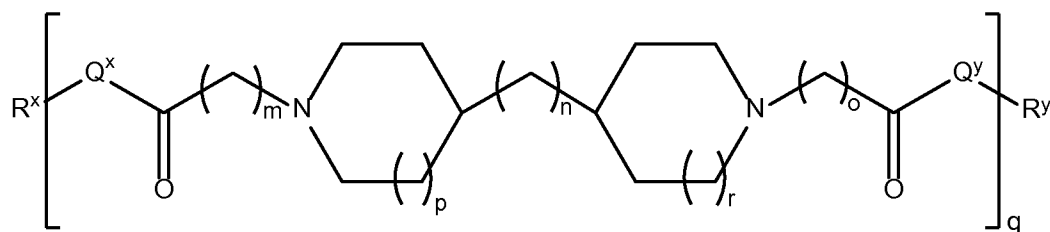
The amine functional polyamides are water soluble.

This invention relates to pharmaceutical compositions comprising polymers or copolymers of amine functional polyamides. This invention also relates to methods of treating and preventing mucositis and infection, including SSI, lung infections in CF patients, and *C. aeruginosa* lung infections in CF patients with or without biofilm formation, with amine functional polyamides. The amine functional polyamides and the pharmaceutical compositions comprising polymers or copolymers of amine functional polyamides can be administered in multiple dosage forms and for systemic or local administration.

This invention relates to the use of amine functional polyamides and pharmaceutical compositions comprising polymers or copolymers of amine functional polyamides as anti-infective agents. The amine functional polyamides and pharmaceutical compositions comprising polymers or copolymers of amine functional polyamides can be used for the treatment of bacterial, fungal, and viral infections, including mucositis, infections and, specifically, surgical site infections, lung infections associated with CF, and *C. aeruginosa* lung infections in CF patients with or without biofilm formation.

The amine functional polyamides can also be used to coat surfaces of various biomedical devices and other surfaces to prevent infection.

In one aspect of the invention, the amine functional polyamides are a compound comprising the structure of Formula (I):



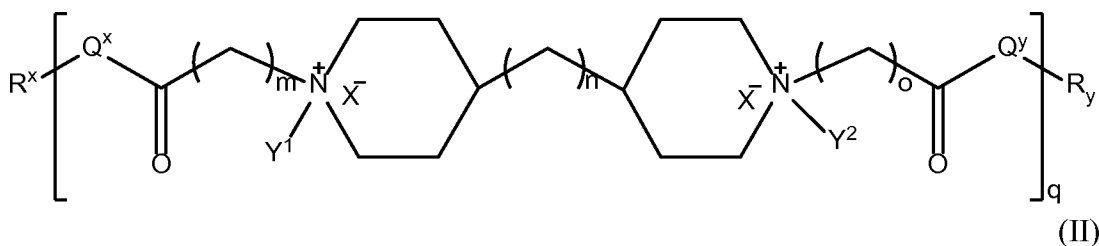
(I)

wherein:

- i) m is 0, 1, 2, or 3;
- ii) n is 0, 1, 2, or 3;
- iii) o is 0, 1, 2, or 3;
- iv) p is 0 or 1;
- v) r is 0 or 1;
- vi) q is an integer from 1 to 400;

- vii) Q^x is NH, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl;
- viii) Q^y is NH-R^w, NH-CH₂-R^w, (C₁-C₁₀)alkyl, or (C₆-C₁₄)aryl, wherein R^w is absent or a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl;
- ix) R^x and R^y are each independently a pharmaceutically acceptable end group.

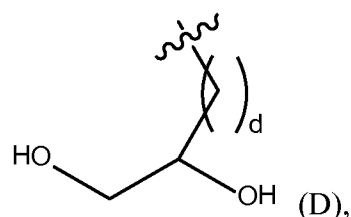
In another aspect of the invention, the amine functional polyamides are a compound comprising the structure of Formula (II):



wherein:

- i) m is 0, 1, 2, or 3;
- ii) n is 0, 1, 2, or 3;
- iii) o is 0, 1, 2, or 3;
- iv) p is 0 or 1;
- v) r is 0 or 1;
- vi) q is an integer from 1 to 400;
- vii) Q^x is NH, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl;
- viii) Q^y is NH-R^w, NH-CH₂-R^w, (C₁-C₁₀)alkyl, or (C₆-C₁₄)aryl, wherein R^w is absent or a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl;
- ix) R^x and R^y are each independently a pharmaceutically acceptable end group;
- x) X⁻ is each independently a halo or any pharmaceutically acceptable anion;
- xi) Y¹ and Y₂ are each independently H or (C₁-C₁₀)alkyl optionally substituted by one or more substituents selected from the group consisting of (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl,

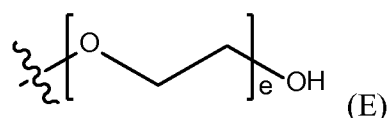
(C₁-C₁₀)alkylamine, -S-O-(C₁-C₁₀)alkyl, -O(O)C-(C₁-C₁₀)alkyl, -(C₁-C₁₀)alkyl-COOH, (C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH, amide, a dihydroxy group, represented by Formula (D),



5

wherein d is an integer from 0 to 25, or

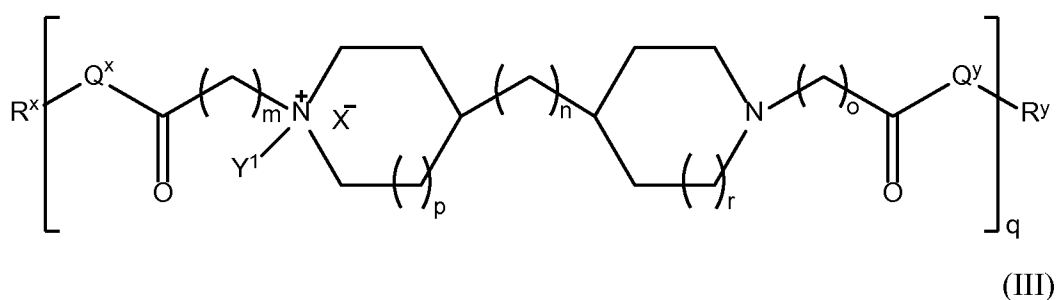
a polyethylene glycol group, represented by Formula (E)



wherein e is an integer from 1 to 25.

In yet another aspect of the invention, the amine functional polyamides are a

10 compound comprising the structure of Formula (III):



wherein:

15

i) m is 0, 1, 2, or 3;

ii) n is 0, 1, 2, or 3;

iii) o is 0, 1, 2, or 3;

iv) p is 0 or 1;

v) r is 0 or 1;

vi) q is an integer from 1 to 400;

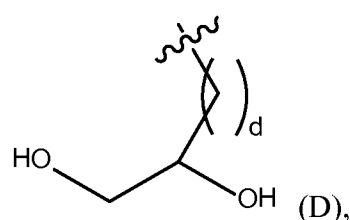
20

vii) Q^x is NH, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl;

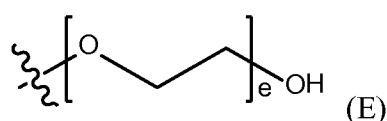
viii) Q^y is NH-R^w, NH-CH₂-R^w, (C₁-C₁₀)alkyl, or (C₆-C₁₄)aryl,

wherein R^w is absent or a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl;

- ii) R^x and R^y are each independently a pharmaceutically acceptable end group;
- ix) X^- is a halo or any pharmaceutically acceptable anion;
- x) Y^1 is H or (C_1-C_{10}) alkyl optionally substituted by one or more substituents selected from the group consisting of (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_3-C_{10}) cycloalkyl, (C_2-C_9) heterocycloalkyl, (C_6-C_{14}) aryl, (C_2-C_9) heteroaryl, (C_1-C_{10}) alkylamine, $-S-O-(C_1-C_{10})$ alkyl, $-O(O)C-(C_1-C_{10})$ alkyl, $-(C_1-C_{10})$ alkyl-COOH, (C_3-C_{10}) cycloalkyl-COOH, $-(O)CH_3$, $-OH$, amide, a dihydroxy group, represented by Formula (D),



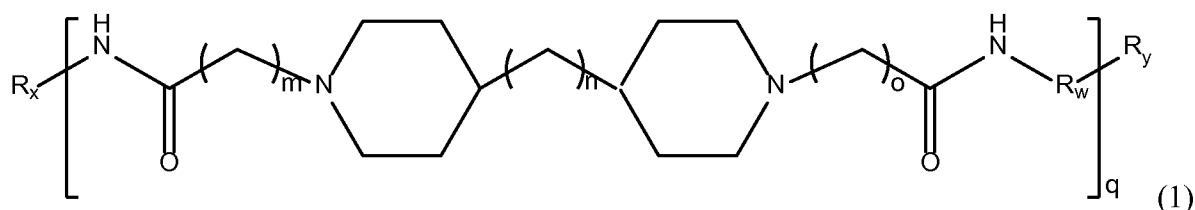
wherein d is an integer from 0 to 25, or
a polyethylene glycol group, represented by Formula (E),



wherein e is an integer from 1 to 400.

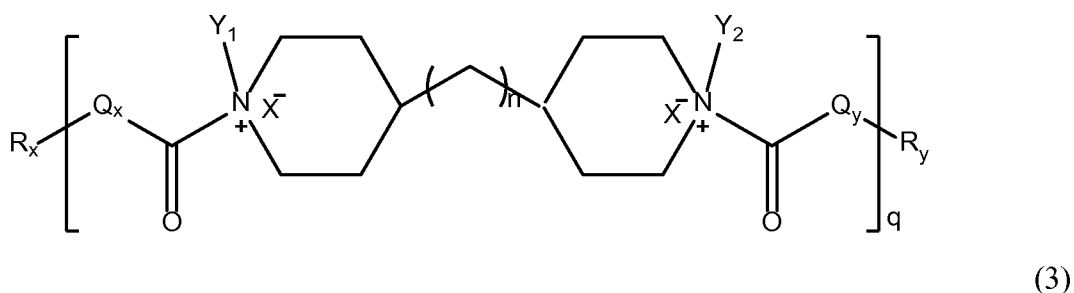
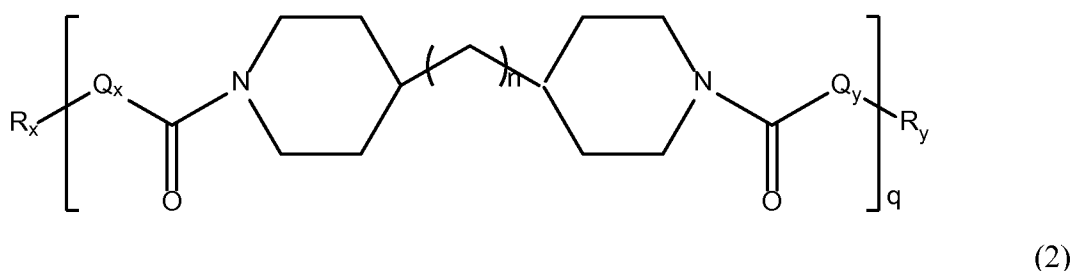
In preferred embodiments of the invention, the amine functional polyamides of are compounds of Formula (I), Formula (II) or Formula (III) where p and r are both 0 and p and r are both 1. In other preferred embodiments of the invention, the amine functional polyamides of are compounds of Formula (I), Formula (II) or Formula (III) where n , p and r are all 0, n is 0 and p and r are both 1, and n is 3 and p and r are both 1.

In a preferred embodiment of the invention, the amide functional polyamides are a compound comprising the structure of Formula (1).



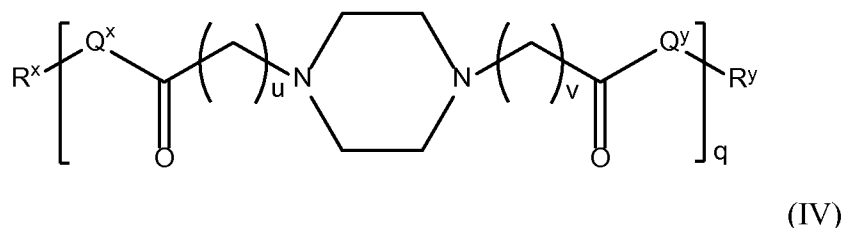
wherein R^w is a (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_6-C_{14}) aryl, or (C_2-C_9) heteroaryl.

In another preferred embodiment of the invention, the amide functional polyamides are a compound comprising the structure of Formula (2) or Formula (3):



5 In yet another preferred embodiment of the invention, the amine functional polyamides are comprised of a compound comprising the structure of Formula (I), Formula (II), Formula (III), Formula (1), Formula (2) or Formula (3), wherein R^x and R^y are independently selected from a methoxy group, a guanidino group, or a guanidinobenzene group.

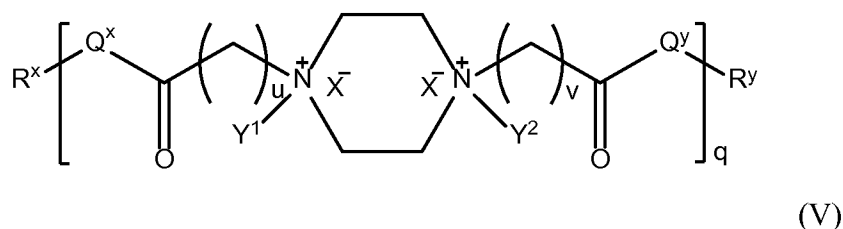
10 In another aspect of the invention, the amine functional polyamides are a compound comprising the structure of Formula (IV):



wherein:

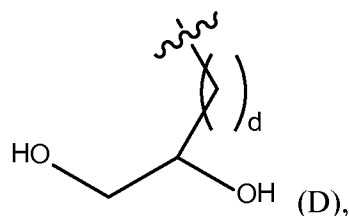
- 15 i) u is 0, 1, 2, or 3;
- ii) v is 0, 1, 2, or 3;
- iii) q is an integer from 1 to 400;
- iv) Q^x is NH, $(\text{C}_1\text{-C}_{10})$ alkyl, $(\text{C}_2\text{-C}_9)$ heteroalkyl, $(\text{C}_3\text{-C}_{10})$ cycloalkyl, $(\text{C}_2\text{-C}_9)$ heterocycloalkyl, $(\text{C}_6\text{-C}_{14})$ aryl, $(\text{C}_2\text{-C}_9)$ heteroaryl;
- 20 v) Q^y is NH-R^w , $\text{NH-CH}_2\text{-R}^w$, $(\text{C}_1\text{-C}_{10})$ alkyl, or $(\text{C}_6\text{-C}_{14})$ aryl, wherein R^w is absent or a $(\text{C}_1\text{-C}_{10})$ alkyl, $(\text{C}_2\text{-C}_9)$ heteroalkyl, $(\text{C}_6\text{-C}_{14})$ aryl, or $(\text{C}_2\text{-C}_9)$ heteroaryl;
- vii) R^x and R^y are each independently a pharmaceutically acceptable end group.

In yet another aspect of the invention, the amide functional polyamides are a compound comprising the structure of Formula (V):

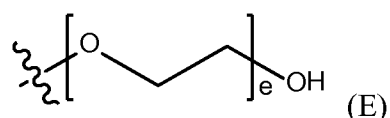


wherein:

- i) u is 0, 1, 2, or 3;
- ii) v is 0, 1, 2, or 3;
- iii) q is an integer from 1 to 400;
- iv) Q^x is NH, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl;
- v) Q^y is NH-R^w, NH-CH₂-R_w, (C₁-C₁₀)alkyl, or (C₆-C₁₄)aryl, wherein R^w is absent or a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl;
- vii) R^x and R^y are each independently a pharmaceutically acceptable end group;
- ix) X⁻ is independently a halo or any pharmaceutically acceptable anion,
- x) Y¹ and Y² are independently H or (C₁-C₁₀)alkyl optionally substituted by one or more substituents selected from the group consisting of (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl, (C₁-C₁₀)alkylamine, -S-O-(C₁-C₁₀)alkyl, -O(O)C-(C₁-C₁₀)alkyl, -(C₁-C₁₀)alkyl-COOH, (C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH, amide, a dihydroxy group, represented by Formula (D),



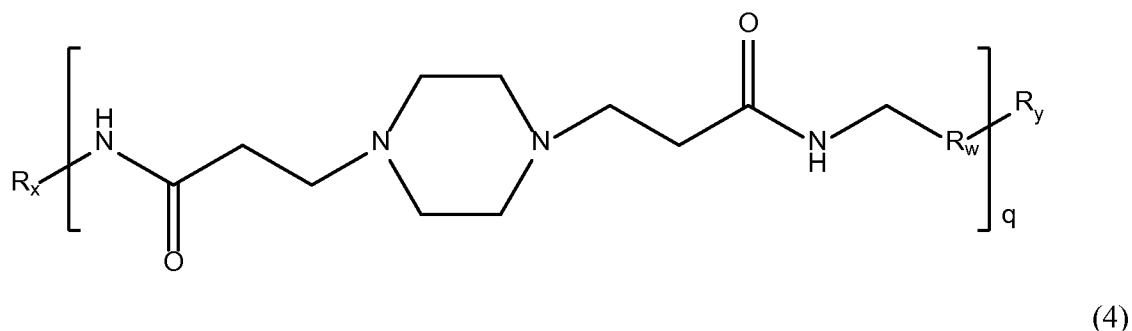
wherein d is an integer from 0 to 25, or
a polyethylene glycol group, represented by Formula (E)



wherein e is an integer from 1 to 400.

In a preferred embodiment of the invention, the amine functional polyamides of are compounds of Formula (IV) or Formula (V) where u and v are both 2.

5 In a preferred embodiment of the invention, the amide functional polyamides are a compound comprising the structure of Formula (4).



In one aspect of the invention, the amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (I). In another aspect of the invention, amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (II). In yet another aspect of the invention, amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (III). In preferred embodiments of the invention, the amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (1), Formula (2), or Formula (3). In yet another preferred embodiment of the invention, the amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (I), Formula (II), Formula (III), Formula (1), Formula (2) or Formula (3), wherein R_x and R_y are independently selected from a methoxy group, a guanidino group, or a guanidinobenzene group.

In another aspect of the invention, amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (IV). In another aspect of the invention, amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (V). In a preferred embodiment of the invention, the amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (4).

In another preferred embodiment of the invention, the amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (I), (II), (III), (IV), (V), (1), (2), (3) or (4) for use in the treatment or prevention of a condition selected from mucositis, oral mucositis and infection. In yet another preferred
5 embodiment, the amine functional polyamides are a pharmaceutical composition comprising a compound comprising the structure of Formula (I), (II), (III), (IV), (V), (1), (2), (3) or (4) for use in the treatment or prevention of a surgical site infection, a lung infection associated with cystic fibrosis, a *Pseudomonas aeruginosa* lung infection, and a *Pseudomonas aeruginosa* lung infection where biofilms are present.

10 In one embodiment of the invention, the amine functional polyamides are polymers. In some embodiments, the polymers may comprise a monomer comprising a compound having a repeat unit according to any of Formulas (I), (II), (III), (IV), (V), (1), (2), (3) or (4).

In one embodiment of the invention, the amine functional polyamides are copolymers. In some embodiments, the copolymers may comprise a monomer comprising a compound
15 having at least one unit according to any of Formulas (I), (II), (III), (IV), (V), (1), (2), (3) or (4) which is copolymerized with one or more other comonomers or oligomers or other polymerizable groups. Non-limiting examples of suitable comonomers which may be used alone or in combination with at least one unit according to any of Formulas (I), (II), (III), (IV), (V), (1), (2), (3) or (4) to form the amine functional polyamides are presented in Table

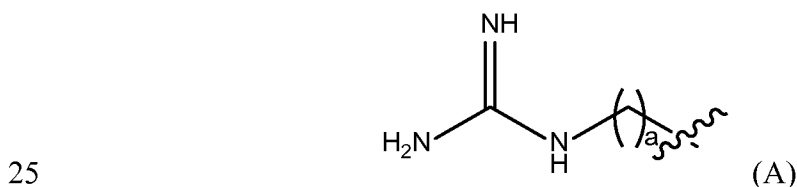
20 1.

In one embodiment of the invention, the amine functional polyamides are polymers or copolymers comprised of about 1 to about 400 repeat units according to any of Formulas (I), (II), (III), (IV), (V), (1), (2), (3) or (4). In one aspect of the invention, the amine functional polyamides are polymers or copolymers comprised of about 1 to about 200 repeat units
25 according to any of Formulas (I), (II), (III), (IV), (V), (1), (2), (3) or (4). In another aspect of the invention, the amine functional polyamides are polymers or copolymers comprised of about 1 to about 100 repeat units according to any of Formulas (I), (II), (III), (IV), (V), (1), (2), (3) or (4). In some embodiments, the amine functional polyamides are polymers or copolymers comprised of about 1 to about 50 repeat units according to any of Formulas (I),
30 (II), (III), (IV), (V), (1), (2), (3) or (4). In an additional embodiment, the amine functional polyamides are polymers or copolymers comprised of about 1 to about 25 repeat units according to any of Formulas (I), (II), (III), (IV), (V), (1), (2), (3) or (4). In yet another embodiment, the amine functional polyamides are polymers or copolymers comprised of

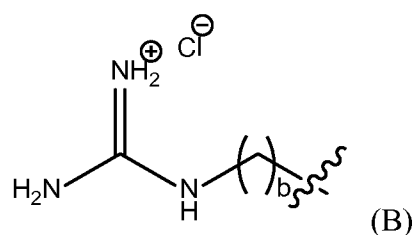
about 1 to about 10 repeat units according to any of Formulas (I), (II), (III), (IV), (V), (1), (2), (3) or (4). In other embodiments, the amine functional polyamides are polymers or copolymers comprised of about 5 to about 40 repeat units according to any of Formulas (I), (II), (III), (IV), (V), (1), (2), (3) or (4). In one embodiment, the amine functional polyamides are polymers or copolymers comprised of about 5 to about 30 repeat units according to any of Formulas (I), (II), (III), (IV), (V), (1), (2), (3) or (4). In another embodiment, the amine functional polyamides are polymers or copolymers comprised of about 5 to about 25 repeat units according to any of Formulas (I), (II), (III), (IV), (V), (1), (2), (3) or (4). In yet another embodiment, the amine functional polyamides are polymers or copolymers comprised of about 5 to about 10 repeat units according to any of Formulas (I), (II), (III), (IV), (V), (1), (2), (3) or (4).

In one aspect of the invention, the amine functional polyamides have a molecular weight less than about 10,000 g/mol. In another aspect of the invention, the amine functional polyamides have a molecular weight less than about 9,000 g/mol. In an additional aspect of the invention, the amine functional polyamides have a molecular weight less than about 8,000 g/mol. In yet another aspect of the invention, the amine functional polyamides have a molecular weight less than about 7,000 g/mol.

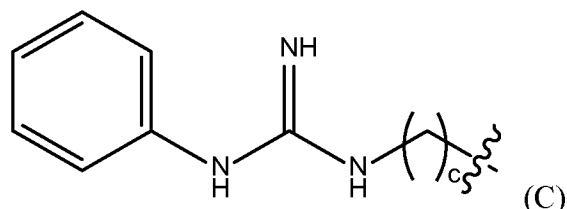
In one aspect of the invention, the amine functional polyamides are optionally, independently terminated (R^x and R^y) with a pharmaceutically acceptable end group. Representative examples of pharmaceutically acceptable end groups will be obvious to one of skill in the art, including H, (C₁-C₁₀)alkyl, (C₂-C₁₀)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl, (C₁-C₁₀)alkylamine, -O(O)C-(C₁-C₁₀)alkyl, (C₁-C₁₀)alkyl-COOH, (C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH, amide, a guanidino group represented by Formula (A)



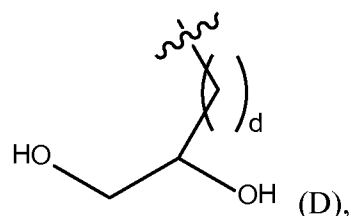
wherein a is an integer from 0 to 25, a guanidinium chloride group represented by Formula (B),



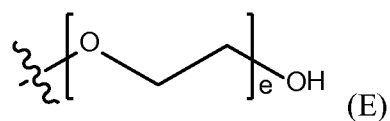
wherein b is an integer from 0 to 25, a guanidinobenzene group represented by Formula (C),



wherein c is an integer from 0 to 25, a dihydroxy group, represented by Formula (D),



wherein d is an integer from 0 to 25, or a polyethylene glycol group, represented by Formula (E)



wherein e is an integer from 1 to 400

The number of repeat units and the molecular weight of the amine functional polyamides are controlled by synthesis of the compound. Methods of preparing preferred amine functional polyamides of the invention and controlling for the number of repeat units and molecular are described in Example 3.

Table 1: Amine Functional Polyamides

Polymer Description	Structure
Poly (4,4'-trimethylene dipiperidine bispropanoic acid--N,N'-dimethyl-1,3-diaminopropane)	

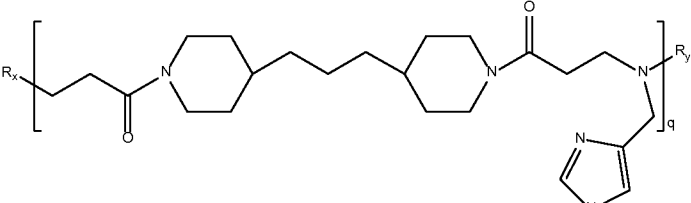
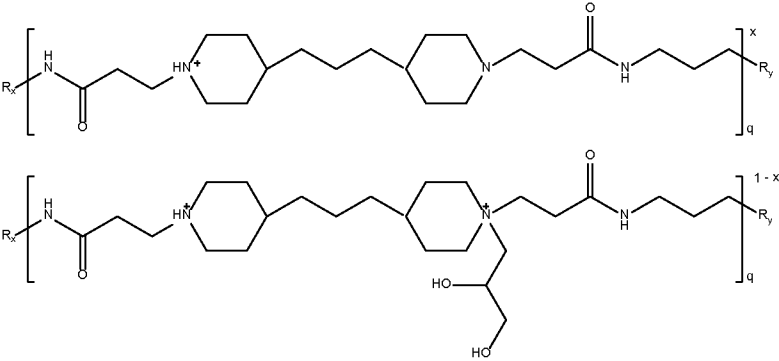
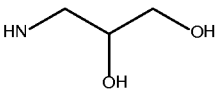
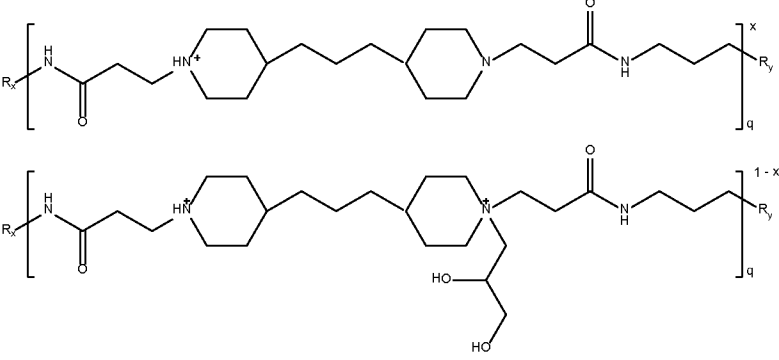
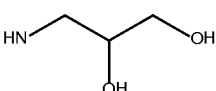
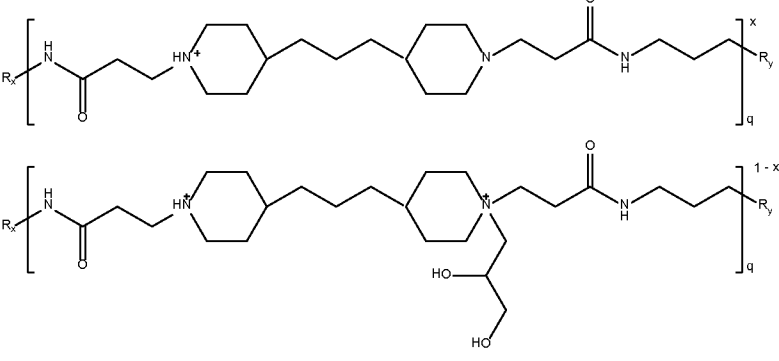
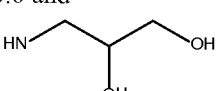
Poly (4,4'-trimethylene dipiperidine bispropanoic acid-4,4'-trimethylene dipiperidine)	
Poly (4,4'-trimethylene dipiperidine bispropanoic acid-piperazine)	
Poly(4,4'-trimethylene dipiperidine bispropanoic acid-diaminoethane) Mw <10K	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane) Mw <10K	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane) Mw >10K	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane) Mw 1650	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane) Mw 7.7K	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane) Mw 3K	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane) Mw 5K	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane) , Mw 3250	

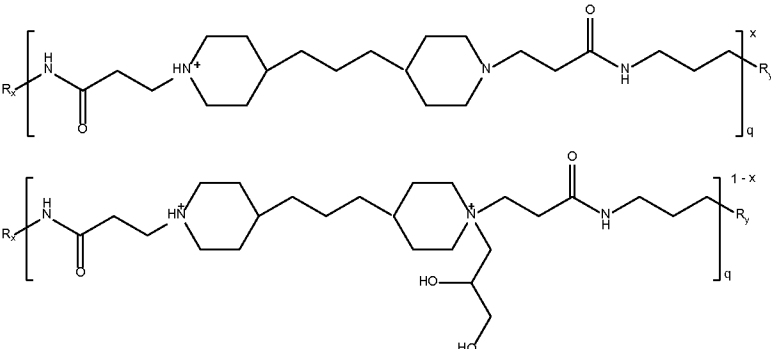
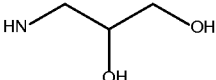
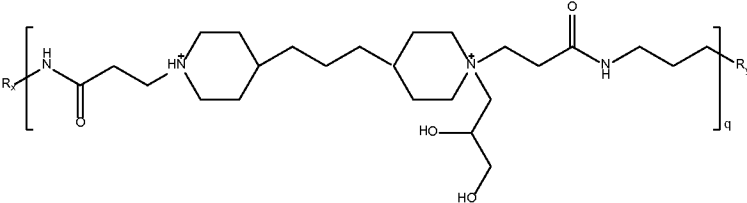
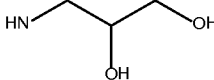
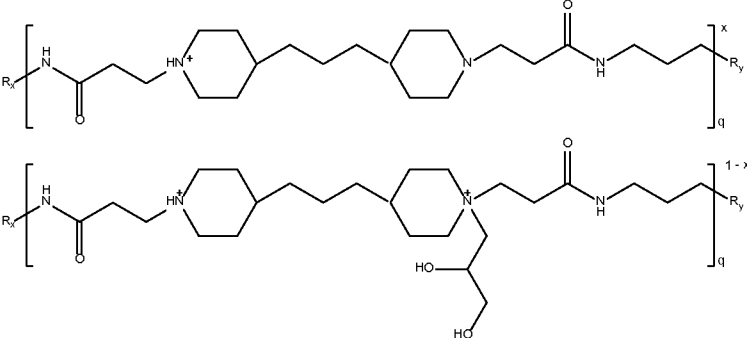
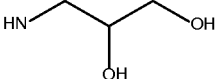
Poly(4,4-trimethylene dipiperidinebispropionic acid-diaminopropane) , Mw 4700	
Poly(4,4-trimethylene dipiperidinebispropionic acid-diaminopropane) , Mw 2500	
Poly(4,4-trimethylene dipiperidinebispropionic acid-diaminopropane)	
Poly(4,4-trimethylene dipiperidinebispropionic acid-diaminopropane) Mw 1400	
Poly(4,4-trimethylene dipiperidinebispropionic acid-diaminobutane) Mw <10K	
Poly(4,4-trimethylene dipiperidinebispropionic acid-diaminobutane) Mw >10K	
Poly(4,4-trimethylene dipiperidinebispropionic acid-diaminotriPEG) Mw <10K	
Poly(4,4-trimethylene dipiperidinebispropionic acid-diaminotriPEG) Mw >10K	
Poly(4,4-trimethylene dipiperidinebispropionic acid-N(2-aminoethyl)-diaminoethane)	
Poly(4,4'-trimethylenedipiperidinebispropionic acid-2,2'-diamino diethylamine) Mw 5.5K	

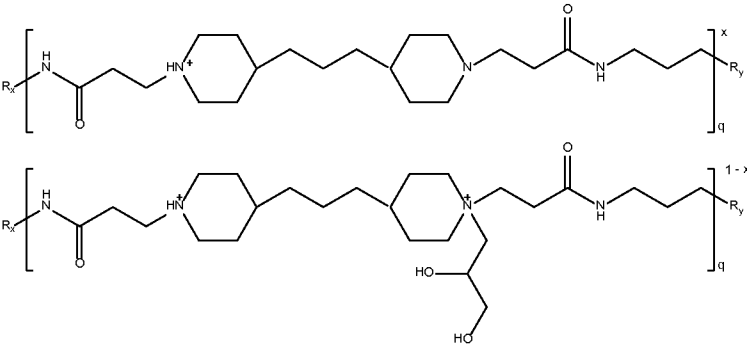
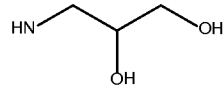
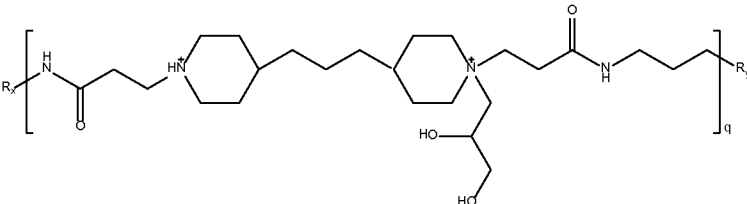
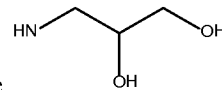
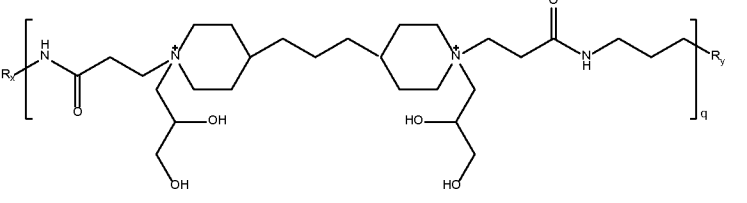
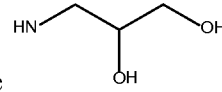
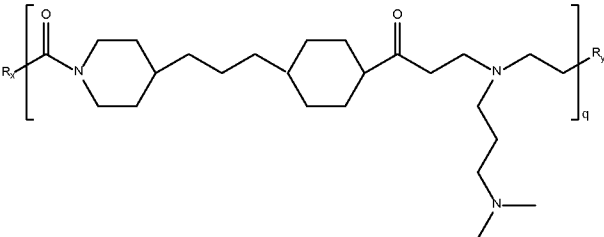
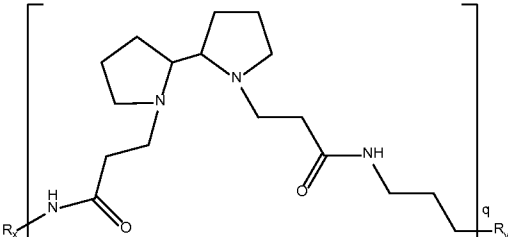
Poly(4,4'-trimethylenedipiperidinebispropanoic acid-2,2'-diamino diethylamine) Mw ~14,000	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-1,4-benzyl diamine) Mw <10K	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-N(3-aminopropyl)1,3-propane diamine) Mw <10K	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-3,3'-diamino-N-methyl-dipropylamine) Mw <10K	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-2,2'-diamino-N-methyl-diethylamine) Mw <10K	
Poly(piperazinebispropanoic acid-1,2-bis(2-aminoethoxy)ethane) Mw <10K	
Poly(piperazinebispropanoic acid-2,2'-diaminodiethylamine) Mw <10K	
Poly(piperazinebispropanoic acid-N-methyl-2,2'-diaminodiethylamine) Mw <10K	
Poly(piperazinebispropanoic acid-N(3-aminopropyl)1,3-propane diamine) Mw <10K	

Poly(piperazinebispropanoic acid-3,3'-diamino-N-methyl-dipropylamine) Mw <10K	
Poly(piperazinebispropanoic acid-1,3-diaminopropane) Mw ~3,700	
Poly(piperazinebispropanoic acid-1,4-diaminobutane) Mw ~4,400	
Poly(4,4'-dipiperidinebispropanoic acid-2,2'-diamino diethylamine) Mw <5K	
Poly(4,4'-dipiperidinebispropanoic acid-2,2'-diamino diethylamine) Mw 5.1K	
Poly(4,4'-dipiperidinebispropanoic acid-2,2'-diamino N-methyl diethylamine) Mw <5K	
Poly(4,4'-dipiperidinebispropanoic acid-2,2'-diamino N-methyl diethylamine) Mw <5K	
Poly(4,4'-dipiperidinebispropanoic acid-3,3'-diamino-dipropylamine) Mw <5K	
Poly(4,4'-dipiperidinebispropanoic acid-3,3'-diamino-dipropylamine) Mw <5K	
Poly(4,4'-dipiperidinebispropanoic acid-3,3'-diamino-N-methyl-dipropylamine) Mw <5 k	

Poly(4,4'-dipiperidinebispropionic acid-3,3'-diamino-N-methyl-dipropylamine) Mw ~5.5K	
Poly(4,4'-dipiperidinebispropionic acid-ethylenediamine) Mw <5K	
Poly(4,4'-dipiperidinebispropionic acid-1,3-diaminopropane) Mw <5K	
Poly(4,4'-dipiperidinebispropionic acid-1,4-diaminobutane) Mw <5K	
Poly(4,4'-trimethylenedipiperidinebispropionic acid-bis(4-aminobutyl)ether) Mw <5K	
Poly(4,4'-trimethylenedipiperidinebispropionic acid-2-hydroxy 1,3-diaminopropane) Mw <5K	
Poly(4,4'-trimethylenedipiperidine-1,3-diaminopropane-N,N'-di-3-propionic acid) Mw <5K	
Poly (4,4'-trimethylene dipiperidine bispropionic acid--N,N'-dimethyl-1,3-diaminopropane) , Mw 1K	
Poly(4,4'-trimethylene dipiperidinebispropionic acid-4,4'-dipiperidine), Mw 10631	

<p>Poly(4,4-trimethylene dipiperidinebispropionic acid-histamine), Mw 2.3K</p>	
<p>40 mol% glycidol modified poly(4,4-trimethylene dipiperidinebispropionic acid-diaminopropane), Mw 8000</p>	 <p>Wherein x is 0.6 and</p> <p>R_x and R_y are  or NH_2.</p>
<p>40 mol% glycidol modified poly(4,4-trimethylene dipiperidinebispropionic acid-diaminopropane), Mw 4700</p>	 <p>Wherein x is 0.6 and</p> <p>R_x and R_y are  or NH_2.</p>
<p>40 mol% glycidol modified Poly(4,4-trimethylene dipiperidinebispropionic acid-diaminopropane) Mw 5000</p>	 <p>Wherein x is 0.6 and</p> <p>R_x and R_y are  or NH_2.</p>

<p>40 mol% glycidol modified Poly(4,4- trimethylene dipiperidinebispr opanoic acid- diaminopropane) Mw 5000</p>	 <p>Wherein x is 0.6 and</p> <p>R_x and R_y are  or NH_2.</p>
<p>100 mol% glycidol modified Poly(4,4- trimethylene dipiperidinebispr opanoic acid- diaminopropane) , Mw 8K</p>	 <p>Wherein R_x and R_y are  or NH_2.</p>
<p>25 mol% glycidol modified Poly(4,4- trimethylene dipiperidinebispr opanoic acid- diaminopropane) , Mw 7800</p>	 <p>Wherein x is 0.75 and</p> <p>R_x and R_y are  or NH_2.</p>

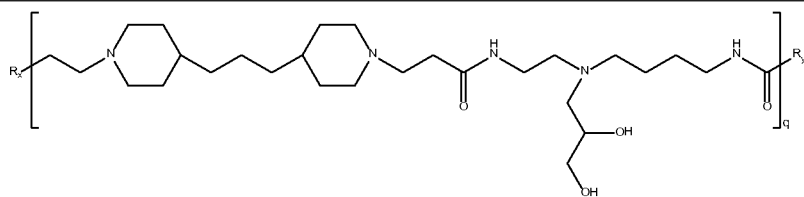
<p>50 mol% glycidol modified Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane), Mw 7800</p>	 <p>Wherein x is 0.5 and</p> <p>R_x and R_y are  or NH_2.</p>
<p>150 mol% glycidol modified Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane), Mw 7800</p>	 <p>Wherein R_x and R_y are  or NH_2.</p>
<p>Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane) modified with 200mol% of glycidol, Mw 7800</p>	 <p>Wherein R_x and R_y are  or NH_2.</p>
<p>Poly (4,4'-trimethylene dipiperidine bispropanoic acid--3-(dimethylamino) 1-propylamine), Mw 1K</p>	
<p>Poly(2,2'-bipyrrolidine bispropanoic acid-diaminopropane), Mw 2.5K</p>	

Poly(2,2'-bipyrrolidine bispropanoic acid-butyl diamine)	
Poly(2,2'-bipyrrolidine bispropanoic acid-penta diamine)	
Poly(2,2'-bipyrrolidine bispropanoic acid-ethyl diamine)	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-aminomethyl benzene)	
Poly[4,4'-trimethylene dipiperidinebispropanoic acid-(1-aminomethyl-4-guanidinomethyl benzene)]	
Carboxy terminated Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane)	
Methyl ester terminated Poly(4,4'-trimethylene)	

dipiperidinebispr opanoic acid- diaminopropane)	
Guanidine terminated Poly(4,4- trimethylene dipiperidinebispr opanoic acid- diaminopropane)	
Guanidine terminated Poly(4,4- trimethylene dipiperidinebispr opanoic acid- diaminopropane) Mw 4700	
Guanidine terminated poly(4,4- trimethylene dipiperidinebispr opanoic acid- diaminopropane) Mw 7700	
Poly(4,4- trimethylene dipiperidinebispr opanoic acid- diaminopropane) (guanidine ended)	
4,4'-trimethylene dipiperidinebispr opanoic acid- diaminopropane	
4- guanidinobenzen e terminated Poly(4,4- trimethylene dipiperidinebispr opanoic acid- diaminopropane)	
Poly(4,4'- trimethylenedipi peridine bisethylacrylami de-co-1,3- diamine propane)	

Poly(4,4'-trimethylenedipiperidine bisethylacrylamide-co-1-amino-3-guanidine propane)	
4,4'-trimethylene dipiperidinebispropanoic acid-1-amino-3-guanidine propane	
Poly(4,4'-trimethylenedipiperidine bisethylacrylamide-1,3-diamine propane)-co-Poly(4,4'-trimethylenedipiperidine bisethylacrylamide-1-aminobutyl-3-carbamoylpyridinium)	
Poly(4,4'-trimethylenedipiperidine bisethylacrylamide-1-aminobutyl-3-carbamoylpyridinium)	
4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane pentamer	
4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane heptamer	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-N-glycidol dipropylene triamine)	
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-N-glycidol diethylene triamine)	

Poly(4,4'-
trimethylene
dipiperidinebispr
opanoic acid-N-
glycidol
diethylene
triamine)



In an embodiment of the invention, the amine functional polyamides are administered as a pharmaceutical composition. In another embodiment of the invention, the amine functional polyamides are administered in an effective amount to achieve the desired therapeutic effect. The skilled artisan will be able to determine the effective amount of the amine functional polyamides depending on the individual and the condition being treated.

In one embodiment of the invention, the amine functional polyamides are used in the treatment all forms of mucositis, and are particularly effective when used to treat oral mucositis. Treatment includes prophylactic and therapeutic uses of the disclosed amine functional polyamides and uses of the disclosed pharmaceutical compositions comprising amine functional polyamides. Desired prophylactic effects include prevention and inhibition of mucositis, reduction in severity of mucositis, reduction in size of mucositis lesions and reduction in likelihood of developing mucositis through the application or administration of amine functional polyamides or pharmaceutical compositions comprising amine functional polyamides. Desired therapeutic effects include amelioration of the discomfort associated with the mucositis, and/or increased rate of healing of mucositis lesion.

In one embodiment, the amine functional polyamides and pharmaceutical compositions comprising amine functional polyamides can be used to treat all forms of infection, including but not limited to SSI, lung infection in CF patients, and *C. aeruginosa* lung infection in CF patients with or without biofilm formation. The amine functional polyamides and pharmaceutical compositions comprising amine functional polyamides can be used in prophylactic and therapeutic applications to treat and prevent infection.

In another embodiment, the amine functional polyamides and pharmaceutical compositions comprising amine functional polyamides can be used to treat all forms of SSIs. Treatment includes prophylactic and therapeutic uses of the disclosed amine functional polyamides and uses of the disclosed pharmaceutical compositions comprising amine functional polyamides. A desired prophylactic use is the immediate administration of amine functional polyamides or pharmaceutical compositions comprising amine functional polyamides to the surgical wound post-surgery in order to prevent and/or reduce the likelihood of developing a SSI. Another desired prophylactic use is the administration of amine functional polyamides or pharmaceutical compositions comprising amine functional polyamides prior to surgery in order to prevent and/or reduce the likelihood of developing a SSI. Desired therapeutic effects include the treatment of an existing SSI through the

application or administration of amine functional polyamides or pharmaceutical compositions comprising an amine functional polyamide.

In another embodiment, the amine functional polyamides and pharmaceutical compositions comprising amine functional polyamides can be used to treat all forms of lung infections and chronic lung infections associated with CF, including *C. aeruginosa* lung infections in CF patients with or without biofilm formation. Treatment includes prophylactic and therapeutic uses of the disclosed amine functional polyamides and uses of the disclosed pharmaceutical compositions comprising amine functional polyamides. Desired therapeutic effects include the treatment of an existing lung infection or chronic lung infection through the administration of amine functional polyamides or pharmaceutical compositions comprising an amine functional polyamide. In one embodiment, the amine functional polyamides and pharmaceutical compositions comprising amine functional polyamides are used to treat *P. aeruginosa* infections associated with CF without biofilm formation. In another embodiment, the amine functional polyamides and pharmaceutical compositions comprising amine functional polyamides are used to treat *P. aeruginosa* infections associated with CF with biofilm formation. A desired prophylactic use is the administration of amine functional polyamides or pharmaceutical compositions comprising amine functional polyamides to the CF patient in order to prevent and/or reduce the likelihood of developing a lung infection, including *C. aeruginosa* lung infections. Desired therapeutic effects include the treatment of an existing lung infection or chronic lung infection through the administration of amine functional polyamides or pharmaceutical compositions comprising an amine functional polyamide.

The amine functional polyamides of the present invention may be administered alone or in a pharmaceutical composition comprising amine functional polyamides. Suitable pharmaceutical compositions may comprise an amine functional polyamide and one or more pharmaceutically acceptable excipients. The form in which the polymers are administered, for example, powder, tablet, capsule, solution, or emulsion, depends in part on the route by which it is administered. The amine functional polyamides can be administered, for example, topically, orally, intranasally, by aerosol or rectally. Suitable excipients include, but are not limited to, are inorganic or organic materials such as gelatin, albumin, lactose, starch, stabilizers, melting agents, emulsifying agents, salts and buffers. Suitable pharmaceutically acceptable excipients for topical formulations such as ointments, creams and gels include, but

are not limited to, commercially available inert gels or liquids supplemented with albumin, methyl cellulose, or a collagen matrix.

The amine functional polyamides and pharmaceutical compositions comprising amine functional polyamides can be administered alone or in combination with one or more additional drugs. Additional drugs administered in combination with the amine functional polyamides and pharmaceutical compositions comprising amine functional polyamides of the present invention include antibiotics and other compounds, including those used prophylactically and/or therapeutically for the treatment or prevention of mucositis and infection, including SSI and lung infection and chronic lung infection associated with CF, especially *P. aeruginosa* infection, with or without biofilm formation. The additional drugs may be administered concomitantly with the amine functional polyamide or pharmaceutical compositions comprising amine functional polyamides. The additional drugs may also be administered in series with the amine functional polyamide or pharmaceutical compositions comprising amine functional polyamides. The pharmaceutical composition comprising amine functional polyamides may also further comprise a drug used prophylactically and/or therapeutically for the treatment or prevention of mucositis and infection, including SSI and lung infection and chronic lung infection associated with CF, especially *P. aeruginosa* infection, with or without biofilm formation.

Examples

Example 1: *In vitro* Studies

Example 1- 1: Cytotoxicity Assay, RPTEC Cells and NHDF Cells

Mammalian cell cytotoxicity assays were performed using two primary human cell lines: renal proximal tubule epithelial cells (RPTEC – Cambrex CC-2553) and normal human dermal fibroblasts (NHDF – Cambrex CC-2509). Cells were plated at 3,000 cells/well (RPTEC) or 5,000 cells/well (NHDF) in 96-well plates and incubated overnight at 37°C. The compounds were added to the wells, and the cells were incubated for 4 days. Alomar Blue was added to one set of plates and incubated for 4 hours. The plates were read when the compound was added (time zero) and at the end of the study. Fluorescence was read using 530 nm (excitation) and 590 nm (emission) according to the manufacturer's instructions. The 50% inhibitory concentration (IC₅₀) was calculated as 50% of the maximum signal minus the value at time zero. The 50% lethal concentration (LC₅₀) was calculated as 50% of the time zero value minus the minimum signal.

Table 2 displays the renal proximal tubule epithelial cells and normal human dermal fibroblasts IC₅₀ and LC₅₀ for selected compounds.

Example 1- 2: Cytotoxicity Assay, Human Lung Epithelial Cells

Cytotoxicity of the polymers towards human lung epithelial cells was performed using human lung epithelial Carcinoma cell line (A 549 –ATCC # CCL-185). The cells were incubated for 96 hours at 37°C with 5% CO₂ in a 96-well plate. CellTiter-Glo[®] (Promega) reagent was added to the plates. The plates were read by measuring the luminescence arising from luciferase catalyzed reaction of luciferin with ATP according to the manufacturer's suggested protocol. The concentration of ATP is directly proportional to cell viability; accordingly, higher luminescence measures high cell viability.

Table 2 displays the human lung epithelial cells IC₅₀ for selected compounds.

Example 1- 3: Erythrocyte Lysis Assay

The compounds were incubated overnight at 37°C in Dulbecco's phosphate-buffered saline containing fresh washed erythrocytes at a hematocrit of 1%. After incubation, the plates were centrifuged and the supernatant transferred to flat-bottomed 96-well plates. The supernatant was assayed using the QuantiChrom Hemoglobin kit according to the manufacturer's instructions. The IC₅₀ values were calculated using GraphPad Prism.

Table 2 displays the IC₅₀ values for selected compounds.

Example 1- 4: Minimum Inhibitory Concentration Assay

The minimum inhibitory concentration (MIC) assay determines the lowest concentration of an antimicrobial agent required to inhibit the growth of test organisms after incubation. MIC assays were performed against an internal standard panel of organisms to identify compounds with antimicrobial activity. The MIC assay was subsequently repeated against other specialized microbial panels. Assays were conducted against the following clinically relevant microorganisms: *Staphylococcus aureus subsp. aureus*, *Staphylococcus epidermis*, *Escherichia coli*, *Pseudomonas aeruginosa*, *Haemophilus influenzae*. The compounds were tested for bacteriocidal activity, time course of killing, toxicity against tissue culture cells grown *in vitro*, and in some cases were tested for antimicrobial activity *in vivo*.

The MIC assays were performed according to the Performance Standards for Antimicrobial Susceptibility Testing, 2006, vol. M100-S15, Fifteenth Informational Supplement, NCCLS, 940 West Valley Road, Suite 1400, Wayne, PA 19087.

The polymers tested were dissolved in 0.85% saline to a final concentration of either 830 or 1000 $\mu\text{g/mL}$ and the pH was adjusted to 7.0. The solution was then filter-sterilized through a 0.22 μm filter. Two-fold serial dilutions of polymer were prepared in Mueller-Hinton broth with cations aliquotted into 96-well microtiter plates. The plates were then
5 inoculated with 5×10^5 cells/mL of target organism and incubated 18-24 hours at 35°C. The optical density (OD) was read at 590 nm, and microorganism growth was scored (OD > 0.1 is considered to be growth; OD < 0.1 is considered to be growth inhibition). The MIC value is the lowest concentration of compound that inhibits growth; accordingly, a higher MIC value indicates less potency where a lower MIC valued indicated more potency.

10 MIC values of representative amine functional polyamides against clinically relevant microorganisms are presented in **Table 2**.

Table 2: *In vitro* Results of Representative Amine Functional Polyamides
Cytotoxicity Assay [Kidney Epithelial and Human Dermal Fibroblast IC₅₀ and LC₅₀],
Erythrocyte Lysis Assay [Hemolysis IC₅₀],
and MIC values against Clinically Relevant Microorganisms

5

Polymer Description	Cytotoxicity (Kidney Epithelial IC₅₀)	Cytotoxicity (Kidney Epithelial LC₅₀)	Cytotoxicity (Human Dermal Fibroblast IC₅₀)	Cytotoxicity (Human Dermal Fibroblast LC₅₀)	Lung Epi IC₅₀	Erythrocyte Lysis (Hemolysis IC₅₀)	Staphylococcus aureus subsp. aureus (MIC)	Staphylococcus epidermis (MIC)	Escherichia coli (MIC)	Pseudomonas aeruginosa (MIC)	Haemophilus influenzae (MIC)
Poly(4,4-trimethylene dipiperidinebi soproanoic acid-diaminopropane) <10K	32.2	55.7	141.3	189.3	391	>6400	1.0	0.3	2.0	32	16.0
Poly(4,4-trimethylene dipiperidinebi soproanoic acid-diaminobutane) >10K	5.3	8.7	16.3	22.1	74.1	>6400	4.0	1.0	4.0	32.0	16.0
Poly(4,4-trimethylene dipiperidinebi soproanoic acid-diaminobutane) <10K	28.6	61.9	138.3	196.2	377.9	>6400	4.0	1.0	4.0	128.0	16.0
Poly(4,4-trimethylene dipiperidinebi soproanoic acid-diaminoethane) <10K	33	78	178	273	799	>6400	32.0	4.0	16.0	128.0	128.0
Poly(4,4-trimethylene dipiperidinebi soproanoic acid-diaminotriPE G) <10K	45.6	133.3	218.4	436.5	427.1	2066	128.0	128.0	32.0	128.0	128.0
Poly(4,4-trimethylene dipiperidinebi soproanoic acid-diaminotriPE G) >10K	2.7	6.9	8.5	18.8	29.0	1792	128.0	8.0	8.0	64.0	128.0
Poly(4,4-trimethylene dipiperidinebi soproanoic acid-diaminopropane) <10K	32.2	55.7	141.3	189.3	--	>6400	1.0	0.5	4.0	64.0	16.0

Polymer Description	Cytotoxicity (Kidney Epithelial IC₅₀)	Cytotoxicity (Kidney Epithelial LC₅₀)	Cytotoxicity (Human Dermal Fibroblast IC₅₀)	Cytotoxicity (Human Dermal Fibroblast LC₅₀)	Lung Epi IC₅₀	Erythrocyte Lysis (Hemolysis IC₅₀)	Staphylococcus aureus subsp. aureus (MIC)	Staphylococcus epidermis (MIC)	Escherichia coli (MIC)	Pseudomonas aeruginosa (MIC)	Haemophilus influenzae (MIC)
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-N(2-aminoethyl)-diaminoethane)	<1.46 ₃	<1.463	1.7	2.0	6.5	9	4.0	0.5	8.0	16.0	64.0
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-N(3-aminopropyl) 1,3-propane diamine) <10K	<1.46 ₃	<1.463	<1.463	<1.463	13.9	2220.0	1.0	0.5	8.0	64.0	32.0
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane) <10K	10.8	25.6	62.6	99.7	333.0	>6400	1.0	0.5	4.0	16.0	16.0
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-3,3'-diamino-N-methyl-dipropylamine) <10K	<1.46 ₃	<1.463	<1.463	1.9	65.0	>6400	8.0	1.0	16.0	64.0	64.0
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-2,2'-diamino-N-methyl-diethylamine)	2.9	7.2	9.3	20.1	125.5	>6400	4.0	1.0	8.0	64.0	64.0
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-1,4-benzylidiamine) <10K	2.0	2.9	4.703	5.1	31.3	87.0	32.0	8.0	16.0	128.0	128.0
Poly(piperazinebispropanoic acid-1,2-bis(2-aminoethoxy)ethane) <10K	>3200	>3200	>3200	>3200	>6400	>6400	128.0	128.0	128.0	128.0	128.0

Polymer Description	Cytotoxicity (Kidney Epithelial IC₅₀)	Cytotoxicity (Kidney Epithelial LC₅₀)	Cytotoxicity (Human Dermal Fibroblast IC₅₀)	Cytotoxicity (Human Dermal Fibroblast LC₅₀)	Lung Epi IC₅₀	Erythrocyte Lysis (Hemolysis IC₅₀)	Staphylococcus aureus subsp. aureus (MIC)	Staphylococcus epidermis (MIC)	Escherichia coli (MIC)	Pseudomonas aeruginosa (MIC)	Haemophilus influenzae (MIC)
Poly(piperazin ebispropanoic acid-2,2- diaminodiethy lamine) <10K	212.1	838.5	1111.6	1999.8	>6400	>6400	128.0	64.0	128.0	128.0	128.0
Poly(piperazin ebispropanoic acid-3,3'- diamino-N- methyl- dipropylamine) <10K	5.3	33.7	94.7	246.2	4164.4	>6400	128.0	32.0	128.0	128.0	128.0
Poly(piperazin ebispropanoic acid-N(3- aminopropyl) 1,3-propane diamine) <10K	<1.46 3	3.0	15.3	35.0	497.0	>6400	8.0	4.0	128.0	128.0	128.0
Poly(piperazin ebispropanoic acid-N- methyl-2,2- diaminodiethy lamine) <10K	1204.2	>3200	3068.6	>3200	>6400	>6400	128.0	128.0	128.0	128.0	128.0
Poly(4,4'- trimethylenedi piperidinebi spropanoic acid- 2,2'-diamino diethylamine) 5.5K	<1.46 3	1.5	2.0	3.4	14.6	88.0	4.0	0.3	4.0	16.0	64.0
Poly(4,4'- dipiperidinebi spropanoic acid-2,2'- diamino diethylamine) 5.1K	<1.5	2	2	4	50.0	1284.0	1.0	0.5	16.0	64.0	128.0
Poly(4,4'- dipiperidinebi spropanoic acid-2,2'- diamino N- methyl diethylamine) <5K	<1.5	2	4	7	93.0	>6400	2.0	0.5	32.0	64.0	128.0
Poly(4,4'- dipiperidinebi spropanoic acid-2,2'- diamino N- methyl diethylamine) <5K	5	9	20	30	294.0	>6400	8.0	0.5	32.0	32.0	128.0

Polymer Description	Cytotoxicity (Kidney Epithelial IC₅₀)	Cytotoxicity (Kidney Epithelial LC₅₀)	Cytotoxicity (Human Dermal Fibroblast IC₅₀)	Cytotoxicity (Human Dermal Fibroblast LC₅₀)	Lung Epi IC₅₀	Erythrocyte Lysis (Hemolysis IC₅₀)	Staphylococcus aureus subsp. aureus (MIC)	Staphylococcus epidermis (MIC)	Escherichia coli (MIC)	Pseudomonas aeruginosa (MIC)	Haemophilus influenzae (MIC)
Poly(4,4'-dipiperidinebispropanoic acid-2,2'-diamino N-methyl diethylamine) ~5K	11	21	61	88	823.0	>6400	8.0	1.0	64.0	64.0	128.0
Poly(4,4'-dipiperidinebispropanoic acid-3,3'-diamino-dipropylamine) <5K	<1.5	<1.5	<1.5	<1.5	10.0	300.0	1.0	0.3	16.0	32.0	128.0
Poly(4,4'-dipiperidinebispropanoic acid-3,3'-diamino-dipropylamine) ~5K	<1.5	<1.5	2	3	30.0	>6400	1.0	0.5	16.0	64.0	128.0
Poly(4,4'-dipiperidinebispropanoic acid-3,3'-diamino-N-methyl-dipropylamine) ~5K	<1.5	2	3	5	81.0	>6400	8.0	1.0	32.0	64.0	128.0
Poly(4,4'-dipiperidinebispropanoic acid-3,3'-diamino-N-methyl-dipropylamine) >5K	<1.5	<1.5	2	3	26.0	3622.0	2.0	0.5	16.0	32.0	128.0
Poly(4,4'-trimethylenedipiperidinebispropanoic acid-2,2'-diamino diethylamine) ~14,000	<1.46 3	<1.463	2.1	2.8	6.8	9.0	4.0	1.0	8.0	16.0	128.0
Poly(piperazinebispropanoic acid-1,3-diaminopropane) ~3,700	924	>3200	>3200	>3200	>6400	>6400	128.0	128.0	128.0	128.0	128.0
Poly(piperazinebispropanoic acid-1,4-diaminobutane) ~4,400	541	1946	1905	3080	4539.0	3388.0	128.0	64.0	128.0	128.0	128.0

Polymer Description	Cytotoxicity (Kidney Epithelial IC₅₀)	Cytotoxicity (Kidney Epithelial LC₅₀)	Cytotoxicity (Human Dermal Fibroblast IC₅₀)	Cytotoxicity (Human Dermal Fibroblast LC₅₀)	Lung Epi IC₅₀	Erythrocyte Lysis (Hemolysis IC₅₀)	Staphylococcus aureus subsp. aureus (MIC)	Staphylococcus epidermis (MIC)	Escherichia coli (MIC)	Pseudomonas aeruginosa (MIC)	Haemophilus influenzae (MIC)
Poly(4,4'-dipiperidinebispropionic acid-1,3-diaminopropane) <5K	6	11	7	16	22.0	>6400	1.0	0.1	16.0	1.0	32.0
Poly(4,4'-dipiperidinebispropionic acid-1,4-diaminobutane) <5K	4	7	5	17	22.0	>6400	2.0	0.3	8.0	8.0	32.0
Poly(4,4'-dipiperidinebispropionic acid-ethylenediamine) <5K	8	21	19	29	77.0	>6400	4.0	0.5	16.0	32.0	128.0
Poly(4,4'-trimethylenedipiperidine-1,3-diaminopropane-N,N'-di-3-propionic acid) <5K	2	3	2	3	4.0	6.3	16.0	4.0	8.0	16.0	128.0
Poly(4,4'-trimethylenedipiperidinebispropionic acid-2-dihydroxy 1,3-diaminopropane) <5K	15	80	50	127	63.0	>6400	4.0	1.0	8.0	128.0	32.0
Poly(4,4'-trimethylenedipiperidinebispropionic acid-bis(4-aminobutyl)ether) <5K	2	5	3	5	8.0	165.0	128.0	8.0	8.0	128.0	16.0
Poly (4,4'-trimethylene dipiperidine bispropanoic acid-4,4'-trimethylene dipiperidine)	<1.46 3	3	3	4	2	5.6	16.0	4.0	16.0	32.0	128.0
Poly (4,4'-trimethylene dipiperidine bispropanoic acid-N,N'-dimethyl-1,3-diaminopropane)	<1.46 3	2	5	11	6	11.3	128.0	4.0	4.0	16.0	128.0

Polymer Description	Cytotoxicity (Kidney Epithelial IC₅₀)	Cytotoxicity (Kidney Epithelial LC₅₀)	Cytotoxicity (Human Dermal Fibroblast IC₅₀)	Cytotoxicity (Human Dermal Fibroblast LC₅₀)	Lung Epi IC₅₀	Erythrocyte Lysis (Hemolysis IC₅₀)	Staphylococcus aureus subsp. aureus (MIC)	Staphylococcus epidermis (MIC)	Escherichia coli (MIC)	Pseudomonas aeruginosa (MIC)	Haemophilus influenzae (MIC)
Poly (4,4'-trimethylene dipiperidine bispropanoic acid-piperazine)	12	22	53	75	21	59.0	128.0	128.0	128.0	128.0	128.0
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane) modified with 40mol% of glycidol, 8K	2	3	5	8	6.0	636.0	0.5	0.3	2.0	4.0	8.0
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane), 1650	3.1	5.5	9.4	15.5	21.5	>3200	0.5	0.1	2.0	4.0	8.0
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane), 5K	1.6	2.9	6.8	9.9	10.8	651.0	1.0	0.3	2.0	8.0	8.0
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane), 7.7K	<1.5	<1.463	2.0	2.8	3.7	50.0	2.0	0.3	2.0	4.0	8.0
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane), 3K	8.7	12.3	20.7	32.8	17	1260.0	1.0	0.3	4.0	16.0	4.0
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane) modified with 100mol% of glycidol, 8K	3.0	5.0	7.0	11.0	7.0	500.0	8.0	0.5	8.0	16.0	16.0

Polymer Description	Cytotoxicity (Kidney Epithelial IC₅₀)	Cytotoxicity (Kidney Epithelial LC₅₀)	Cytotoxicity (Human Dermal Fibroblast IC₅₀)	Cytotoxicity (Human Dermal Fibroblast LC₅₀)	Lung Epi IC₅₀	Erythrocyte Lysis (Hemolysis IC₅₀)	Staphylococcus aureus subsp. aureus (MIC)	Staphylococcus epidermis (MIC)	Escherichia coli (MIC)	Pseudomonas aeruginosa (MIC)	Haemophilus influenzae (MIC)
Poly(4,4-trimethylene dipiperidinebi spropanoic acid-4,4'-dipiperidine), 10,631	<1.5	2.0	2.0	4	<1.5	3.2	16.0	4.0	8.0	16.0	128.0
Poly(4,4-trimethylene dipiperidinebi spropanoic acid-histamine), 2,3K	7.0	16.0	18.0	27.0	9.0	19.0	128.0	32.0	32.0	128.0	128.0
Poly(4,4-trimethylene dipiperidinebi spropanoic acid-diaminopropane), 3250	5.8	10.3	--	--	--	>3200	0.5	0.1	2.0	8.0	8.0
Poly(4,4-trimethylene dipiperidinebi spropanoic acid-diaminopropane), 4700	1.9	3.3	--	--	--	176.497	1.0	0.3	1.0	4.0	8.0
Poly(4,4-trimethylene dipiperidinebi spropanoic acid-diaminopropane) modified with 200mol% of glycidol, 7800	2	3	--	--	--	96.0	2.0	0.5	4.0	8.0	16.0
Poly(4,4-trimethylene dipiperidinebi spropanoic acid-diaminopropane) modified with 150mol% of glycidol, 7800	2	4	--	--	--	152.0	2.0	1.0	8.0	16.0	16.0
Poly(4,4-trimethylene dipiperidinebi spropanoic acid-diaminopropane) modified with 50mol% of glycidol, 7800	2	4	--	--	--	110.0	4.0	0.5	8.0	16.0	16.0

Polymer Description	Cytotoxicity (Kidney Epithelial IC₅₀)	Cytotoxicity (Kidney Epithelial LC₅₀)	Cytotoxicity (Human Dermal Fibroblast IC₅₀)	Cytotoxicity (Human Dermal Fibroblast LC₅₀)	Lung Epi IC₅₀	Erythrocyte Lysis (Hemolysis IC₅₀)	Staphylococcus aureus subsp. aureus (MIC)	Staphylococcus epidermis (MIC)	Escherichia coli (MIC)	Pseudomonas aeruginosa (MIC)	Haemophilus influenzae (MIC)
Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane), 2500	8.7	51.2	--	--	--	>3200	0.5	0.3	8.0	64.0	32.0
Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane) modified with 25mol% of glycidol, 7800	2	2	--	--	--	55.0	2.0	0.5	4.0	8.0	16.0
Poly (4,4'-trimethylene dipiperidine bispropanoic acid--3-(dimethylamino) 1-propylamine), 1K	32	70	--	--	--	182.0	128.0	32.0	128.0	128.0	128.0
Poly (4,4'-trimethylene dipiperidine bispropanoic acid--N,N'-dimethyl-1,3-diaminopropane), 1K	4	7	--	--	--	76.0	128.0	8.0	16.0	32.0	128.0
Poly(2,2-bipyrrolidine bispropanoic acid-diaminopropane), 2.5K	28	164	--	--	--	>3200	128.0	16.0	128.0	128.0	128.0
Poly(2,2'-bipyrrolidine bispropanoic acid-butyl diamine)	12	314	--	--	--	>3200	128.0	128.0	128.0	128.0	128.0
Poly(2,2'-bipyrrolidine bispropanoic acid-ethyl diamine)	73	292	--	--	--	>3200	128.0	128.0	128.0	128.0	128.0
Poly(2,2'-bipyrrolidine bispropanoic acid-penta diamine)	21	297	--	--	--	>3200	128.0	128.0	128.0	128.0	128.0

Polymer Description	Cytotoxicity (Kidney Epithelial IC₅₀)	Cytotoxicity (Kidney Epithelial LC₅₀)	Cytotoxicity (Human Dermal Fibroblast IC₅₀)	Cytotoxicity (Human Dermal Fibroblast LC₅₀)	Lung Epi IC₅₀	Erythrocyte Lysis (Hemolysis IC₅₀)	Staphylococcus aureus subsp. aureus (MIC)	Staphylococcus epidermis (MIC)	Escherichia coli (MIC)	Pseudomonas aeruginosa (MIC)	Haemophilus influenzae (MIC)
4,4'-trimethylene dipiperidinebi sopropanoic acid-diaminopropane pentamer	>512	>512	--	--	259.0	>3200	16.0	1.0	16.0	>128	64.0
4,4'-trimethylene dipiperidinebi sopropanoic acid-diaminopropane heptamer	190	>512	--	--	418.0	>3200	2.0	1.0	8.0	>128	128.0
Poly(4,4'-trimethylene dipiperidinebi sopropanoic acid-diaminopropane) (guanidine ended)	2	4	--	--	10	738.0	2.0	0.3	2.0	8.0	8.0
40mol% modified Poly(4,4'-trimethylene dipiperidinebi sopropanoic acid-diaminopropane)	3	5	--	--	7	757.0	0.5	0.1	1.0	8.0	4.0
40mol% modified Poly(4,4'-trimethylene dipiperidinebi sopropanoic acid-diaminopropane)	2	6	--	--	8	118.0	1.0	0.3	2.0	8.0	4.0
Poly(4,4'-trimethylene dipiperidinebi sopropanoic acid-N-glycidol diethylene triamine)	>512	>512	--	--	>512	>3200	>128	>128	>128	>128	>128
Poly(4,4'-trimethylene dipiperidinebi sopropanoic acid-N-glycidol diethylene triamine)	72	178	--	--	>512	>3200	>128	32.0	>128	>128	>128

Polymer Description	Cytotoxicity (Kidney Epithelial IC₅₀)	Cytotoxicity (Kidney Epithelial LC₅₀)	Cytotoxicity (Human Dermal Fibroblast IC₅₀)	Cytotoxicity (Human Dermal Fibroblast LC₅₀)	Lung Epi IC₅₀	Erythrocyte Lysis (Hemolysis IC₅₀)	Staphylococcus aureus subsp. aureus (MIC)	Staphylococcus epidermis (MIC)	Escherichia coli (MIC)	Pseudomonas aeruginosa (MIC)	Haemophilus influenzae (MIC)
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-N-glycidol dipropylene triamine)	70	161	--	--	>512	>3200	>128	32.0	>128	>128	>128
Poly(4,4'-trimethylenedi piperidine bisethylacrylamide-co-1,3-diamine propane)	20	32	--	--	--	267.0	>128	8.0	64.0	128.0	>128
Poly(4,4'-trimethylenedi piperidine bisethylacrylamide-co-1-amino-3-guanidine propane)	22	34	--	--	--	122.0	16.0	8.0	16.0	64.0	>128
4,4'-trimethylene dipiperidinebispropanoic acid-1-amino-3-guanidine propane	>512	>512	--	--	--	>3200	>128	>128	>128	>128	>128
Poly(4,4'-trimethylene dipiperidinebispropanoic acid-diaminopropane), 1400	82	499	--	--	--	>3200	4.0	4.0	16.0	>128	128.0
Poly(4,4'-trimethylenedi piperidine bisethylacrylamide-1,3-diamine propane)-co-Poly(4,4'-trimethylenedi piperidine bisethylacrylamide-1-aminobutyl-3-carbamoyl-pyridinium)	5	10	--	--	--	44.0	32.0	2.0	8.0	16.0	>128

Polymer Description	Cytotoxicity (Kidney Epithelial IC₅₀)	Cytotoxicity (Kidney Epithelial LC₅₀)	Cytotoxicity (Human Dermal Fibroblast IC₅₀)	Cytotoxicity (Human Dermal Fibroblast LC₅₀)	Lung Epi IC₅₀	Erythrocyte Lysis (Hemolysis IC₅₀)	Staphylococcus aureus subsp. aureus (MIC)	Staphylococcus epidermis (MIC)	Escherichia coli (MIC)	Pseudomonas aeruginosa (MIC)	Haemophilus influenzae (MIC)
Poly(4,4'-trimethylenedi piperidine bisethylacrylamine-1-aminobutyl-3-carbamoyl-pyridinium)	11	16	--	--	--	83.0	>128	32.0	64.0	>128	>128
40 mol% glycidol modified poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane), 4700	1	2	--	--	--	287.0	1.0	0.5	2.0	4.0	8.0
Guanidine terminated Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane), 4700	1	4	--	--	--	334.0	1.0	0.5	2.0	4.0	8.0
Guanidine terminated poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane), 7700	1	1	--	--	--	18.0	1.0	0.5	2.0	4.0	8.0
Carboxy terminated Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane)	43	>512	--	--	--	>3200	64.0	32.0	128.0	>128	>128
Methyl ester terminated Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane)	1.5	4	--	--	--	152.0	4.0	0.5	4.0	16.0	8.0

Polymer Description	Cytotoxicity (Kidney Epithelial IC₅₀)	Cytotoxicity (Kidney Epithelial LC₅₀)	Cytotoxicity (Human Dermal Fibroblast IC₅₀)	Cytotoxicity (Human Dermal Fibroblast LC₅₀)	Lung Epi IC₅₀	Erythrocyte Lysis (Hemolysis IC₅₀)	Staphylococcus aureus subsp. aureus (MIC)	Staphylococcus epidermis (MIC)	Escherichia coli (MIC)	Pseudomonas aeruginosa (MIC)	Haemophilus influenzae (MIC)
Guanidine terminated Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane), 2200	93	169	--	--	--	>3200	1.0	0.3	2.0	64.0	16.0
Poly(4,4-trimethylene dipiperidinebispropanoic acid-aminomethyl benzene)	4	10	--	--	--	14.0	64.0	8.0	16.0	16.0	>128
4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane	>512	>512	--	--	--	>3200	>128	>128	>128	>128	>128
Poly[4,4-trimethylene dipiperidinebispropanoic acid-(1-aminomethyl-4-guanidinomethyl benzene)]	3	5	--	--	--	52.0	4.0	64.0	8.0	64.0	>128
4-guanidinobenzene terminated Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane)	6	12	--	--	--	1142	4.0	2.0	4.0	64.0	32.0
Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane), <10K	29.6	110.1	--	--	--	>3200	1.0	0.5	4.0	64.0	16.0

-- indicates not tested.

Example 1- 5: Inhibition of *Pseudomonas aeruginosa* in Cystic Fibrosis Bronchial Epithelial Cells

Cystic fibrosis bronchial epithelial (CFBE) cells were grown in 12-well plates for 7–9 days. The cells were washed twice with imaging medium before *Pseudomonas aeruginosa* (mucoid strain, SMC 1585) was inoculated into each well at a multiplicity of infection (MOI) of ~30 (~ 6×10^6 cfu/well). The plates were incubated at 37°C, 5% CO₂ for 1 hour to allow bacterial attachment to the airway cells. The supernatant was then replaced with imaging medium containing 0.4% arginine and then incubated for 5 hours to form biofilms on CFBE cells. To estimate the efficacy of antimicrobial polymer treatment in preformed biofilms, the plates were washed twice with imaging medium and antimicrobial agent (antimicrobial polymer or tobramycin [positive control]) were applied at designated concentrations to disrupt established biofilms for 16 hours. The supernatant was then removed and washed twice with imaging medium. CFBE cells were lysed with 0.1% Triton X-100 for approximately 15 minutes. The lysate was vortexed for 3 minutes before serial dilution and spot titration onto LB plates to determine the cfu/well. The bacterial strain was defined as ‘susceptible’ to the antibiotic treatment in the static co-culture model if the CFBE monolayers were not disrupted after overnight antibiotic treatment and there was more than a 2 log₁₀ difference in cfu recovery between no treatment and antibiotic antimicrobial agent treatment.

To test the ability of antibiotics to prevent biofilm formation, these compounds were applied after the 1 hour period for bacterial attachment. The plates were incubated for 5 hours, and cfu/well was determined as described above. The detection limit of the static co-culture assay was 200 cfu/well. All experiments were performed at least three times. The susceptibility of *Pseudomonas aeruginosa* biofilm to Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane) <10K is displayed in Figure I below.

Example 2: *In vivo* Studies**Example 2- 1: Toxicity – Maximum Tolerated Dose**

Acute, 24 hour, toxicity studies to determine the maximum tolerated dose of a compound were carried out in male rats and mice of approximately 8-10 weeks of age. Animals were housed singly in standard polycarbonate cages and fed normal chow diets. Following one week of acclimation, compounds were administered in a single intraperitoneal (I.P.) or intravenous (I.V.) dose, typically in a PBS vehicle. The doses generally ranged from 1 mg/kg to as high as 400 mg/kg. Animals were observed for signs of pain, distress, and local or systemic signs of toxicity for one hour post-dosing, and then in 1 hour intervals for 6 hours after dosing. The following day at 24 hours post-dose, the animals were sacrificed and blood removed for serum chemistry analysis. Serum chemistry analyses performed include: ALT, AST, Creatinine and Urea Nitrogen. Major organs were also examined for abnormal signs.

Table 3 displays the Maximum Tolerated Dose (MTD) for select test compounds at select routes of administration.

Table 3: Maximum Tolerated Dose (MTD)

Treatment	Animal Model	Route of Administration	MTD
Poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane), MW = 4,700	Rat	I.P.	5 mg/kg
Poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane), MW = 2,500	Rat	I.P.	5 mg/kg
Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane), MW < 10K	Mice	I.P.	5 mg/kg
Poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane), MW = 2,500	Mice	I.V.	40 mg/kg

Example 2- 2: Efficacy – Surgical Site Infection

The test compound, poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane) modified with 40 mol% of glycidol, was evaluated for anti-infective activity against *Staphylococcus aureus*, Methicillin Resistant (MRSA) and *Escherichia coli* (*E. coli*)

in mice. Male ICR mice weighing approximately 22 g were used to evaluate the anti-infective activity against each bacterium.

Example 2- 2(a): MRSA

Five groups of 10 male mice were inoculated intraperitoneally with a LD₉₀₋₁₀₀ of MRSA (1.90 x 10⁸ CFU/mouse) suspended in 0.5 mL of brain heart infusion (BHI) broth containing 5% mucin. One hour after bacteria inoculation, groups of 10 animals were intraperitoneally administered one of the following:

- 0.2 mg/kg poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane) modified with 40 mol% of glycidol suspended in 0.9% NaCl,
- 5 mg/kg poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane) modified with 40 mol% of glycidol suspended in 0.9% NaCl,
- 1 mg/kg ofloxacin,
- 3 mg/kg ofloxacin, and
- 5 mL/kg vehicle (0.9% NaCl).

Mortality was recorded once daily for 7 days and an increase of survival relative to vehicle control group was evaluated.

Table 4 displays the results against MRSA for the test compounds.

Table 4: MRSA Activity

Treatment	Dose	Animals Dosed (N)	Survival (n/N)	Survival Increase (%)
Vehicle	5 mL/kg	10	0/10	---
Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane) modified with 40 mol% of glycidol	0.2 mg/kg	10	1/10	10%
	5 mg/kg	10	6/10	60%*

* Survival increase $\geq 50\%$ indicates significant anti-microbial effect

Example 2- 2(b): *E. coli*

Five groups of 10 male mice were inoculated intraperitoneally with a LD₉₀₋₁₀₀ of *E. coli* (2.20 x 10⁵ CFU/mouse) suspended in 0.5 mL of BHI broth containing 5% mucin. One hour after bacteria inoculation, groups of 10 animals were intraperitoneally administered one of the following:

- 0.2 mg/kg poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane) modified with 40 mol% of glycidol suspended in 0.9% NaCl,

- 5 mg/kg poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane) modified with 40 mol% of glycidol suspended in 0.9% NaCl,
- 0.3 mg/kg gentamicin,
- 1 mg/kg gentamicin, and
- 5 mg/kg vehicle (0.9% NaCl).

Mortality was recorded once daily for 7 days and an increase of survival relative to vehicle control group was evaluated.

Table 5 displays the results against *E. coli* for the test compounds.

Table 5: *E. coli* Activity

Treatment	Dose	Animals Dosed (N)	Survival (n/N)	Survival Increase (%)
Vehicle	5 mL/kg	10	0/10	---
Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane) modified with 40 mol% of glycidol	0.2 mg/kg	10	0/10	0%
	5 mg/kg	10	8/10	80%*

* Survival increase $\geq 50\%$ indicates significant anti-microbial effect

The test compound, poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane) modified with 40 mol% of glycidol, afforded significant anti-microbial protection, exhibiting 60% and 80% increase in survival rate in MRSA and *E. coli* infected mouse models.

Example 2- 3: Efficacy – Mucositis

The goal of this study was to examine the role of schedule and route of administration on the observed efficacy of poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane) (1 mg/mL) on the frequency, severity and duration of oral mucositis induced by acute radiation. Male LVG Syrian Golden Hamsters, aged 5 to 6 weeks with an average body weight of 86.3 g at study commencement were used to evaluate the activity of each compound against radiation induced oral mucositis. Study endpoints were mucositis score, weight change and survival.

Male Syrian Golden Hamsters were randomly and prospectively divided into treatment groups of seven (7) animals per group (test article) and one group of ten (10) animals (control).

On day 0, all animals were given an acute radiation dose of 40 Gy directed to their left buccal cheek pouch. On day 0, animals were dosed topically once. From day 0 to day 20, 0.5 mL doses were applied topically to the left buccal pouch three times per day.

To evaluate mucositis severity, animals were anesthetized with an inhalation anesthetic, and the left cheek pouch everted. Mucositis was scored visually by comparison to a validated photographic scale; the scale ranges from 0 for normal, to 5 for severe ulceration. A descriptive version of the mucositis scoring scale used in this study is presented in **Table 6**.

Table 6: Mucositis Scoring Scale

Score	Description:
0	Pouch completely healthy. No erythema or vasodilation.
1	Light to severe erythema and vasodilation. No erosion of mucosa.
2	Severe erythema and vasodilation. Erosion of superficial aspects of mucosa leaving denuded areas. Decreased stippling of mucosa.
3	Formation of off-white ulcers in one or more places. Ulcers may have a yellow/gray appearance due to pseudomembrane formation. Cumulative size of ulcers should equal about $\frac{1}{4}$ of the pouch. Severe erythema and vasodilation.
4	Cumulative size of ulcers should equal about $\frac{1}{2}$ of the pouch. Loss of pliability. Severe erythema and vasodilation.
5	Virtually all of pouch is ulcerated. Loss of pliability (pouch can only partially be extracted from mouth).

A score of 1 or 2 represent a mild stage of injury, a score of 3, 4 or 5 indicates moderate to severe mucositis. Following visual scoring, a digital photograph was taken of each animal's mucosa using a standardized technique. At the conclusion of the experiment, all images were randomly numbered and graded in blinded fashion by at least two independent trained observers using the above-described scale (blinded scoring).

Animal deaths were evaluated during the course of the study. In the model, deaths are most commonly attributable to adverse effects associated with anesthesia typically occurring at the time of radiation, or toxicity of the experimental compound. There were no deaths associated with the experimental compounds.

Weight change was also evaluated as it represents a secondary method of examining potential toxicities of experimental treatments. Animals were weighed daily throughout the study; weight changes were similar in all groups. The mean percent weight gain during the study is provided in **Table 7**.

To evaluate the significance of these differences, the mean area under the curve (AUC) was calculated for each animal from the percent weight gain data, and the means and

standard errors were plotted. Using a one way ANOVA, no statistically significant difference in weight change was observed in any of the groups.

To evaluate efficacy, the mean group mucositis scores were compared to the control group in each experiment. A clinical mucositis score of 3 in hamsters indicates the presence of an ulcer. Ulceration is the point in the development of mucositis where the physical integrity of the oral mucosa is breached. In the clinic, a patient presenting with severe oral ulcerations may require hospitalization for analgesic, narcotic and/or antibiotic therapies or fluid support. The average cost to the healthcare system is significant. Advanced mucositis in humans (with ulcerative sores, correlating to a score of 3 or greater) often requires the interruption of therapy for patients receiving radiation and, if sepsis occurs, these patients risk death. A therapeutic that significantly reduces the time that a patient with oral mucositis had ulcers would be of great value to the clinician. The cumulative number of days that an animal had a score of 3 or greater was determined due to its clinical significance.

The significance of group differences in scores of 3 or greater was determined using Chi-squared (χ^2) difference analysis for the total number of animal days with a score of 3 or higher over the course of the entire study; these results are presented in **Table 7**. The severity and course of mucositis was favorably attenuated in the poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane) group.

Table 7: Mucositis Safety and Efficacy

Group	Dose (Concentration)	% Weight Gain (Days 0 to 20)	% Animal Days with Mucositis Score ≥ 3
Control <i>topical, tid</i>	0.5 mL	41.6	41.3
Poly(4,4-trimethylene dipiperidinebispropanoic acid-diaminopropane), <i>topical, tid</i>	0.5 mL (1 mg/mL)	34.2	28.6

Example 3: Synthesis of Amine Functional Polyamides

Example 3- 1: Synthesis of 4,4'-trimethylene dipiperidine bispropanoic acetate

To 5.0 g of 4,4'- trimethylene dipiperidine in 20 mL of methanol solution (20 mL), 4.6 g of methyl acrylate was added drop-wise. The resulting reaction mixture was stirred at room temperature for 16 hours. The solvent was removed under reduced pressure and the residue was purified by column chromatography using a gradient solvent system comprising

of from 100% hexane to 100% ethyl acetate. Removal of the solvent under reduced pressure yielded 7 g of the desired product as a white solid.

Example 3- 2: Synthesis of 4,4'-dipiperidine bispropanoic acetate

To 10.0 g of 4,4'- dipiperidine HCl dissolved in 80 mL of methanol was added 12.6 g of potassium carbonate. The reaction mixture was stirred at room temperature for 3 hours, at which time 8.03 g of methyl acrylate was added slowly. The resulting reaction solution was then stirred at room temperature for 18 hours. The reaction mixture was filtered and the filtrate was evaporated to dryness under reduced pressure. The residue was treated with 300 mL of ethyl acetate. The resulting suspension was stirred at room temperature for 2 hours followed by filtration. The filtrate was evaporated to dryness under reduced pressure. The resulting mass was dried at room temperature under the vacuum to give 11.34 g of the desired product as an off white solid.

Example 3- 3: Synthesis of piperazine bispropanoic acetate

To 10 g of piperazine hexahydrate dissolved in 40 mL of methanol was added 9.97 g of methyl acrylate in a drop-wise manner. The reaction mixture was stirred at room temperature for 18 hours. At the end of this time, the reaction mixture was evaporated to dryness under reduced pressure. The residue was recrystallized from hexane/methylene chloride (1:1 v/v). After filtration and drying at room temperature under reduced pressure, 12.2 g of the desired product was obtained as a white solid.

Example 3- 4: Synthesis of 1,1'-diacryl-4,4'-trimethylene dipiperidine

To 3.8 g of acryloyl chloride dissolved in 50 mL of dichloromethane was added a solution of 4.0 g of 4, 4-trimethylene dipiperidine dissolved in 20 mL dichloromethane in a drop-wise manner at 0°C. To this solution was added 4.23 g of triethyl amine slowly with a syringe. The resulting reaction mixture was stirred for 18 hours and was allowed to warm to room temperature. The reaction mixture was filtered and the filtrate was collected. After removing the solvent under reduced pressure, the residue was treated with 100 mL of ethyl acetate. The solution was extracted with 1M HCl (1 x 100 mL), saturated NaHCO₃ (2 x 100 mL), and finally with brine (2 x 100 mL). The organic layer was collected and dried over Na₂SO₄. After filtration, the filtrate was evaporated to dryness under reduced pressure. The residue was purified by column chromatography using a gradient solvent system from 100% hexane to 100% ethyl acetate. Upon removal of the solvent, 3 g of the desired product was obtained as viscous oil.

Example 3- 5: Synthesis of 2,2'-bipyrrolidine bispropanoic acetate

To a solution of 5 g of 2,2'-bipyrrolidine in 20 mL of methanol was added 6.9 g of methyl acrylate (6.9 g, 80 mmol) in a drop-wise manner. The resulting reaction mixture was stirred at room temperature for 16 hours. The reaction mixture was evaporated to dryness yielding 10 g of the desired product as viscous oil.

5 **Example 3- 6: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane)**

The reaction mixture consisting of 1 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate (**Example 3- 1**) and 0.387 g of 1, 3-diamino propane was heated at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of
10 dichloromethane and was poured into 50 mL of ethyl acetate. After filtering off the solvent, the residue was dissolved in 20 mL of deionized (DI) water. The pH of the solution was brought to 2 by addition of HCl. The resulting solution was dialyzed against DI water for 24 hours using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis bag was dried by lyophilization yielding 90 mg of the desired
15 product as a light yellow solid.

Example 3- 7: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-diamino ethane)

The reaction mixture containing 0.5 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate (**Example 3- 1**) and 0.157 g of diamino ethane was stirred at 100°C under nitrogen
20 atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 50 mg of the
25 desired product as a light yellow solid.

Example 3- 8: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,4-diamino butane)

The reaction mixture containing 0.5 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate and 0.23 g of 1, 4-diamino butane was stirred at 100°C under nitrogen atmosphere for
30 18 hours. The resulting product was dissolved in 5 mL of CH₂Cl₂ and then precipitated in 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the

dialysis membrane was lyophilized to dryness yielding 60 mg of the desired product as a light yellow solid.

Example 3- 9: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,2-bis (2-aminoethoxy) ethane

5 The reaction mixture containing 0.5 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate and 0.26 g of 1, 2-bis(2-aminoethoxy) ethane was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed
10 against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 60 mg of the desired product as a light yellow solid.

Example 3- 10: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,4-bis(aminomethyl) benzene

15 The reaction mixture containing 0.5 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate and 0.7 g of 1,4-bis(aminomethyl) benzene (0.7 g, 5.1mmol) was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2,
20 it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 40 mg of the desired product as a light yellow solid.

Example 3- 11: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-2,2'diamino diethylamine

25 The reaction mixture containing 0.5 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate and 0.35 g of 2,2'diamino diethylamine was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed
30 against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 63 mg of the desired product as a light yellow solid.

Example 3- 12: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-N-methyl-2,2'diamino diethylamine

The reaction mixture containing 1 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate and 0.61 g of N-methyl-2,2'diamino diethylamine (0.61 g, 5.2mmol) was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 130 mg of the desired product as a light yellow solid.

Example 3- 13: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-N-(3-aminopropyl)-1,3-propane diamine

The reaction mixture containing 1 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate and 0.68 g of N-(3-aminopropyl)-1,3-propane diamine (0.68 g, 5.2 mmol) was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 180 mg of the desired product as a light yellow solid.

Example 3- 14: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-3,3'-diamino-N-methyl dipropylamine

The reaction mixture containing 1 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate and 0.76 g of 3,3'-diamino-N-methyl dipropylamine was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 110 mg of the desired product as a light yellow solid.

Example 3- 15: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino-2-propanol

The reaction mixture containing 1 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate and 0.47 g of 1,3-diamino-2-propanol (0.47 g, 5.2 mmol) was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 60 mg of the desired product as a light yellow solid.

Example 3- 16: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-4-(4-amino-butoxyl)-butyl amine

4-(4-amino-butoxyl)-butyl amine HCl salt (1 g) was dissolved in 20 mL of methanol. To this solution 0.72 g of aqueous sodium hydroxide solution (50% w/w)) was added. The reaction mixture was stirred at room temperature for 1 hour. After filtering off the solids, the filtrate was evaporated to dryness. The residue was treated with 20 mL of ethanol. The reaction mixture was filtered and the filtrate was evaporated to dryness yielding 0.55 g of an off white solid. This solid was combined with 0.75 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate and the resulting reaction mixture was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 90 mg of the desired product as a light yellow solid.

Example 3- 17: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-3,5-diamino-1,2,4-triazol

The reaction mixture containing 1 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate and 0.31 g of 3,5-diamino-1,2,4-triazole was treated with 1 mL of DMSO. The resulting reaction mixture was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis

membrane was lyophilized to dryness yielding 10 mg of the desired product as a light yellow solid.

Example 3- 18: Synthesis of poly (piperazine bispropanoic acid-co-diamino ethane)

The reaction mixture containing 1 g of piperazine bispropanoic acetate (**Example 3- 3**) and 0.47 g of diamino ethane was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 10 mg of the desired product as a light yellow solid.

Example 3- 19: Synthesis of poly(piperazine bispropanoic acid-co-1,3-diamino propane)

The reaction mixture containing 1 g of piperazine bispropanoic acetate (**Example 3- 3**) and 0.5 g of 1,3-diamino propane was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 30 mg of the desired product as a light yellow solid.

Example 3- 20: Synthesis of poly (piperazine bispropanoic acid-co-1,4-diamino butane)

The reaction mixture containing 1 g of piperazine bispropanoic acetate (**Example 3- 3**) and 0.6 g of 1,4-diamino butane was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 60 mg of the desired product as a light yellow solid.

Example 3- 21: Synthesis of poly (piperazine bispropanoic acid-co-1,2-bis (2-aminoethoxy) ethane)

The reaction mixture containing 1 g of piperazine bispropanoic acetate (**Example 3- 3**) and 1.15 g of 1,2-bis (2-aminoethoxy) ethane was stirred at 100°C under nitrogen

atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 30 mg of the desired product as a light yellow solid.

Example 3- 22: Synthesis of poly (piperazine bispropanoic acid-co-2,2'diamino diethylamine

The reaction mixture containing 1 g of piperazine bispropanoic acetate (**Example 3- 3**) and 0.8 g of 2,2'-diamino diethylamine was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 60 mg of the desired product as a light yellow solid.

Example 3- 23: Synthesis of poly (piperazine bispropanoic acid- co-N-methyl-2,2'diamino diethylamine

The reaction mixture containing 1 g of piperazine bispropanoic acetate (**Example 3- 3**) and 0.9 g of N-methyl-2,2'-diamino diethylamine was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 50 mg of the desired product as a light yellow solid.

Example 3- 24: Synthesis of poly (piperazine bispropanoic acid-co-N-(3-aminopropyl)-1,3-propane diamine

The reaction mixture containing 1 g of piperazine bispropanoic acetate (**Example 3- 3**) and 1.02 g of N-(3-aminopropyl)-1,3-propane diamine was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed

against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 90 mg of the desired product as a light yellow solid.

Example 3- 25: Synthesis of poly (piperazine bispropanoic acid-co-3,3'-diamino-N-methyl dipropylamine

The reaction mixture containing 1 g of piperazine bispropanoic acetate (**Example 3- 3**) and 1.12 g of 3,3'-diamino-N-methyl dipropylamine was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 120 mg of the desired product as a light yellow solid.

Example 3- 26: Synthesis of poly (4,4'-dipiperidine bispropanoic acid-co-diamino ethane

The reaction mixture containing 1 g of 4,4'-dipiperidine bispropanoic acetate (**Example 3- 2**) and 0.31 g of diamino ethane was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 90 mg of the desired product as a light yellow solid.

Example 3- 27: Synthesis of poly (4,4'-dipiperidine bispropanoic acid-co-1,3-diamino propane

The reaction mixture containing 1 g of 4,4'-dipiperidine bispropanoic acetate (**Example 3- 2**) and 0.38 g of 1,3-diamino propane was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 60 mg of the desired product as a light yellow solid.

Example 3- 28: Synthesis of poly (4,4'-dipiperidine bispropanoic acid-co-1,4-diamino butane

The reaction mixture containing 1 g of 4,4'-dipiperidine bispropanoic acetate (**Example 3- 2**) and 0.45 g of 1,4-diamino butane was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 90 mg of the desired product as a light yellow solid.

Example 3- 29: Synthesis of poly (4,4'-dipiperidine bispropanoic acid-co-1,2-bis (2-aminoethoxy) ethane

The reaction mixture containing 1 g of 4,4'-dipiperidine bispropanoic acetate (**Example 3- 2**) and 0.76 g of 1,2-bis (2-aminoethoxy) ethane was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 100 mg of the desired product as a light yellow solid.

Example 3- 30: Synthesis of poly (4,4'-dipiperidine bispropanoic acid-co-2,2'diamino diethylamine

The reaction mixture containing 1 g of 4,4'-dipiperidine bispropanoic acetate and 0.45 g of 2,2'diamino diethylamine was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 310 mg of the desired product as a light yellow solid.

Example 3- 31: Synthesis of poly (4,4'-dipiperidine bispropanoic acid-co-N-methyl-2,2'diamino diethylamine

The reaction mixture containing 1 g of 4,4'-dipiperidine bispropanoic acetate and 0.52 g of N-methyl-2,2'-diamino diethylamine was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 480 mg of the desired product as a light yellow solid.

Example 3- 32: Synthesis of poly (4,4'-dipiperidine bispropanoic acid-co-N-(3-aminopropyl)-1,3-propane diamine

The reaction mixture containing 1 g of 4,4'-dipiperidine bispropanoic acetate and 0.58 g of N-(3-aminopropyl)-1,3-propane diamine was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 540 mg of the desired product as a light yellow solid.

Example 3- 33: Synthesis of poly (4,4'-dipiperidine bispropanoic acid-co-3,3'-diamino-N-methyl dipropylamine

The reaction mixture containing 1 g of 4,4'-dipiperidine bispropanoic acetate and 0.64 g of 3,3'-diamino-N-methyl dipropylamine was stirred at 100°C under nitrogen atmosphere for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 420 mg of the desired product as a light yellow solid.

Example 3- 34: Synthesis of Poly (1,1'-diacryl-4,4'-trimethylene dipiperidine –co-1,3-diaminopropane

The reaction mixture containing 1 g of 1,1'-diacryl-4,4'-trimethylene dipiperidine, 0.35 g 1,3-diamino propane and 1 mL of methanol was stirred at room temperature for 18 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50

mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 640 mg of the desired product as a light yellow solid.

Example 3- 35: Synthesis of poly (1,1'-diacryl-4,4'-trimethylene dipiperidine –co-N,N'-dimethyl-1,3-propanediamine

The reaction mixture containing 1 g of 1,1'-diacryl-4,4'-trimethylene dipiperidine, 0.36 g of N,N'-dimethyl-1,3-propanediamine and 1 mL of methanol was stirred at 60°C for 24 hours. The solvent was removed under reduced pressure and the residue was dissolved in 20 mL of DI water. The pH of the solution was adjusted to 2 by adding HCl. The polymer solution dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 180 mg of the desired product as a light yellow solid.

Example 3- 36: Synthesis of poly (1,1'-diacryl-4,4'-trimethylene dipiperidine –co-4,4'-trimethylene dipiperidine

The reaction mixture containing 1 g of 1,1'-diacryl-4,4'-trimethylene dipiperidine, 0.99 g of 4,4'-trimethylene dipiperidine, 1 mL of methanol was stirred at 60°C for 12 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 220 mg of the desired product as a light yellow solid.

Example 3- 37: Synthesis of poly (1,1'-diacryl-4,4'-trimethylene dipiperidine –co-piperazine

The reaction mixture containing 1 g of 1,1'-diacryl-4,4'-trimethylene dipiperidine, 0.91 g of piperazine hexahydrate and 1 mL of methanol was stirred at 60°C for 12 hours. The resulting product was dissolved in 5 mL of dichloromethane and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis

membrane was lyophilized to dryness yielding 80 mg of the desired product as a light yellow solid.

Example 3- 38: Synthesis of poly (1,1'-diacryl-4,4'-trimethylene dipiperidine –co-4,4'-bipiperidine

5 A solution containing 1.14 g of 4,4'- dipiperidine HCl and 5 mL of methanol was treated with 1.14 g of potassium carbonate. The reaction mixture was stirred at room temperature for 2 hours. The reaction mixture was filtered and the filtrate was combined with 1g of 1,1'-diacryl-4,4'-trimethylene dipiperidine dissolved in 3 mL of methanol. The resulting reaction mixture was stirred at 60°C for 15 hours. The resulting product was poured
10 into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 140 mg of the desired product as a light yellow solid.

Example 3- 39: Synthesis of poly (1,1'-diacryl-4,4'-trimethylene dipiperidine –co-histamine)

The reaction mixture containing 1 g of 1,1'-diacryl-4,4'-trimethylene dipiperidine, 0.5 g of histamine and 1 mL of methanol was stirred at 60°C for 18 hours. The resulting product was poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was
20 dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 120 mg of the desired product as a light yellow solid.

Example 3- 40: Synthesis of poly (1,1'-diacryl-4,4'-trimethylene dipiperidine –co-3-(dimethylamino)-1-propylamine)

25 The reaction mixture containing 1 g of 1,1'-diacryl-4,4'-trimethylene dipiperidine, 0.53 g of 3-(dimethylamino)-1-propylamine and 1 mL of methanol was stirred at 50°C for 10 hours. The resulting product was poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the
30 solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 1 g of the desired product as a light yellow solid.

Example 3- 41: Synthesis of poly (1,1'-diacryl-4,4'-trimethylene dipiperidine –co-propyl amine

The reaction mixture containing 0.64 g of 1,1'-diacryl-4,4'-trimethylene dipiperidine, 0.35 g of propyl amine, and 1 mL methanol was stirred at 60°C for 20 hours. The resulting product was poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 740 mg of the desired product as a light yellow solid.

Example 3- 42: Synthesis of poly (1,1'-diacryl 4,4'-trimethylene dipiperidine –co-1-aminobutyl-3-carbamoyl pyridinium

The reaction mixture containing 0.5 g of 1,1'-diacryl-4,4'-trimethylene dipiperidine, 0.35 g of 1-aminobutyl-3-carbamoyl pyridinium, and 3 mL of methanol was stirred at 50°C for 20 hours. The resulting product was poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 20 mg of the desired product as a light yellow solid.

Example 3- 43: Synthesis of poly (1,1'-diacryl-4,4'-trimethylene dipiperidine –co-1-aminobutyl-3-carbamoyl pyridinium)-co-4,4'-trimethylene dipiperidine bispropanoic acid-2-dihydroxy-1,3-diamino propane)

The reaction mixture containing 1.0 g of 1,1'-diacryl-4,4'-trimethylene dipiperidine, 0.36 g of 1-aminobutyl-3-carbamoyl pyridinium, 0.27g of mono N-boc-1,3-diaminopropane, and 3 mL of methanol stirred at 50°C for 20 hours. The reaction mixture was poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration. The residue was washed with ethyl acetate (3 x 50 mL) and dried under reduced pressure.

Above product was dissolved in 5 mL of methanol and mixed with 0.5 g of 4,4'-trimethylene dipiperidine bispropanoic acid and 0.25 mL of concentrated HCl. The resulting reaction mixture was stirred at 50°C for 6 hours. The resulting product was poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the

dialysis membrane was lyophilized to dryness yielding 210 mg of the desired product as a light yellow solid.

Example 3- 44: Synthesis of poly (2,2'-bipyrrolidine bispropanoic acid-co-diamino ethane)

5 The reaction mixture containing 1.0 g of 2,2'-bipyrrolidine bispropanoic acetate and 0.38 g diamino ethane was stirred at 100°C under nitrogen atmosphere for 20 hours. The resulting product was dissolved in 3 mL of methanol and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis
10 membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 10 mg of the desired product as a light yellow solid.

Example 3- 45: Synthesis of poly (2,2'-bipyrrolidine bispropanoic acid-co-1,3-diamino propane)

15 The reaction mixture containing 1.0 g of 2,2'-bipyrrolidine bispropanoic acetate and 0.47 g of 1, 3-diamino propane was stirred at 100°C under nitrogen atmosphere for 20 hours. The resulting product was dissolved in 3 mL of methanol and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis
20 membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 540 mg of the desired product as a light yellow solid.

Example 3- 46: Synthesis of poly (2,2'-bipyrrolidine bispropanoic acid-co-1,3-diamino butane)

25 The reaction mixture containing 1.0 g of 2,2'-bipyrrolidine bispropanoic acetate and 0.56 g of 1, 4-diamino butane was stirred at 100°C under nitrogen atmosphere for 20 hours. The resulting product was dissolved in 3 mL of methanol and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis
30 membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 380 mg of the desired product as a light yellow solid.

Example 3- 47: Synthesis of poly (2,2'-bipyrrolidine bispropanoic acid-co-1,5-diamino pentane)

The reaction mixture containing 1.0 g of 2,2'-bipyrrolidine bispropanoic acetate and 0.65 g of 1, 5-diamino pentane was stirred at 100°C under nitrogen atmosphere for 20 hours.

5 The resulting product was dissolved in 3 mL of methanol and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 10 mg of the desired product as a light yellow
10 solid.

Example 3- 48: Synthesis of poly (2,2'-bipyrrolidine bispropanoic acid-co-1,6-diamino hexane)

The reaction mixture containing 1.0 g of 2,2'-bipyrrolidine bispropanoic acetate and 0.74 g of 1, 6-diamino hexane was stirred at 100°C under nitrogen atmosphere for 20 hours.

15 The resulting product was dissolved in 3 mL of methanol and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 10 mg of the desired product as a light yellow
20 solid.

Example 3- 49: Synthesis of poly (4,4-Trimethylene dipiperidine bispropanoic acid-co-4-(1,2-diol)-1,4,7-triazaheptane)**Example 3- 49(a): Synthesis of 4-(1,2-diol)-1,4,7-triazaheptane**

In 5 mL of ethanol 1 g of 1,7-bis-Boc-1,4,7-triazaheptane and 0.3 g of glycidol were
25 added and the reaction mixture was refluxed for 15 hours. The resulting product was purified by column chromatography using gradient solvent system in the range of 100% hexane to 100% yielding 0.4g of 1, 7-bis-boc-4-(1,2-diol)-1,4,7-triazaheptane. To 0.4g of 1, 7-bis-boc-4-(1,2-diol)-1,4,7-triazaheptane dissolved in 2mL of methanol was added 0.3 mL of concentrated HCl. The reaction mixture was stirred at 50°C for 24 hours. After removing the
30 solvent under reduced pressure, the residue was dissolved in 10 mL of methanol:water (1:1 v/v). To this solution was added 5.0 g of Amberlyst OH 26 resin. After stirring at room temperature for 3 hours, the resin was filtered off. The solvent was evaporated under reduced

pressure. The resulting oil was lyophilized to dry to give 0.15 g of the desired product as a viscous liquid.

Example 3- 49(b): Synthesis of poly(4,4-Trimethylene dipiperidine bispropanoic acid-co-4-(1,2-diol)-1,4,7-triazaheptane)

5 The reaction mixture containing 0.288 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate and 0.15 g of 4-(1,2-diol)-1,4,7-triazaheptane (**Example 3- 49(a)**) stirred at 100°C for 18 hours. The resulting product was dissolved in 3 mL of methanol and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2, it was dialyzed against DI
10 water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 160 mg of the desired product as a light yellow solid.

Example 3- 50: Synthesis of poly (4,4-trimethylene dipiperidine bispropanoic acid-co-4-(1,2-diol)-1,4,7-triazaheptane-co-1,3-diamino propane)

15 The reaction mixture containing 0.25 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate, 0.09 g of 4-(1,2-diol)-1,4,7-triazaheptane (**Example 3- 49(a)**) and 0.05 g of 1,3-diamino propane stirred at 100°C for 18 hours. The resulting product was dissolved in 3 mL of methanol and poured into 50 mL of ethyl acetate. The precipitate was isolated by filtration and was dissolved in 20 mL of DI water. After adjusting the pH of the solution to 2,
20 it was dialyzed against DI water using a dialysis membrane of molecular weight cut off of 1000 Dalton. The solution remaining in the dialysis membrane was lyophilized to dryness yielding 150 mg of the desired product as a light yellow solid.

Example 3- 51: Synthesis of poly (4,4-Trimethylene dipiperidine bispropanoic acid-co-5-(1,2-diol)-1,5,9-triazanonane)

Example 3- 51(a): Synthesis of 5-(1,2-diol)-1,5,9-triazanonane

25 The reaction mixture containing 1.5 g of 1,9-Bis-BOC-1,5,9-triazanonane, 0.34 g of glycidol, and 10mL of ethanol was refluxed for 15 hours. After removal of the solvent, the residue was purified by column chromatography using a gradient solvent system ranging from 100% hexane to 100% ethyl acetate) yielding 0.7g of 1, 9-bis-boc-5-(1,2-diol)-1,5,9-triazanonane. To 0.7g of 1, 9-bis-boc-5-(1,2-diol)-1,5,9-triazanonane dissolved in 2 mL of
30 methanol was added 0.25 mL of concentrated HCl and the reaction mixture stirred at 50°C for 24 hours. After removal of the solvent under reduced pressure, the residue was dissolved in 10mL of methanol/water (1:1) mixture and 5 g of Amberlyst OH 26 resin was added it. After

stirring at room temperature for 3 hours, the resin was filtered off. The solvent was removed under reduced pressure and the residue was lyophilized to dryness yielding 0.28g of the desired product as light yellow oil.

Example 3- 51(b): Synthesis of poly(4,4'-Trimethylene dipiperidine bispropanoic acid-co-5-(1,2-diol)-1,5,9-triazanonane)

The reaction mixture containing 0.23 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate and 0.15 g of 5-(1,2-diol)-1,5,9-triazanonane was stirred at 100°C for 18 hours. The resulting reaction mixture was dissolved in 5 mL of methanol and poured into 50 mL of ethyl acetate. After filtering off the solvent, the residue was dissolved in 20 mL of DI water. The pH of the solution was adjusted to 2 by adding dilute HCl and the solution subjected to centrifugation using with Microsep membrane filter with a molecular weight cut off of 1000 Dalton. The fraction with molecular weight higher than 1000 Dalton was collected and lyophilized to dryness yielding 100 mg of the desired product as a light yellow solid.

Example 3- 52: Synthesis of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-5-(1,2-diol)-1,5,9-triazanonane-co-1,3-diamino propane).

The reaction mixture containing 0.125 g 4, 4'-trimethylene dipiperidine bispropanoic acetate (**Example 3- 1**), 0.05 g of 5-(1,2-diol)-1,5,9-triazanonane (**Example 3- 51(a)**), and 0.3 g of 1,3-diamino propane was stirred at 100°C for 18 hours. The resulting reaction mixture was dissolved in 5 mL of methanol poured into 50 mL of ethyl acetate. After filtering off the solvent, the residue was dissolved in 20 mL of DI water. The pH of the solution was adjusted to 2 by adding dilute HCl and the solution subjected to centrifugation using with Microsep membrane filter with a molecular weight cut off of 1000 Dalton. The fraction with molecular weight higher than 1000 Dalton was collected and lyophilized to dryness yielding 90 mg of the desired product as a light yellow solid.

Example 3- 53: Synthesis of glycidol modified poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane)

To 0.26 g poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane) (**Example 3- 6**) dissolved in 2 mL of ethanol was added 16.5 mg of glycidol. The reaction mixture at 140°C for 30minutes using a microwave reactor. The resulting reaction mixture was poured into 50 mL of ethyl acetate. After filtration, the residue was washed with ethyl acetate (3 x 50 mL). Subsequently, it was dissolved in 10 mL of DI water and was subjected to centrifugation using with Microsep membrane filter with a molecular weight cut

off of 1000 Dalton. The fraction with molecular weight higher than 1000 Dalton was collected and lyophilized to dryness yielding 126 mg of the desired product as a light yellow solid.

Example 3- 54: Synthesis of Guanidine terminated poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane)

To 0.3g of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane) (**Example 3- 6**) dissolved in 2 mL of methanol was added 0.1 g of 1H-pyrazole-1-carboxamide and 0.11 g of N,N'-diisopropylethylamine. The reaction mixture was stirred at 60°C for 8 hours. The resulting reaction mixture was poured into 50 mL of ethyl acetate. After filtration, the residue was washed with ethyl acetate (3 x 50 mL). The resulting solid was dissolved in 2 mL of DI water and was passed through a PD-10 Sephadex column. The desired fractions were collected, lyophilized to dryness yielding 0.19 g of the polymer as a light yellow solid.

Example 3- 55: Synthesis of Polyethylene glycol (PEG-4) terminated poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane).

To 0.128 g of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane) (**Example 3- 6**) dissolved in 5 mL of methanol solution was added 0.2 mL of triethyl amine followed by 0.075 g of m-dPEG4-NHS ester. The reaction solution was stirred at room temperature for 22 hours. The resulting reaction mixture was poured into 50 mL of ethyl acetate. After filtration, the residue was washed with ethyl acetate (5 x 50 mL). The residue was subsequently dissolved in 2 mL of DI water and the pH of the resulting solution was adjusted to 2 using dilute HCl was subjected to centrifugation using a Microsep membrane filter with a molecular weight cut off of 1000 Dalton. The fraction with molecular weight higher than 1000 Dalton was collected and lyophilized to dryness yielding 50 mg of the desired product as a light yellow solid.

Example 3- 56: Synthesis of Polyethylene glycol (PEG-12) terminated poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane)

To 0.1 g of poly (4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane) (**Example 3- 6**) dissolved 5 mL of methanol was added 0.2 mL of triethyl amine followed by 0.12 g of m-dPEG12-NHS ester. The reaction solution was stirred at room temperature for 22 hours. The resulting reaction mixture was poured into 50 mL of ethyl acetate. After filtration, the residue was washed with ethyl acetate (5 x 50 mL). The residue was subsequently dissolved in 2 mL of DI water and the pH of the resulting solution was

adjusted to 2 using dilute HCl. was subjected to centrifugation using with Microsep membrane filter with a molecular weight cut off of 1000 Dalton. The fraction with molecular weight higher than 1000 Dalton was collected and lyophilized to dryness yielding 60 mg of the desired product as a light yellow solid.

5 **Example 3- 57: Synthesis of monodispersed polymer (heptamer) of poly(4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane)**

Example 3- 57(a): Synthesis of 4,4'-trimethylene dipiperidine bispropanoic acid-1,3-diamino propane trimer

10 The reaction mixture containing 3 g of 4, 4'-trimethylene dipiperidine bispropanoic acetate (**Example 3- 2**) and 4.1 g of mono N-boc-1,3-diamino propane was stirred at 100°C for 18 hours. The resulting reaction mixture was purified by column chromatography using an amine modified silica column and the gradient solvent system ranging from 100% hexane to ethyl acetate/hexane (50/50)). The appropriate fraction was collected and removal of the solvent under reduced pressure produced 2.6 g of 4, 4'-trimethylene dipiperidine
15 bispropanoic acid- bis-BOC-1,3-diamino propane.

To 0.55 g of 4,4'-trimethylene dipiperidine bispropanoic acid- bis-boc-1,3-diamino propane dissolved in 5 mL of methanol was added 0.5 mL of concentrated HCl and the reaction mixture was stirred at 50°C for 10 hours. After removal of the solvent under reduced pressure, the residue was dissolved in 10 mL of methanol/water (1:1) and was treated with 5
20 g of Amberlyst OH 26 resin. After stirring at room temperature for 3 hours, the resin was filtered off. The filtrate was evaporated dryness and the residue was lyophilized yielding 0.5 g of the product as a white solid.

Example 3- 57(b): Synthesis of 1-BOC-4,4'-trimethylene-1'-propanoic acid

To 2 g of 1-BOC-4,4'-trimethylene-1'propanoic methyl ester, 0.9 g of 50 wt%
25 solution of aqueous sodium hydroxide was added and the reaction mixture was stirred at 60°C for 15 hours. To this reaction mixture was added concentrated HCl until pH of the reaction reached 7.5. The reaction mixture was evaporated to dryness and residue was lyophilized to complete dryness. To this dry residue was added 10 mL of dichloromethane and the resulting mixture was stirred at room temperature for 30 minutes. After filtering off the insoluble
30 particles, the filtrate was evaporated to dryness to give 0.7 g of a white solid product.

Example 3- 57(c): Synthesis of bis-boc-4,4'-trimethylene dipiperidine bispropanoic acid-1,3-diamino propane pentamer

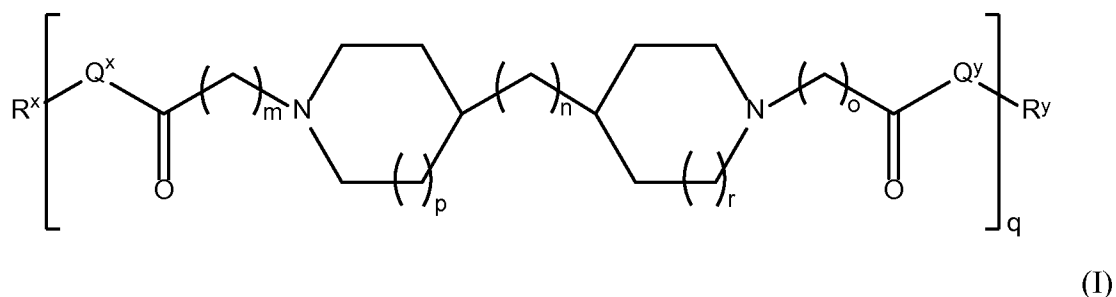
To 90 mg of 1-boc-4,4'-trimethylene-1'-propanoic acid (**Example 3- 57(b)**) dissolved in 2 mL of dichloromethane/DMF (1:1 v/v) was added 38 mg of 1,1-carbonyl diimidazole. After stirring at room temperature for 1 hour, 0.05 g of 4,4'-trimethylene dipiperidine bispropanoic acid-1,3-diamino propane trimer (**Example 3- 57(a)**) was added to reaction mixture. The resulting reaction mixture was stirred at room temperature for 20 hours. After removing the solvent under reduced pressure, the residue was purified by column chromatography using an amine modified silica column using a gradient solvent system ranging from 100% ethyl acetate to ethyl acetate/methanol (95/5)) yielding 80 mg of the product as a colorless oil. This oil was dissolved 2 mL of methanol followed by addition of 0.5 mL of concentrated HCl. The reaction mixture was stirred at 50°C for 10 hours. The solvent was evaporated removed under reduced pressure and the residue was lyophilized to dry to yield 60 mg of the desired product as yellow viscous oil.

Example 3- 57(d): Synthesis of poly(4,4'-trimethylene dipiperidine bispropanoic acid-co-1,3-diamino propane) heptamer

To 35 mg of 4,4'-trimethylene dipiperidine bispropanoic acid-1,3-diamino propane pentamer (**Example 3- 57(c)**) dissolved in 1 mL of methanol was added 0.08 mL of triethyl amine and 24 mg of boc-(3-acrylamido) propyl amine. The reaction mixture was stirred at room temperature overnight. The reaction mixture was poured into 10 mL of ethyl acetate. The residue was isolated by filtration and was washed with ethyl acetate (3 x 10 mL). The residue was dried at room temperature under reduced pressure yielding 40 mg of a white solid. To this solid residue was added 2 mL of methanol and 0.5 mL of concentrated HCl. The resulting reaction mixture was added stirred at 50°C for 10 hours. After removing the solvent under reduced pressure, residue was purified by preparative HPLC yielding 10 mg of the desired product as light yellow viscous oil.

CLAIMS

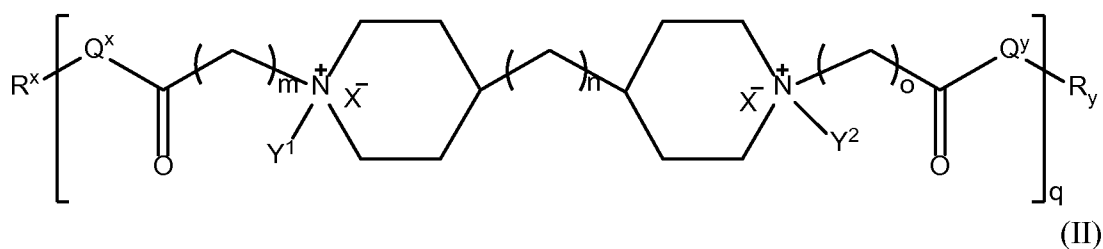
1. A compound comprising the structure of Formula (I):



wherein:

- i) m is 0, 1, 2, or 3;
- ii) n is 0, 1, 2, or 3;
- iii) o is 0, 1, 2, or 3;
- iv) p is 0 or 1;
- v) r is 0 or 1;
- vi) q is an integer from 1 to 400;
- vii) Q^x is NH, (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_3-C_{10}) cycloalkyl, (C_2-C_9) heterocycloalkyl, (C_6-C_{14}) aryl, (C_2-C_9) heteroaryl;
- viii) Q^y is $NH-R^w$, $NH-CH_2-R^w$, (C_1-C_{10}) alkyl, or (C_6-C_{14}) aryl, wherein R^w is absent or a (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_6-C_{14}) aryl, or (C_2-C_9) heteroaryl;
- ix) R^x and R^y are each independently a pharmaceutically acceptable end group.

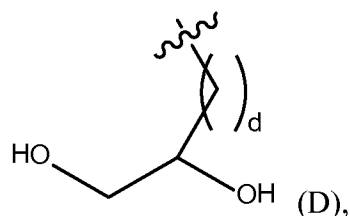
2. A compound comprising the structure of Formula (II):



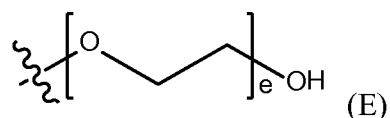
wherein:

- i) m is 0, 1, 2, or 3;
- ii) n is 0, 1, 2, or 3;
- iii) o is 0, 1, 2, or 3;
- iv) p is 0 or 1;

- v) r is 0 or 1;
- vi) q is an integer from 1 to 400;
- vii) Q^x is NH, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl;
- viii) Q^y is NH-R^w, NH-CH₂-R^w, (C₁-C₁₀)alkyl, or (C₆-C₁₄)aryl, wherein R^w is absent or a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl;
- ix) R^x and R^y are each independently a pharmaceutically acceptable end group;
- x) X⁻ is each independently a halo or any pharmaceutically acceptable anion;
- xi) Y¹ and Y₂ are each independently H or (C₁-C₁₀)alkyl optionally substituted by one or more substituents selected from the group consisting of (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl, (C₁-C₁₀)alkylamine, -S-O-(C₁-C₁₀)alkyl, -O(O)C-(C₁-C₁₀)alkyl, -(C₁-C₁₀)alkyl-COOH, (C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH, amide, a dihydroxy group, represented by Formula (D),

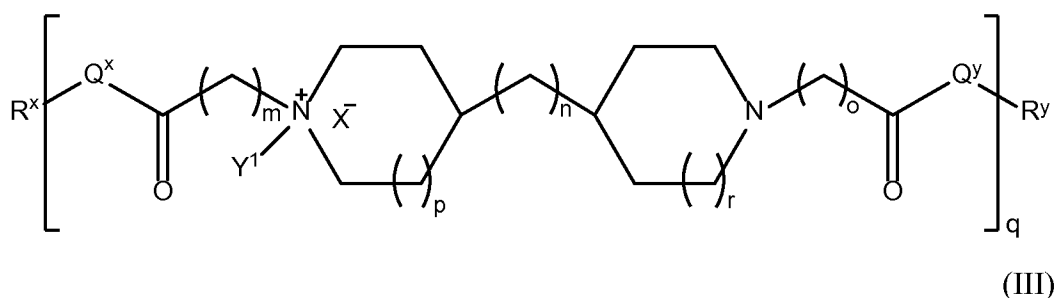


wherein d is an integer from 0 to 25, or a polyethylene glycol group, represented by Formula (E)



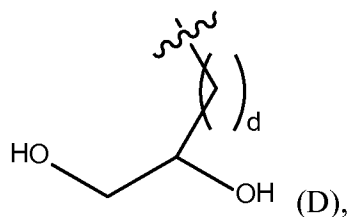
wherein e is an integer from 1 to 25.

3. A compound comprising the structure of Formula (III):



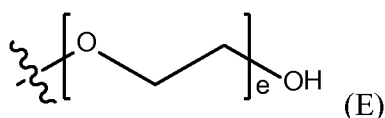
wherein:

- i) m is 0, 1, 2, or 3;
- ii) n is 0, 1, 2, or 3;
- iii) o is 0, 1, 2, or 3;
- iv) p is 0 or 1;
- v) r is 0 or 1;
- vi) q is an integer from 1 to 400;
- vii) Q^x is NH, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl;
- viii) Q^y is NH-R^w, NH-CH₂-R^w, (C₁-C₁₀)alkyl, or (C₆-C₁₄)aryl, wherein R^w is absent or a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl;
- iii) R^x and R^y are each independently a pharmaceutically acceptable end group;
- ix) X⁻ is a halo or any pharmaceutically acceptable anion;
- x) Y¹ is H or (C₁-C₁₀)alkyl optionally substituted by one or more substituents selected from the group consisting of (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl, (C₁-C₁₀)alkylamine, -S-O-(C₁-C₁₀)alkyl, -O(O)C-(C₁-C₁₀)alkyl, -(C₁-C₁₀)alkyl-COOH, (C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH, amide, a dihydroxy group, represented by Formula (D),



wherein d is an integer from 0 to 25, or

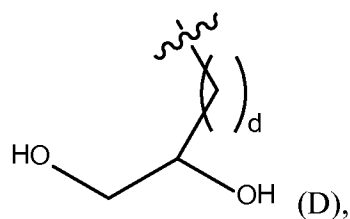
a polyethylene glycol group, represented by Formula (E),



wherein e is an integer from 1 to 400.

4. The compound according to claim 1 wherein n is 0.
5. The compound according to claim 2 wherein n is 0.
6. The compound according to claim 3 wherein n is 0.
7. The compound according to claim 1 wherein n is 1.
8. The compound according to claim 2 wherein n is 1.
9. The compound according to claim 3 wherein n is 1.
10. The compound according to claim 1 wherein n is 2.
11. The compound according to claim 2 wherein n is 2.
12. The compound according to claim 3 wherein n is 2.
13. The compound according to claim 1 wherein n is 3.
14. The compound according to claim 2 wherein n is 3.
15. The compound according to claim 3 wherein n is 3.
16. The compound according to claim 1 wherein p is 0.
17. The compound according to claim 2 wherein p is 0.
18. The compound according to claim 3 wherein p is 0.
19. The compound according to claim 1 wherein p is 1.
20. The compound according to claim 2 wherein p is 1.
21. The compound according to claim 3 wherein p is 1.
22. The compound according to claim 1 wherein r is 0.
23. The compound according to claim 2 wherein r is 0.
24. The compound according to claim 3 wherein r is 0.
25. The compound according to claim 1 wherein r is 1.
26. The compound according to claim 2 wherein r is 1.
27. The compound according to claim 3 wherein r is 1.
28. The compound according to claim 1 wherein p is 0 and r is 0.
29. The compound according to claim 2 wherein p is 0 and r is 0.
30. The compound according to claim 2 wherein p is 0 and r is 0.
31. The compound according to claim 1 wherein p is 1 and r is 1.
32. The compound according to claim 2 wherein p is 1 and r is 1.

33. The compound according to claim 3 wherein p is 1 and r is 1.
34. The compound according to claim 1 wherein n is 3; p is 1; and r is 1.
35. The compound according to claim 2 wherein n is 3; p is 1; and r is 1.
36. The compound according to claim 3 wherein n is 3; p is 1; and r is 1.
37. The compound according to claim 1 wherein n is 0; p is 1; and r is 1.
38. The compound according to claim 2 wherein n is 0; p is 1; and r is 1.
39. The compound according to claim 3 wherein n is 0; p is 1; and r is 1.
40. The compound according to claim 1 wherein n is 0; p is 0; and r is 0.
41. The compound according to claim 2 wherein n is 0; p is 0; and r is 0.
42. The compound according to claim 3 wherein n is 0; p is 0; and r is 0.
43. The compound according to claim 31 wherein:
 - i) Q^x is $-NH$; and
 - ii) Q^y is $NH-R^w$ -, wherein R^w is a (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_6-C_{14}) aryl, or (C_2-C_9) heteroaryl.
44. The compound according to claim 32 wherein:
 - i) Q^x is $-NH$; and
 - ii) Q^y is $NH-R^w$ -, wherein R^w is a (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_6-C_{14}) aryl, or (C_2-C_9) heteroaryl.
45. The compound according to claim 33 wherein:
 - i) Q^x is $-NH$; and
 - ii) Q^y is $NH-R^w$ -, wherein R^w is a (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_6-C_{14}) aryl, or (C_2-C_9) heteroaryl.
46. The compound according to claim 34 wherein:
 - i) m is 2;
 - ii) o is 2;
 - iii) Q^x is NH ; and
 - iv) Q^y is $NH-R^w$ -, wherein R^w is a (C_3) alkyl.
47. The compound according to claim 35 wherein:
 - i) m is 2;
 - ii) o is 2;
 - iii) Y^1 is H; and
 - iv) Y^2 is a dihydroxy group, represented by Formula (D),



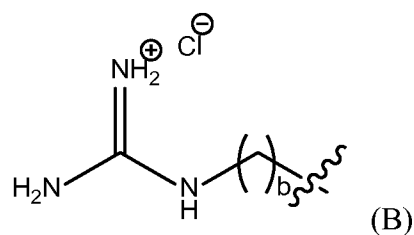
wherein d is 1.

48. The compound according to claim 36 wherein:
 - i) m is 2;
 - ii) o is 2;
 - iii) Q^x is NH; and
 - iv) Q^y is $NH-R^w$ wherein R^w is a (C_3) alkyl.
49. The compound according to claim 47, wherein the compound is a polymer.
50. The compound according to claim 48, wherein the compound is a polymer.
51. The compound according to claim 49, wherein the polymer is a copolymer with the compound according to claim 50.
52. The compound according to claim 31 wherein:
 - i) m is 0; and
 - ii) o is 0.
53. The compound according to claim 32 wherein:
 - i) m is 0; and
 - ii) o is 0.
54. The compound according to claim 33 wherein:
 - i) m is 0; and
 - ii) o is 0.
55. The compound according to claim 40 wherein:
 - i) Q^x is NH; and
 - ii) Q^y is $N-R^w$, wherein R^w is absent or a (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_6-C_{14}) aryl, or (C_2-C_9) heteroaryl.
56. The compound according to claim 41 wherein:
 - i) Q^x is NH; and
 - ii) Q^y is $N-R^w$, wherein R^w is absent or a (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_6-C_{14}) aryl, or (C_2-C_9) heteroaryl.
57. The compound according to claim 42 wherein:
 - i) Q^x is NH; and

- ii) Q^y is $N-R^w$, wherein R^w is absent or a (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_6-C_{14}) aryl, or (C_2-C_9) heteroaryl.

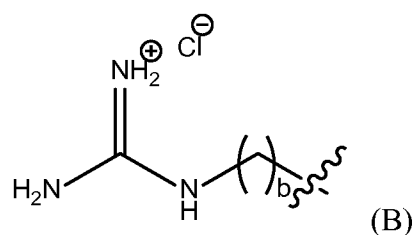
58. The compound according to claim 34 wherein:

- i) m is 2;
 ii) o is 2;
 iii) Q^x is NH ;
 iv) Q^y is $NH-R_w$, wherein R_w is a (C_3) alkyl
 v) R^x is a guanidinium chloride group represented by Formula (B),



wherein b is 3; and

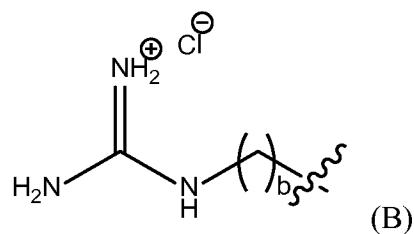
- vi) R^y is a guanidinium chloride group represented by Formula (B),



wherein b is 0.

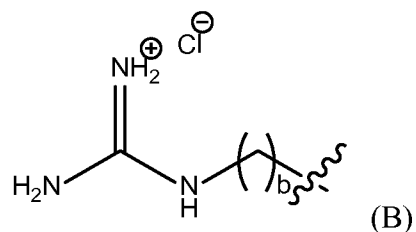
59. The compound according to claim 35 wherein:

- i) m is 2;
 ii) o is 2;
 iii) Q^x is NH ;
 iv) Q^y is $NH-R_w$, wherein R_w is a (C_3) alkyl
 v) R^x is a guanidinium chloride group represented by Formula (B),



wherein b is 3; and

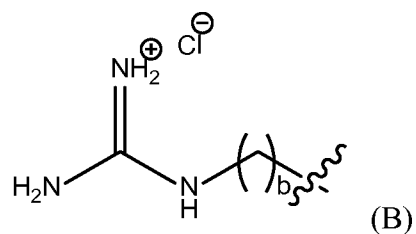
- vi) R^y is a guanidinium chloride group represented by Formula (B),



wherein b is 0.

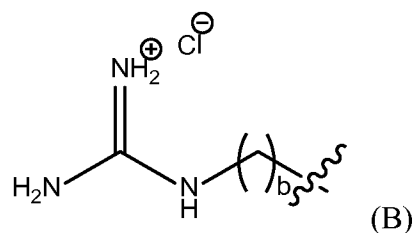
60. The compound according to claim 36 wherein:

- i) m is 2;
- ii) o is 2;
- iii) Q^x is NH;
- iv) Q^y is $NH-R_w$, wherein R_w is a (C_3) alkyl
- v) R^x is a guanidinium chloride group represented by Formula (B),



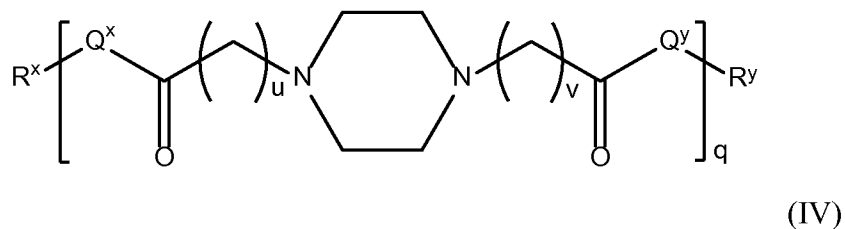
wherein b is 3; and

- vi) R^y is a guanidinium chloride group represented by Formula (B),



wherein b is 0.

61. A compound comprising the structure of Formula (IV):

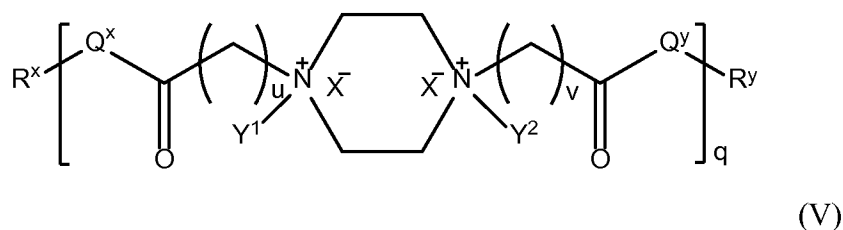


wherein:

- i) u is 0, 1, 2, or 3;
- ii) v is 0, 1, 2, or 3;
- iii) q is an integer from 1 to 400;

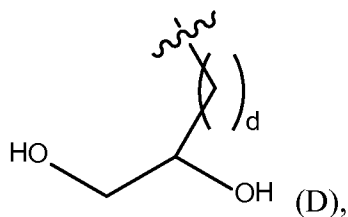
- iv) Q^x is NH, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl;
- v) Q^y is NH-R^w, NH-CH₂-R^w, (C₁-C₁₀)alkyl, or (C₆-C₁₄)aryl, wherein R^w is absent or a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl;
- vi) R^x and R^y are each independently a pharmaceutically acceptable end group.

62. A compound comprising the structure of Formula (V):

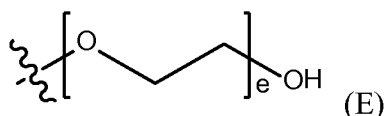


wherein:

- i) u is 0, 1, 2, or 3;
- ii) v is 0, 1, 2, or 3;
- iii) q is an integer from 1 to 400;
- iv) Q^x is NH, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl;
- v) Q^y is NH-R^w, NH-CH₂-R^w, (C₁-C₁₀)alkyl, or (C₆-C₁₄)aryl, wherein R^w is absent or a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl;
- vi) R^x and R^y are each independently a pharmaceutically acceptable end group;
- vii) X⁻ is independently a halo or any pharmaceutically acceptable anion,
- viii) Y¹ and Y² are independently H or (C₁-C₁₀)alkyl optionally substituted by one or more substituents selected from the group consisting of (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl, (C₁-C₁₀)alkylamine, -S-O-(C₁-C₁₀)alkyl, -O(O)C-(C₁-C₁₀)alkyl, -(C₁-C₁₀)alkyl-COOH, (C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH, amide, a dihydroxy group, represented by Formula (D),



wherein d is an integer from 0 to 25, or
a polyethylene glycol group, represented by Formula (E)



wherein e is an integer from 1 to 400.

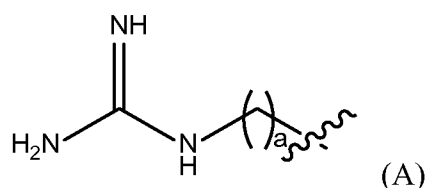
63. The compound according to claim 61 wherein u is 0.
64. The compound according to claim 62 wherein u is 0.
65. The compound according to claim 61 wherein u is 1.
66. The compound according to claim 62 wherein u is 1.
67. The compound according to claim 61 wherein u is 2.
68. The compound according to claim 62 wherein u is 2.
69. The compound according to claim 61 wherein u is 3.
70. The compound according to claim 62 wherein u is 3.
71. The compound according to claim 61 wherein v is 0.
72. The compound according to claim 62 wherein v is 0.
73. The compound according to claim 61 wherein v is 1.
74. The compound according to claim 62 wherein v is 1.
75. The compound according to claim 61 wherein v is 2.
76. The compound according to claim 62 wherein v is 2.
77. The compound according to claim 61 wherein v is 3.
78. The compound according to claim 62 wherein v is 3.
79. The compound according to claim 61 wherein u is 2 and v is 2.
80. The compound according to claim 62 wherein u is 2 and v is 2.
81. The compound according to claim 79 wherein:
 - i) Q_x is NH ;
 - ii) Q_y is $NH-CH_2-R_w$,

wherein R_w is a (C_1-C_{10}) alkyl, (C_2-C_9) heteroalkyl, (C_6-C_{14}) aryl,
or (C_2-C_9) heteroaryl.
82. The compound according to claim 80 wherein:

- i) Q_x is NH;
 ii) Q_y is NH-CH₂-R_w,

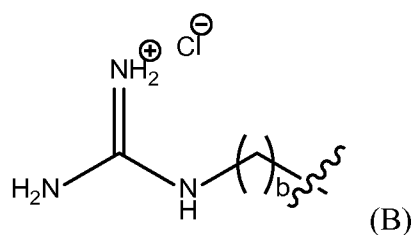
wherein R_w is a (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₆-C₁₄)aryl, or (C₂-C₉)heteroaryl.

83. The compound according to claim 1, wherein R^x and R^y are each independently H, (C₁-C₁₀)alkyl, (C₂-C₉)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl, (C₁-C₁₀)alkylamine, -O(O)C-(C₁-C₁₀)alkyl, (C₁-C₁₀)alkyl-COOH, (C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH, amide, a guanidino group represented by Formula (A)



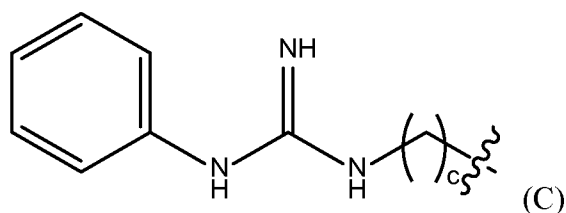
wherein a is an integer from 0 to 25,

a guanidinium chloride group represented by Formula (B),



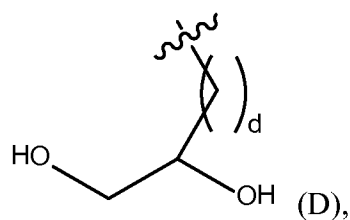
wherein b is an integer from 0 to 25,

a guanidinobenzene group represented by Formula (C),



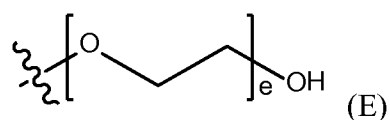
wherein c is an integer from 0 to 25,

a dihydroxy group, represented by Formula (D),



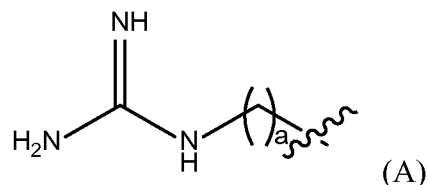
wherein d is an integer from 0 to 25, or

a polyethylene glycol group, represented by Formula (E)



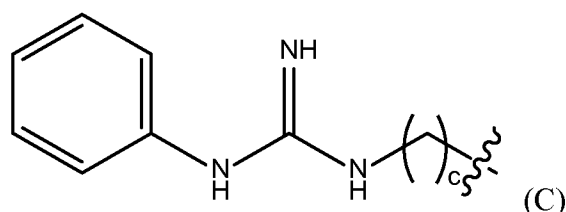
wherein e is an integer from 1 to 400.

84. The compound according to claim 83, wherein R^x and R^y are each independently selected from $-(O)CH_3$, a guanidino group represented by Formula (A)



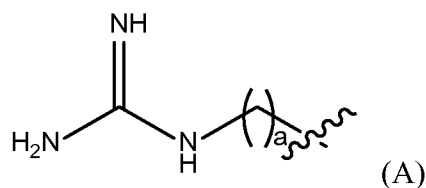
wherein a is an integer from 0 to 25, or

a guanidinobenzene group represented by Formula (C),



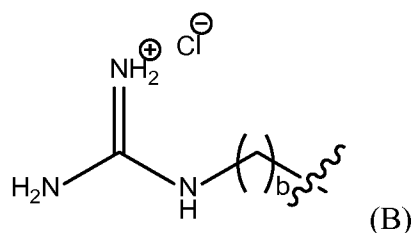
wherein c is an integer from 0 to 25.

85. The compound according to claim 2, wherein R^x and R^y are each independently H, (C_1-C_{10}) alkyl, (C_2-C_0) heteroalkyl, (C_3-C_{10}) cycloalkyl, (C_2-C_9) heterocycloalkyl, (C_6-C_{14}) aryl, (C_2-C_9) heteroaryl, (C_1-C_{10}) alkylamine, $-O(O)C-(C_1-C_{10})$ alkyl, (C_1-C_{10}) alkyl-COOH, (C_3-C_{10}) cycloalkyl-COOH, $-(O)CH_3$, $-OH$, amide, a guanidino group represented by Formula (A)



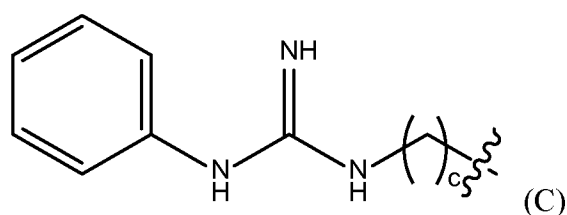
wherein a is an integer from 0 to 25,

a guanidinium chloride group represented by Formula (B),

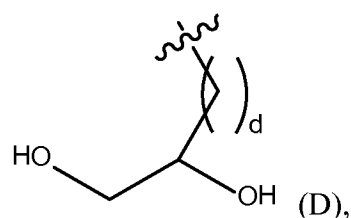


wherein b is an integer from 0 to 25,

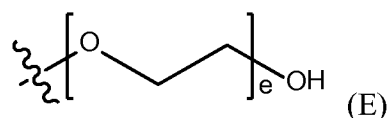
a guanidinobenzene group represented by Formula (C),



wherein c is an integer from 0 to 25,
a dihydroxy group, represented by Formula (D),

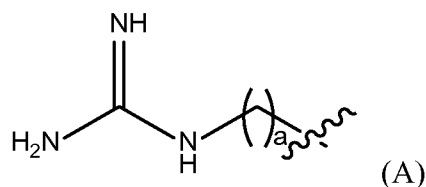


wherein d is an integer from 0 to 25, or
a polyethylene glycol group, represented by Formula (E)

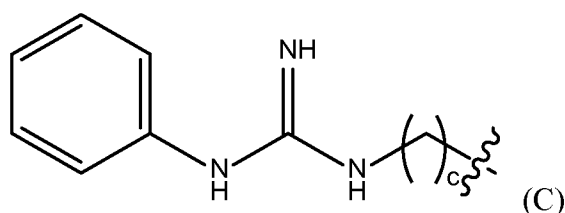


wherein e is an integer from 1 to 400.

86. The compound according to claim 85, wherein R^x and R^y are each independently selected from $-(O)CH_3$, a guanidino group represented by Formula (A)

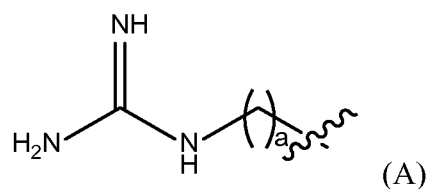


wherein a is an integer from 0 to 25, or
a guanidinobenzene group represented by Formula (C),

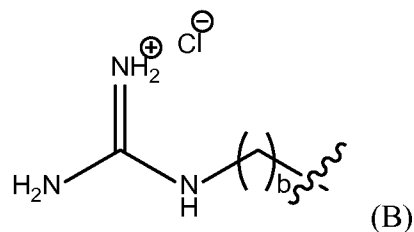


wherein c is an integer from 0 to 25.

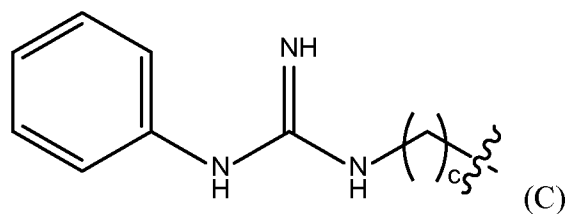
87. The compound according to claim 3, wherein R^x and R^y are each independently H, (C_1-C_{10}) alkyl, (C_2-C_0) heteroalkyl, (C_3-C_{10}) cycloalkyl, (C_2-C_9) heterocycloalkyl, (C_6-C_{14}) aryl, (C_2-C_9) heteroaryl, (C_1-C_{10}) alkylamine, $-O(O)C-(C_1-C_{10})$ alkyl, (C_1-C_{10}) alkyl-COOH, (C_3-C_{10}) cycloalkyl-COOH, $-(O)CH_3$, $-OH$, amide, a guanidino group represented by Formula (A)



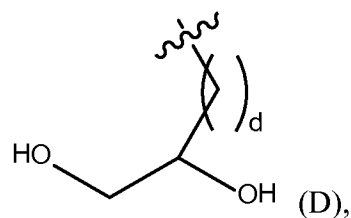
wherein a is an integer from 0 to 25,
a guanidinium chloride group represented by Formula (B),



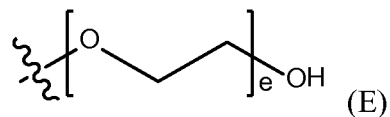
wherein b is an integer from 0 to 25,
a guanidinobenzene group represented by Formula (C),



wherein c is an integer from 0 to 25,
a dihydroxy group, represented by Formula (D),

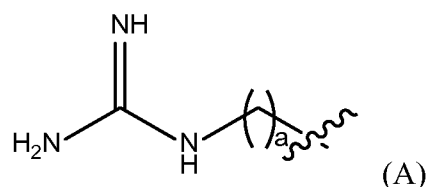


wherein d is an integer from 0 to 25, or
a polyethylene glycol group, represented by Formula (E)

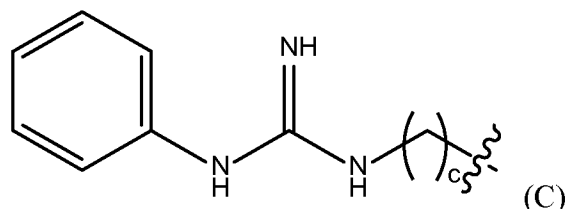


wherein e is an integer from 1 to 400.

88. The compound according to claim 87, wherein R^x and R^y are each independently selected from $-(O)CH_3$, a guanidino group represented by Formula (A)

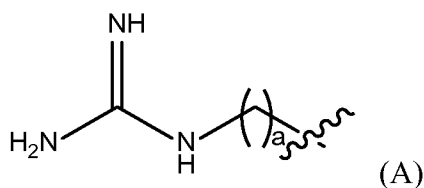


wherein a is an integer from 0 to 25, or
a guanidinobenzene group represented by Formula (C),

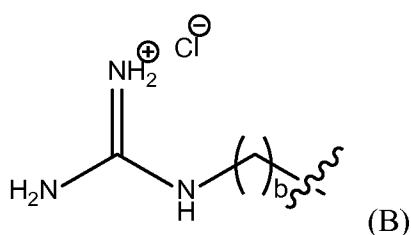


wherein c is an integer from 0 to 25.

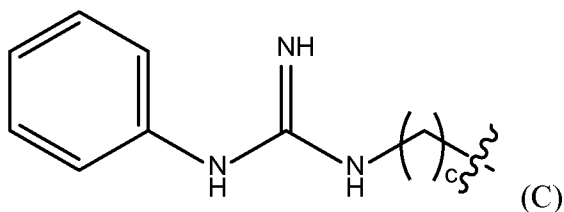
89. The compound according to claim 61, wherein R^x and R^y are each independently H, (C₁-C₁₀)alkyl, (C₂-C₀)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl, (C₁-C₁₀)alkylamine, -O(O)C-(C₁-C₁₀)alkyl, (C₁-C₁₀)alkyl-COOH, (C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH, amide, a guanidino group represented by Formula (A)



wherein a is an integer from 0 to 25,
a guanidinium chloride group represented by Formula (B),

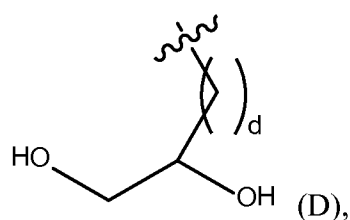


wherein b is an integer from 0 to 25,
a guanidinobenzene group represented by Formula (C),



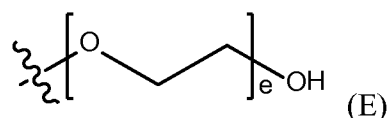
wherein c is an integer from 0 to 25,

a dihydroxy group, represented by Formula (D),



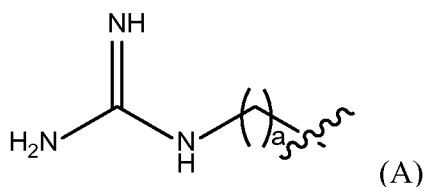
wherein d is an integer from 0 to 25, or

a polyethylene glycol group, represented by Formula (E)



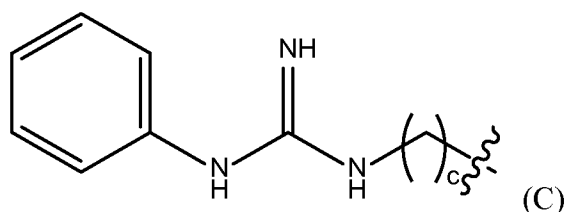
wherein e is an integer from 1 to 400.

90. The compound according to claim 89, wherein R^x and R^y are each independently selected from -(O)CH₃, a guanidino group represented by Formula (A)



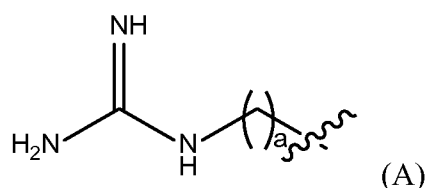
wherein a is an integer from 0 to 25, or

a guanidinobenzene group represented by Formula (C),



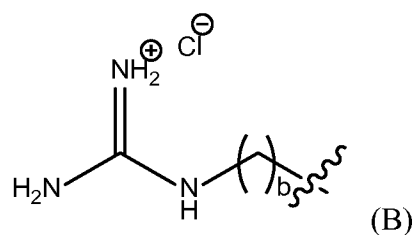
wherein c is an integer from 0 to 25.

91. The compound according to claim 62, wherein R^x and R^y are each independently H, (C₁-C₁₀)alkyl, (C₂-C₀)heteroalkyl, (C₃-C₁₀)cycloalkyl, (C₂-C₉)heterocycloalkyl, (C₆-C₁₄)aryl, (C₂-C₉)heteroaryl, (C₁-C₁₀)alkylamine, -O(O)C-(C₁-C₁₀)alkyl, (C₁-C₁₀)alkyl-COOH, (C₃-C₁₀)cycloalkyl-COOH, -(O)CH₃, -OH, amide, a guanidino group represented by Formula (A)



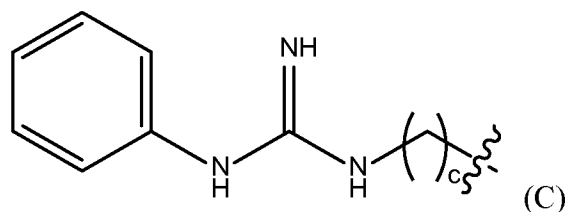
wherein a is an integer from 0 to 25,

a guanidinium chloride group represented by Formula (B),



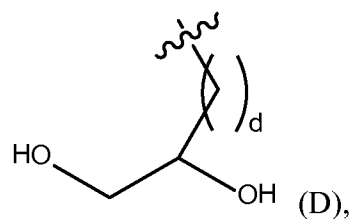
wherein b is an integer from 0 to 25,

a guanidinobenzene group represented by Formula (C),



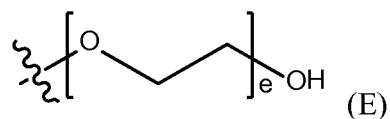
wherein c is an integer from 0 to 25,

a dihydroxy group, represented by Formula (D),



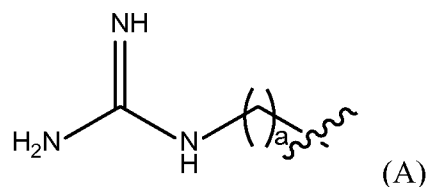
wherein d is an integer from 0 to 25, or

a polyethylene glycol group, represented by Formula (E)



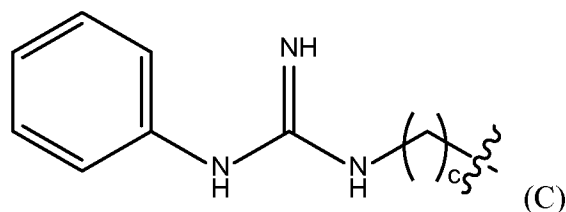
wherein e is an integer from 1 to 400.

92. The compound according to claim 91, wherein R^x and R^y are each independently selected from -(O)CH₃, a guanidino group represented by Formula (A)



wherein a is an integer from 0 to 25, or

a guanidinobenzene group represented by Formula (C),



wherein c is an integer from 0 to 25.

93. A pharmaceutical composition comprising a compound according to claim 1.
94. A pharmaceutical composition comprising a compound according to claim 2.
95. A pharmaceutical composition comprising a compound according to claim 3.
96. A pharmaceutical composition comprising a compound according to claim 61.
97. A pharmaceutical composition comprising a compound according to claim 62.
98. The pharmaceutical composition according to claim 93 for use in the treatment of mucositis.
99. The pharmaceutical composition according to claim 94 for use in the treatment of mucositis.
100. The pharmaceutical composition according to claim 95 for use in the treatment of mucositis.
101. The pharmaceutical composition according to claim 96 for use in the treatment of mucositis.
102. The pharmaceutical composition according to claim 97 for use in the treatment of mucositis.
103. The pharmaceutical composition according to claim 93 for use in the treatment of oral mucositis.
104. The pharmaceutical composition according to claim 94 for use in the treatment of oral mucositis.
105. The pharmaceutical composition according to claim 95 for use in the treatment of oral mucositis.
106. The pharmaceutical composition according to claim 96 for use in the treatment of oral mucositis.
107. The pharmaceutical composition according to claim 97 for use in the treatment of oral mucositis.
108. The pharmaceutical composition according to claim 93 for use in the treatment of infection.

109. The pharmaceutical composition according to claim 94 for use in the treatment of infection.
110. The pharmaceutical composition according to claim 95 for use in the treatment of infection.
111. The pharmaceutical composition according to claim 96 for use in the treatment of infection.
112. The pharmaceutical composition according to claim 97 for use in the treatment of infection.
113. The pharmaceutical composition according to claim 108, wherein the infection is a surgical site infection.
114. The pharmaceutical composition according to claim 108, wherein the infection is a lung infection associated with cystic fibrosis.
115. The pharmaceutical composition according to claim 114, wherein the infection is a *Pseudomonas aeruginosa* lung infection.
116. The pharmaceutical composition according to claim 115, wherein biofilms are present in the *Pseudomonas aeruginosa* lung infection.
117. The pharmaceutical composition according to claim 109, wherein the infection is a surgical site infection.
118. The pharmaceutical composition according to claim 109, wherein the infection is a lung infection associated with cystic fibrosis.
119. The pharmaceutical composition according to claim 118, wherein the infection is a *Pseudomonas aeruginosa* lung infection.
120. The pharmaceutical composition according to claim 119, wherein biofilms are present in the *Pseudomonas aeruginosa* lung infection.
121. The pharmaceutical composition according to claim 110, wherein the infection is a surgical site infection.
122. The pharmaceutical composition according to claim 110, wherein the infection is a lung infection associated with cystic fibrosis.
123. The pharmaceutical composition according to claim 122, wherein the infection is a *Pseudomonas aeruginosa* lung infection.
124. The pharmaceutical composition according to claim 123, wherein biofilms are present in the *Pseudomonas aeruginosa* lung infection.

125. The pharmaceutical composition according to claim 111, wherein the infection is a surgical site infection.
126. The pharmaceutical composition according to claim 111, wherein the infection is a lung infection associated with cystic fibrosis.
127. The pharmaceutical composition according to claim 126, wherein the infection is a *Pseudomonas aeruginosa* lung infection.
128. The pharmaceutical composition according to claim 127, wherein biofilms are present in the *Pseudomonas aeruginosa* lung infection.
129. The pharmaceutical composition according to claim 112, wherein the infection is a surgical site infection.
130. The pharmaceutical composition according to claim 112, wherein the infection is a lung infection associated with cystic fibrosis.
131. The pharmaceutical composition according to claim 130, wherein the infection is a *Pseudomonas aeruginosa* lung infection.
132. The pharmaceutical composition according to claim 131, wherein biofilms are present in the *Pseudomonas aeruginosa* lung infection.
133. A method of treating a condition selected from mucositis, oral mucositis, and infection comprising administering a compound according to claim 1.
134. A method of treating a condition selected from mucositis, oral mucositis, and infection comprising administering a compound according to 2.
135. A method of treating a condition selected from mucositis, oral mucositis, and infection comprising administering a compound according to 3
136. A method of treating a condition selected from mucositis, oral mucositis, and infection comprising administering a compound according to 61.
137. A method of treating a condition selected from mucositis, oral mucositis, and infection comprising administering a compound according to 62.
138. The method according to claim 133, wherein the infection is a surgical site infection.
139. The method according to claim 133, wherein the infection is a lung infection associated with cystic fibrosis.
140. The method according to claim 139, wherein the infection is a *Pseudomonas aeruginosa* lung infection.
141. The method according to claim 140, wherein biofilms are present in the *Pseudomonas aeruginosa* lung infection.

142. The method according to claim 134, wherein the infection is a surgical site infection.
143. The method according to claim 134, wherein the infection is a lung infection associated with cystic fibrosis.
144. The method according to claim 143, wherein the infection is a *Pseudomonas aeruginosa* lung infection.
145. The method according to claim 144, wherein biofilms are present in the *Pseudomonas aeruginosa* lung infection.
146. The method according to claim 135, wherein the infection is a surgical site infection.
147. The method according to claim 135, wherein the infection is a lung infection associated with cystic fibrosis.
148. The method according to claim 147, wherein the infection is a *Pseudomonas aeruginosa* lung infection.
149. The method according to claim 148, wherein biofilms are present in the *Pseudomonas aeruginosa* lung infection.
150. The method according to claim 136, wherein the infection is a surgical site infection.
151. The method according to claim 136, wherein the infection is a lung infection associated with cystic fibrosis.
152. The method according to claim 151, wherein the infection is a *Pseudomonas aeruginosa* lung infection.
153. The method according to claim 152, wherein biofilms are present in the *Pseudomonas aeruginosa* lung infection.
154. The method according to claim 137, wherein the infection is a surgical site infection.
155. The method according to claim 137, wherein the infection is a lung infection associated with cystic fibrosis.
156. The method according to claim 155, wherein the infection is a *Pseudomonas aeruginosa* lung infection.
157. The method according to claim 156, wherein biofilms are present in the *Pseudomonas aeruginosa* lung infection.

Figure I: Susceptibility of *Pseudomonas aeruginosa* Present in Biofilm of Cystic Fibrosis-Derived Human Airway Epithelial Cells of Representative Amine Functional Polyamide

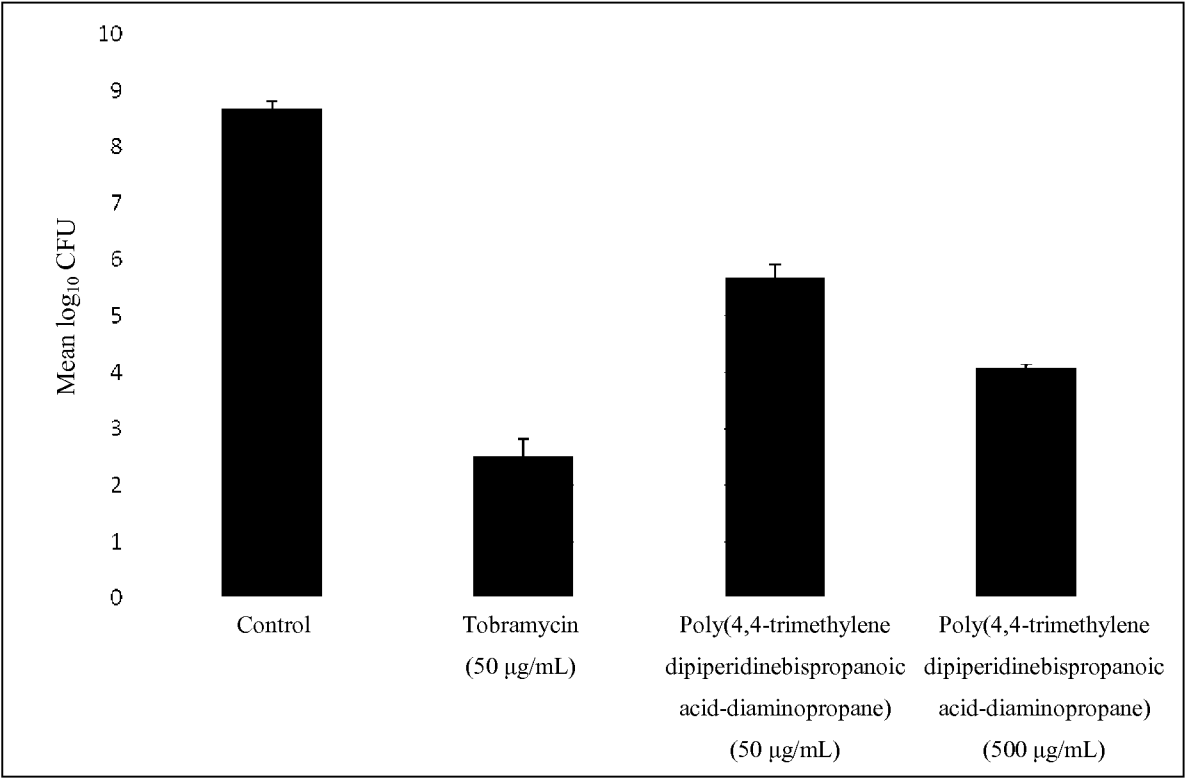


Figure 1 of 1

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2014/022984

A. CLASSIFICATION OF SUBJECT MATTER
INV. C08G69/00 A61K31/00
ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
C08G A61K C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 2012/295922 A1 (SCOTT RICHARD W [US] ET AL) 22 November 2012 (2012-11-22) paragraphs [0002], [0013]; compound Formula III -----	1-157
A	WO 2012/151554 A1 (HARVARD COLLEGE [US]; LOSICK RICHARD [US]; KOLODKIN-GAL ILLANA [US]; C) 8 November 2012 (2012-11-08) the whole document -----	1-157
A	WO 01/93878 A2 (GELTEX PHARMA INC [US]) 13 December 2001 (2001-12-13) the whole document -----	1-157
A	WO 2010/054269 A1 (NOVABAY PHARMACEUTICALS INC [US]; JAIN RAKESH K [US]; LOW EDDY [US]; F) 14 May 2010 (2010-05-14) the whole document -----	1-157

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Further documents are listed in the continuation of Box C.

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See patent family annex.

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"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

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"&" document member of the same patent family

Date of the actual completion of the international search

23 July 2014

Date of mailing of the international search report

25/08/2014

Name and mailing address of the ISA/

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Authorized officer

Zellner, Armin

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US2014/022984

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 2012295922 A1	22-11-2012	CA 2836099 A1	22-11-2012
		EP 2709619 A2	26-03-2014
		KR 20140078582 A	25-06-2014
		TW 201247654 A	01-12-2012
		US 2012295922 A1	22-11-2012
		WO 2012158672 A2	22-11-2012

WO 2012151554 A1	08-11-2012	US 2014056951 A1	27-02-2014
		US 2014056952 A1	27-02-2014
		WO 2012151554 A1	08-11-2012
		WO 2012151555 A1	08-11-2012

WO 0193878 A2	13-12-2001	AU 7545001 A	17-12-2001
		EP 1286676 A1	05-03-2003
		PT 1395289 E	16-03-2011
		US 2002081340 A1	27-06-2002
		WO 0193878 A2	13-12-2001

WO 2010054269 A1	14-05-2010	CA 2741667 A1	14-05-2010
		EP 2341886 A1	13-07-2011
		JP 2012508251 A	05-04-2012
		US 2010158818 A1	24-06-2010
		WO 2010054269 A1	14-05-2010

摘要

胺官能性聚酰胺沿聚合物链方向包含胺和铵基团。胺官能性聚酰胺可用作药物和在药物组合物中使用。胺官能性聚酰胺具体地用于治疗或预防粘膜炎和感染，尤其是口腔粘膜炎、手术部位感染和与囊性纤维化有关的肺部感染。