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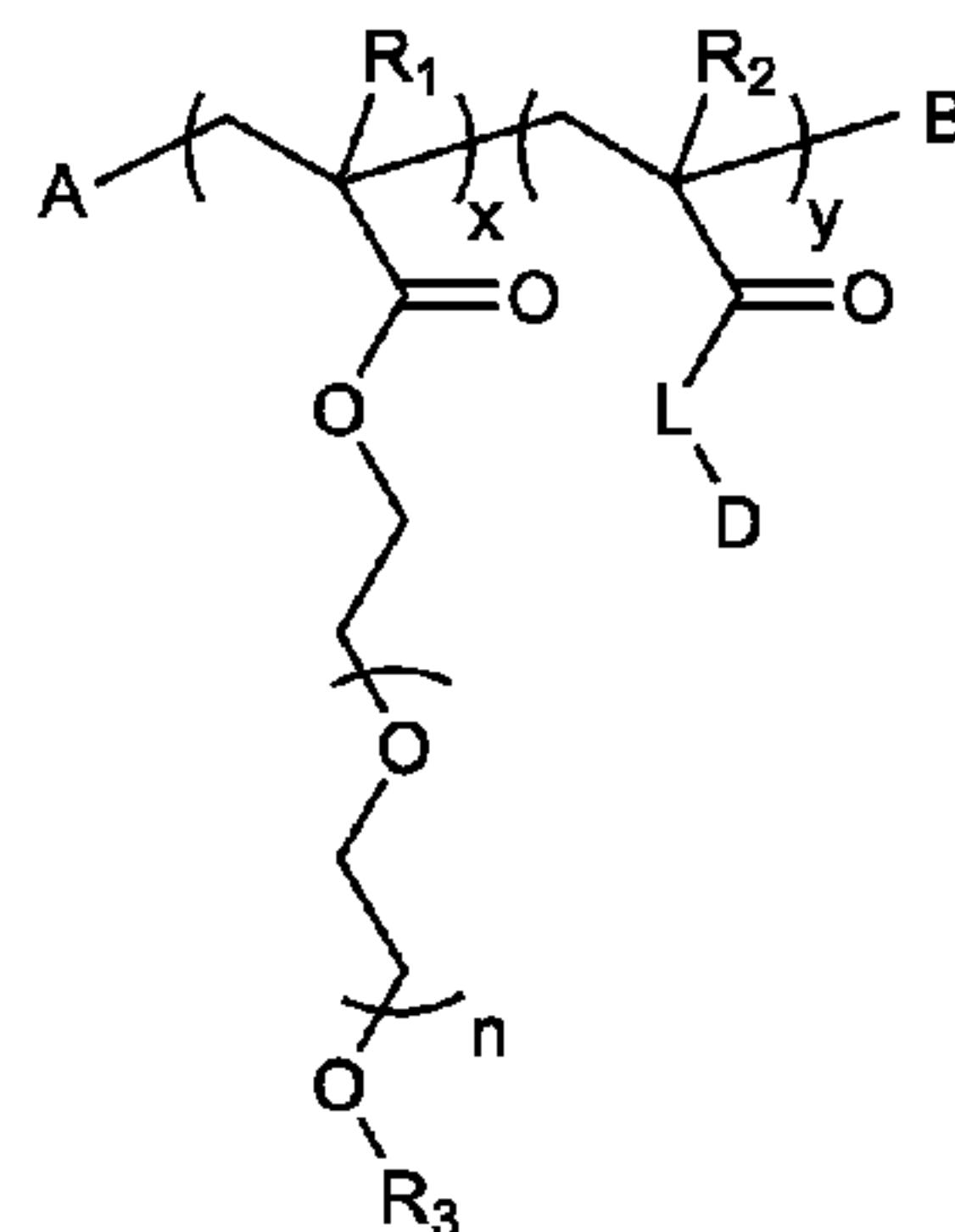
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(54) Titre : COPOLYMERES DIBLOCS AUTO-ASSEMBLÉS COMPOSÉS DE PEGMEMA ET DE SEGMENTS
POLYMERES PORTANT UN MÉDICAMENT

(54) Title: SELF-ASSEMBLED DIBLOCK COPOLYMERS COMPOSED OF PEGMEMA AND DRUG BEARING
POLYMERIC SEGMENTS



Formula I

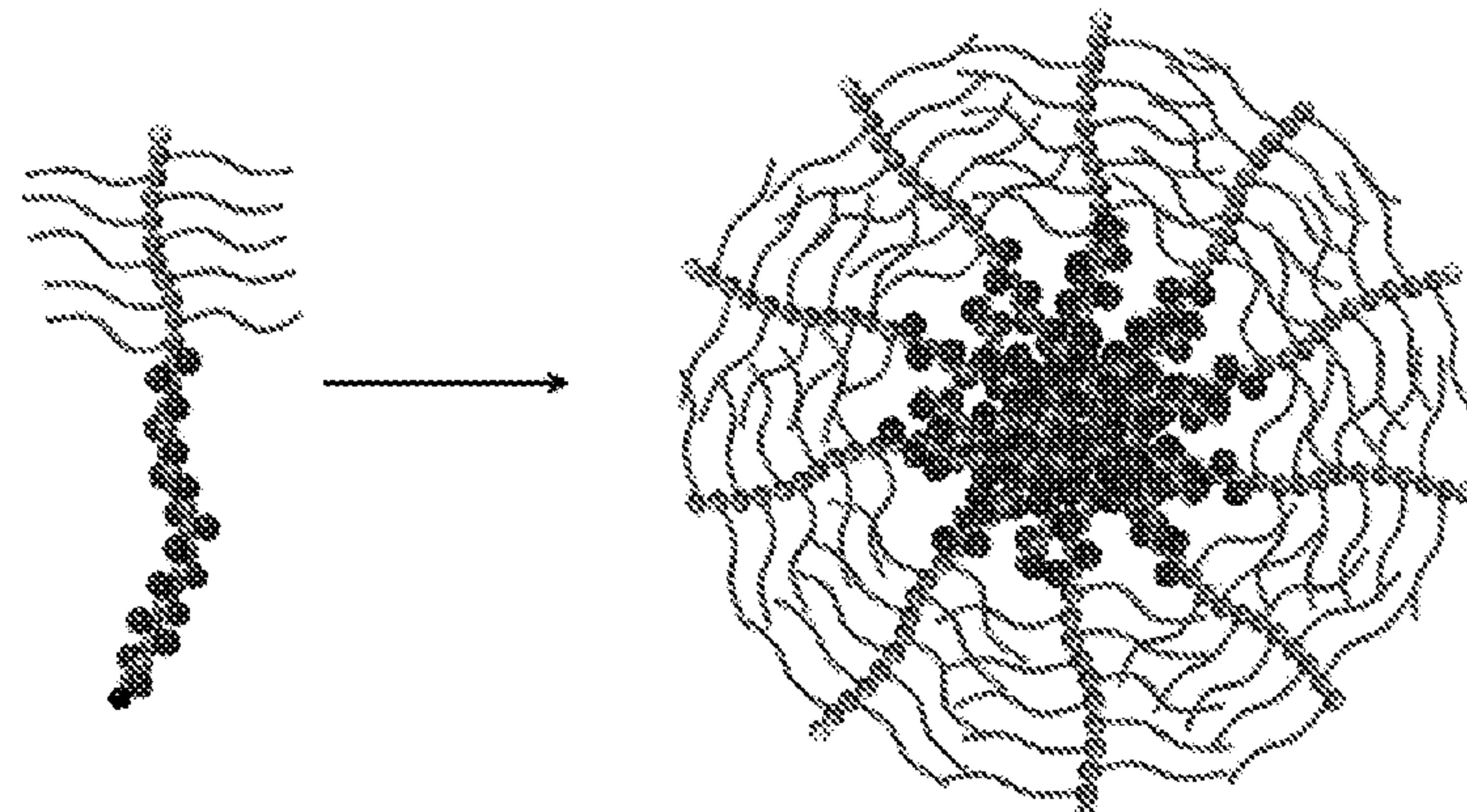


Figure 2

(57) Abrégé/Abstract:

This invention relates to polymer drug conjugates according to formula I, and their use for treatment of diseases such as cancer.

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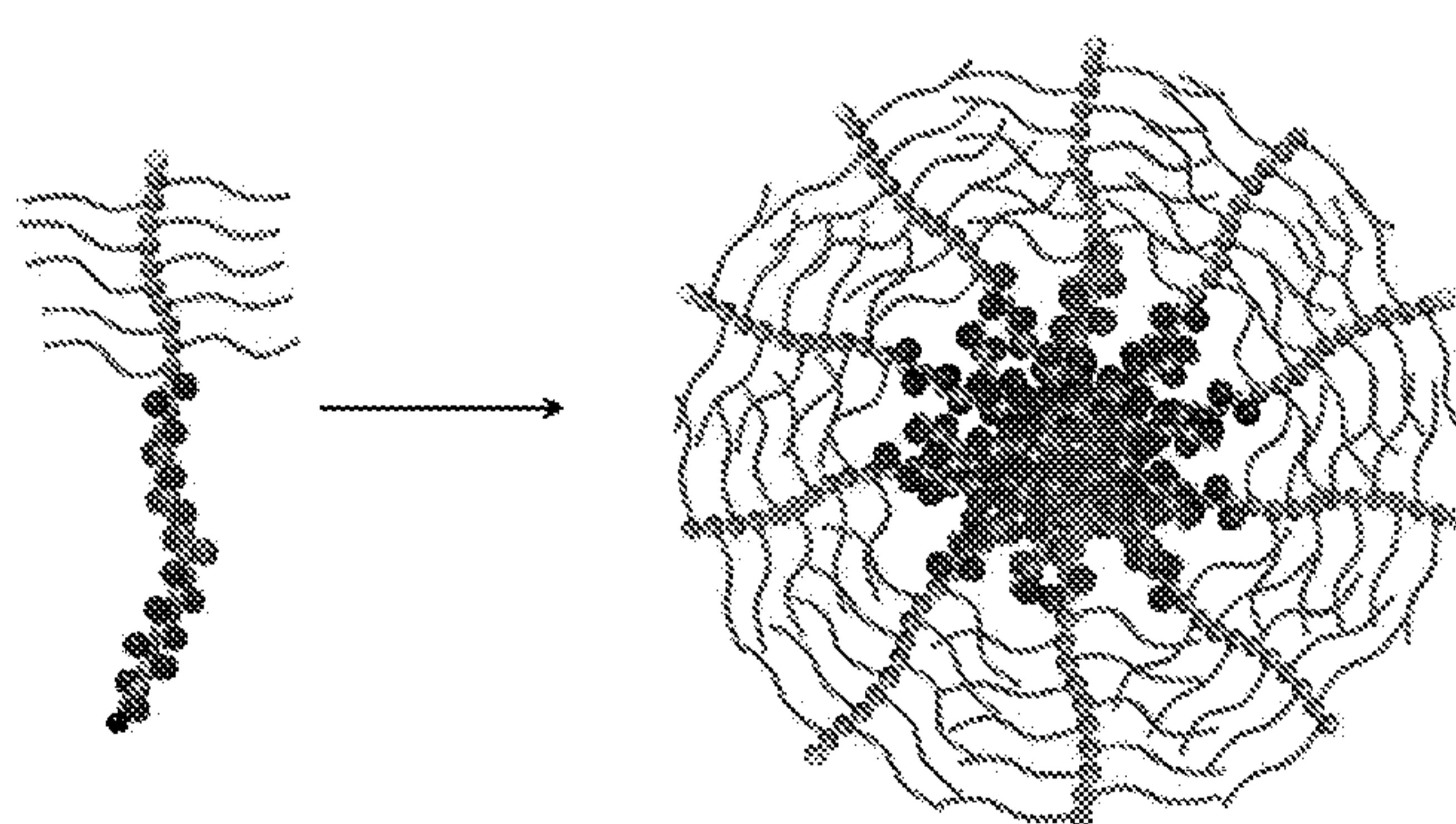
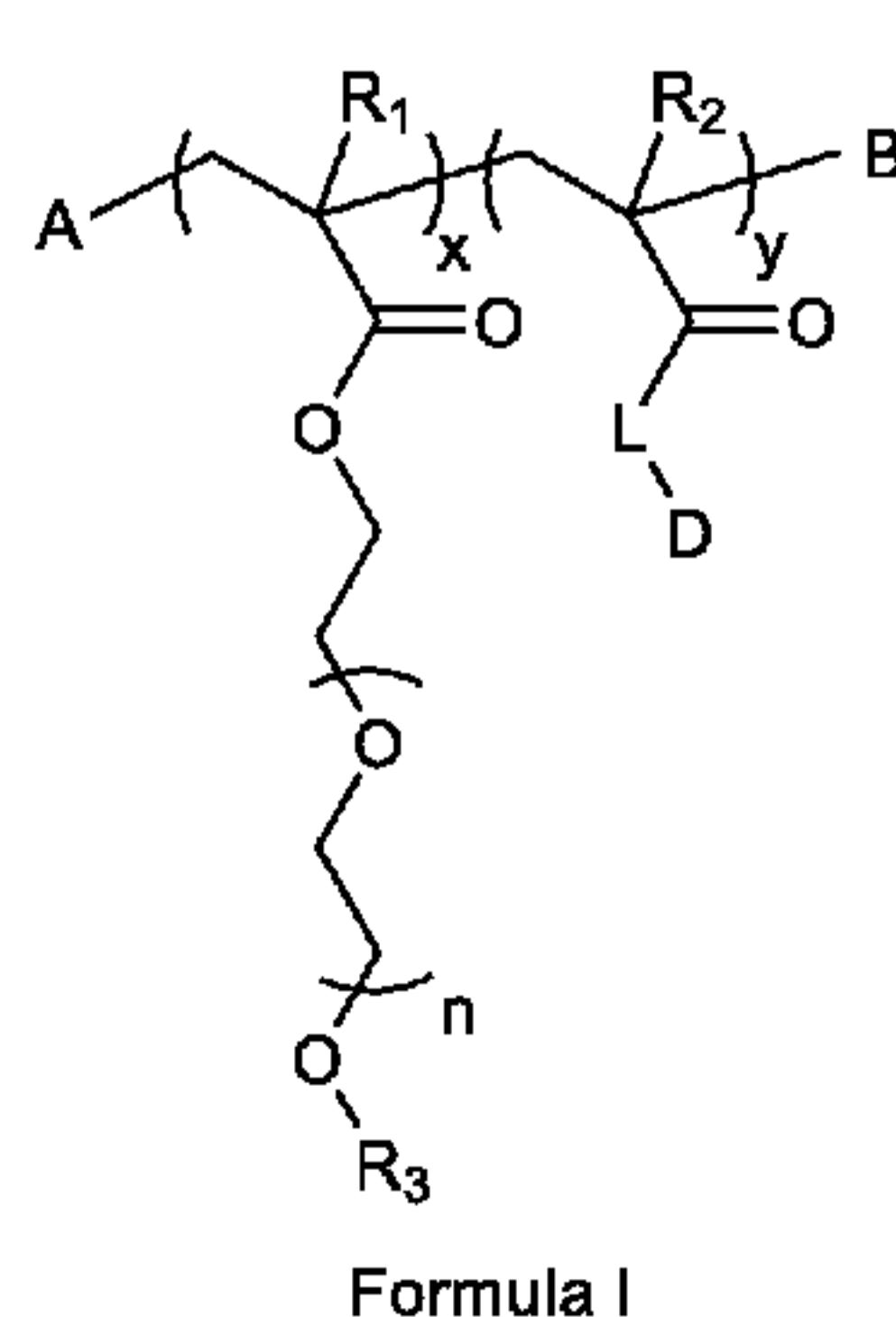


Figure 2

WO 2018/092048 A1

(57) **Abstract:** This invention relates to polymer drug conjugates according to formula I, and their use for treatment of diseases such as cancer.

SELF-ASSEMBLED DIBLOCK COPOLYMERS COMPOSED OF PEGMEMA AND DRUG BEARING POLYMERIC SEGMENTS

This invention relates to polymer drug conjugates according to formula I, assemblies made up of polymer drug conjugates of formula I, methods of preparing said polymer-drug conjugates and 5 assemblies and their use for treatment of diseases such as cancer.

BACKGROUND OF THE INVENTION

Chemotherapy agents used for treatment of cancer are mostly cytotoxic. These agents may accumulate in the tissues of the body in addition to the targeted area, which in turn causes lowered therapeutic benefit and undesired distribution of the drug throughout healthy body 10 tissues. The uncontrolled distribution of these agents throughout the body causes severe side effects to the patient.

In order to address the abovementioned problems, drug delivery systems, which can deliver the drug to targeted areas of the body, are developed. For example in some approaches of cancer treatment, these systems make use of the enhanced permeability and retention effect (EPR) 15 which implies that drug carriers having high molecular weight and large hydrodynamic volume accumulate in solid tumors and this, in turn leads to passive targeting of the drug molecule to the tumor tissue and minimize the damage of the chemotherapy agents to healthy tissue.

Since the discovery of the enhanced permeability and retention (EPR) effect, macromolecules with long blood circulation have become the primary substances for drug delivery due to 20 improved accumulation in tumors through fenestrated blood vessels.

There have been great interest on polymeric assemblies in drug delivery systems for cancer disease mainly due to their body distribution properties. Present invention also relates to assemblies that are suitable for use in drug delivery, specifically anticancer agents for treatment of cancer.

TECHNICAL FIELD OF THE INVENTION

There are various types of techniques for the preparation of polymeric assemblies known in the present state of the art. For example, self-assembly of polymeric compounds is an attractive method for polymeric assembly formation. Eisenberg and coworkers demonstrated the first 5 example of self-assembly of asymmetric copolymers in solutions in the 1990s. Then, there have been focused attention on amphiphilic block copolymers having polymer blocks with different physical and chemical properties to generate polymeric structures via self-assembly

Drug loading in such polymeric assemblies can be achieved by covalent attachment or physical encapsulation via hydrophobic interactions. Considering the very low total volume of these 10 assemblies in their colloidal suspensions, high drug loading capacity is of crucial importance to achieve effective drug formulations. Low drug loading capacity of polymeric assemblies mainly in micellar formulations is still a major drawback in the field.

Several researchers addressed this problem by taking advantage of π - π interactions through modifying the drug molecules or using an additive compound rather than providing covalent 15 attachment. However, non-capsulated drug aggregates in the micellar type polymeric nanoparticle formulations is another drawback to provide a safe formulation for their clinical applications.

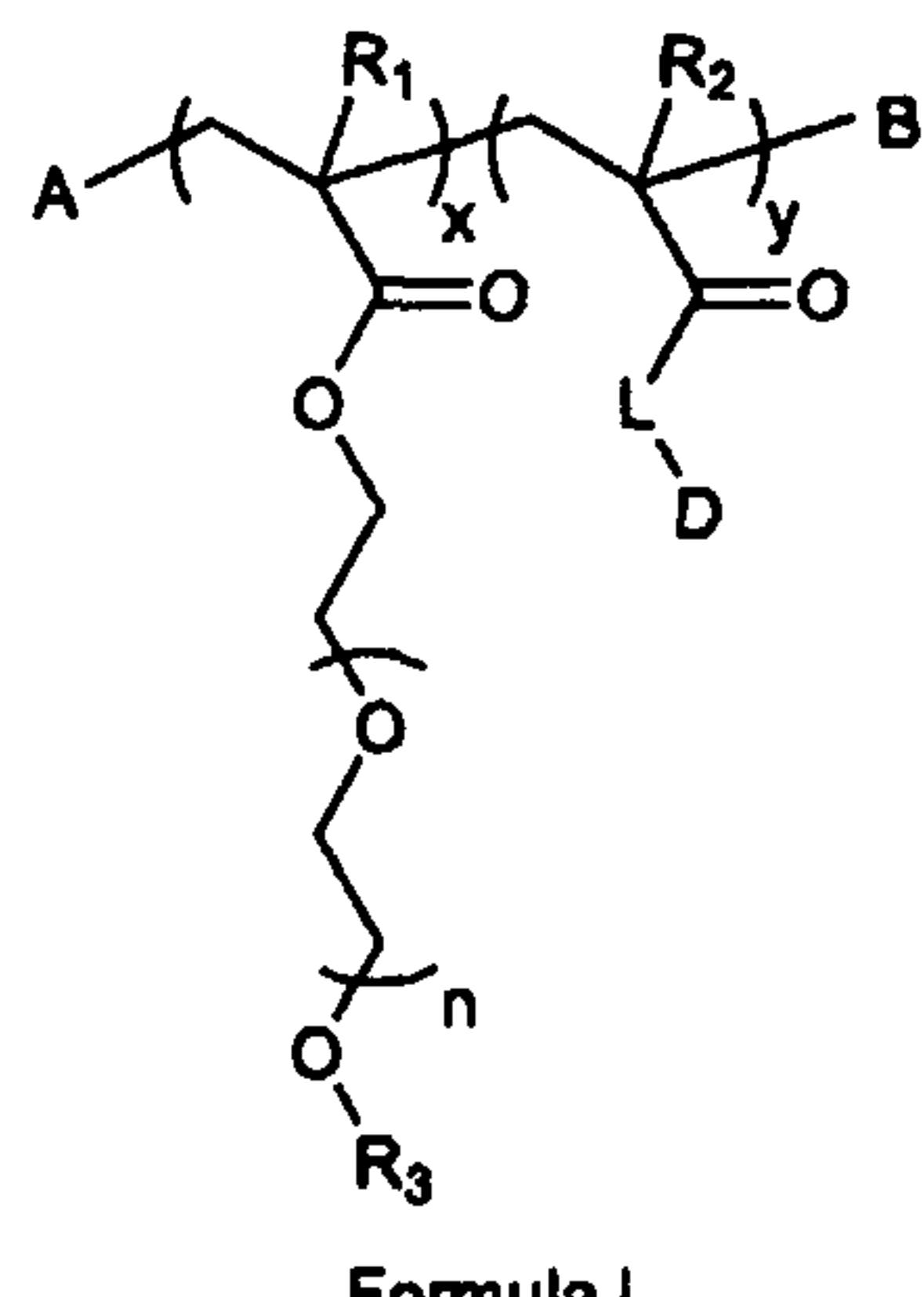
In another approach, Stenzel and co-workers have utilized drug attachable amphiphilic block copolymers to form hydrophobic segment of the block copolymers for the formation of 20 polymeric assemblies. This strategy has the lack of complete conjugation of the drug to the polymer backbone due to steric hindrance which is resulted in not well-defined final structure. Moreover, remaining reactive groups on the polymer backbone due to incomplete conjugation may be the most important drawback of this kind of post-polymerization conjugation strategies due to possible undesirable biological interactions *in vivo*.

25 In light of the current state of the art summarized above there is need for block copolymers and assemblies formed thereof that are well defined by means of the content of the drug, that provide efficient conjugation of the drug without any remaining reactive sites and which has a high drug loading capacity.

The inventors have found that the amphiphilic block copolymers according to present invention provide formation of micellar type assemblies that has high drug content and that are well defined by means of drug attachment.

BRIEF DESCRIPTION OF THE INVENTION

5 Present invention relates to a polymer-drug conjugate in the form of a block co-polymer for delivery of therapeutic agents as shown in Formula I



In other words, the invention relates to a polymer-drug conjugate of Formula I in the form of a

10 block co-polymer and assemblies made from polymer-drug conjugates of Formula I, wherein

- R₁ and R₂ are independently selected from H or -CH₃
- R₃ is selected from -H or -CH₃
- x is a natural number between 1-100
- y is a natural number between 1-100

15

- n is a natural number between 1-50 and
- L is a cleavable linker or L may be null and
- D is a therapeutic agent that is selected from a group comprising combretastatin, 5-Fluorouracil, gemcitabine, chloroquine and doxorubicine
- A is an end group or A may be null

20

- B is an end group or B may be null

The inventors have found that in addition to overcoming the disadvantages of the prior art polymeric assemblies; the polymeric assemblies of present invention that are made up of polymer-drug conjugates of formula I also reduces the amount of polymer given to a patient due to high drug loading.

5 DETAILED DESCRIPTION OF THE INVENTION

The term “polymer-drug conjugate” refers to a polymeric structure having a therapeutic agent covalently attached to the polymer.

The term “polymeric assembly” refers to a structure having a diameter in between 1-100 nm. Said assemblies are made of self-assembled polymer chains and they may or may not have 10 hollow cavities of various sizes.

The terms “polymeric backbone” and “polymer backbone” can be used interchangeably and refer to a polymer chain having side chains or pendant groups. For example, a side chain may have an oligo ethylene glycol unit and a pendant group may be bearing one therapeutic agent or any other group that can be utilized to attach a therapeutic and/or diagnostic agent or a targeting group.

15 Throughout the text, the term “the polymer-drug conjugate of the invention” should be construed to mean “a polymer-drug conjugate according to formula I” or “a polymer-drug conjugate of formula I” or “formula I” and these terms can be used interchangeably.

Throughout the text, the term “polymeric assemblies” should be construed to mean an 20 “nanoparticle made up of polymer-drug conjugate of formula I” or “nanoparticle made of a co-polymer as shown in formula I” or “nanoparticle that formed as a result of the self assembly of a polymer-drug conjugate of formula I” or “a micellar assembly made of a co-polymer as shown in formula I” and these terms can be used interchangeably.

The term “PEG” refers to a polyether compound having the structure of H-(O-CH₂-CH₂)_n-OR₃, n being a natural number between 1-200 and R₃ selected from H or -CH₃. PEG is defined as an 25 oligomer or polymer of ethylene oxide. The terms “PEG”, “polyethylene glycol”, “polyethylene oxide”, “PEO”, “polyoxyethylene” and “POE” refer to the same structure and may be used interchangeably within this text.

In an embodiment of the invention R_1 , R_2 and R_3 are independently selected from H or $-CH_3$

In one embodiment $R_1=H$, $R_2=H$, $R_3=H$ or $R_1=H$, $R_2=H$, $R_3= -CH_3$ or $R_1=H$, $R_2= -CH_3$, $R_3=H$ or $R_1=H$, $R_2= -CH_3$, $R_3= -CH_3$ or $R_1= -CH_3$, $R_2= H$, $R_3=H$ or $R_1= -CH_3$, $R_2= H$, $R_3= -CH_3$ or $R_1= -CH_3$, $R_2= -CH_3$, $R_3=H$ or $R_1= -CH_3$, $R_2= -CH_3$, $R_3= -CH_3$.

5 The term "block copolymer" refers to a copolymer wherein all of one type of monomer is grouped together and the all of the other type of monomers are grouped together. The polymer-drug conjugates of the invention are in the form of block copolymer. The fact that the polymer-drug conjugates of the invention are in the form of a block copolymer allows the formation of the polymeric assembly of the invention.

10 The therapeutic agent is attached to the polymer via a cleavable linker so that the therapeutic agent can be released, for example, under reducing conditions, oxidizing conditions or by hydrolysis of an ester, amide, hydrazide.

Said cleavable linker can be any hydrocarbon or substituted hydrocarbon based compound which is capable of dissociating under physiological conditions. In a preferred embodiment the linker 15 can be selected from compounds that are cleaved under the acidic conditions of the tumor (such as any C_1 - C_{10} substituted or unsubstituted and/or linear and/or cyclic hydrocarbon comprising an acetal or an ester functional group) or with the help of the overexpressed enzymes present in the intercellular or intracellular matrix of the tumor cells.

The linker can be any sort of entity capable of binding to both the polymer backbone and to the 20 drug, such as, a poly(ethylene glycol), an amino acid, poly(amino acid) (e.g. a peptide or oligopeptide), or polypeptide (e.g. a protein), such that one end of it is capable of forming a covalent bond with the polymer backbone and the other end of it is capable of forming a covalent bond with the therapeutic agent. The linkers may also include short peptides with specific peptide sequences that are cathepsin B labile, such as Gly-Phe-Leu-Gly (SEQ ID NO: 1) also 25 denoted as GFLG or Val-Cit or Phe-Lys or Val-Ala or Ala-Leu-Ala-Leu (SEQ ID NO: 2)

The linker can also be a C_1 - C_{10} hydrocarbon or a C_1 - C_{10} substituted or hetero substituted hydrocarbon such that it comprises a functional group that dissociates under physiological conditions, such as an acetal, ester, imine, amide, disulfide, carbonate, carbamate, hydrazone.

In an embodiment of the invention, the linker (L) is GFLG or Val-Cit or Phe-Lys or Val-Ala or Ala-Leu-Ala-Leu.

In an embodiment of the invention, the linker is a C₁-C₁₀ hetero substituted hydrocarbon comprising at least one disulfide functional group or a C₁-C₁₀ hetero substituted hydrocarbon comprising at least one acetal functional group or a C₁-C₁₀ hetero substituted hydrocarbon comprising at least one ester functional group or a C₁-C₁₀ hetero substituted hydrocarbon comprising at least one imine functional group or a C₁-C₁₀ hetero substituted hydrocarbon comprising at least one amide functional group or a C₁-C₁₀ hetero substituted hydrocarbon comprising at least one carbonate functional group or a C₁-C₁₀ hetero substituted hydrocarbon comprising at least one carbamate functional group or a C₁-C₁₀ hetero substituted hydrocarbon comprising at least one hydrazone functional group.

In another embodiment of the invention the linker may comprise a C₁-C₁₀ substituted or hetero substituted hydrocarbon comprising two or more functional groups selected from the group comprising acetal, ester, imine, amide, disulfide, carbonate, carbamate, hydrazone.

15 In another embodiment of the invention the linker may be null, which means that the therapeutic agent D is attached directly to the polymer chain. The direct attachment of the therapeutic agent to the polymer chain can be through an ester, imine, amide, disulfide, carbonate, carbamate, hydrazine bond.

20 In a preferred embodiment of the invention the drug molecule is attached to the polymer via a direct ester bond that dissociates under physiological conditions of the tumor. In another embodiment of the invention the linker may be a combination of a C₁-C₁₀ substituted hydrocarbon comprising at least one functional group selected from the group comprising acetal, ester, imine, amide, disulfide, carbonate, carbamate, hydrazone and a peptide chain selected from the group comprising GFLG, Val-Cit or Phe-Lys or Val-Ala or Ala-Leu-Ala-Leu. The 25 therapeutic agent selected from combretastatin, 5-Fluorouracil, gemcitabine, chloroquine and doxorubicine can be present in an amount in between 5% to 50% by weight of the polymer-drug conjugate, preferably in an amount between 6% to 48% by weight of the drug-polymer conjugate and most preferably in an amount between 10% to 45% by weight of the drug-polymer

conjugate. The therapeutic agent can be present in an amount in the range of for example; 10% to 48% or 15% to 46% or 20% to 45% or 25% to 44% by weight of the drug-polymer conjugate.

The term "end group" refers to functionalities or constitutional units that are at the extremity of a polymer. The end groups shown as A and B can be identical to or different from one another.

5 In an embodiment A is an end group that is optionally a fragment of a chain transfer agent or initiator or a fragment of a chain transfer agent or initiator conjugated with a targeting moiety or a fragment of a chain transfer agent or initiator having a reactive functional group.

In an embodiment B is an end group that is optionally a fragment of a chain transfer agent or initiator.

10 The term "chain transfer agent" refers to chemical compounds that are capable of regulating the molecular weight of polymers through interrupting the uncontrolled growth of the polymer chain. The chain transfer agents are well established in the literature and their structures are well known to those skilled in the art. The terms "chain transfer agent" and its abbreviation "CTA" refer to the same group of compounds and can be used interchangeably throughout the text.

15 A and B can optionally be a fragment of a chain transfer agent. Particularly when a CTA is used in the polymerization reaction, the CTA fragment may stay as an end group to the polymer. The CTA used herein, may be any material suitable for initiating the polymerization reaction known in the art. In one embodiment, A and/or B is selected from the group consisting of 3,5-Bis(2-dodecylthiocarbonothioylthio-1-oxopropoxy)benzoic acid, 3-Butenyl 2-

20 (dodecylthiocarbonothioylthio)-2-methylpropionate, 2-Cyanobutan-2-yl 4-chloro-3,5-dimethyl-1H-pyrazole-1-carbodithioate, 2-Cyanobutanyl-2-yl 3,5-dimethyl-1H-pyrazole-1-carbodithioate, 4-Cyano-4-[(dodecylsulfanylthiocarbonyl)sulfanyl]pentanoic acid, 4-Cyano-4-[(dodecylsulfanylthiocarbonyl)sulfanyl]pentanol, Cyanomethyl (3,5-Dimethyl-1H-pyrazole)-carbodithioate, Cyanomethyl dodecyl trithiocarbonate, Cyanomethyl [3-(trimethoxysilyl)propyl]

25 trithiocarbonate, 2-Cyano-2-propyl dodecyl trithiocarbonate, S,S-Dibenzyl trithiocarbonate, 2-(Dodecylthiocarbonothioylthio)-2-methylpropionic acid, 2-(Dodecylthiocarbonothioylthio)-2-methylpropionic acid 3-azido-1-propanol ester, 2-(Dodecylthiocarbonothioylthio)-2-methylpropionic acid N-hydroxysuccinimide ester, 2-(Dodecylthiocarbonothioylthio)-2-methylpropionic acid pentafluorophenyl ester, 2-(Dodecylthiocarbonothioylthio)propionic acid,

Methyl 2-(dodecylthiocarbonothioylthio)-2-methylpropionate, Pentaerythritol tetrakis[2-(dodecylthiocarbonothioylthio)-2-methylpropionate], Phthalimidomethyl butyl trithiocarbonate, 1,1,1-Tris[(dodecylthiocarbonothioylthio)-2-methylpropionate]ethane, Benzyl 1H-pyrrole-1-carbodithioate, Cyanomethyl diphenylcarbamodithioate, Cyanomethyl 5 methyl(phenyl)carbamodithioate, Cyanomethyl methyl(4-pyridyl)carbamodithioate, 2-Cyanopropan-2-yl N-methyl-N-(pyridin-4-yl)carbamodithioate, Methyl 2-[methyl(4-pyridinyl)carbamothioylthio]propionate, 1-Succinimidyl-4-cyano-4-[N-methyl-N-(4-pyridinyl)carbamothioylthio]pentanoate, Benzyl benzodithioate, Cyanomethyl benzodithioate, 4-Cyano-4-(phenylcarbonothioylthio)pentanoic acid, 4-Cyano-4-(phenylcarbonothioylthio)pentanoic acid N-succinimidyl ester, 2-Cyano-2-propyl benzodithioate, 2-Cyano-2-propyl 4-cyanobenzodithioate, Ethyl 2-(4-methoxyphenylcarbonothioylthio)acetate, Ethyl 2-methyl-2-(phenylthiocarbonylthio)propionate, Ethyl 2-(phenylcarbonothioylthio)-2-phenylacetate, Ethyl 2-(phenylcarbonothioylthio)propionate, 1-(Methoxycarbonyl)ethyl benzodithioate, 2-(4-Methoxyphenylcarbonothioylthio)ethanoic acid, 2-Nitro-5-(2-propynyl)benzyl 4-cyano-4-(phenylcarbonothioylthio)pentanoate 2-(Phenylcarbonothioylthio)propanoic acid, 2-Phenyl-2-propyl benzodithioate, Cyanomethyl methyl(4-pyridyl)carbamodithioate, 2-Cyanopropan-2-yl N-methyl-N-(pyridin-4-yl)carbamodithioate, Methyl 2-[methyl(4-pyridinyl)carbamothioylthio]propionate, 1-Succinimidyl-4-cyano-4-[N-methyl-N-(4-pyridinyl)carbamothioylthio]pentanoate or any fragment of the initiators listed herein as itself or as conjugated with a targeting moiety before or after the polymerization reaction or as modified with a reactive group.

The term "fragment" as used herein refers to compounds that form due to breaking of one or more of the covalent bonds forming the initiator molecule.

25 Fragmentation of the CTAs listed herein and the structure of the formed fragments are well established in the art, thus the structure of A and B can be determined without use of inventive skills by making use of current state of the art.

A and/or B is optionally null.

In an embodiment of the invention A is null and B is a fragment of a CTA.

In another embodiment B is null and A is a fragment of a CTA.

In another embodiment A and B are both fragments of a CTA however they are structurally different from one another. In other words, A and B are different fragments of the same CTA.

In another embodiment A and B are both fragments of a CTA and they have the same chemical
5 structure.

The term “targeting moiety” refers to molecules which has a tendency to bind to specific target sites in the body. In other words targeting moiety are molecules that bind specifically to the cell that has the complimentary receptor.

Targeting moiety can be selected from a group comprising antibodies; antibody fragments, or
10 peptides such as Cyclo (Arg-Gly-Asp-D-Phe-Lys) (SEQ ID No: 3) (cRGDfK). In a preferred embodiment cRGDfK is used as the targeting moiety.

The term “reactive group” refers to an atom or associated group of atoms in a chemical substance that is intended or can be reasonably anticipated to undergo facile chemical reaction.

The reactive group can be an acetal, , hemiacetal, carboxylic acid, alcohol, amide, imide, ,
15 anhydride, aryl halide, azo compound, diazo compound, hydrazine, azide, , carbonate, chlorosilane, cyanide, ester, sulfate ester, phosphate ester, thiophosphate ester, isocyanate, isothiocyanate, thiocarbamate ester, dithiocarbamate ester. Preferably the reactive group is an ester, imide or carbonate. The reactive group can for example be N-hydroxysuccinimide.

Depending on the polymerization techniques employed in preparation of the polymer conjugates
20 of the invention A and/or B can be a fragment of an initiator.

An initiator as used herein refers to a chemical compound that reacts with a monomer to form an intermediate compound capable of linking successively with a large number of other monomers into a polymeric compound. The terms “initiator” and “polymerization initiator” can be used interchangeably within the context of this application.

25 Depending on the polymerization techniques used for preparing the polymer conjugates of the invention, different initiator agents can be used.

In an embodiment A and/or B is a fragment of an initiator that can be selected from the group comprising 4,4'-Azobis(4-cyanovaleric acid), 4,4'-Azobis(4-cyanovaleric acid), 1,1'-Azobis(cyclohexanecarbonitrile), 2,2'-Azobis(2-methylpropionamide) dihydrochloride, 2,2'-Azobis(2-methylpropionitrile) (also known as AIBN), Ammonium persulfate, 5 hydroxymethanesulfinic acid monosodium salt dihydrate, potassium persulfate, sodium persulfate, tert-Butyl hydroperoxide, tert-Butyl peracetate, Cumenehydroperoxide, 2,5-Di(tert-butylperoxy)-2,5-dimethyl-3-hexyne, Dicumyl peroxide, 2,5-Bis(tert-butylperoxy)-2,5-dimethylhexane, 1,1-Bis(tert-butylperoxy)-3,3,5-trimethylcyclohexane, 1,1-Bis(tert-amylperoxy)cyclohexane, Benzoyl peroxide, 2-Butanone peroxide, tert-Butyl peroxide, Di-tert-10 amyl peroxide, Lauroyl peroxide, tert-Butyl peroxybenzoate, tert-Butylperoxy 2-ethylhexyl carbonate, tert-Butyl hydroperoxide, 2-Azidoethyl 2-bromoisobutyrate, Bis[2-(2-bromoisobutyryloxy)undecyl] disulfide, Bis[2-(2'-bromoisobutyryloxy)ethyl]disulfide, 2-Bromoisobutanoic acid N-hydroxysuccinimide ester, 2-Bromoisobutyric anhydride, α -Bromoisobutyryl bromide, 2-(2-Bromoisobutyryloxy)ethyl methacrylate, tert-Butyl α -bromoisobutyrate, 3-Butynyl 2-bromoisobutyrate, Dipentaerythritolhexakis(2-bromoisobutyrate), 15 Dodecyl 2-bromoisobutyrate, Ethyl α -bromoisobutyrate, Ethylene bis(2-bromoisobutyrate), 2-Hydroxyethyl 2-bromoisobutyrate, 1-(DL-1,2-Isopropylideneglycyl) 2-bromoisobutyrate, Methyl α -bromoisobutyrate, Octadecyl 2-bromoisobutyrate, Pentaerythritoltetrakis(2-bromoisobutyrate), 1-(Phthalimidomethyl) 2-bromoisobutyrate, Poly(ethylene glycol) bis(2-20 bromoisobutyrate), Propargyl 2-bromoisobutyrate, 1,1,1-Tris(2-bromoisobutyryloxymethyl)ethane, 10-Undecenyl 2-bromoisobutyrate, N-tert-Butyl-O-[1-[4-(chloromethyl)phenyl]ethyl]-N-(2-methyl-1-phenylpropyl)hydroxylamine, N-tert-Butyl-N-(2-methyl-1-phenylpropyl)-O-(1-phenylethyl)hydroxylamine, TEMPO, TEMPO methacrylate, 2,2,5-Trimethyl-4-phenyl-3-azahexane-3-nitroxide, 3,5-Bis(2-dodecylthiocarbonothiylthio-1-25 oxopropoxy)benzoic acid, 3-Butenyl 2-(dodecylthiocarbonothiylthio)-2-methylpropionate, 4-Cyano-4-[(dodecylsulfanylthiocarbonyl)sulfanyl]pentanoic acid, 4-Cyano-4-[(dodecylsulfanylthiocarbonyl)sulfanyl]pentanol, Cyanomethyl dodecyl, Cyanomethyl [3-(trimethoxysilyl)propyl] trithiocarbonate, 2-Cyano-2-propyl dodecyl trithiocarbonate, S,S-Dibenzyl trithiocarbonate, 2-(Dodecylthiocarbonothiylthio)-2-methylpropionic acid, 2-(Dodecylthiocarbonothiylthio)-2-methylpropionic acid, 3-azido-1-propanol ester, 2-(Dodecylthiocarbonothiylthio)-2-methylpropionic acid N-hydroxysuccinimide ester, 2-

(Dodecylthiocarbonothioylthio)-2-methylpropionic acid pentafluorophenyl ester, 2-(Dodecylthiocarbonothioylthio)propionic acid, Methyl 2-(dodecylthiocarbonothioylthio)-2-methylpropionate, Pentaerythritol tetrakis[2-(dodecylthiocarbonothioylthio)-2-methylpropionate], Phthalimidomethyl butyl trithiocarbonate, 1,1,1-5 Tris[(dodecylthiocarbonothioylthio)-2-methylpropionate]ethane, benzyl benzodithioate, Cyanomethyl benzodithioate, 4-Cyano-4-(phenylcarbonothioylthio)pentanoic acid, 4-Cyano-4-(phenylcarbonothioylthio)pentanoic acid N-succinimidyl ester, 2-Cyano-2-propyl benzodithioate, 2-Cyano-2-propyl 4-cyanobenzodithioate, Ethyl 2-(4-methoxyphenylcarbonothioylthio)acetate, Ethyl 2-methyl-2-(phenylthiocarbonylthio)propionate, 10 Ethyl 2-(phenylcarbonothioylthio)-2-phenylacetate, Ethyl 2-(phenylcarbonothioylthio)propionate, 1-(Methoxycarbonyl)ethyl benzodithioate, 2-(4-Methoxyphenylcarbonothioylthio)ethanoic acid, 2-Nitro-5-(2-propynyloxy)benzyl, 4-cyano-4-(phenylcarbonothioylthio)pentanoate, 2-(Phenylcarbonothioylthio)propanoic acid, 2-Phenyl-2-propyl benzodithioate, Cyanomethyl methyl(4-pyridyl)carbamodithioate, Cyanopropan-2-yl N-15 methyl-N-(pyridin-4-yl)carbamodithioate, Methyl 2-[methyl(4-pyridinyl)carbamothioylthio]propionate, 1-Succinimidyl-4-cyano-4-[N-methyl-N-(4-pyridinyl)carbamothioylthio]pentanoate or a combination thereof as itself or as **conjugated with a targeting moiety before or after the polymerization reaction or as modified with a reactive group.**

The definition and examples of targeting moiety are as described above.

20 The definition and examples of the reactive group are as described above.

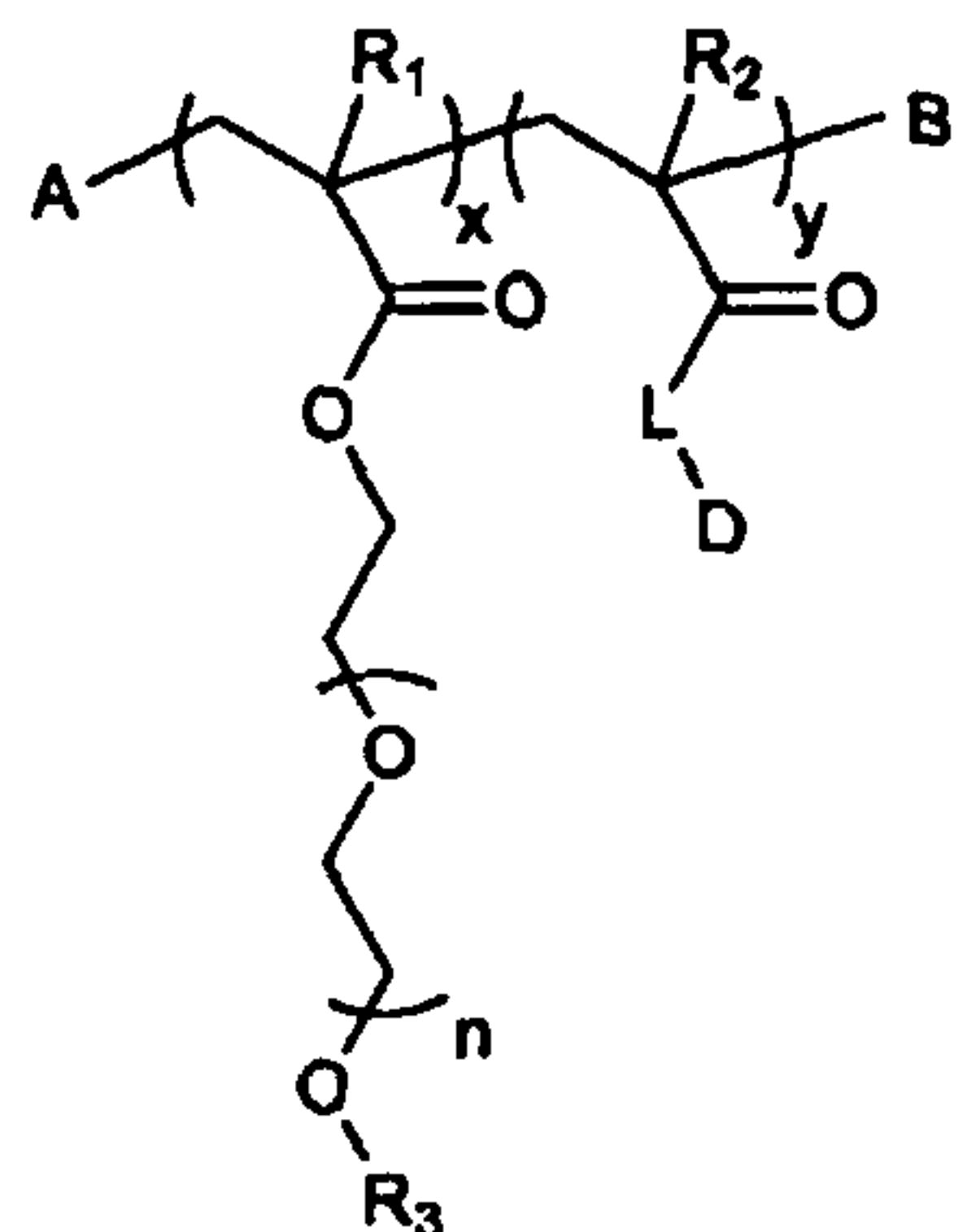
The chain transfer agents and the initiators listed within this application are given as example for illustration of the invention, any chemical compound published before or after the filing of this document that can act as a CTA and/or initiator are included within the scope of the invention.

25 The measurement of the amount of drug in the polymer conjugate of the invention is made by using conventional techniques well-known in the art, for example by calculation of the drug ratio from a ¹H-NMR of the polymer-drug conjugate or by determination of the amount by forced release of the drug.

In another embodiment the polymer-drug conjugate of the invention has an average molecular weight in between 5 kDa to 60 kDa. In a preferred embodiment the polymer-drug conjugate of the invention has an average molecular weight in between, 6 Da to 50 kDa and in a most preferred embodiment the polymer-drug conjugate of the invention has an average molecular weight in between 7 kDa to 40 kDa.

Molecular weight of the polymer-drug conjugate of the invention is determined by using conventional techniques known in the art for example by using gel permeation chromatography (GPC).

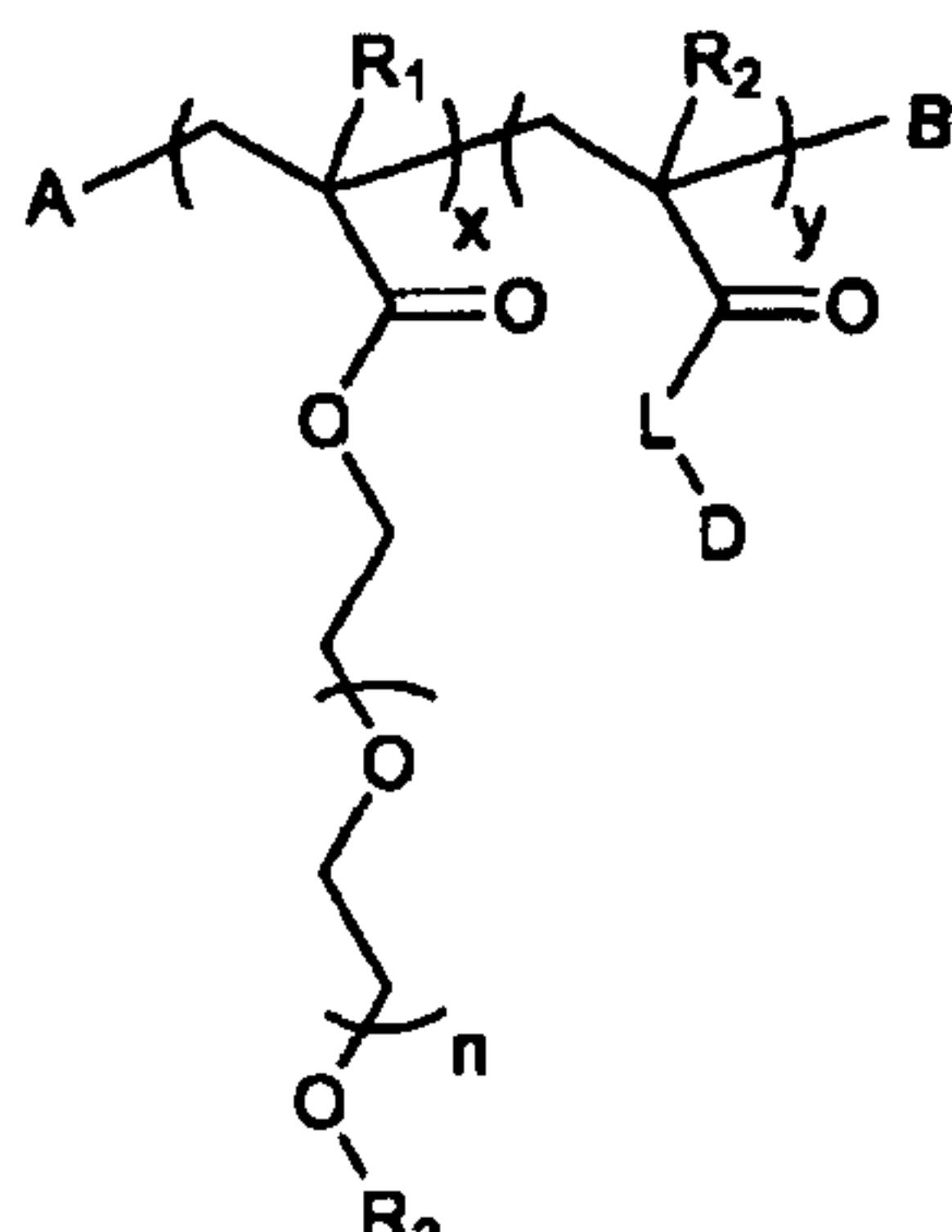
Another embodiment of the invention is polymeric assemblies formed with polymer-drug conjugates shown with formula I wherein;



Formula I

- R₁ and R₂ are independently selected from H or -CH₃
- R₃ is selected from -H or -CH₃
- x is a natural number between 1-100
- y is a natural number between 1-100
- n is a natural number between 1-50 and
- L is a cleavable linker or L may be null and
- D is a therapeutic agent that is combretastatin or 5-Fluorouracil or gemcitabine or chloroquine or doxorubicine
- A is an end group or A may be null
- B is an end group or B may be null

Another embodiment of the invention is polymeric micelles formed with polymer-drug conjugates shown with formula I wherein;



Formula I

- R₁ and R₂ are independently selected from H or -CH₃
- 5 • R₃ is selected from -H or -CH₃
- x is a natural number between 1-100
- y is a natural number between 1-100
- n is a natural number between 1-50 and
- L is a cleavable linker or L may be null and
- 10 • D is a therapeutic agent that is combretastatin or 5-Fluorouracil or gemcitabine or chloroquine or doxorubicine
- A is an end group or A may be null
- B is an end group or B may be null

15 In another embodiment, assemblies according to present invention are used to encapsulate therapeutic molecules, in that sense polymeric assemblies made up of polymer-drug conjugates of formula I encapsulating therapeutic agents other than those attached to the polymer chain are another embodiment of this invention.

The term "encapsulating" refers to confinement of a guest molecule, for example a therapeutic agent inside the cavity of a host molecule, for example a polymeric assembly made up of the polymer-drug conjugate of formula I. The encapsulation preferably takes place through non-covalent interactions of the therapeutic molecule with the polymeric assembly of the invention.

The term "therapeutic agent" refers to any compound that is suitable for use in treatment of a disease. The terms "therapeutic agent", "chemotherapy agent", "anticancer agent" and "antineoplastic agent" all refer to the compounds suitable for use in treatment of a disease and these terms can be used interchangeably. In one embodiment, the disease is cancer.

5 Additionally, a "therapeutic agent" also refers to any agent that is suitable for use in treating of a disease, for example cancer. Any therapeutic agent which can be directly or indirectly attached to the polymer-drug conjugate of the invention can be used. U.S. Patent Nos 6 6,342,221 also describe agents related to anticancer agents and this document is incorporated herein by reference. Anticancer agents can be classified as but are not limited to, chemotherapeutic agents, 10 cytotoxins, antimetabolites, alkylating agents, protein kinase inhibitors, anthracyclines, antibiotics, antimitotic agents (e.g. antitubulin agents), corticosteroids, radiopharmaceuticals, and proteins (e.g. cytokines, enzymes, or interferons). Specific examples of anticancer agents are for example, docetaxel, gemcitabine, imatinib, 5-fluorouracil, 9-aminocamptothecin, amine-modified geldanamycin, doxorubicin, paclitaxel, procarbazine, hydroxyurea, meso e-chlorin, 15 cisplatin and radionuclides (e.g 1-131, Y-90, In-111, and Tc-99m). There are many other anticancer agents known in the art and many continue to be developed, those agents are also included within the scope of this invention.

The therapeutic agent can also be selected from a sub group comprising, but not limited to, nucleoside analogs, antifolates, other metabolites, topoisomerase I inhibitors, anthracyclines, 20 podophyllotoxins, taxanes, vinca alkaloids, alkylating agents, platinates, antihormones, radiopharmaceutics, monoclonal antibodies, tyrosine kinase inhibitors, mammalian target of rapamycin (mTOR) inhibitors, retinoids, immunomodulatory agents, histone deacetylase inhibitors and other agents.

Nucleoside analogs can be selected from a group comprising, but not limited to, azacitidine, 25 cladribine, clofarabine, cytarabine, decitabine, floxuridine, fludarabine, gemcitabine, mercaptopurine, nelarabine, pentostatin, tioguanine, trifluridine, tipiracil.

Antifolates can be selected from a group comprising, but not limited to, methotrexate, pemetrexed, pralatrexed, raltitrexed.

Other metabolites can be selected from a group comprising, but not limited to, hydroxycarbamide.

Topoisomerase I inhibitors can be selected from a group comprising, but not limited to, irinotecan and topotecan.

5 Anthracyclines can be selected from a group comprising, but not limited to, daunorubicin, doxorubicin, epirubicin, idarubicin, mitoxantrone, valrubicin.

Podophyllotoxins can be selected from a group comprising, but not limited to, etoposide and teniposide.

10 Taxanes can be selected from a group comprising, but not limited to, cabazitaxel, docetaxel, paclitaxel.

Vinca alkaloids can be selected from a group comprising, but not limited to, vinblastine, vincristine, vindesine, vinflunine, vinorelbine.

Alkylating agents can be selected from a group comprising, but not limited to, bendamustine, chlorambucil, dacarbazine, melphalan, streptozotocin, trabectedin.

15 Antihormone compounds can be selected from a group comprising, but not limited to, abiraterone, bicalutamide, cyproterone, degarelix, exemestane, fulvestrant, goserelin, histrelin, leuprolide, mifepristone, triptorelin.

Tyrosine kinase inhibitors can be selected from a group comprising, but not limited to, afatinib, axitinib, bosutinib, cobimetinib, crizotinib, dasatinib, erlotinib, gefitinib, imatinib, lapatinib, 20 nilotinib, osimertinib, pazopanib, ruxolitinib, sunitinib, vandetanib.

Mammalian target of rapamycin (mTOR) inhibitors can be selected from a group comprising, but not limited to everolimus, temsirolimus.

Retinoids can be selected from a group comprising, but not limited to, alitretinoin, bexarotene, isotretinoin, tamibarotene, tretinoin.

Immunomodulatory agents can be selected from a group comprising, but not limited to, lenalidomide, pomalidomide, thalidomide.

Histone deacetylase inhibitors can be selected from a group comprising, but not limited to, belinostat, panobinostat, valproate, vorinostat.

5 Platinates can be selected from a group comprising, but not limited to, cisplatin, carboplatin, oxaliplatin, nedaplatin.

Other agents can be selected from a group comprising, but not limited to, anagrelide, ceritinib, dabrafenib, idelalisib, ibrutinib, palbociclib, vemurafenib, bleomycin, bortezomib, dactinomycin, eribulin, estramustine, ixabepilone, mitomycin, procarbazine, alectinib, fluxymesterone, 10 iobenguane, imiquimod, interferon, ixazomib, lanreotide, lentinan, octreotide, omacetaxine, tegafur, gimerazil, oteracil, uracil, combretastatin, chloroquine.

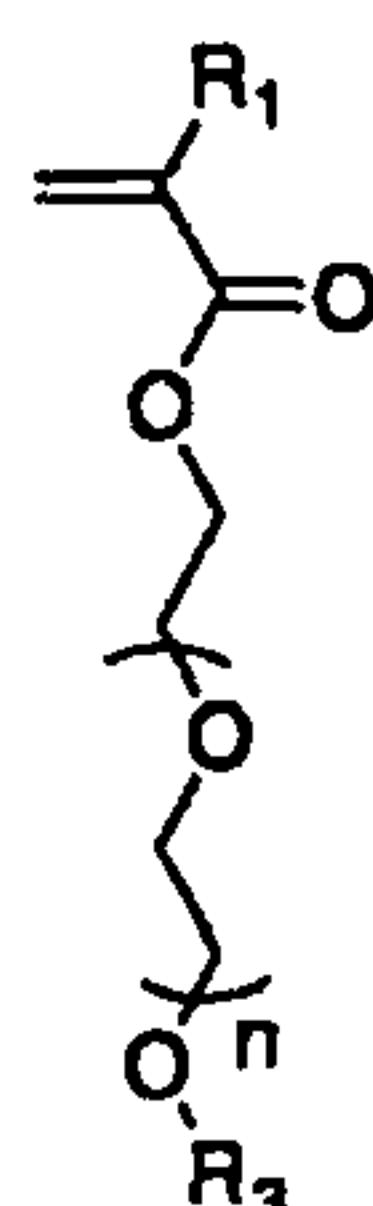
In a preferred embodiment of the invention the therapeutic agent is selected from taxanes, antifolates, tyrosine kinase inhibitors, anthracyclines, nucleoside analogs or other agents. Most preferably the therapeutic agent is selected from a group comprising docetaxel, pemetrexed, 15 chloroquine, combretastatin, gemcitabine, doxorubicine, Fluorouracil (5-FU), 5'-Deoxy 5-Fluorocytidine (5'-DFCR), lapatinib.

In an embodiment of the invention the therapeutic agent is docetaxel.

In an embodiment of the invention the therapeutic agent is carboplatin.

In an embodiment of the invention the therapeutic agent is doxorubicine.

20 Another embodiment of the invention is a method (Method I) for preparation of the polymer-drug conjugate of the invention (formula I) which comprises (i) polymerization of PEG (meth)acrylate monomer (formula II)



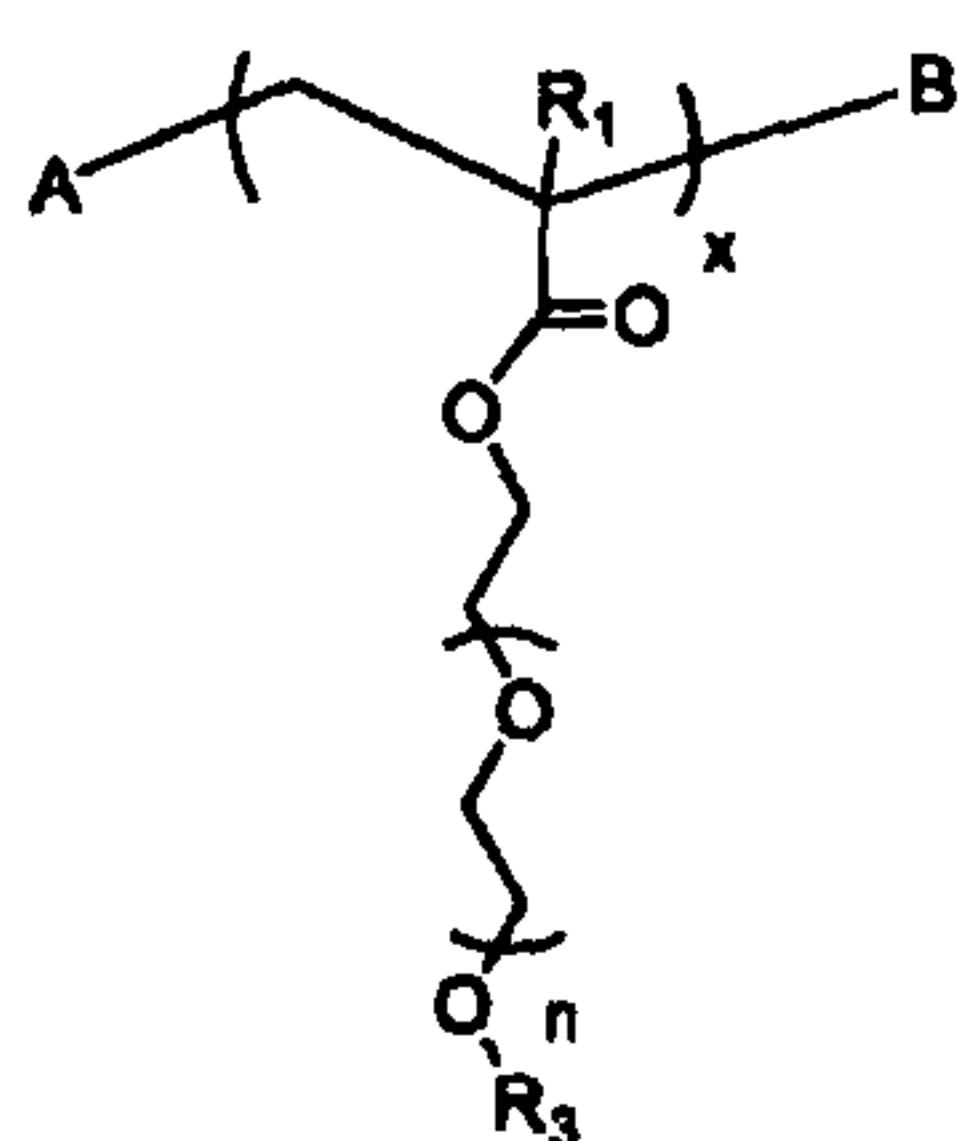
Formula II

wherein

- R1 is selected from H or -CH₃
- R3 is selected from -H or -CH₃

5 • n is a natural number between 1-50

to give a polymer of formula IIa



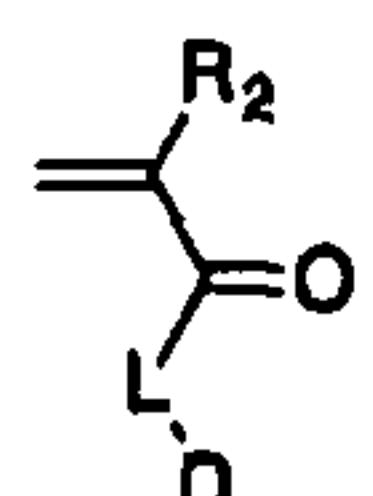
Formula IIa

wherein

10 • A is an end group or A may be null

 • B is an end group or B may be null

And then (ii) further reacting formula IIa with a (meth)acrylate-L-D monomer (Formula IIIa)

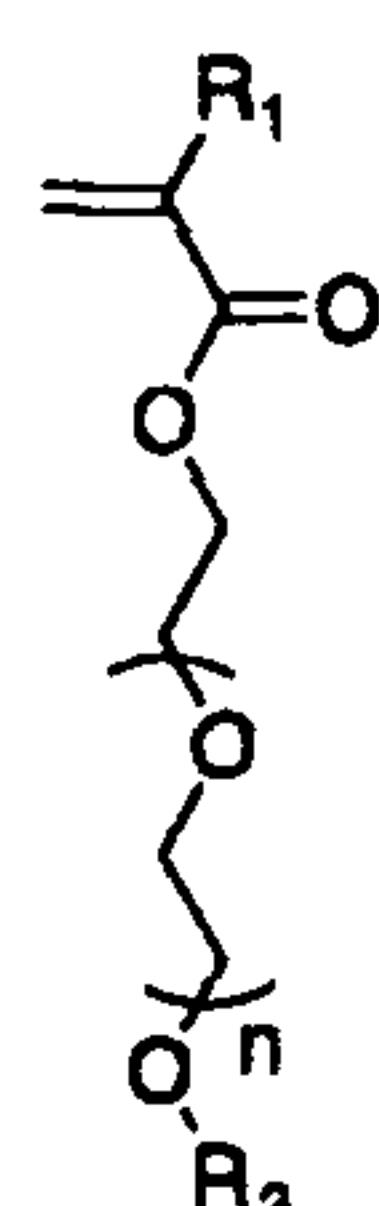


Formula IIIa

- wherein R_2 is selected from H or $-CH_3$
- L is a cleavable linker or L may be null
- D is a therapeutic agent selected from a group comprising combretastatin, 5-FU, gemcitabine, chloroquine, doxorubicine to give polymer-drug conjugate of formula I.

5

Another embodiment of the invention is a method (Method III) for preparation of the polymer-drug conjugate of the invention (formula I) which comprises (i) polymerization of PEG (meth)acrylate monomer (Formula II)

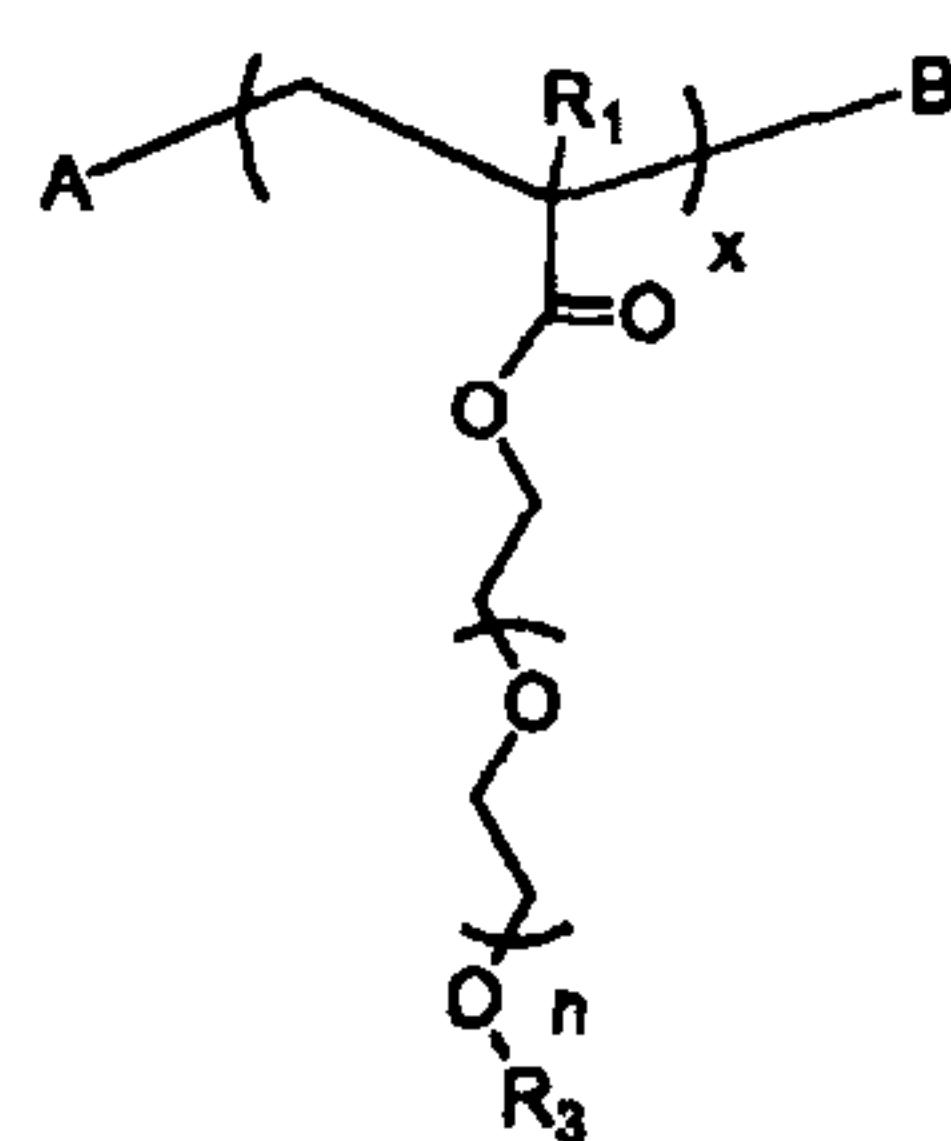


Formula II

10 wherein

- R_1 is selected from H or $-CH_3$
- R_3 is selected from $-H$ or $-CH_3$
- n is a natural number between 1-50

to give a polymer of formula IIa



15

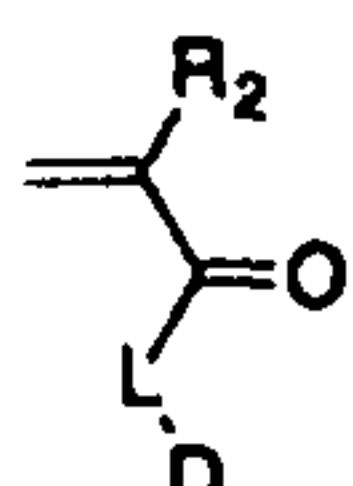
Formula IIa

18

wherein

- A is an end group or A may be null
- B is an end group or B may be null

And then (ii) further reacting formula IIa with a (meth)acrylate-L-D monomer (Formula IIIa)



5 Formula IIIa

- wherein R₂ is selected from H or -CH₃
- L is a cleavable linker or L may be null
- D is a therapeutic agent selected from a group comprising combretastatin, 5-FU, gemcitabine, chloroquine, doxorubicine to give a block copolymer

10 And then (iii) reacting formed block co-polymer with a targeting moiety to give a polymer-drug conjugate shown in formula I.

The term “acrylate” refers to derivatives of acrylic acids. These derivatives include the parent acid (CH₂CHCO₂H) and esters, thus the term “acrylate based” defines functional groups having any of the abovementioned acrylate derivatives.

15 The term “methacrylate” refers to derivatives of methacrylic acids. These derivatives include the parent acid (CH₂C(CH₃)CO₂H) and esters. Thus the term “methacrylate based” defines functional groups having any of the abovementioned methacrylate derivatives.

20 The term “(meth)acrylate” refers to the terms “acrylate” and “methacrylate”. Thus, the term “(meth)acrylate” can be used interchangeably with “acrylate” and “methacrylate” and comprises all features of these terms as described above. The term “(meth)acrylate” should be construed to mean “methacrylate and/or acrylate”

Step (i) of methods I, II and III may further comprise use of a chain transfer agent and/or an initiator.

25 In another aspect, the invention relates to polymer-drug conjugate of formula I prepared by any one of the methods I, II or III.

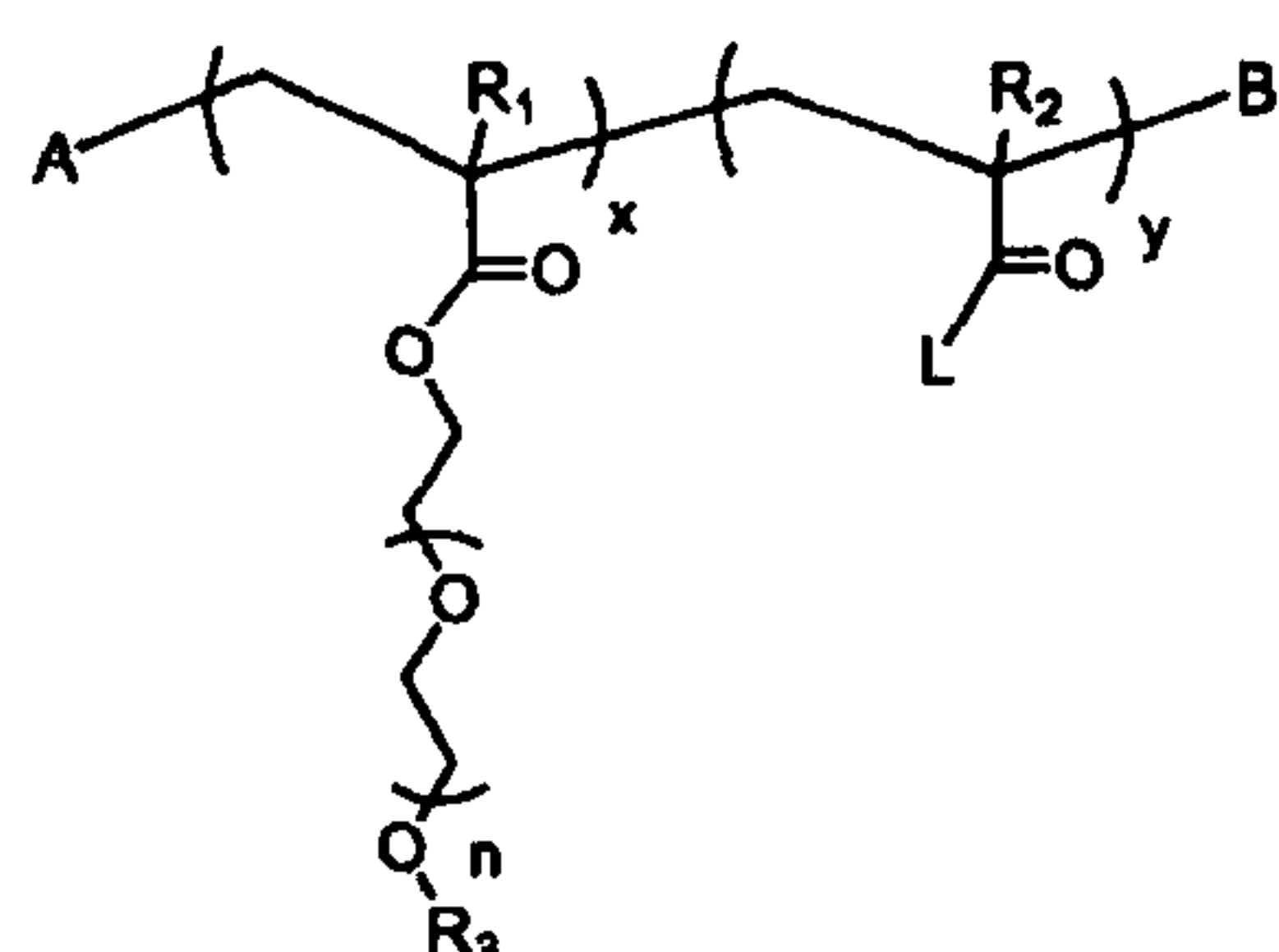
In another embodiment, invention relates to polymer-drug conjugates of formula I prepared by method I as this method provides a polymer-drug conjugate with a highly well defined polymer structure and drug content. Also, polymer-drug conjugates prepared with this method do not have free reactive groups on the side chain which remain unconjugated to a drug molecule.

5 In one embodiment, PEG (meth)acrylate (Formula II) is preferably selected from a group comprising; polyethylene glycol methyl ether methacrylate (CAS No: 26915-72-0), polyethylene glycol methacrylate (CAS No: 25736-86-1), polyethylene glycol methyl ether acrylate (CAS No: 32171-39-4), and poly ethylene glycol acrylate (CAS No: 9051-31-4), see table 1 for structures of the compounds. In a preferred embodiment of the invention, polyethylene glycol methyl ether 10 methacrylate, wherein R_1 and R_3 are both- CH_3 , is used.

In an embodiment of the invention PEG (meth)acrylate (Formula II) has an average molecular weight in between 50-2000 g/mol is used. In a preferred embodiment PEG (meth)acrylate (Formula II) has an average molecular weight in between 100-1500 g/mol and in a most preferred embodiment PEG (meth)acrylate (Formula II) has an average molecular weight in 15 between 150-1000 g/mol. PEG (meth)acrylate (Formula II) of the invention can have an average molecular weight of for example 60, 70, 80, 80, 100, 150, 200, to 250, 300, 400, 500, 600, 700, 800, 900, 1000 g/mol.

The cleavable linker according to present invention can be obtained from commercial sources or may be prepared according to known methods provided in literature.

20 In another aspect, present invention relates to a block copolymer of formula IIb for use in synthesis of polymer drug conjugate of formula I



Formula IIb

wherein;

- x is a natural number between 1-100,
- y is a natural number between 1-100,
- R₁ and R₂ are independently selected from H or -CH₃,
- 5 - R₃ is selected from -H or -CH₃
- n is a natural number between 1-50 and,
- L is a cleavable linker or L may be null
- A is an end group or A may be null
- B is an end group or B may be null.

10

As disclosed above, methods for preparing the polymer drug conjugate of formula I (Method I, II and III) comprise at least two polymerization steps, wherein monomers formula II and monomers of formula IIIa or IIIb are polymerized. In an embodiment of the invention, these polymerization steps may further comprise an initiator and/or a polymerization agent and/or a solvent.

15 The polymer-drug conjugate of the invention can be prepared by any of the known polymerization methods. Any suitable initiators and/or catalysts known in the art can be used for the preparation of the polymer-drug conjugate of the present invention. Where a polymerization initiator is used, the initiator or a fragment thereof may be present in the resulting polymer-drug conjugate.

20 The polymer backbone of the polymer-drug conjugate of the invention can be obtained by for example bulk polymerization, solution polymerization and/or suspension polymerization techniques known in the art.

25 The polymerization technique used for the preparation of the polymer-drug conjugate of the present invention may propagate through free-radical polymerization or controlled/living free radical polymerization. Herein the term "controlled/living free radical polymerization" refers to atom transfer radical polymerization (ATRP), Reversible addition fragmentation chain transfer (RAFT) polymerization, iodine transfer polymerization (ITP), selenium centered radical mediated polymerization, telluride mediated polymerization (TERP), nitroxide mediated

polymerization (NMP). In a preferred embodiment of the invention RAFT polymerization is used to prepare the polymer-drug conjugates of the invention.

A suitable polymerization initiator can be selected from the group given in this document.

In case of employing RAFT polymerization a CTA and an initiator can be used together for 5 preparing the polymer durg conjugates of the invention according to the Methods I, II or III provided herein.

In an embodiment of the invention the polymer-drug conjugate of the invention and polymeric assemblies made from them can further comprise targeting groups. Herein the term "targeting group" refers to tumor specific ligands that bind specifically to the cell, preferably the tumor cell 10 that has a complimentary receptor.

The term "targeting group" means a molecule which serves to deliver the polymer-drug conjugate of the invention to a specific site for the desired activity, i.e. it provides localization of the compound. The localization is mediated by specific recognition of molecular determinants, molecular size of the targeting agent or conjugate, ionic interactions, hydrophobic interactions, 15 and the like. Other mechanisms of targeting an agent to a particular tissue or region are known to those of skill in the art. Targeting ligands include, for example, molecules that bind to molecules on a targeted cell surface. Exemplary targeting ligands include antibodies, antibody fragments, small organic molecules, peptides, peptoids, proteins, polypeptides, oligosaccharides, transferrin, HS- glycoprotein, coagulation factors, serum proteins, beta-glycoprotein, G-CSF, GM-CSF, M- 20 CSF, EPO, and the like. In exemplary embodiments of the present invention, the targeting system includes covalently attaching a targeting ligand such as RGDfK, EPPT1 peptide, bisphosphonic acid or folate to the carrier molecule or linker.

In certain embodiments, the present invention is characterized by polymer-drug conjugates with or without a targeting ligand. In some embodiments the targeting ligand can be RGDfK, EPPT1, 25 bisphosphonic acid or folate.

Another embodiment of the present invention provides a method for delivering a therapeutic agent, comprising administering to a subject an effective amount of polymer-drug conjugate shown with formula I.

Another embodiment of the present invention provides a method for delivering a therapeutic agent, comprising administering to a subject an effective amount of a polymeric assembly made up of polymer-drug conjugate shown with formula I.

Another embodiment of the invention is the polymer-drug conjugate of the invention for use in
5 treatment of a variety of disorders that require the delivery of anticancer or similar agents.

Another embodiment of the invention is the polymeric assemblies made up of polymer-drug conjugates of the invention for use in treatment of a variety of disorders that require the delivery of anticancer or similar agents.

In a preferred embodiment, the invention is related to a polymer-drug conjugate shown with
10 formula I for use as a medicament for treatment of cancer.

In a yet another preferred embodiment, the invention is related to a polymeric assembly made up of polymer-drug conjugate of formula I for use as a medicament for treatment of cancer.

As used herein, "treat" or "treating" means to inhibit, reduce, modulate, ameliorate, or block at least one symptom that characterizes a pathologic condition, in a subject threatened by, or 15 afflicted with, the condition. A non-limiting list of different types of cancers is as follows: carcinomas, carcinomas of solid tissues, squamous cell carcinomas, adenocarcinomas, sarcomas, gliomas, high grade gliomas, blastomas, neuroblastomas, plasmacytomas, histiocytomas, melanomas, adenomas, hypoxic tumours, myelomas, metastatic cancers, or cancers in general. Specific examples of cancers that the disclosed compositions can be used to treat include B 20 cell lymphoma, T cell lymphoma, mycosis fungoides, Hodgkin's Disease, bladder cancer, brain cancer, nervous system cancer, head and neck cancer, squamous cell carcinoma of head and neck, kidney cancer, lung cancers such as small cell lung cancer and non-small cell lung cancer, neuroblastoma/glioblastoma, ovarian cancer, pancreatic cancer, prostate cancer, skin cancer, liver cancer, melanoma, squamous cell carcinomas of the mouth, throat, larynx, colon cancer, 25 cervical cancer, cervical carcinoma, breast cancer, and epithelial cancer, renal cancer, genitourinary cancer, pulmonary cancer, esophageal carcinoma, head and neck carcinoma, large bowel cancer, hematopoietic cancers; testicular cancer; colon and rectal cancers, prostatic cancer, or pancreatic cancer.

Polymer-drug conjugates of invention and/or polymeric assemblies made up of the polymer-drug conjugates of the invention may also be used for the treatment of precancer conditions such as cervical and anal dysplasias, other dysplasias, severe dysplasias, hyperplasias, atypical hyperplasias, and neoplasias.

5 The terms "cancer" and "cancerous" as used herein refers to malignant tumors or describe the physiological condition characterized by unregulated cell growth.

As discussed herein, polymer-drug conjugates of invention and/or polymeric assemblies made up of the polymer-drug conjugates of the present invention find use in the treatment and/or prophylaxis of cancer. For such use the polymer-drug conjugate of invention and/or polymeric 10 assemblies made up of the polymer-drug conjugates of the present invention will generally be administered in the form of a pharmaceutical composition.

Thus, according to the invention there is provided a pharmaceutical composition comprising a polymer-drug conjugate according to formula I and at least one pharmaceutically acceptable diluent, excipient and/or carrier. The term "treatment" includes either therapeutic or prophylactic 15 therapy.

Moreover, according to the invention there is provided a pharmaceutical composition comprising a polymeric assembly made up of a polymer-drug conjugate according to formula I and at least one pharmaceutically acceptable diluent, excipient and/or carrier.

The composition comprising the polymer-drug conjugate of the invention and/or polymeric 20 assemblies made up of the polymer-drug conjugates of the present invention may be in any suitable form depending upon the desired method of administering it to a patient. The composition comprising polymer drug conjugates of the invention and/or polymeric assemblies made up of the polymer-drug conjugates of the invention can be formulated to be administered orally, e.g. in the form of liquid dispersions or aqueous or oily suspensions or they can 25 formulated for parenteral administration, for example for subcutaneous, intravenous, intramuscular, intrasternal, intraperitoneal, intradermal, transdermal or other infusion techniques. The composition comprising the polymer drug conjugates of the invention and/or polymeric assemblies made up of the polymer-drug conjugates of the present invention can also be formulated for administration by inhalation in form of an aerosol or solution for administration

with and inhaler or nebulizer. The polymer-drug conjugates of the invention and/or polymeric assemblies made up of the polymer-drug conjugates of the invention are preferably administered to a subject transdermally, subcutaneously, intranasally, intravenously, intramuscularly, intratumorally or via inhalation. The most suitable route for administration in any given case will 5 depend on the particular therapeutic agent present in the polymer-drug conjugate of the present invention, the subject, and the nature and severity of the disease and the physical condition of the subject.

The polymer-drug conjugates of the invention and/or polymeric assemblies made up of the polymer-drug conjugates of the invention may be administered in combination, e.g. 10 simultaneously, sequentially or separately, with one or more other therapeutically active compounds, which may be an anti-cancer agent or it is an immunomodulatory, antiviral, antiinfective, antimicrobial, antiinfective or anesthetic agent or combinations thereof.

Polymeric assemblies of the invention may be encapsulating a second therapeutic agent selected 15 from the list provided above and additionally may further be administered in combination, e.g. simultaneously, sequentially or separately, with one or more other therapeutically active compounds, which may be an anti-cancer agent or it is an immunomodulatory, antiviral, antiinfective, antimicrobial, antiinfective or anesthetic agent or combinations thereof.

Said second therapeutic agent can be selected from the therapeutic agents listed above on the condition that it is different from the one present in the polymer-drug conjugate of the invention. 20 Comprising in the context of the present specification is intended to mean including.

Where technically appropriate, embodiments of the invention may be combined.

Embodiments are described herein as comprising certain features/elements. The disclosure also extends to separate embodiments consisting or consisting essentially of said features/elements.

Technical references such as patents and applications are incorporated herein by reference.

25 Any embodiments specifically and explicitly recited herein may form the basis of a disclaimer either alone or in combination with one or more further embodiments.

The invention will now be described with reference to the following examples, which are merely illustrative and should not in any way be construed as limiting the scope of the present invention.

EXAMPLES

Below examples provide step-by-step preparation of the polymer-drug conjugates of formula 1 and polymeric assemblies thereof.

Example 1A: Synthesis of Polymerizable Combretastatin-A4 Monomer (CombMA)

5 Combretastatin-A4 (300 mg, 0.95 mmol), triethyl amine (TEA, 191 mg, 1.89 mmol), methacryloyl chloride (198 mg, 1.89 mmol) were dissolved in dry dichloromethane (DCM, 10 mL) under N₂ in a 25 mL round bottom flask. Reaction was stirred for 16 h at room temperature. The crude was extracted with saturated NaHCO₃ (20 mL x 2) and distilled water (20 mL x 2). Organic layer was dried with Na₂SO₄ and the solvent was evaporated. The CombMA monomer
10 was purified using column chromatography on silica with hexanes.

Example 1B: Preparation of chain transfer agent (CTA) having a reactive functional group

4-cyanopentanoic acid dithiopentanoate (CPDB) is a CTA and this CTA was modified with N-hydroxysuccinimide (NHS) which is a reactive functional group in accordance with the below procedure;

15 Briefly, CPADB (200 mg, 0.72 mmol) and N-hydroxysuccinimide (125 mg, 1.07 mmol) were dissolved in anhydrous DCM (4 mL). Dicyclohexylcarbodiimide (DCC) (177 mg, 0.86 mmol) was dissolved in anhydrous DCM (1 mL). Two solution mixtures were then mixed and the reaction mixture was stirred at room temperature in the dark for 16 h. The insoluble white by-product dicyclohexylurea (DCU) was removed by filtration. Obtained solution was dried under
20 vacuo and the crude product was purified by column chromatography on silica with hexane and EtOAc.

Example 1C: Synthesis of POEGMEMA and NHS-POEGMEMA Homopolymers

Reversible addition-fragmentation chain transfer (RAFT) polymerization was used for the synthesis of POEGMEMA and NHS activated POEGMEMA (NHS-POEGMEMA) homopolymers. For the synthesis of POEGMEMA polymer, to a solution of OEGMA (600 mg, 2.0 mmol) and CPADB (20.12 mg, 0.072 mmol) in DMF (3 mL), was added AIBN (1.31 mg, 0.008 mmol). The mixture was purged with N₂ to remove O₂ and polymerization was stirred for at 70 °C. Polymerization was stopped by cooling and air exposure. POEGMEMA polymer was

purified by precipitation in diethyl ether. The polymer precipitate was dried under vacuo to give approximately 35 OEGMEMA repeating units (460 mg, 77 % yield). For the synthesis of NHS-POEGMEMA homopolymer same procedure was applied using SCPDB (27.10 mg, 0.072 mmol) as chain transfer agent to give approximately 35 OEGMEMA repeating units (480 mg, 80 %
5 yield)

Example 1D: Synthesis of POEGMEMA-co-CombMA and NHS-POEGMEMA-co-CombMA Block Copolymers

RAFT polymerization was used for the synthesis of POEGMEMA-*b*-PCombMA and NHS-POEGMEMA-*b*-PCombMA block copolymers. To a solution of CombMA (100 mg, 0.26 mmol)
10 and POEGMEMA polymer as macro chain transfer agent (50 mg, 0.005 mmol) in DMF (0.75 mL), was added AIBN (0.15 mg, 0.00092 mmol). The mixture was purged with N₂ to remove O₂ and polymerization was stirred at 65 °C. Polymerization was stopped by cooling and air exposure. The crude was precipitated in diethyl ether. The polymer precipitate was dried under vacuo to give approximately 32 CombMA repeating units (98 mg, 65 % yield). Same procedure
15 was applied for the synthesis of NHS-POEGMEMA-*b*-PCombMA block copolymer using NHS-POEGMEMA (50 mg, 0.005 mmol) as macro chain transfer agent to give approximately 33 CombMA repeating units (105 mg, 70 % yield).

Example 1E: Synthesis of Targeted Block Copolymer

NHS-POEGMEMA-*b*-PCombMA (50 mg, 0.002 mmol) and cRGDfK (6.4 mg, 0.01 mmol)
20 were co-dissolved in DMF (0.25 mL) and *N,N*-Diisopropylethylamine (6.5 mg, 0.05 mmol) was added to this reaction mixture. The reaction mixture was stirred at 30 °C for 24 h. The crude was precipitated in diethyl ether. The polymer precipitate was dried under reduced pressure to give cRGDfK-POEGMEMA-*b*-CombMA (41 mg, 82 % yield). An overall scheme showing the preparation of the cRGDfK-POEGMEMA-*b*-CombMA Block Copolymer is provided as Figure
25 1.

Example 1F: Preparation of Targeted and Non-Targeted Polymeric Assemblies

Herein “targeted” refers to polymeric assemblies made up of cRGDfK-POEGMEMA-*b*-CombMA block copolymers and “non-targeted” refers to polymeric assemblies made up of POEGMEMA-*b*-PCombMA block copolymers.

Two solutions were prepared to obtain targeted and non-targeted assemblies. For the formation 5 of non-targeted assemblies 2.3 mg POEGMEMA-*b*-PCombMA block copolymer was dissolved in 500 μ L THF in a glass vial and 3 mL water was added to this solution. THF in the vials was then evaporated at room temperature for 24 h. at open atmosphere to give micelles. Targeted assemblies were prepared in the same manner using cRGDfK-POEGMEMA-*b*-PCombMA and POEGMEMA-*b*-PCombMA mixture (1/5, w/w). A schematic representation of the preparation 10 of non-targeted and targeted assemblies are provided in Figure 2 and Figure 3.

Example 1G: Critical Micelle Concentration (CMC) Measurements

The fluorescence probe method was utilized for the determination of CMC values of targeted and non-targeted assemblies. Block copolymer solutions (450 μ L) were prepared in THF using serial dilution in vials as above. 50 μ L solution of Nile Red in THF (0.03 mg/mL) and then 3 mL water 15 was added to each vial. THF was evaporated completely at open atmosphere and final concentration values for 16 samples ranging between 1×10^{-9} and 1×10^{-5} M in 3 mL water were obtained. Fluorescence measurements were recorded by fluorescence spectrophotometer at an excitation wavelength of 550 nm and the emission was monitored from 580 to 660 nm.

For the determination of the minimum required concentration of the amphiphilic polymers to 20 form the micelle type polymeric structures via self-assembly, the CMC values of the block copolymers were calculated by tracking the fluorescence intensity of Nile Red as function of polymer concentration.

Fluorescence emission spectra of Nile Red in non-targeted and targeted assemblies were obtained at an excitation wavelength of 550 nm and the emission monitored from 580 to 660 nm 25 (Figure 4 and Figure 5 respectively)

For the determination CMC values, plots of the emission intensity at 612 nm versus the log of copolymer concentration were obtained. The CMC values for the non-targeted and targeted assembly formation were calculated by the intersection of the trendlines belonging to intensity

ratios with relatively constant values and the rapidly increased intensity ratios (Figure 6 and Figure 7 respectively). The CMC values for non-targeted and targeted assembly formation was found to be 1.796×10^{-6} M and 1.566×10^{-6} M, respectively.

Example 2A: Synthesis of Polymerizable 5FU Monomer (5FU-MA)

5 5-fluorouracil (250 mg, 1.92 mmol) and 37% formalin (244 μ L) was reacted at 60 °C for 2h. Then the mixture was cooled down and lyophilized completely to yield a viscous compound. As the second step, this product (315 mg, 1.97 mmol) was dissolved in anhydrous acetonitrile (5.8 mL) and then triethylamine (422 μ L, 3.03 mmol) was added into this solution. After cooling the reaction mixture down to 0 °C, methacryloyl chloride (232 μ L, 2.38 mmol) was added dropwise.

10 The reaction was let to run for overnight at room temperature and then white precipitate was filtered out. All volatiles were evaporated under vacuo and then the crude was dissolved with dichloromethane (20 mL), followed by extraction with 1M HCl (2x10 mL), 1M NaHCO₃ (10 mL) and brine (10 mL). After concentrating the organic part, product was isolated in pure form by column chromatography on silica, eluting with ethylacetate: hexane mixture (30:70 v/v). The

15 product was obtained as a viscous colorless oil (305 mg, 68% yield).

Example 2B: Synthesis of POEGMEMA Homopolymer

Reversible addition-fragmentation chain transfer (RAFT) polymerization was used for the synthesis of POEGMEMA homopolymer. For the synthesis of POEGMEMA polymer, to a solution of OEGMA (500 mg, 1.66 mmol) and CPADB (11.97 mg, 42.85 μ mol) in DMF (2.5 mL), was added AIBN (0.78 mg, 4.76 μ mol). The mixture was purged with N₂ to remove O₂ and polymerization was stirred for at 70 °C for 16h. Polymerization was stopped by cooling and air exposure. POEGMEMA polymer was purified by precipitation in diethyl ether. The polymer precipitate was dried under vacuo to give 95 mg polymer with 19 % yield.

Example 2C: Synthesis of POEGMEMA-co- 5FU-MA Block Copolymer

25 RAFT polymerization was used for the synthesis of POEGMEMA-b-P5FU- MA block copolymer. To a solution of 5FU-MA (17.9 mg, 78.6 μ mol) and POEGMEMA polymer as macro chain transfer agent (40 mg, 3.93 μ mol) in DMF (0.30 mL), was added AIBN (0.13 mg, 0.786 μ mol). The mixture was purged with N₂ to remove O₂ and polymerization was stirred at 70

°C for 16h. Polymerization was stopped by cooling and air exposure. The crude was precipitated in diethyl ether. The polymer precipitate was dried under vacuo to give 16 mg polymer with 28 % yield).

Example 3A: Synthesis of Polymerizable Combretastatin-A4 Monomer (Comb-ss-MA)

5 2-hydroxethyl methacrylate (500 mg, 3.8 mmol), 4,4'-dithiodibutyric acid (1.83 g, 7.6 mmol), N,N'-dicyclohexylcarbodiimide (1.59 mg, 7.8 mmol), 4-(Dimethylamino)pyridine (281 mg, 2.3 mmol) were dissolved in anhydrous dichloromethane (DCM) (54 mL) under N₂ in a 100 mL round bottom flask. Reaction was stirred for 16 h at room temperature. The crude was precipitated in cold diethyl ether and was left for 20 min under -20 °C. The product was purified
10 by using column chromatography on silica with ethyl acetate and hexane (20:80 v/v) (426 mg, 32 % yield).

Then, the obtained product (300 mg, 0.85 mmol) and Combretastatin-A4 (270 mg, 0.85 mmol) were dissolved in anhydrous dichloromethane (DCM) (6 mL) with DIPC (129 mg, 1.02 mmol) and DMAP (104 mg, 0.85 mmol) under N₂ in a 25 mL round bottom flask. Reaction was stirred
15 for 16 h at room temperature. The Comb-ss-MA monomer was purified using column chromatography on silica with ethyl acetate and hexane (30:70 v/v). (195 mg, 35 % yield).

Example 3B: Synthesis of POEGMEMA and Comb-ss-MA-POEGMEMA Homopolymers

Reversible addition-fragmentation chain transfer (RAFT) polymerization was used for the synthesis of POEGMEMA. For the synthesis of POEGMEMA polymer, to a solution of
20 OEGMA (1.0 g, 3.3 mmol) and CPADB (36.88 mg, 0.133 mmol) in DMF (5 mL), was added AIBN (2.17 mg, 0.0013 mmol). The mixture was purged with N₂ to remove O₂ and polymerization was stirred for 16h at 70°C. Polymerization was stopped by cooling and air exposure. POEGMEMA polymer was purified by precipitation in diethyl ether. The polymer precipitate was dried under vacuum to give 715 mg polymer with 69 % yield.

25 Example 3C: Synthesis of POEGMEMA-co-Comb-ss-MA Block Copolymer

RAFT polymerization was used for the synthesis of POEGMEMA-*b*-PComb-ss-MA block copolymer. Comb-ss-MA (80 mg, 0.20 mmol) and POEGMEMA polymer as macro chain

transfer agent (134 mg, 0.020 mmol) in DMF (1.07 mL), was added AIBN (0.68 mg, 0.004 mmol). The mixture was purged with N₂ to remove O₂ and polymerization was stirred at 65 °C. Polymerization was stopped by cooling and air exposure. The crude was precipitated in diethyl ether. The polymer precipitate was dried under vacuum to give 98 mg of polymer with 65 % yield.

The examples given above show that the invention is applicable to a range of different linkers such as disulfide and acetal and to different drug molecules such as combretastagtin and 5-FU.

Brief Description of Drawings:

Figure 1: Schematic representation showing synthesis of NHS-modified polymer-drug conjugate (polymer drug conjugate of formula I having a reactive functional group) and cRGDfK modified polymer-drug conjugate (polymer drug conjugate of formula I having a targeting group)

Figure 2: Schematic representation showing preparation of non-targeted polymeric assemblies of invention from polymer drug conjugates of formula I.

Figure 3: Schematic representation showing preparation of targeted polymeric assemblies of invention from polymer-drug conjugates of formula I.

Figure 4: Fluorescence emission spectra of Nile Red in non-targeted polymeric assemblies of invention.

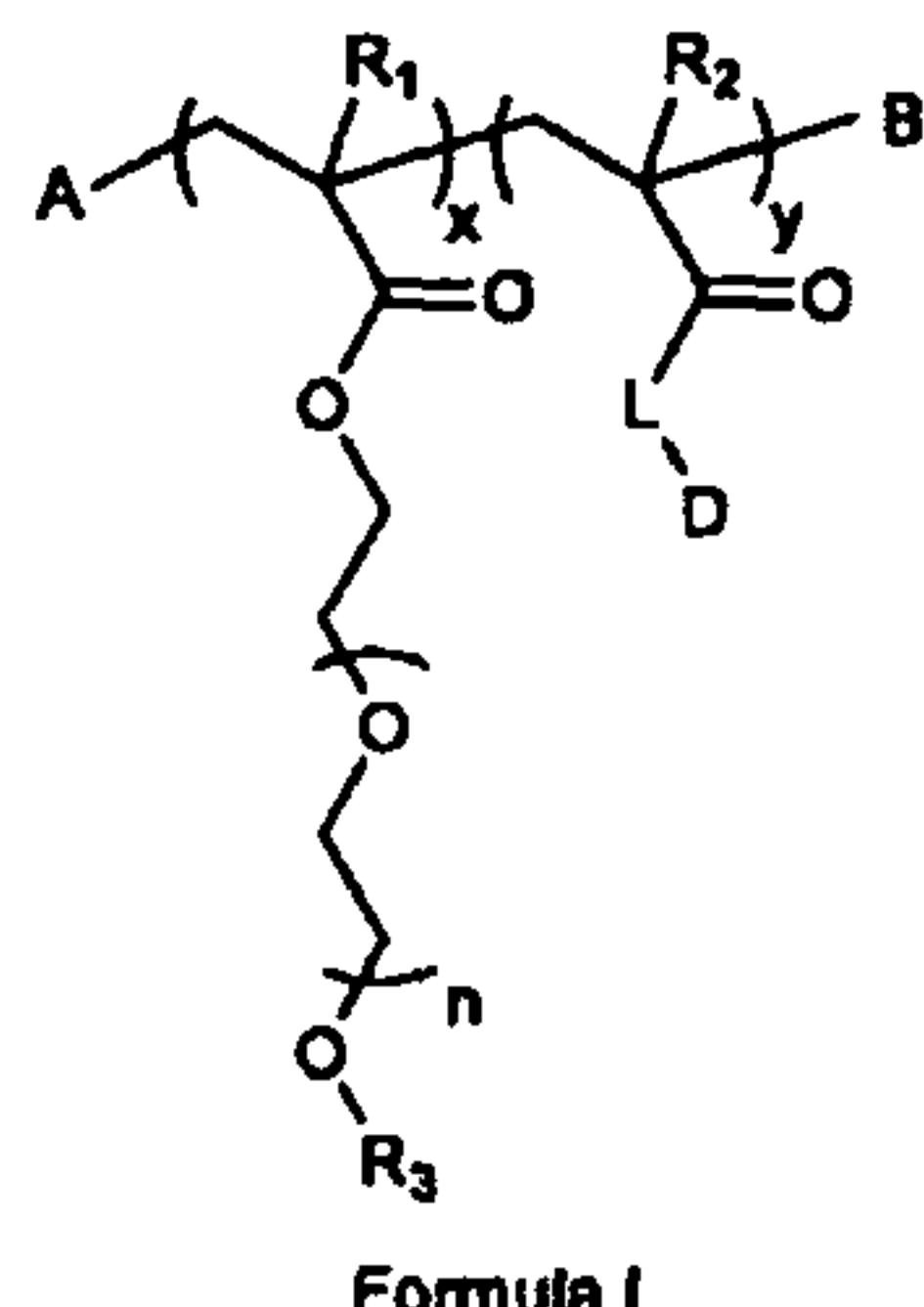
Figure 5: Fluorescence emission spectra of Nile Red in targeted polymeric assemblies of invention.

Figure 6: Plot of the emission intensity at 612 nm versus the log of POEGMEMA-*b*-PCombMA concentration

Figure 7: Plot of the emission intensity at 612 nm versus the log of total POEGMEMA-*b*-PCombMA and cRGDfK-POEGMEMA-*b*-PCombMA concentration

CLAIMS

1. A polymer-drug conjugate of formula I in the form of a block co-polymer for delivery of therapeutic agents



wherein

- R₁, R₂ and R₃ are independently selected from H or -CH₃
- x is a natural number between 1-100
- y is a natural number between 1-100
- n is a natural number between 1-50 and
- L is a cleavable linker and
- D is a therapeutic agent that is combretastatin or 5-Fluorouracil (5-FU) or gemcitabine or chloroquine or doxorubicine
- A is an end group or A may be null
- B is an end group or B may be null

2. A polymer drug conjugate according to claim 1 wherein linker is selected from a group comprising a poly (ethylene glycol), an amino acid, poly (amino acid) and short peptides, preferably short peptide is cathepsin B labile and most preferably short peptide is selected from a group comprising Gly-Phe-Leu-Gly (SEQ ID NO: 1), Val-Cit, Phe-Lys, Val-Ala, Ala-Leu-Ala-Leu (SEQ ID NO: 2).

3. A polymer drug conjugate according to claims 1-2 wherein, the linker is a C₁-C₁₀ hydrocarbon or C₁-C₁₀ substituted or hetero substituted hydrocarbon, which comprises a functional group that dissociates under physiological conditions, preferably said functional group is selected from acetal, ester, imine, amide, disulfide, carbonate, hydrazine, carbamate

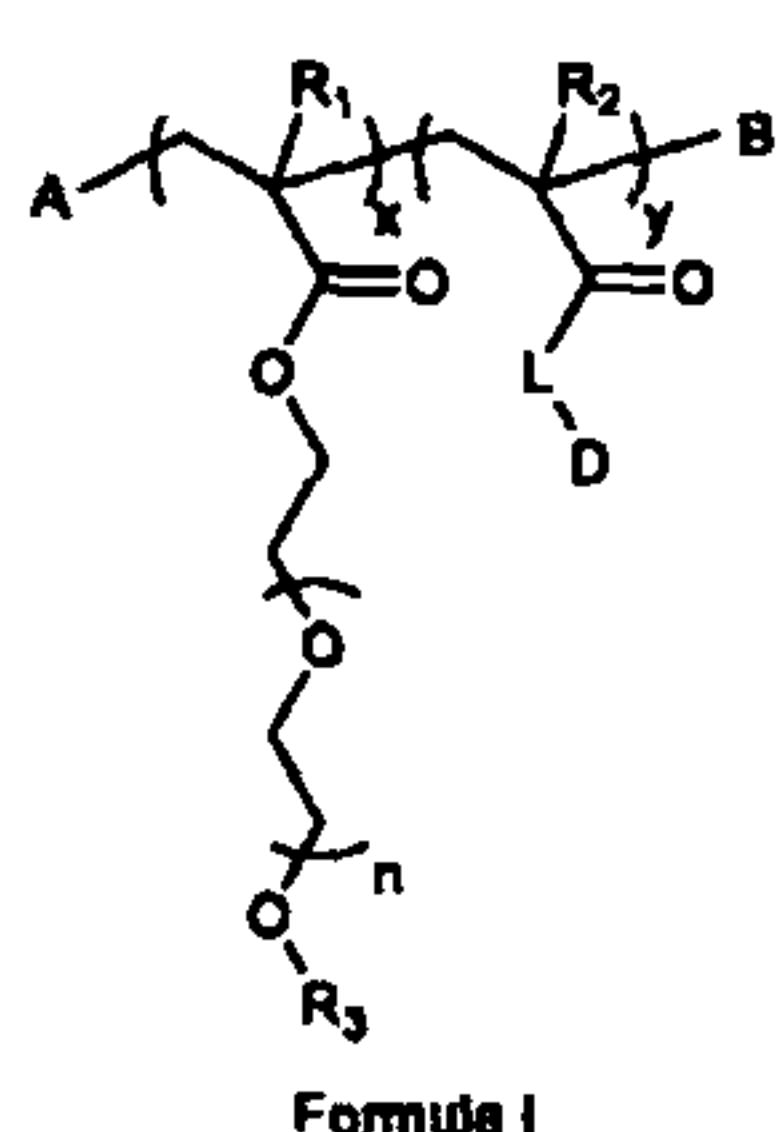
4. A polymer drug conjugate according to claim 1-3 wherein the linker is null.

5. A polymer-drug conjugate according to claims 1-4 wherein A is an end group that is a fragment of a chain transfer agent (CTA) or an initiator or a fragment of a CTA or initiator conjugated with a targeting moiety or a fragment of a CTA or initiator having a reactive functional group, preferably the reactive group is selected from acetal, hemiacetal, carboxylic acid, alcohol, amide, imide, anhydride, aryl halide, azo compound, diazo compound, hydrazine, azide, carbonate, chlorosilane, cyanide, ester, sulfate ester, phosphate ester, thiophosphate ester, isocyanate, isothiocyanate, thiocarbamate ester, dithiocarbamate ester.

6. A polymer-drug conjugate according to claims 1-5 wherein B is an end group that is a fragment of a chain transfer agent or initiator.

7. A polymer-drug conjugate according to claim 5 wherein targeting moiety is selected from a group comprising antibodies; antibody fragments, or peptides such as Cyclo (Arg-Gly-Asp-D-Phe-Lys) (cRGDfK) (SEQ ID No: 3).

8. Polymeric assemblies (nanoparticles or micelles) formed with polymer-drug conjugates of formula I according to claims 1-7

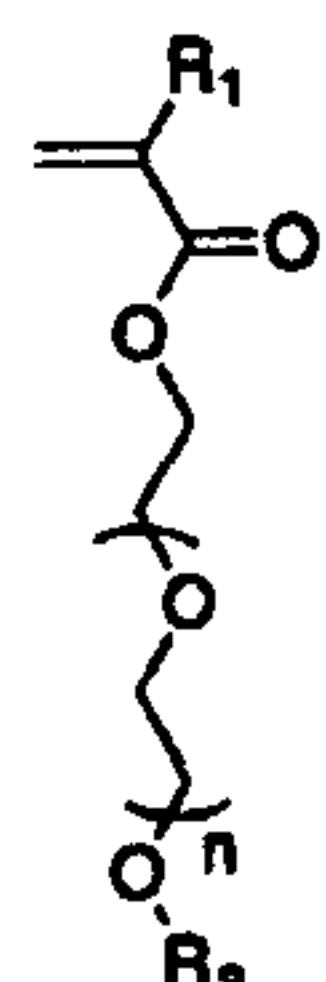


wherein

- R₁, R₂ and R₃ are independently selected from H or -CH₃
- x is a natural number between 1-100
- y is a natural number between 1-100
- n is a natural number between 1-50 and
- L is a cleavable linker or L may be null and
- D is a therapeutic agent that is combretastatin or 5-Fluorouracil or gemcitabine or chloroquine or doxorubicine
- A is an end group or A may be null
- B is an end group or B may be null.

9. Polymeric assemblies (nanoparticles or micelles) according to claim 8 encapsulating therapeutic agents other than those attached to the polymer-drug conjugate of formula I wherein said therapeutic agents can be selected from a group comprising nucleoside analogs, antifolates, other metabolites, topoisomerase I inhibitors, anthracyclines, podophyllotoxins, taxanes, vinca alkaloids, alkylating agents, platinates, antihormones, radiopharmaceutics, monoclonal antibodies, tyrosine kinase inhibitors, mammalian target of rapamycin (mTOR) inhibitors, retinoids, immunomodulatory agents, histone deacetylase inhibitors and other agents.

10. A method for preparation of polymer drug conjugate of formula I comprises polymerization of PEG (meth)acrylate monomer (formula II)

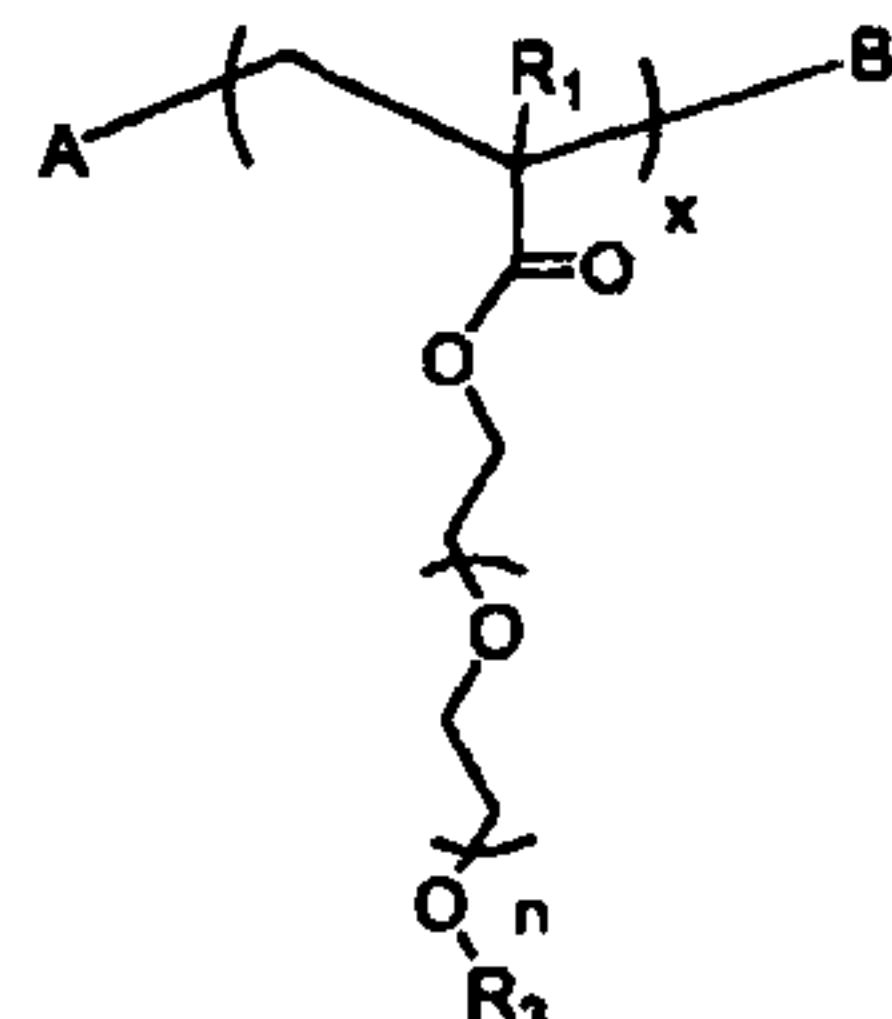


Formula II

wherein

- R_1 and R_3 are independently selected from H or $-CH_3$
- n is a natural number between 1-50

to give a polymer of formula IIa

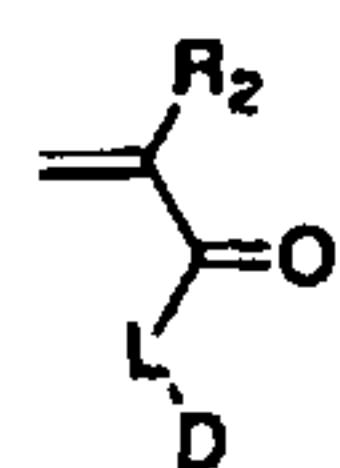


Formula IIa

wherein

- A is an end group or A may be null
- B is an end group or B may be null

and then (ii) further reacting formula IIa with a (meth)acrylate-L-D monomer (formula IIIa)



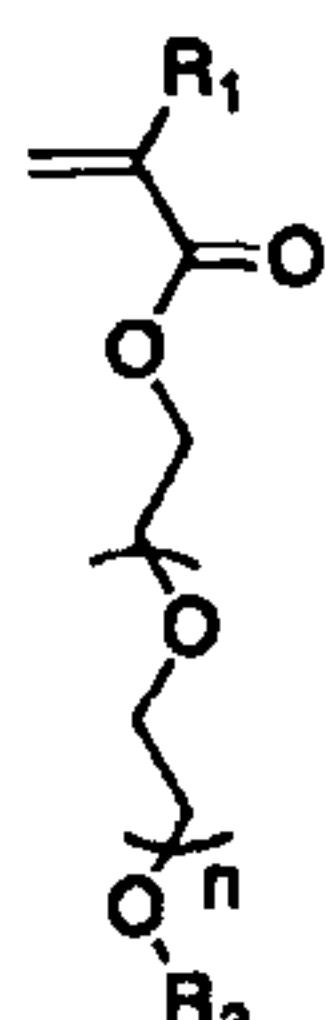
Formula IIIa

wherein

- R_2 is selected from H or $-CH_3$
- L is a cleavable linker or L may be null
- D is a therapeutic agent selected from combretastatin or 5-FU or gemcitabine or chloroquine or doxorubicine

to give polymer-drug conjugate of formula I.

11. A method for preparation of polymer drug conjugate of formula I comprises (i) polymerization of PEG (meth)acrylate monomer (Formula II)

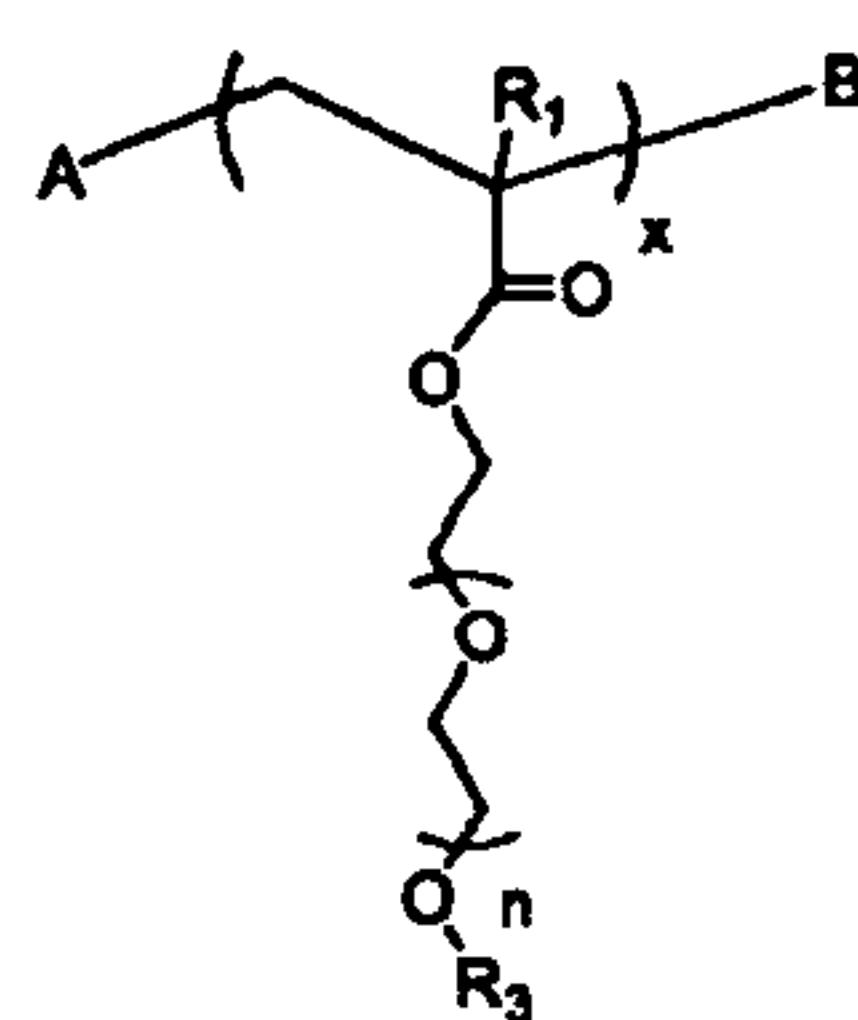


Formula II

wherein

- R_1 and R_3 are independently selected from H or $-CH_3$
- n is a natural number between 1-50

to give a polymer of formula IIa

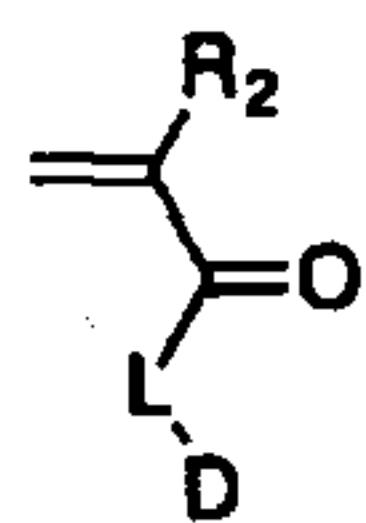


Formula IIa

wherein

- A is an end group or A may be null
- B is an end group or B may be null

And then (ii) further reacting formula IIa with a (meth)acrylate-L-D monomer (Formula IIIa)

**Formula IIIa**

- wherein R_2 is selected from H or $-\text{CH}_3$
- L is a cleavable linker or L may be null
- D is a therapeutic agent selected from combretastatin or 5-FU or gemcitabine or chloroquine or doxorubicine to give a block copolymer

And then (iii) reacting the formed block copolymer with a targeting moiety to give polymer-drug conjugate of formula I.

12. A polymer-drug conjugate of formula I obtainable by a method according to claim 10 or claim 11.
13. A pharmaceutical composition comprising a polymer-drug conjugate of formula I according to claims 1-7 and/or a polymeric assembly according to claims 8-9.
14. A polymer-drug conjugate of formula I according to claims 1-7 for use as a medicament for treatment and/or prophylaxis of cancer.
15. A polymeric assembly according to claims 8-9 for use as a medicament for treatment and/or prophylaxis of cancer.

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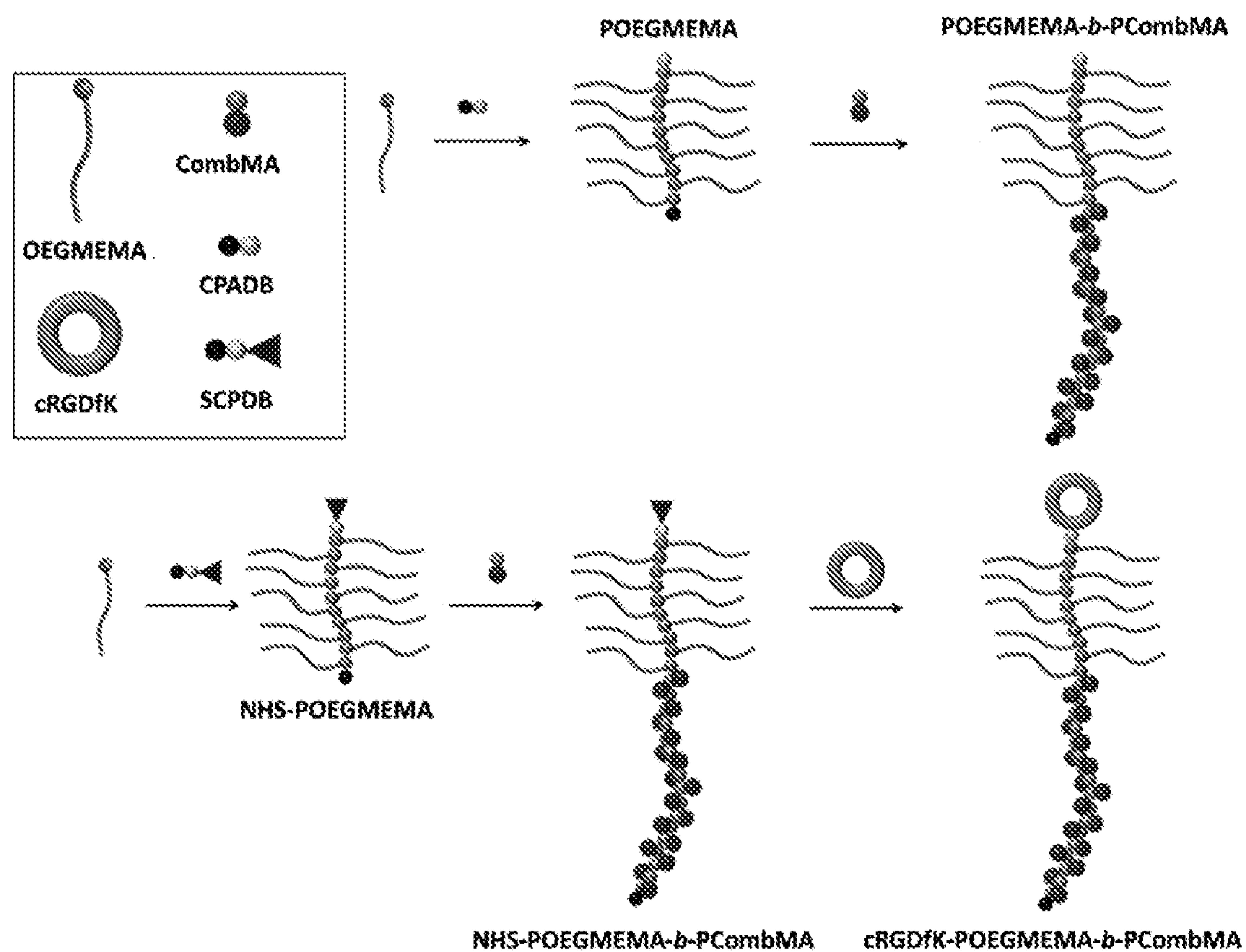


Figure 1

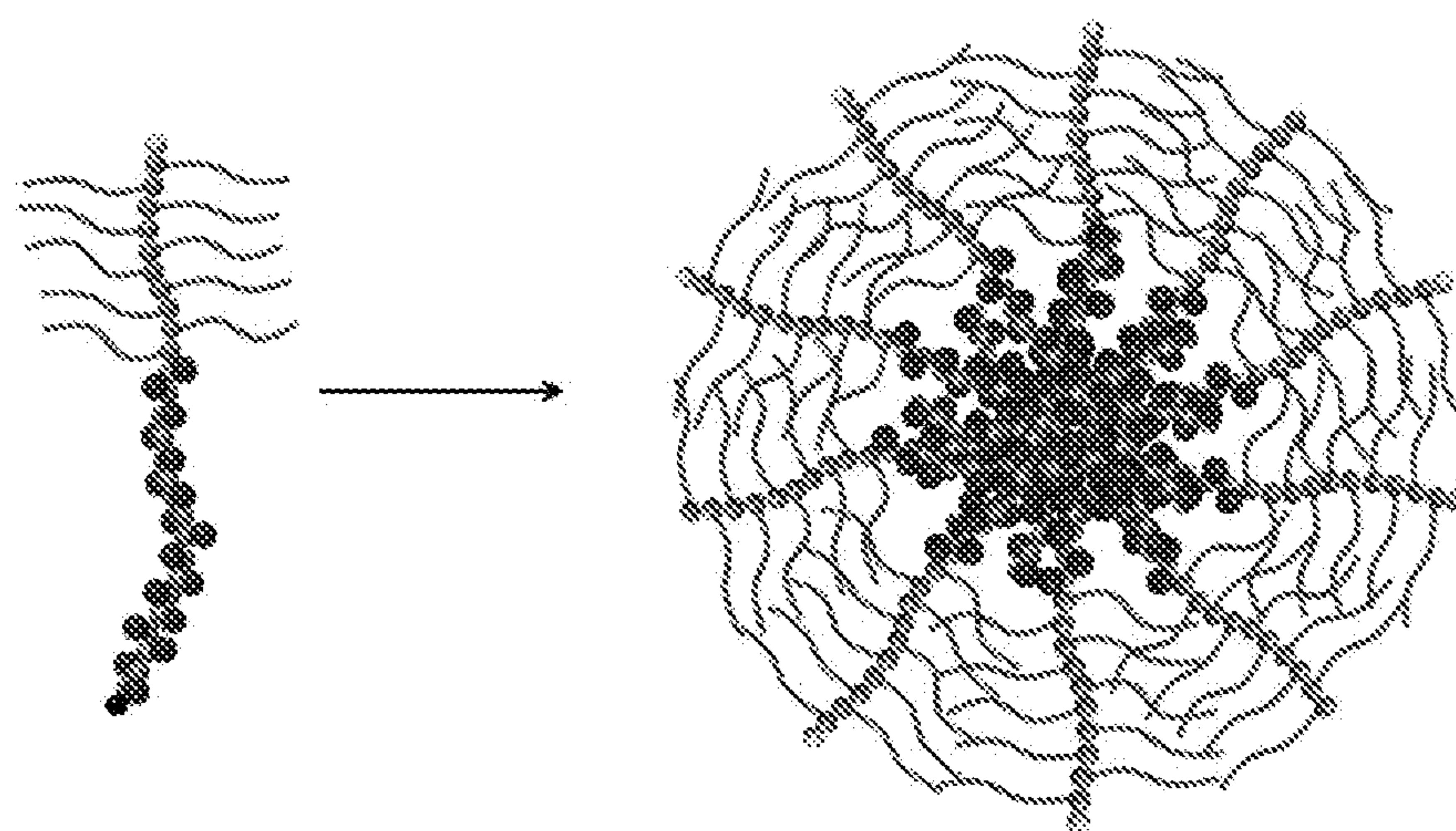


Figure 2

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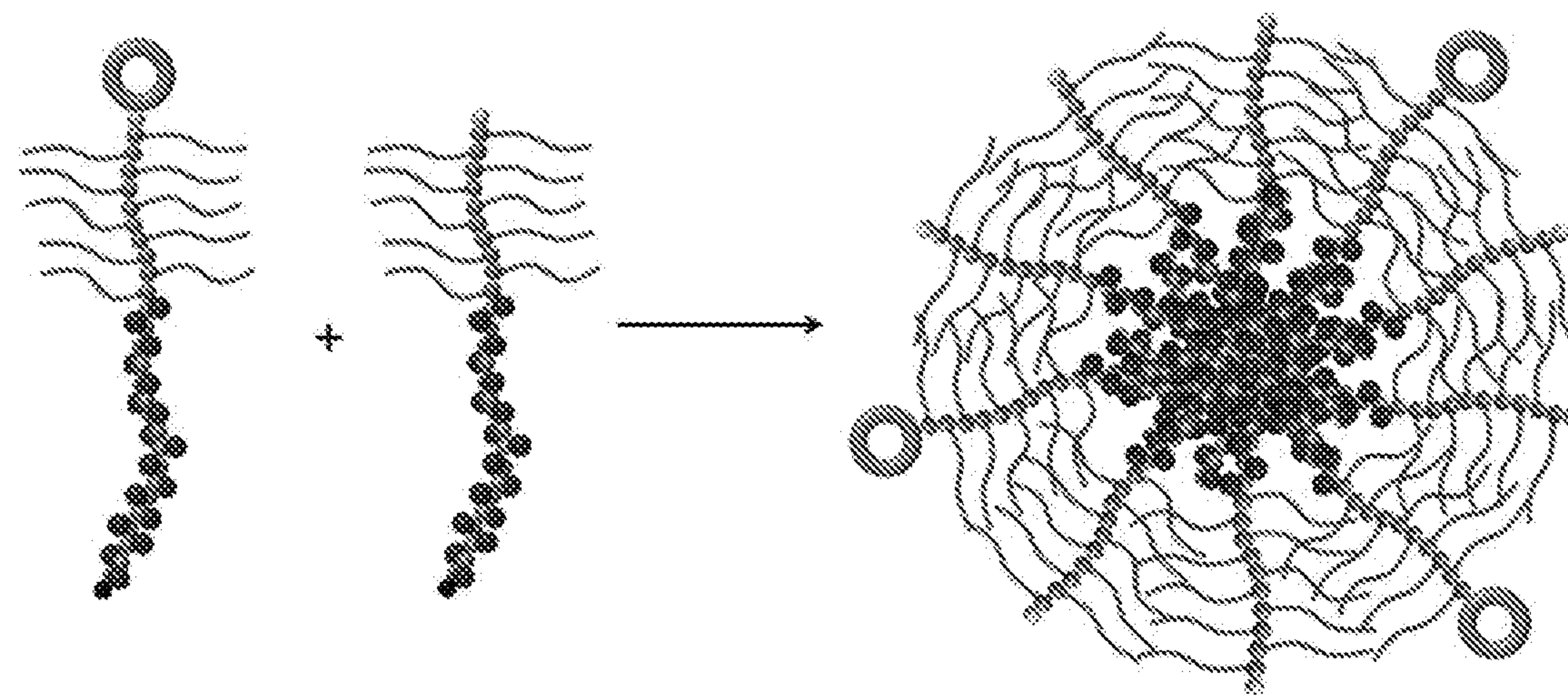


Figure 3

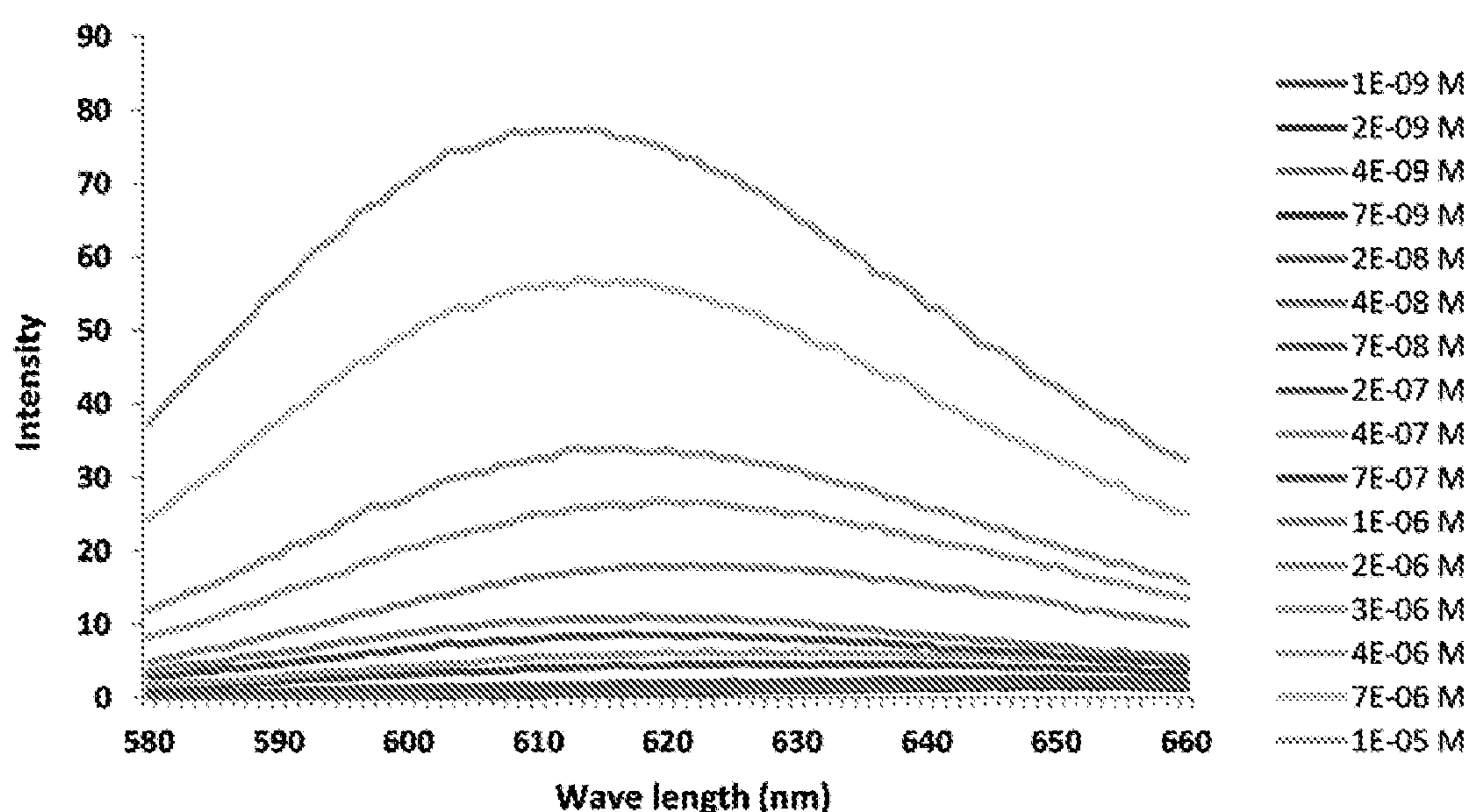


Figure 4

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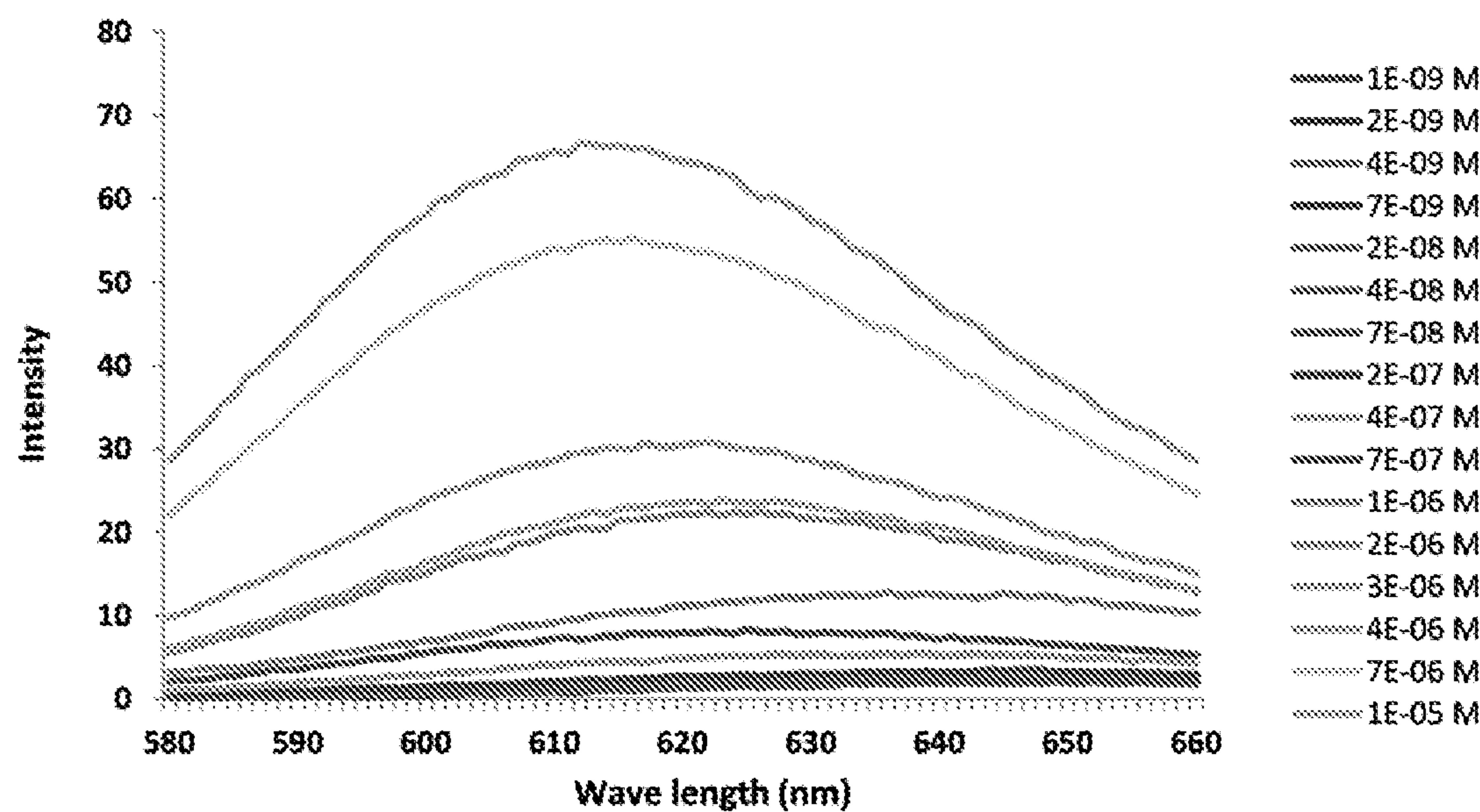


Figure 5

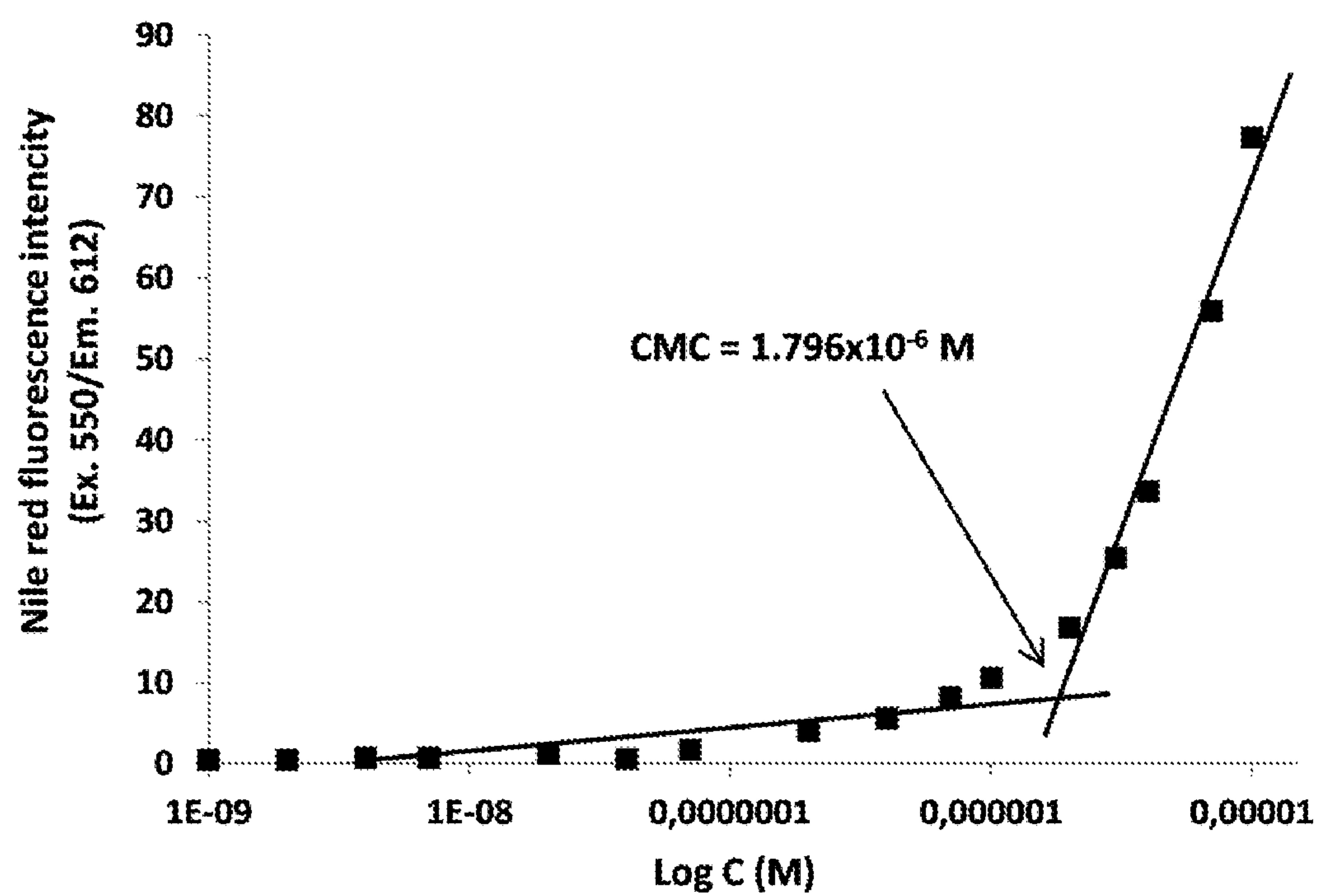
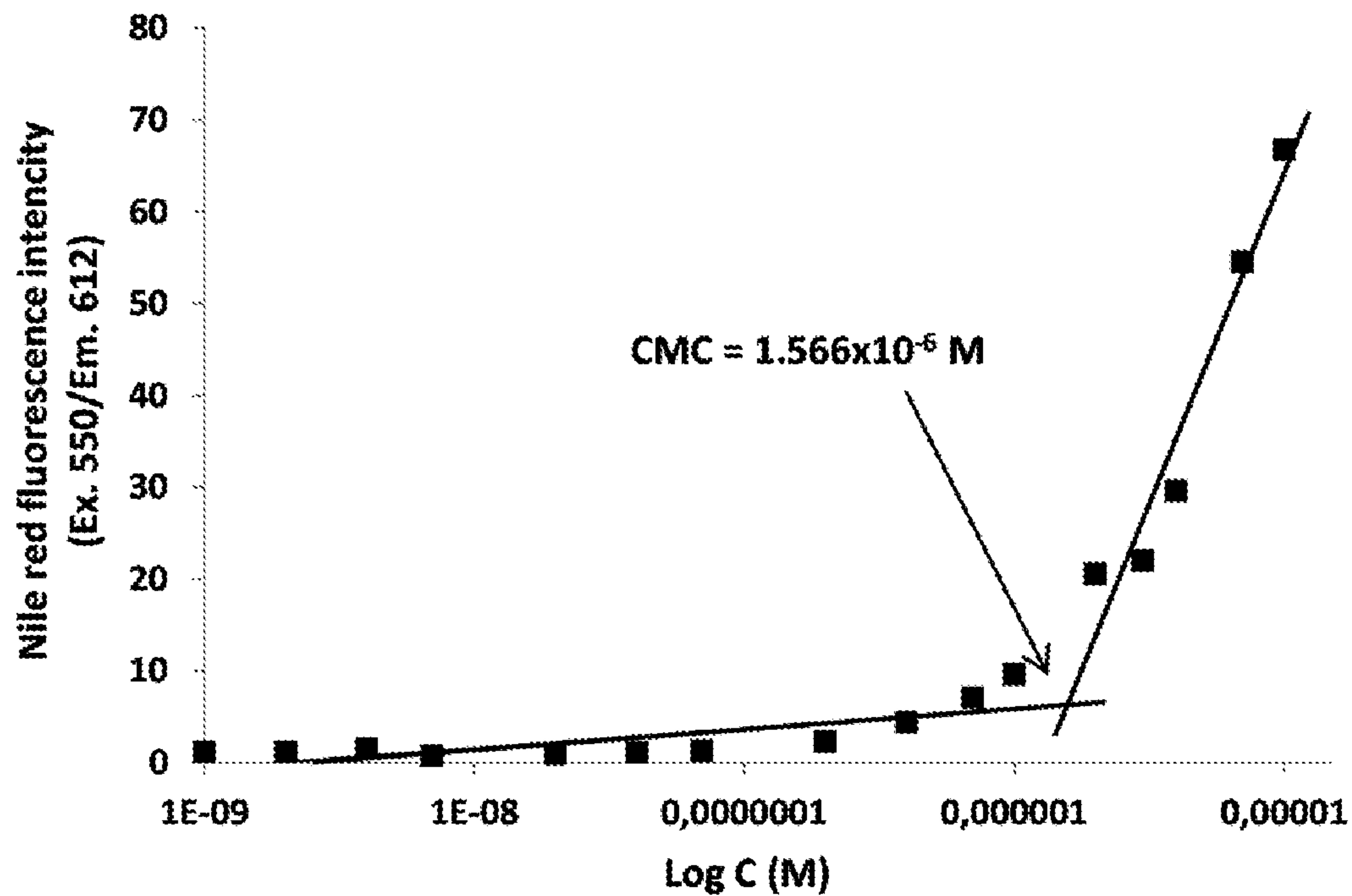
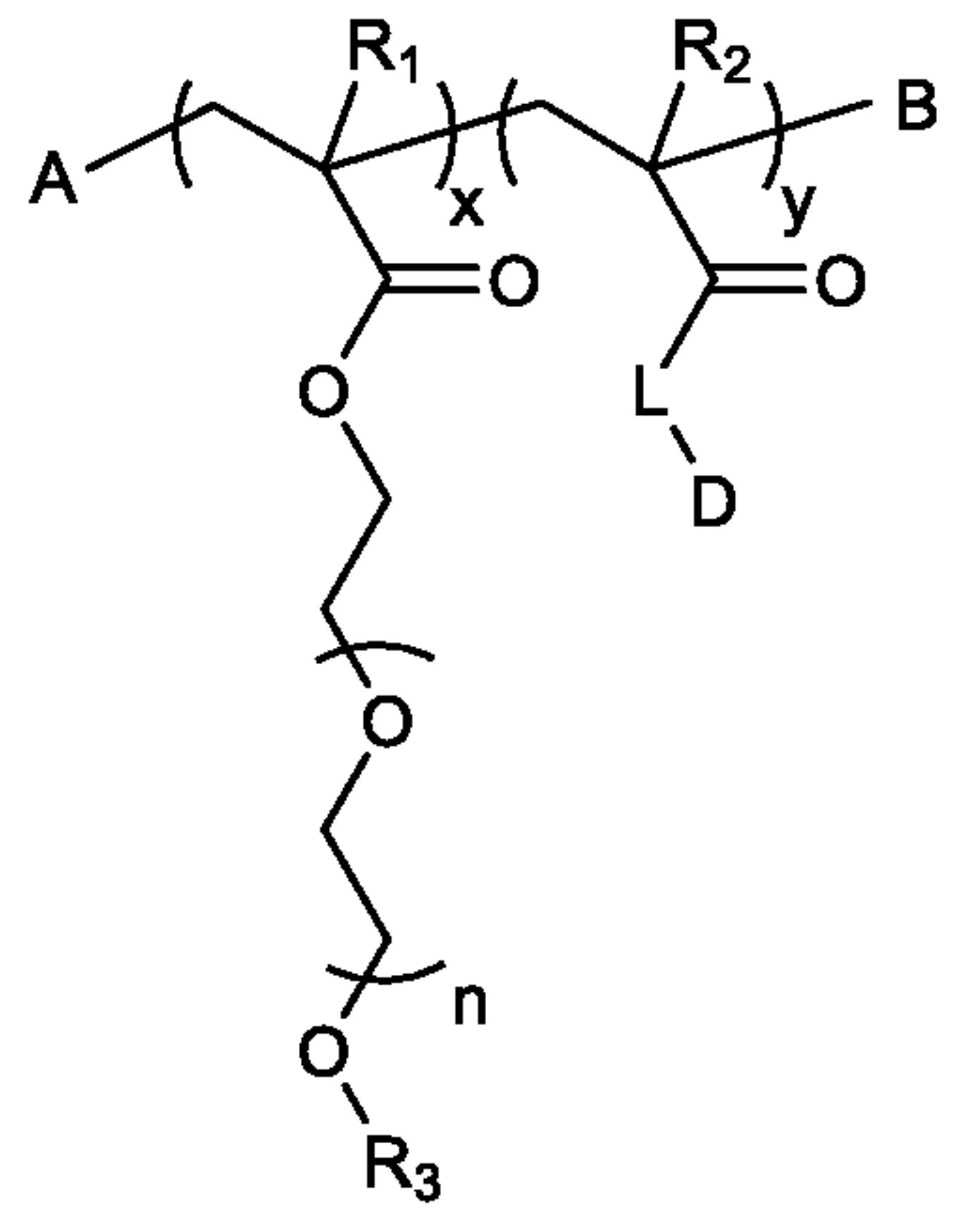


Figure 6

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Formula I

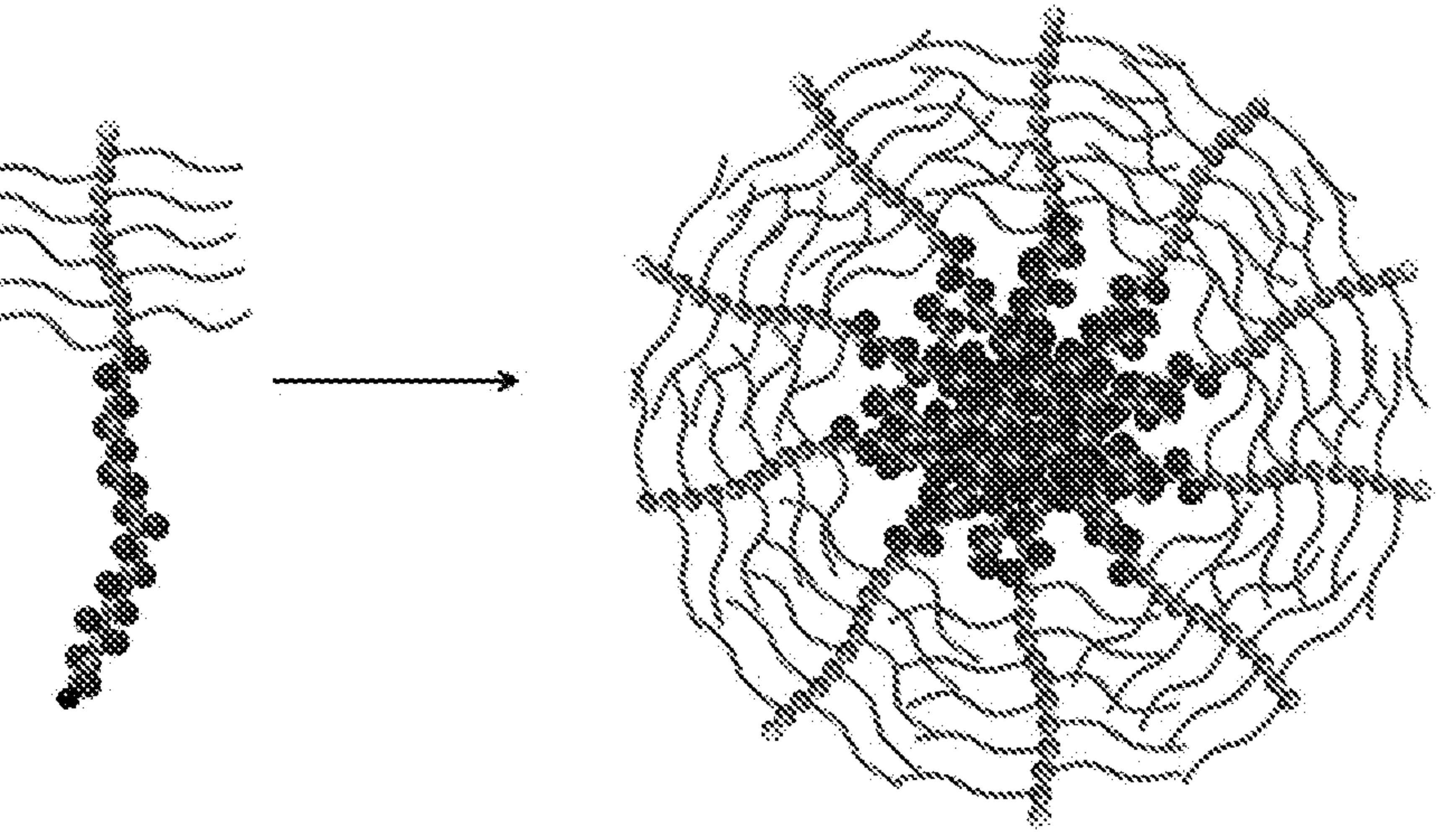


Figure 2