

(S7) Abstract: Improved poly(amido ethylenimine) copolymers for gene delivery are disclosed. One illustrative embodiment includes polyethylene glycol (PEG) covalently bonded to a branched poly (triethylenetetramine/cystamine bisacrylamide) copolymer (poly (TETA/CBA)). The polyethylene glycol can be linear or branched. Another illustrative embodiment includes an RGD peptide covalently bonded to the poly (TETA/CBA) -PEG conjugate. Still another illustrative embodiment includes a method of using these compositions for transfecting a cell with a nucleic acid.

**CLEAVABLE MODIFICATIONS TO REDUCIBLE
POLY(AMIDO ETHYLENIMINE)S TO ENHANCE NUCLEOTIDE DELIVERY**

BACKGROUND OF THE INVENTION

This invention relates to gene delivery. More
5 particularly, this invention relates to nonviral gene
delivery carriers.

Gene therapy has broad potential in treatment of human
genetic and acquired diseases through the delivery and
application of therapeutic gene-based drugs. The use of
10 safe, efficient and controllable gene carriers is a
requirement for the success of clinical gene therapy. R.C.
Mulligan, The basic science of gene therapy, 260 Science
926-932 (1993); I.M. Verma & N. Somia, Gene
therapy-promises, problems and prospects, 389 Nature 239-
15 242 (1997). Although viral vectors are very efficient in
gene delivery, their potential safety and immunogenicity
concerns raise their risk in clinical applications. C.
Baum et al., Mutagenesis and oncogenesis by chromosomal
insertion of gene transfer vectors, 17 Hum. Gene Ther. 253-
20 263 (2006). As an alternative to viral vectors, cationic
polymers such as poly(L-lysine) (PLL), poly(ethylenimine)
(PEI), poly(amidoamine) dendrimers, and cationic liposomes
have been synthesized as gene delivery carriers. The
advantages of these cationic polymer carriers include
25 safety, stability, large DNA and RNA loading capacity, and
easy and large-scale production. S. Li & L. Huang,
Nonviral gene therapy: promises and challenges, 7 Gene
Ther. 31-34 (2000); F. Liu et al., Non-immunostimulatory
nonviral vectors, 18 Faseb J. 1779-1781 (2004); T. Niidome
30 & L. Huang, Gene therapy progress and prospects: nonviral
vectors, 9 Gene Ther. 1647-1652 (2002). The cationic
polymers can condense negatively charged DNA into nanosized
particles through electrostatic interactions, and the
polymer/pDNA polyplexes can enter cells via endocytosis.
35 Y.W. Cho et al., Polycation gene delivery systems: escape
from endosomes to cytosol, 55 J. Pharm. Pharmacol. 721-734

(2003); L. De Laporte et al., Design of modular non-viral gene therapy vectors, 27 Biomaterials 947-954 (2006); E. Piskin et al., Gene delivery: intelligent but just at the beginning, 15 J. Biomater. Sci. Polym. Ed. 1182-1202
5 (2004). As a result, the polymers can protect pDNA from nuclease degradation and facilitate cellular uptake to induce high gene transfection. O. Boussif et al., A versatile vector for gene and oligonucleotide transfer into cells in culture and in vivo: polyethylenimine, 92 Proc.
10 Nat'l Acad. Sci. USA 7297-7301 (1995); D.W. Pack et al., Design and development of polymers for gene delivery, 4 Nat. Rev. Drug. Discov. 581-593 (2005).

The currently available cationic polymers, however, have significant cytotoxicity concerns, mostly due to their
15 poor biocompatibility and non-degradability under physiological conditions.

Therefore, while prior nonviral gene delivery carriers are known and are generally suitable for their limited purposes, they possess certain inherent deficiencies that
20 detract from their overall utility in gene therapy.

In view of the foregoing, it will be appreciated that providing improved carriers to enhance nucleotide delivery would be a significant advancement in the art.

BRIEF SUMMARY OF THE INVENTION

25 An illustrative embodiment of the present invention comprises a composition comprising polyethylene glycol covalently bonded to poly[H₂N-(CH₂-CH₂-N)_n-H/CBA]. In illustrative embodiments of the present invention, n is 1, 2, or 3. In some illustrative embodiments of the present
30 invention, the polyethylene glycol is linear, but in other illustrative embodiments, the polyethylene glycol is branched. The polyethylene glycol typically has a molecular weight of about 1,000 to about 50,000 and, more typically has a molecular weight of about 2,000 to about
35 25,000. In one illustrative embodiment of the invention, the polyethylene glycol has a molecular weight of about

3,400. The poly[H₂N-(CH₂-CH₂-N)_n-H/CBA] typically has a molecular weight of about 1,000 to about 25,000. However, the molecular weights of the poly[H₂N-(CH₂-CH₂-N)_n-H/CBA] and the polyethylene glycol are limited only by their
5 functionality as carriers. In some illustrative embodiments of the present invention, the composition further comprises an RGD peptide covalently bonded to the polyethylene glycol. An illustrative example of an RGD peptide is shown in SEQ ID NO:1.

10 Another illustrative embodiment of the present invention comprises a complex comprising a nucleic acid ionically bonded to a composition comprising polyethylene glycol covalently bonded to poly[H₂N-(CH₂-CH₂-N)_n-H/CBA]. Illustrative nucleic acids comprise plasmids, siRNAs, and
15 oligonucleotides.

Still another illustrative embodiment of the present invention comprises a method of transfecting a cell with a selected nucleic acid, the method comprising contacting the cell with a complex comprising the selected nucleic acid
20 ionically bonded to a composition comprising polyethylene glycol covalently bonded to poly[H₂N-(CH₂-CH₂-N)_n-H/CBA] such that the complex enters the cell.

A still further illustrative embodiment of the invention comprises a solid support coated with a
25 composition comprising polyethylene glycol covalently bonded to poly[H₂N-(CH₂-CH₂-N)_n-H/CBA]. The polyethylene glycol may be linear or branched. In certain illustrative embodiments of the invention, n is 1, 2, or 3. Moreover, in additional illustrative embodiments, an RGD peptide may
30 be bonded to the polyethylene glycol.

BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWINGS

FIG. 1 shows a scheme for synthesis of poly[H₂N-(CH₂-CH₂-N)_n-H/CBA] according to the present invention.

FIG. 2 shows the results of MALDI-TOF analysis showing
35 the degree of branching of poly(TETA/CBA) synthesized at either 30°C or 40°C.

FIG. 3 shows the results of MALDI-TOF analysis showing the degree of branching of poly(TETA/CBA) synthesized at 50°C.

FIG. 4 shows the effect of polymer architecture on transfection efficiency in H9c2 cells. From left to right, results are shown for a no-carrier control (i.e., report plasmid only), a complex of poly(TETA/CBA) and reporter plasmid wherein the poly(TETA/CBA) was synthesized at 30°C and the weight ratio of carrier to reporter plasmid is 6 ("30 Deg w/w 6"), 30 Deg w/w 12, 30 Deg w/w 24, 40 Deg w/w 6, 40 Deg w/w 12, 40 Deg w/w 24, 50 Deg w/w 6, 50 Deg w/w 12, and 50 Deg w/w 24.

FIG. 5 shows the effect of polymer architecture on cell viability in H9c2 cells. Results are presented in the same order as in FIG. 4.

FIG. 6 shows a general synthesis scheme for poly(TETA/CBA) and poly(TETA/CBA)-*g*-PEG₃₄₀₀.

FIG. 7 shows protection of plasmid DNA from serum nucleases in 20% FBS in PBS for up to 24 hours with 1,000 (1K); 5,000 (5K); and 10,000 (10K) molecular weight cut off fractions of poly(TETA/CBA) and the corresponding PEG copolymer (5K-PEG3.4K).

FIG. 8 shows protection of plasmid DNA from serum nucleases by poly(TETA/CBA) synthesized at 30°C and 50°C, as compared to a control having no poly(TETA/CBA).

FIG. 9 shows particle sizes of polyplexes formed from plasmid DNA and poly(TETA/CBA(1k)) (■), poly(TETA/CBA(5k)) (▲), poly(TETA/CBA(10k)) (▼), and poly(TETA/CBA(5k))-PEG₃₄₀₀ (◆).

FIG. 10 shows zeta potentials of polyplexes formed from plasmid DNA and poly(TETA/CBA(1k)) (■), poly(TETA/CBA(5k)) (▲), poly(TETA/CBA(10k)) (▼), and poly(TETA/CBA(5k))-PEG₃₄₀₀ (◆).

FIG. 11 shows particle diameters measured by dynamic light scattering (DSL) of polyplexes formed from plasmid DNA and poly(TETA/CBA(1k)) synthesized at 40°C.

FIG. 12 shows particle diameters measured by dynamic light scattering (DSL) of polyplexes formed from plasmid DNA and poly(TETA/CBA(5k)) synthesized at 40°C.

FIG. 13 shows particle diameters measured by dynamic light scattering (DSL) of polyplexes formed from plasmid DNA and poly(TETA/CBA(10k)) synthesized at 40°C.

FIG. 14 shows particle diameters measured by dynamic light scattering (DSL) of polyplexes formed from plasmid DNA and poly(TETA/CBA(5k))-PEG₃₄₀₀ synthesized at 40°C.

FIG. 15 shows a scheme for synthesis of poly(TETA/CBA)-g-PEG-RGD.

FIG. 16 shows a scheme for synthesis of bPEG-g-poly(TETA/CBA).

FIG. 17 shows particle sizes of poly(TETA/CBA) (◇), poly(TETA/CBA)-PEG₃₄₀₀ (■), and poly(TETA/CBA)-PEG-RGD (▽).

FIG. 18 shows zeta potentials of poly(TETA/CBA) (■), poly(TETA/CBA)-PEG₃₄₀₀ (Δ), and poly(TETA/CBA)-PEG-RGD (▽).

FIG. 19 shows cytotoxicity of poly(TETA/CBA) and bPEG-poly(TETA/CBA) as compared to PEG-PEI and control.

FIG. 20 shows transfection as measured by luciferase expression for poly(TETA/CBA) and bPEG-poly(TETA/CBA) as compared to PEG-PEI and control.

DETAILED DESCRIPTION

Before the present improvement to reducible poly(amido ethylenimine)s (SS-PAEIs) and methods are disclosed and described, it is to be understood that this invention is not limited to the particular configurations, process steps, and materials disclosed herein as such configurations, process steps, and materials may vary somewhat. It is also to be understood that the terminology employed herein is used for the purpose of describing particular embodiments only and is not intended to be limiting since the scope of the present invention will be limited only by the appended claims and equivalents thereof.

The publications and other reference materials referred to herein to describe the background of the invention and to provide additional detail regarding its practice are hereby incorporated by reference. The references discussed herein are provided solely for their disclosure prior to the filing date of the present application. Nothing herein is to be construed as an admission that the inventors are not entitled to antedate such disclosure by virtue of prior invention.

It must be noted that, as used in this specification and the appended claims, the singular forms "a," "an," and "the" include plural referents unless the context clearly dictates otherwise.

Unless defined otherwise, all technical and scientific terms used herein have the same meanings as commonly understood by one of ordinary skill in the art to which this invention belongs.

In describing and claiming the present invention, the following terminology will be used in accordance with the definitions set out below.

As used herein, "comprising," "including," "containing," "characterized by," and grammatical equivalents thereof are inclusive or open-ended terms that do not exclude additional, unrecited elements or method steps. "Comprising" is to be interpreted as including the more restrictive terms "consisting of" and "consisting essentially of." As used herein, "consisting of" and grammatical equivalents thereof exclude any element, step, or ingredient not specified in the claim. As used herein, "consisting essentially of" and grammatical equivalents thereof limit the scope of a claim to the specified materials or steps and those that do not materially affect the basic and novel characteristic or characteristics of the claimed invention.

As used herein, "SS-PAEI" means poly(amido ethylenimine) containing reducible disulfide bonds; "TETA"

means triethylenetetramine; and "CBA" means *N,N'*-cystamine bisacrylamide.

Reducible poly(amido ethylenimine)s (SS-PAEIs) are a class of nonviral carrier used for the delivery of nucleic acids. L.V. Christensen et al., Reducible Poly(amido ethylenimine)s Designed for Triggered Intracellular Gene Delivery, 17 Bioconjugate Chem. 1233-1240 (2006). The synthesis of these types of polymers using Michael addition chemistry between polyamine monomers and cystamine bisacrylamide (CBA) is shown in FIG. 1. The resultant polymers have been shown to mediate effective transfection efficiency of therapeutic plasmid DNA as well as siRNA. L.V. Christensen, C.W. Chang, J.W. Yockman, et al., Reducible poly(amido ethylenimine) for hypoxia-inducible VEGF delivery, 118 J. Control. Rel. 254-261 (2007); J. Hoon Jeong, L.V. Christensen, J.W. Yockman, Z. Zhong, J.F.J. Engbersen, W.J. Kim, J. Feijen, S.W. Kim, Reducible poly(amido ethylenimine) directed to enhance RNA interference, 28 Biomaterials 1912-1917 (2007). These polymers take advantage of the redox potential between the extracellular and the intracellular environment by a triggered release mechanism delivering the nucleic acid of interest into the cytoplasmic compartment of the cell. Such characteristics have a significant advantage over traditional noncleavable or hydrolysable polymers used to date for nonviral gene delivery.

A clear advantage is seen with the delivery of oligonucleotides (such as antisense oligonucleotides and siRNA) that only need to be delivered into the cytoplasm for their effects to be demonstrated. Traditional noncleavable polymers typically remain condensed within the cytoplasm once released from the endosomal compartment due to their high electrostatic interactions. This condensation of the therapeutic nucleotides prevents their use in the host machinery for transcription or translational inhibition. However, the SS-PAEIs are

reduced within the cytoplasm by reductive proteins/enzymes, i.e., glutathione-S-transferase (GSH), releasing the therapeutic nucleotides and significantly increasing the amount of available nucleotides for transcription or
5 inhibition of translation.

Until recently, these polymers have been used for direct delivery via injection *in vivo*. Disadvantages of systemic delivery, including undesired interactions with negatively charged plasma proteins and nonspecific
10 degradation during circulation, has led to the construction of a polymeric system that can overcome the main issues confronted by nonviral gene delivery to date--delivery. Most importantly, reactive linkages between the polymer and the additional polymeric structures an/or targeting
15 moieties are crucial to allow for complete reduction of the nucleotide/polymeric complexes within the cytoplasm. It has been shown that stable linkages of polymeric structures may decrease transfection efficiency within cells, whether by inhibiting release of the nucleotides or hindering
20 movement into the nucleus. M. Meyer & E. Wagner, pH-Responsive shielding of non-viral gene vectors, 3 Expert Opin. Drug Deliv. 563-571 (2006). As this reduction of the SS-PAEIs is crucial for effective nucleotide delivery, effective modifications of SS-PAEIs that protect against
25 negative interactions *in vivo* and deliver nucleic acids to the desired targeted region are disclosed herein.

To improve the poly(triethylenetetramine cystamine-bisacrylamide) (poly(TETA/CBA)) polymer for *in vivo* applications via systemic administration, the polymer may
30 be sterically stabilized using a hydrophilic PEG spacer that may, or may not, contain an accompanying tissue specific targeting moiety, such as RGD. Moreover, previous gene delivery studies have demonstrated that *in vivo* gene transfer is highest with high molecular weight, branched
35 polymers. An example of a co-polymer exhibiting these characteristics is PEG₃₄₀₀-bPEI25k. Therefore, a high

molecular weight, branched poly(TETA/CBA) ("bTETA/CBA") has been synthesized and has been conjugated to PEG₃₄₀₀ and PEG₃₄₀₀-RGD to derive the resulting copolymers termed poly(TETA/CBA)-PEG₃₄₀₀ and poly(TETA/CBA)-PEG₃₄₀₀-RGD.

5 Another illustrative embodiment of the present invention comprises a solid support coated with a composition comprising polyethylene glycol covalently bonded to poly(TETA/CBA). The polyethylene glycol can be linear or branched. In addition, an RGD peptide can be
10 bonded to the polyethylene glycol. Illustratively, the solid support can be a dish bottom, a multiwell plate, or a continuous surface. The solid support can be glass, plastics (such as polytetrafluoroethylene (PTFE), polyvinylidenedifluoride, polystyrene, polycarbonate, or
15 polypropylene), silicon, metal (such as gold), membranes (such as nitrocellulose, methylcellulose, PTFE, or cellulose), paper, biomaterials (such as protein, gelatin, or agar), tissues (such as skin, endothelial tissue, bone, or cartilage), or minerals (such as hydroxyapatite or
20 graphite), slides (such as glass or poly-L-lysine coated slides) or wells of a multiwell or microtiter plate. Typical solid support materials include polystyrene resin, epoxy resin, or glass. The coating amount is typically about 0.1 to about 100 µg/cm². The composition is coated on
25 the surface of the solid support. The composition can be mixed with a matrix, such as proteins, peptides, polysaccharides, or polymers. The protein can be gelatin, collagen, bovine serum albumin, or any other protein that can be used in affixing proteins to a surface. The
30 polymers can be hydrogels, copolymers, non-degradable or biodegradable polymers, and biocompatible materials. The polysaccharide can be any compound that can form a membrane and coat the polymer, such as chitosan. Reagents, such as cytotoxicity reductive reagents, cell binding reagents,
35 cell growing reagents, cell stimulating reagents, or cell inhibiting reagents and compounds for culturing specific

cells, can also be affixed to the solid surface along with the composition.

Illustratively, a gelatin-composition mixture, comprising composition and gelatin in an appropriate solvent, such as water, may be affixed to the solid surface. A cell culture reagent, such as fibronectin, collagen, salts, sugars, protein, or peptides may also be present in the gelatin-polymer mixture. The mixture is spread evenly on a surface of the solid support, such as a slide or a multiwell plate. The resulting product is allowed to dry completely under suitable conditions such that the gelatin-polymer mixture is affixed to the solid support. For example, the resulting product can be dried at selected temperatures or humidities, or in a vacuum desiccator.

Example 1

Poly(TETA/CBA) was synthesized according to the scheme illustrated in FIG. 1 and described in L.V. Christensen et al., *supra*, at 30°C, 40°C, and 50°C.

The degree of branching of poly(TETA/CBA) prepared according to this example was characterized by matrix-assisted laser desorption/ionization time of flight (MALDI-TOF) mass spectrometry. The results show that poly(TETA/CBA) branching was about 90%. Synthesis at 30°C or 40°C (FIG. 2) resulted in about 90% of poly(TETA/CBA) repeat units containing at least one branched chain (arm), with the other 10% being linear. Within this population, about 60% contained one arm, and about 30% contained two arms. Synthesis at 50°C (FIG. 3) resulted in a more dense architecture, with about 7% of the copolymers being linear, about 40% containing one arm, about 40% containing two arms, and about 10% containing three arms.

Example 2

Transfection with poly(TETA/CBA) copolymers was evaluated in H9c2 cells using the plasmid, pBLuc, as a reporter. Cells were maintained in DMEM containing 10%

FBS, streptomycin (100 µg/mL) and penicillin (100 units/mL) at 37°C in a humidified atmosphere with 5% CO₂. Cells were seeded 24 hrs prior to transfection in 24-well plates at initial densities of 4.0×10^4 cells/well. DNA was

5 complexed with the poly(TETA/CBA) polymers at polymer/pBLuc w/w ratios of 6,12, and 24. Controls contained no poly(TETA/CBA) polymer. Complexes were made in HEPES buffer and incubated for 30 min before use. At the time of transfection, the medium in each well was replaced with

10 fresh serum-free medium. Polyplexes (0.5 µg DNA/well) were incubated with the cells for 4 hrs at 37°C. The medium was then replaced with 500 µL of fresh complete medium and cells were incubated for additional 44 hrs. The cells were then washed with pre-warmed PBS, treated with 200 µL cell

15 lysis buffer and subjected to a freezing-thawing cycle. Cellular debris was removed by centrifugation at 14,000 *g* for 5 min. The luciferase activity in cell lysates (25 µL) was measured using a luciferase assay kit (100 µL luciferase assay buffer) on a luminometer (Dynex

20 Technologies Inc., Chantilly, Virginia). The relative luminescence unit (RLU) of luciferase expression was normalized against protein concentration in the cell extracts, measured by a BCA protein assay kit (Pierce, Rockford, Illinois). All transfection assays were carried

25 out in triplicate. Results are shown in FIG. 4. The more highly branched architecture reduced transfection efficiency as compared to the less-branched architecture.

Example 3

H9c2 cells were seeded in a 24-well plate at a density

30 of 4.0×10^4 cells/well and incubated for 24 hrs. Poly(TETA/CBA)/pBLuc complexes of weight ratios (w/w) of 6, 12, and 24 and no polymer (control) were incubated with the cells for 4 hrs in serum-free medium followed by 20 hrs in complete medium. MTT solution (50 µL, 2 mg/mL) was then

35 added and cells were further incubated for 2 hrs. The medium was removed and 300 µL DMSO was then added to each

well. The absorption was measured at 570 nm using a microplate reader (Model 680, Bio-Rad Lab, Hercules, California). The percentage relative cell viability was determined relative to control (untreated) cells, which
5 were not exposed to transfection system and taken as 100% cell viability. All cytotoxicity experiments were performed in triplicate. Results are shown in FIG. 5. Cytotoxicity was essentially nil.

Example 4

10 Poly(TETA/CBA) copolymers prepared according to the procedure of Example 1 were fractionated using ultrafiltration membranes with molecular weight cut offs of 10,000; 5,000; and 1,000. These fractions were termed poly(TETA/CBA(10k)), poly(TETA/CBA(5k)), and
15 poly(TETA/CBA(1k)), respectively.

Polyethylene glycol of molecular weight 3,400 (PEG_{3.4k} or PEG₃₄₀₀) was conjugated in equimolar amounts to poly(TETA/CBA(5k)). DPAS was dissolved in dimethylsulfoxide (DMSO) and added dropwise to NH₂-PEG-COOH
20 with excess pyridine in DMSO and reacted overnight. Next, dithiothreitol (DTT) was added to the reaction mixture and reacted for 4 hr. The resulting PEG product was then purified by ultrafiltration (1000 MWCO). HS-PEG-COOH and excess divinyl sulfone were dissolved in DMSO as a solvent
25 and reacted overnight. The resulting mixture was purified again by ultrafiltration (1000 MWCO). The resulting PEG product was then dissolved in sodium carbonate buffer, pH 9.0, and added dropwise to the poly(TETA/CBA(5k)) carbonate solution and reacted for 16 hr. For visual clarity,
30 poly(TETA/CBA(5k)) appears linear in FIG. 6, however, it should be noted that the poly(TETA/CBA(5k)) was branched. The product resulting from this reaction was termed poly(TETA/CBA(5k))-PEG₃₄₀₀.

Poly(TETA/CBA(1k)), poly(TETA/CBA(5k)),
35 poly(TETA/CBA(10k)), poly(TETA/CBA(5k))-PEG₃₄₀₀, and hyperbranched polyethylenimine (bPEI25k; $M_w = 25,000$;

Aldrich, St. Louis, Missouri) were analyzed by gel permeation chromatography (GPC). Table 1 shows the number average molecular weight (M_n), weight average molecular weight (M_w), and polydispersity index (M_w/M_n ; PDI) derived from GPC measurements. Polymer fraction buffer capacity titrations were determined by the moles of HCl required to shift pH from 7.4 to 5.1 in 0.1 M aqueous NaCl. Degree of branching was determined by reduction of each polymer fraction using TCEP and subsequent free sulfhydryl NEM protection as analyzed by MALDI-TOF.

Table 1					
Polymer	M_n (kDa)	M_w (kDa)	PDI	Titration (mol HCl)	Degree Branching
p(TETA/CBA(1k))	4.2	8.2	1.95	12.1	0.68
p(TETA/CBA(5k))	5.8	8.8	1.53	15.0	0.90
p(TETA/CBA(10k))	12.0	13.1	1.09	13.0	0.91
p(TETA/CBA(5k))-PEG ₃₄₀₀	10.2	11.8	1.16	9.80	--
bPEI25k	16.4	21.0	1.28	35.2	--

Example 5

The ability of poly(TETA/CBA(1k)), poly(TETA/CBA(5k)), poly(TETA/CBA(10k)), and poly(TETA/CBA(5k))-PEG₃₄₀₀ to protect plasmid DNA from serum nucleases was determined.

Polyplexes were prepared as described above, except that poly(TETA/CBA(1k)), poly(TETA/CBA(5k)), poly(TETA/CBA(10k)), and poly(TETA/CBA(5k))-PEG₃₄₀₀ were used instead of unfractionated poly(TETA/CBA). The polyplexes and a plasmid DNA control (not complexed with a poly(TETA/CBA) polymer) were then exposed to serum nucleases (20% FBS; HyClone, Logan, Utah) for 0, 1, 3, 6, 12, 16, or 24 hours. Then, the polyplexes were subjected

to gel electrophoresis (150 ng of pDNA per lane) in TAE buffer (40 mM Tris-HCl, 1% (v/v) acetic acid, 1 mM EDTA). Images were visualized with an image analyzer equipped with UV transilluminator (GelDoc, BioRad, Hercules, California) after ethidium bromide staining. FIG. 7 shows that the polymers successfully protected the plasmid DNA from nuclease degradation for 24 hours.

In another experiment, control plasmid DNA and polyplexes formed with unfractionated poly(TETA/CBA) synthesized at 30°C or 50°C were exposed to serum nucleases in 20% FBS, as described above. FIG. 8 shows that the denser architecture formed at 50°C provides better protection from serum nucleases than does the architecture formed at 30°C.

Example 6

The particle size and zeta potential of polyplexes formed from plasmid DNA and poly(TETA/CBA(1k)), poly(TETA/CBA(5k)), poly(TETA/CBA(10k)), and poly(TETA/CBA(5k))-PEG₃₄₀₀ at various w/w ratios were determined on a Brookhaven Instruments Corp. (Holtsville, New York) ZetaPALS. Values for the particle size are effective mean diameters ($n = 3$ SEM). These results are shown in FIGS. 9 and 10.

Dynamic light scattering (DLS) measurements determined using poly(TETA/CBA(1k)) (FIG. 11), poly(TETA/CBA(5k)) (FIG. 12), poly(TETA/CBA(10k)) (FIG. 13), and poly(TETA/CBA(5k))-PEG_{3.4k} (FIG. 14).

Example 7

Poly(TETA/CBA)-*g*-PEG-RGD was synthesized using poly(TETA/CBA) prepared according to the procedure of Example 1 and commercially available vinyl sulfone-PEG-RGD (VS-PEG-RGD). The reaction scheme is illustrated in FIG. 15.

Example 8

Branched PEG products, such as bPEG2000-COH, have been shown to enhance protection of proteins and polymers similar

to high molecular weight PEG products. These conjugates have achieved similar results, while maintaining low molecular weights. Branched PEG2000 was conjugated to poly(TETA/CBA) to generate an improved copolymer system for gene delivery. The synthesis of bPEG-g-TETA/CBA is shown in FIG. 16.

Example 9

The particle sizes and zeta potentials of poly(TETA/CBA), poly(TETA/CBA)-PEG₃₄₀₀, and poly(TETA/CBA)-PEG-RGD were determined on a Brookhaven Instruments Corp. (Holtsville, New York) ZetaPALS. Values for the particle size are effective mean diameters. The results are shown in FIGS. 17 and 18.

Example 10

PC-3 cells were seeded in a 24-well plate at a density of 4.0×10^4 cells/well and incubated for 24 hrs. Poly(TETA/CBA), poly(TETA/CBA)-PEG₃₄₀₀ and controls PEG-PEI and no polymer were incubated with the cells for 4 hrs in serum-free medium followed by 20 hrs in complete medium. MTT solution (50 μ L, 2 mg/mL) was then added and cells were further incubated for 2 hrs. The medium was removed and 300 μ L DMSO was then added to each well. The absorption was measured at 570 nm using a microplate reader (Model 680, Bio-Rad Lab, Hercules, California). The percentage relative cell viability was determined relative to control (untreated) cells, which were not exposed to transfection system and taken as 100% cell viability. All cytotoxicity experiments were performed in triplicate. Results are shown in FIG. 19.

Example 11

Transfection was evaluated on PC-3 cells using the plasmid, pCMV-Luc, as a reporter. Cells were maintained in DMEM containing 10% FBS, streptomycin (100 μ g/mL) and penicillin (100 units/mL) at 37°C in a humidified atmosphere with 5% CO₂. Cells were seeded 24 hrs prior to transfection in 24-well plates at initial densities of 4.0

× 10⁴ cells/well. DNA was complexed with the poly(TETA/CBA) and PEG-poly(TETA/CBA) polymers. Controls were PEG-PEI and no polymer. Complexes were made in HEPES buffer and incubated for 30 min before use. At the time of

5 transfection, the medium in each well was replaced with fresh serum-free medium. Polyplexes (0.5 µg DNA/well) were incubated with the cells for 4 hrs at 37°C. The medium was then replaced with 500 µL of fresh complete medium and cells were incubated for additional 44 hrs. The cells were

10 then washed with pre-warmed PBS, treated with 200 µL cell lysis buffer and subjected to a freezing-thawing cycle. Cellular debris was removed by centrifugation at 14,000 g for 5 min. The luciferase activity in cell lysates (25 µL) was measured using a luciferase assay kit (100 µL

15 luciferase assay buffer) on a luminometer (Dynex Technologies Inc., Chantilly, Virginia). The relative luminescence unit (RLU) of luciferase expression was normalized against protein concentration in the cell extracts, measured by a BCA protein assay kit (Pierce,

20 Rockford, Illinois). All transfection assays were carried out in triplicate. Results are shown in FIG. 20.

CLAIMS

The subject matter claimed is:

1. A composition comprising polyethylene glycol covalently bonded to poly[H₂N-(CH₂-CH₂-N)_n-H/CBA].
- 5 2. The composition of claim 1 wherein n is 1, 2, or 3.
3. The composition of claim 1 wherein n is 3.
4. The composition of claim 1 wherein the polyethylene glycol is linear.
- 10 5. The composition of claim 4 wherein the polyethylene glycol has a molecular weight of about 1,000 to about 50,000.
6. The composition of claim 5 wherein the polyethylene glycol has a molecular weight of about 2,000
15 to about 25,000.
7. The composition of claim 6 wherein the polyethylene glycol has a molecular weight of about 3,400.
8. The composition of claim 4 wherein the poly[H₂N-(CH₂-CH₂-N)_n-H/CBA] has a molecular weight of about 1,000 to
20 about 25,000.
9. The composition of claim 4, further comprising an RGD peptide covalently bonded to the polyethylene glycol.
10. The composition of claim 9 wherein the RGD peptide comprises SEQ ID NO:1.
- 25 11. The composition of claim 1 wherein the polyethylene glycol is branched.
12. A complex comprising a nucleic acid ionically bonded to a composition comprising polyethylene glycol covalently bonded to poly[H₂N-(CH₂-CH₂-N)_n-H/CBA].
- 30 13. The complex of claim 12 wherein the nucleic acid comprises a plasmid.
14. The complex of claim 12 wherein the nucleic acid comprises an siRNA.
15. The complex of claim 12 wherein the nucleic acid
35 is an oligonucleotide.

16. A method of transfecting a cell with a selected nucleic acid, the method comprising contacting the cell with a complex comprising the selected nucleic acid ionically bonded to a composition comprising polyethylene glycol covalently bonded to poly[H₂N-(CH₂-CH₂-N)_n-H/CBA] such
5 that the complex enters the cell.

17. The method of claim 16 wherein n is 1, 2, or 3.

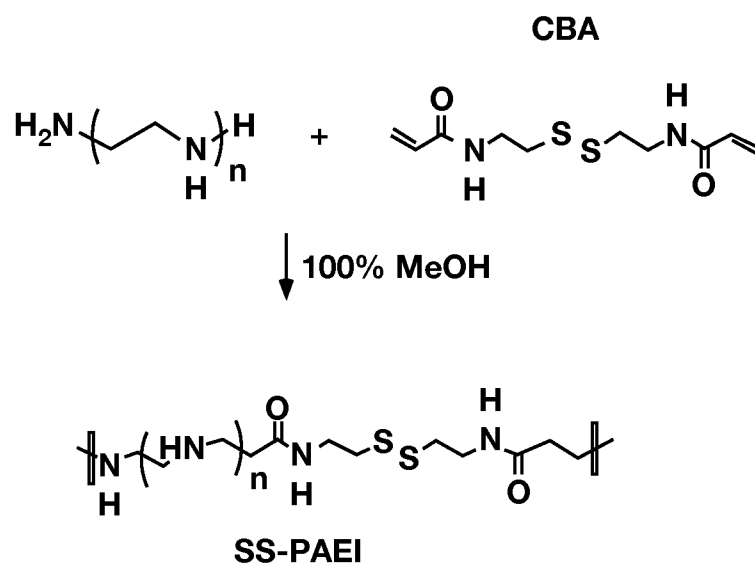
18. The method of claim 16 wherein the selected nucleic acid comprises a plasmid.

10 19. The method of claim 16 wherein the selected nucleic acid comprises an siRNA.

20. The method of claim 16 wherein the selected nucleotide comprises an oligonucleotide.

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**FIG. 1**

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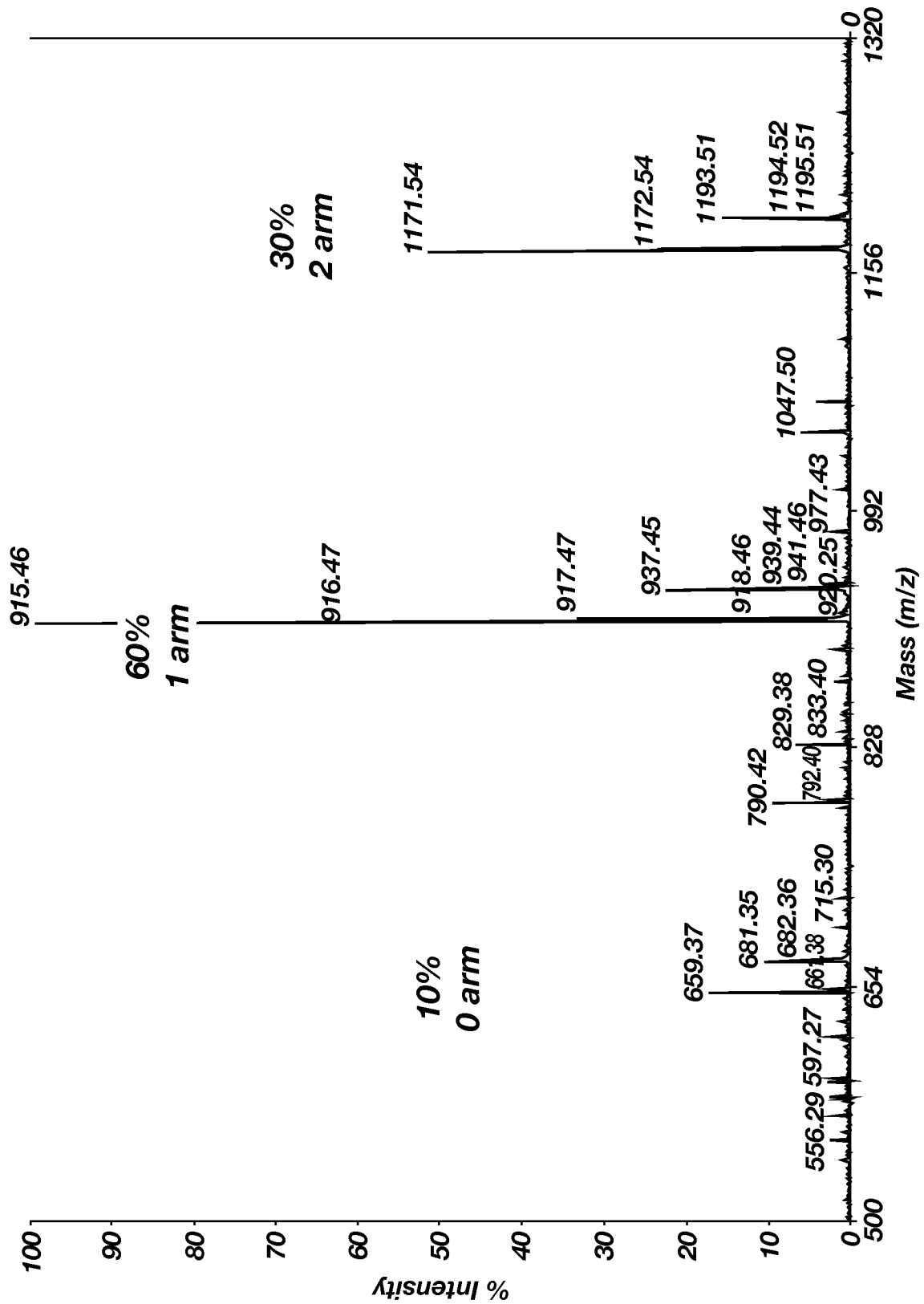


FIG. 2

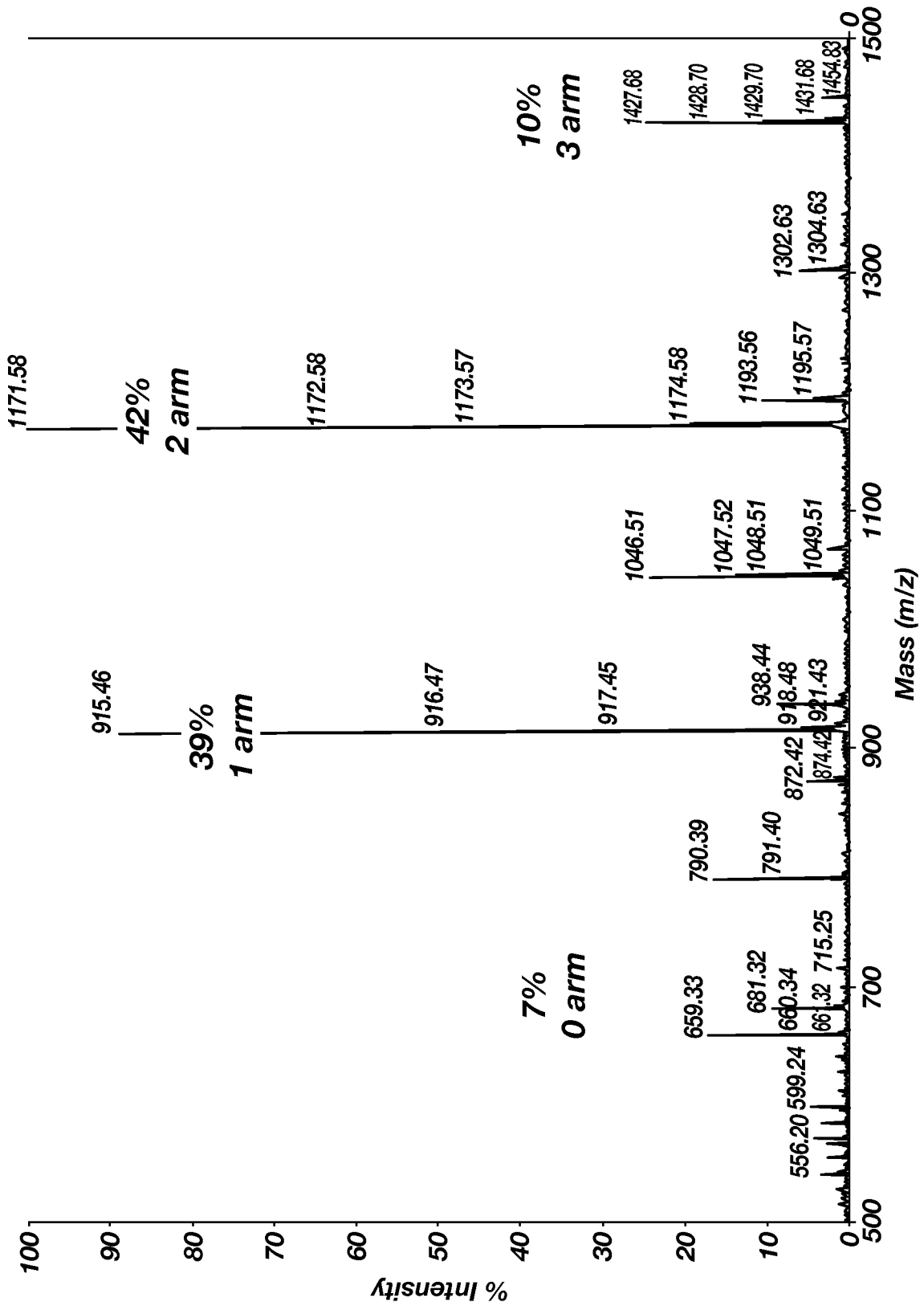


FIG. 3

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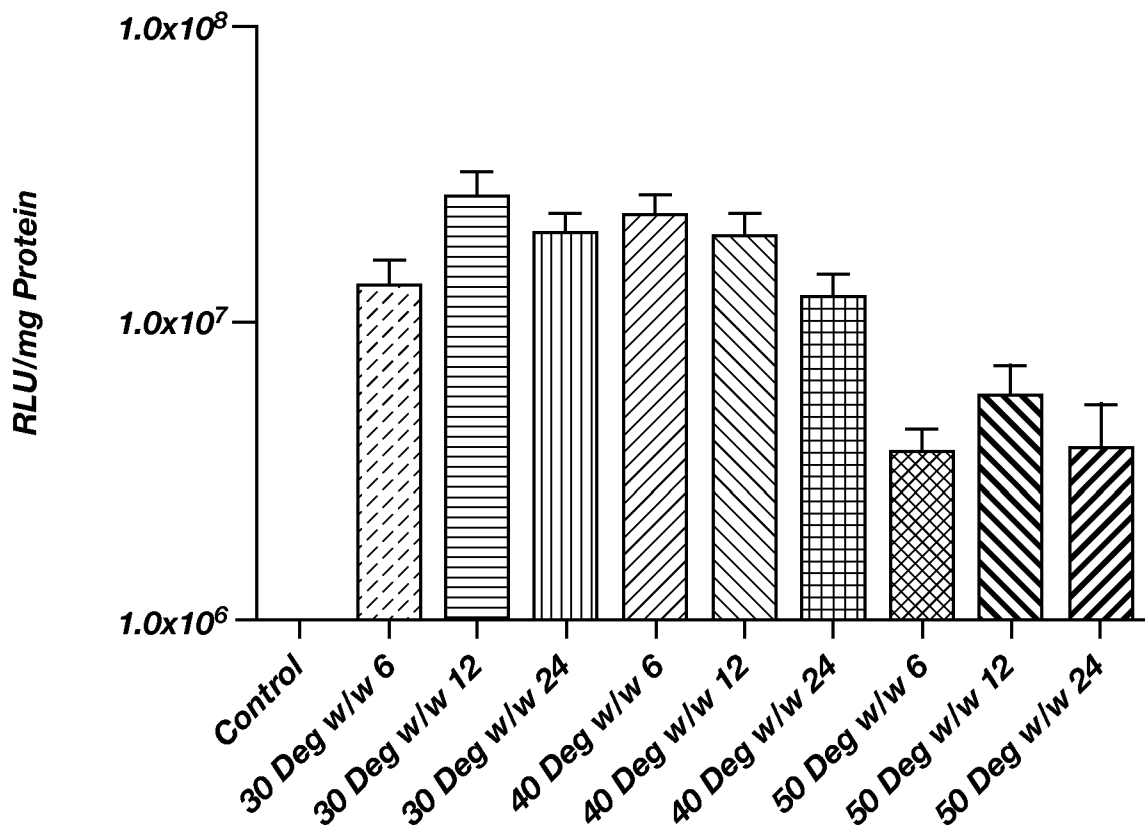


FIG. 4

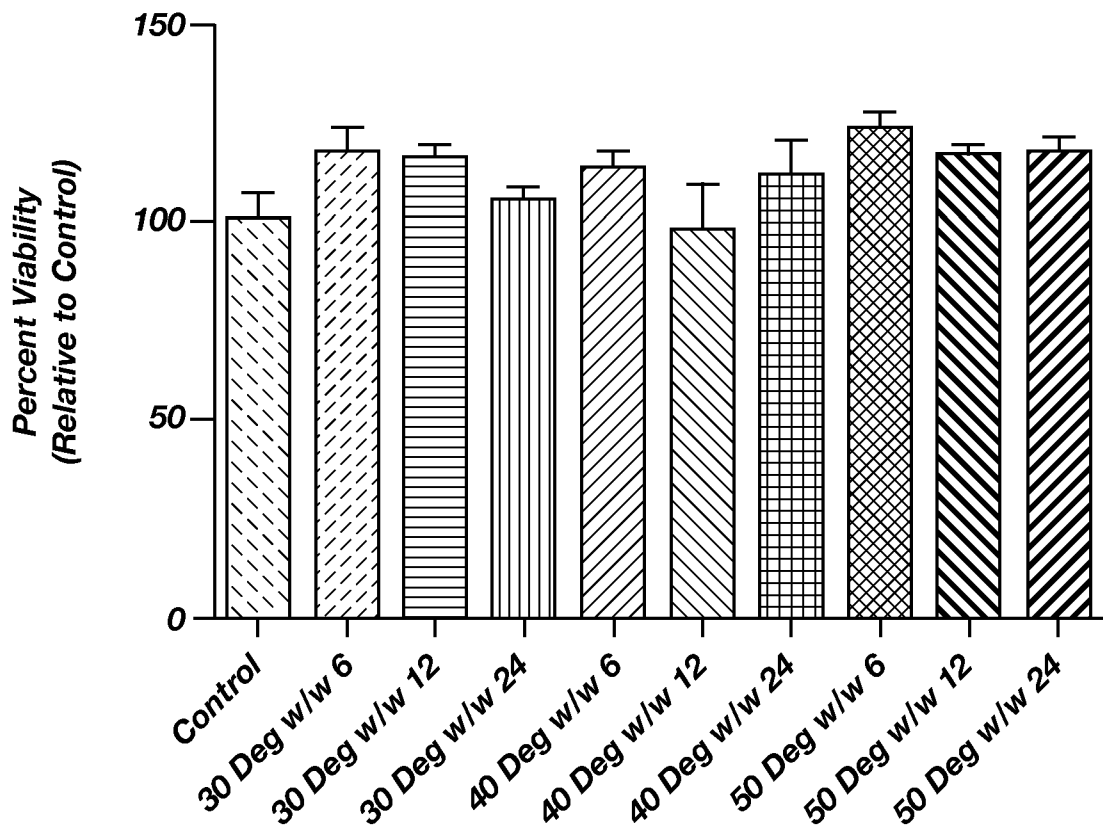


FIG. 5

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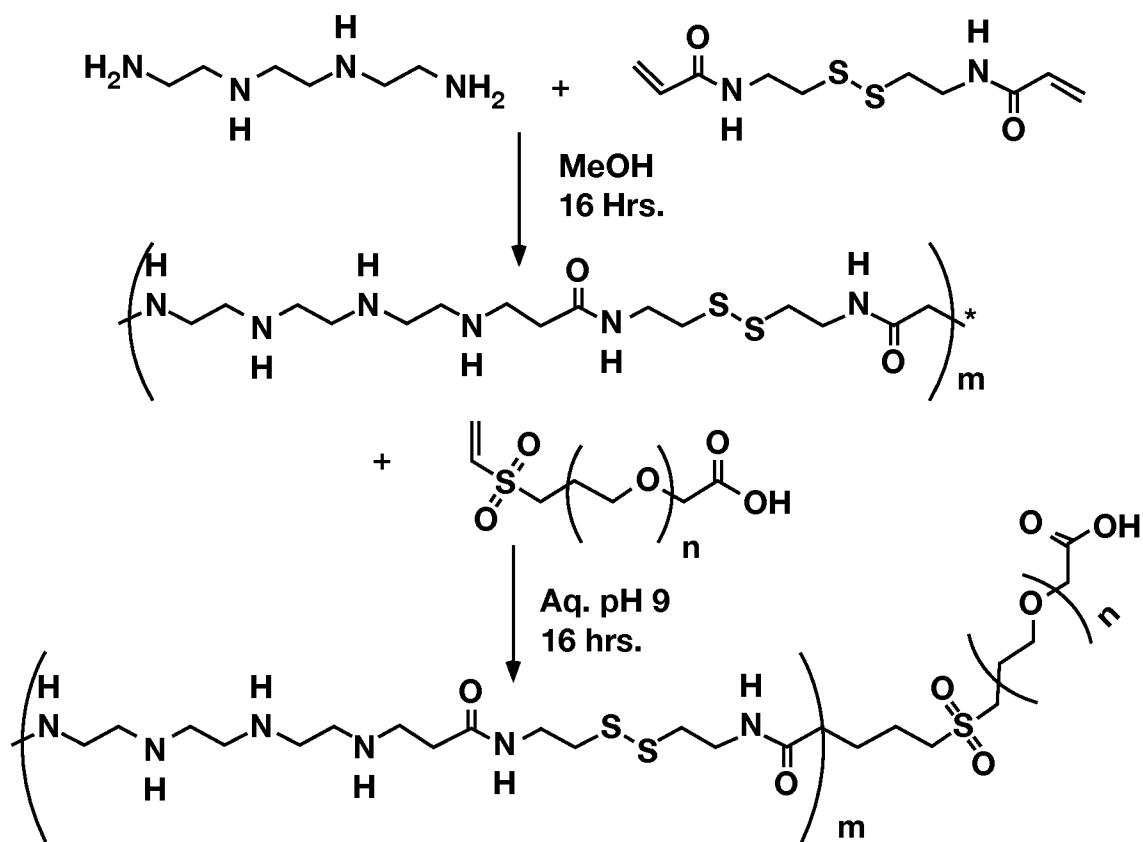


FIG. 6

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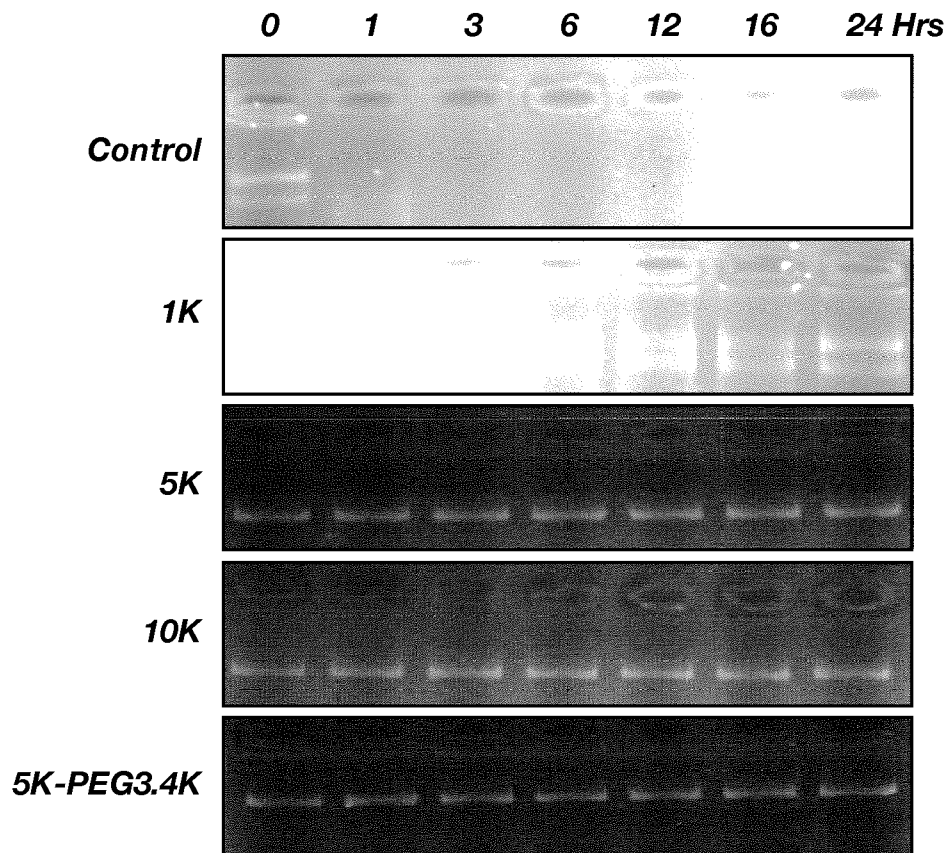


FIG. 7

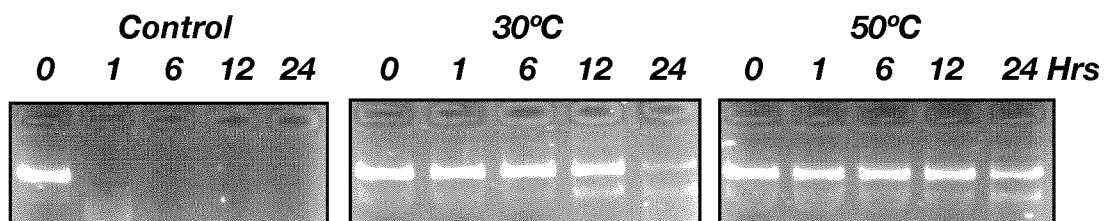


FIG. 8

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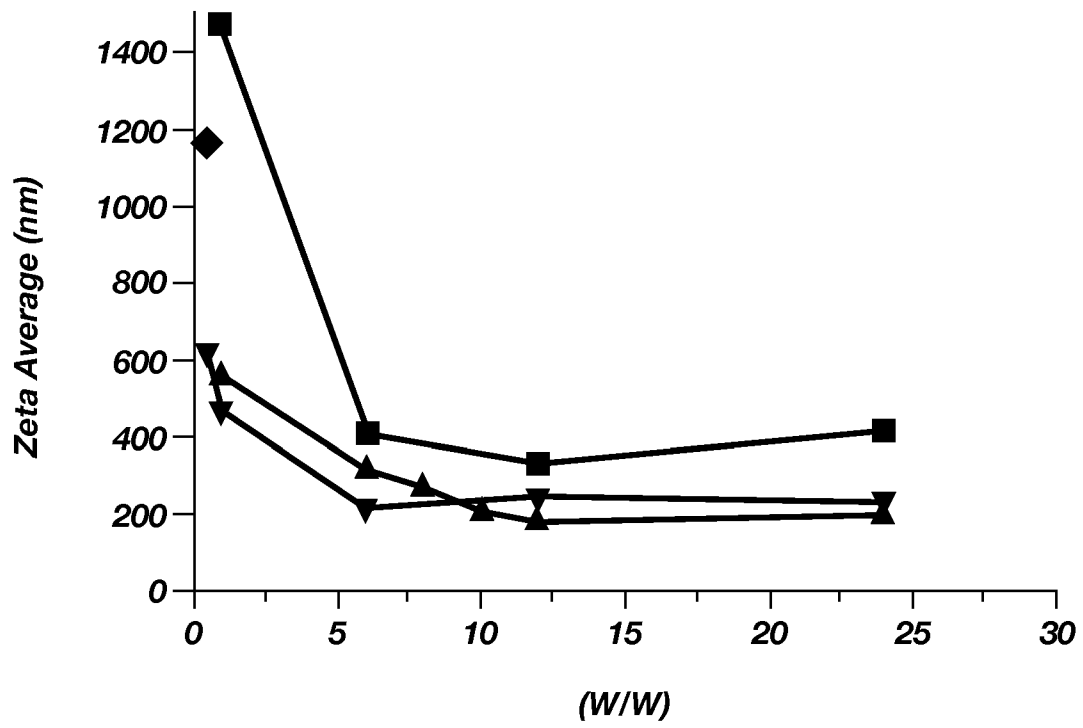


FIG. 9

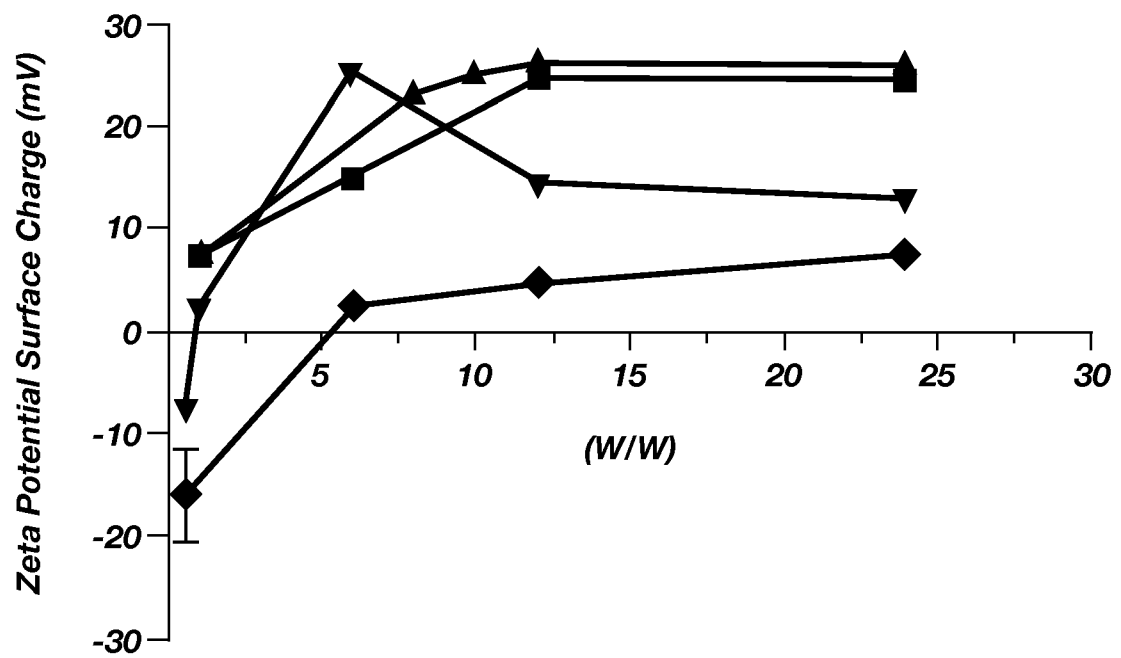


FIG. 10

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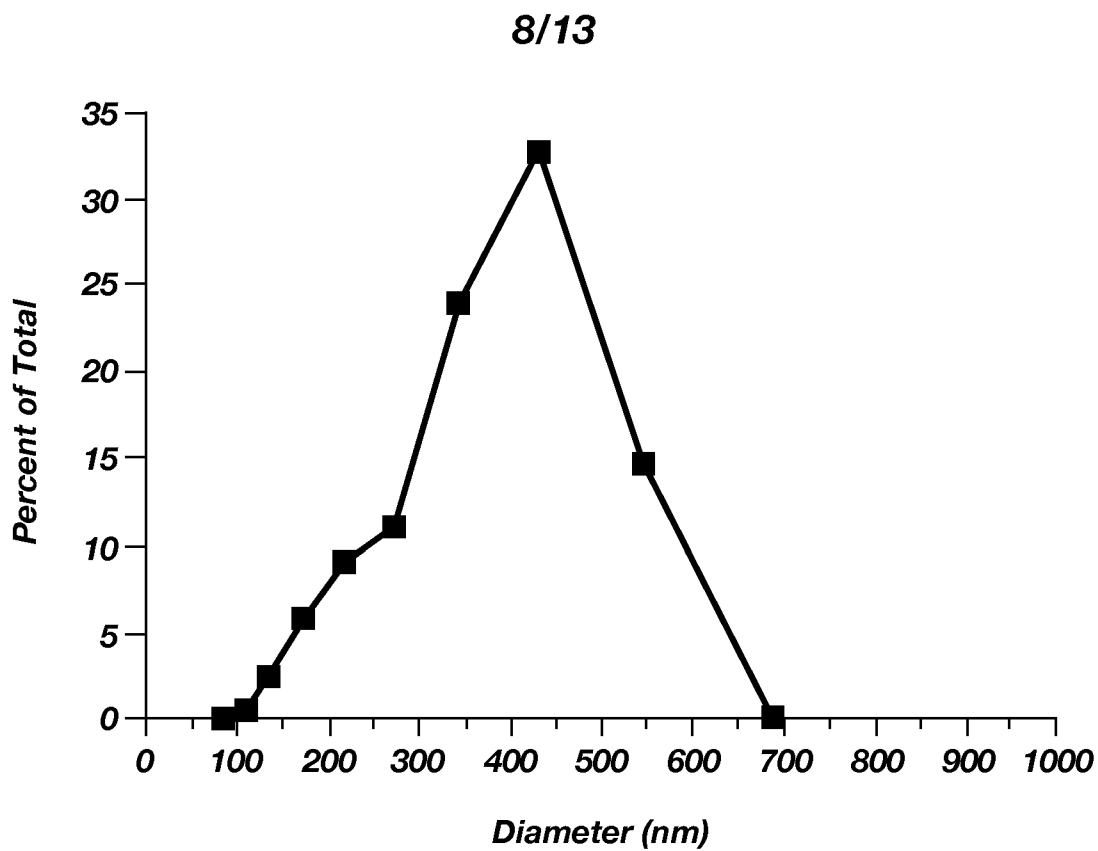


FIG. 11

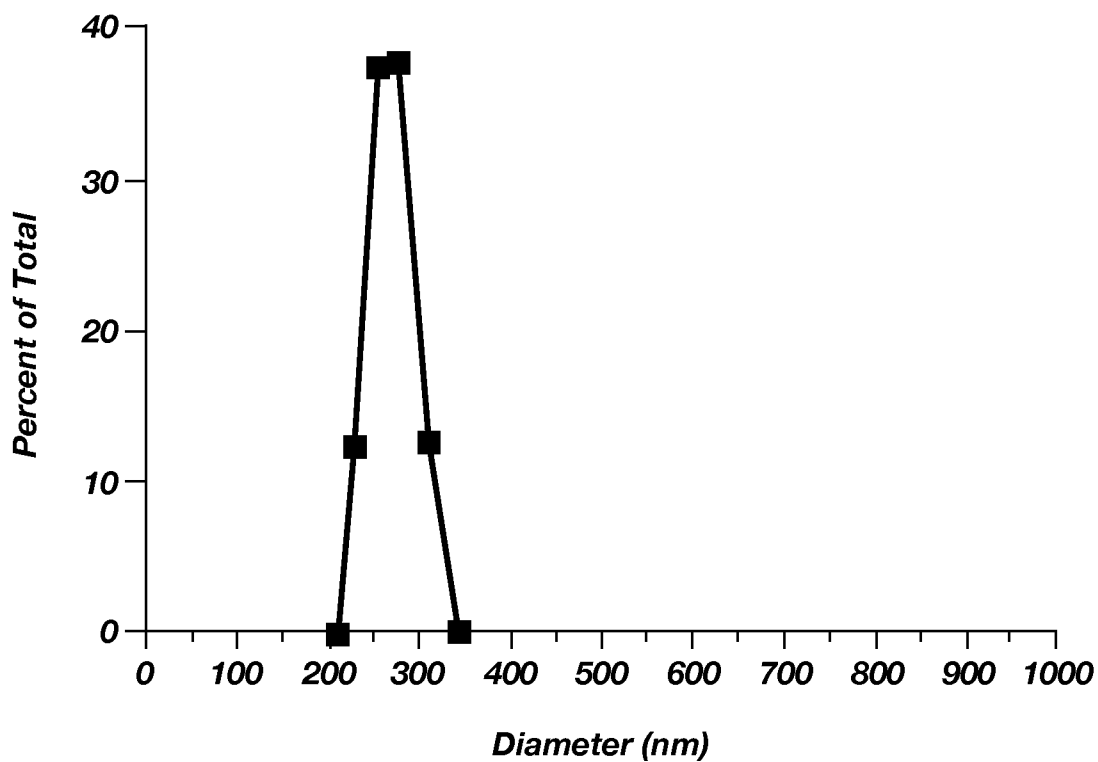


FIG. 12

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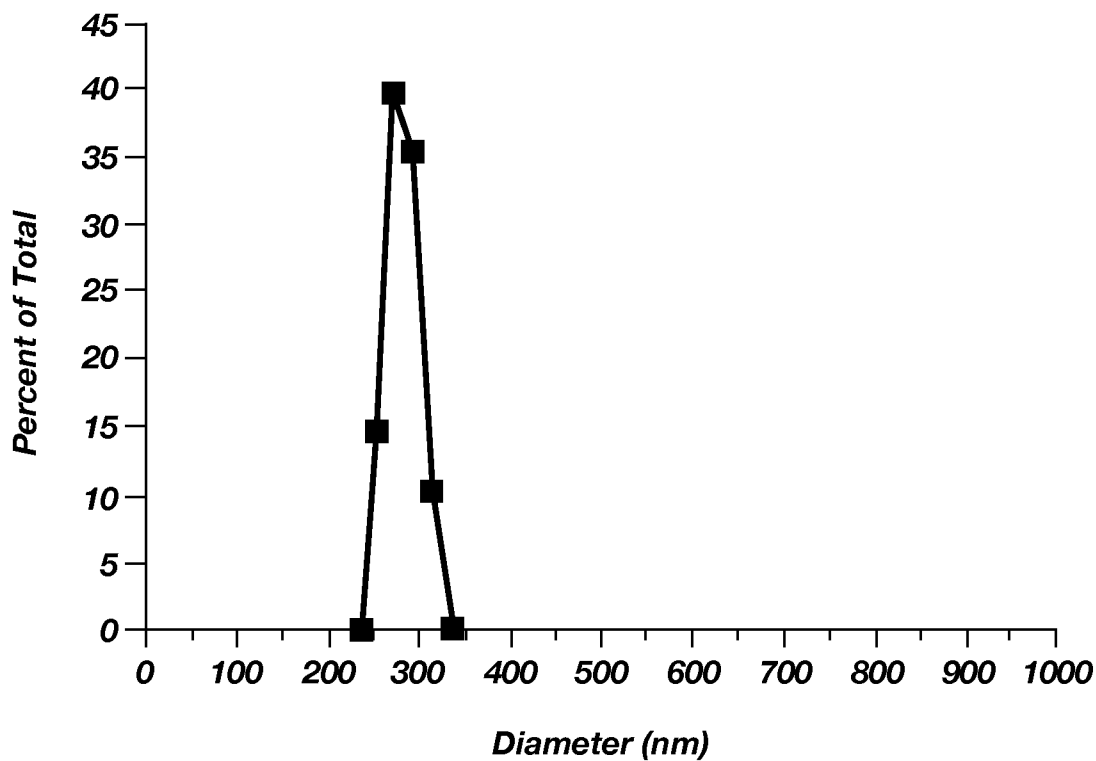


FIG. 13

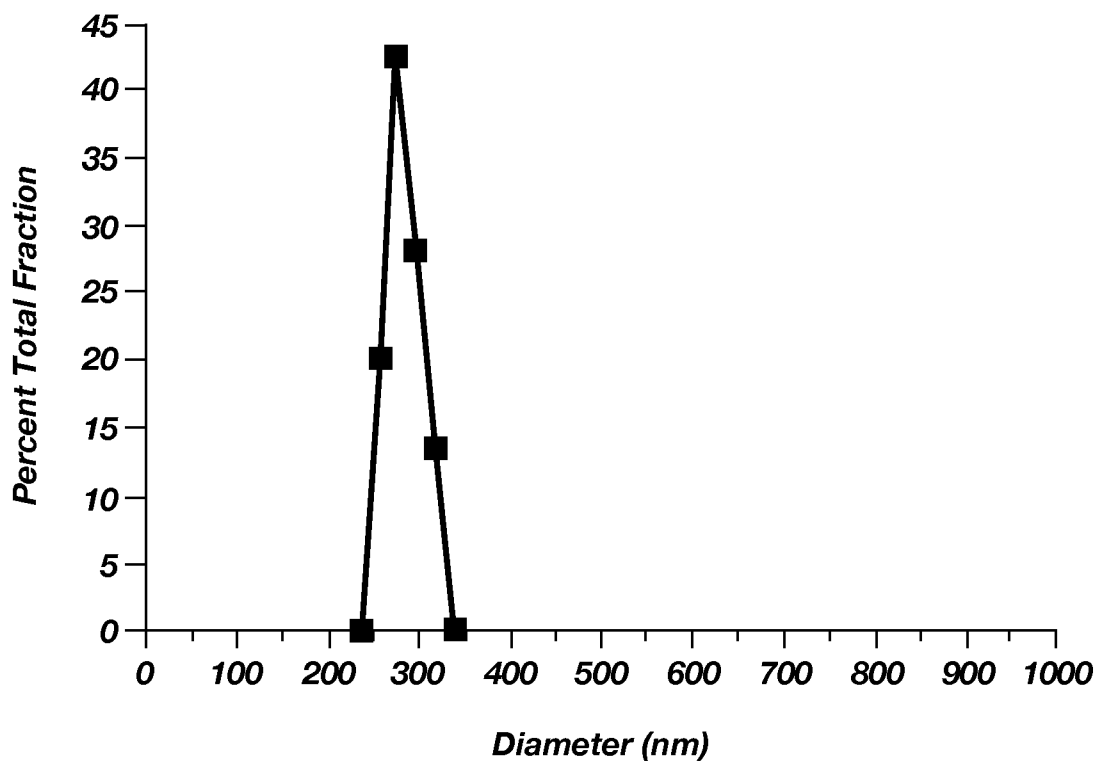


FIG. 14

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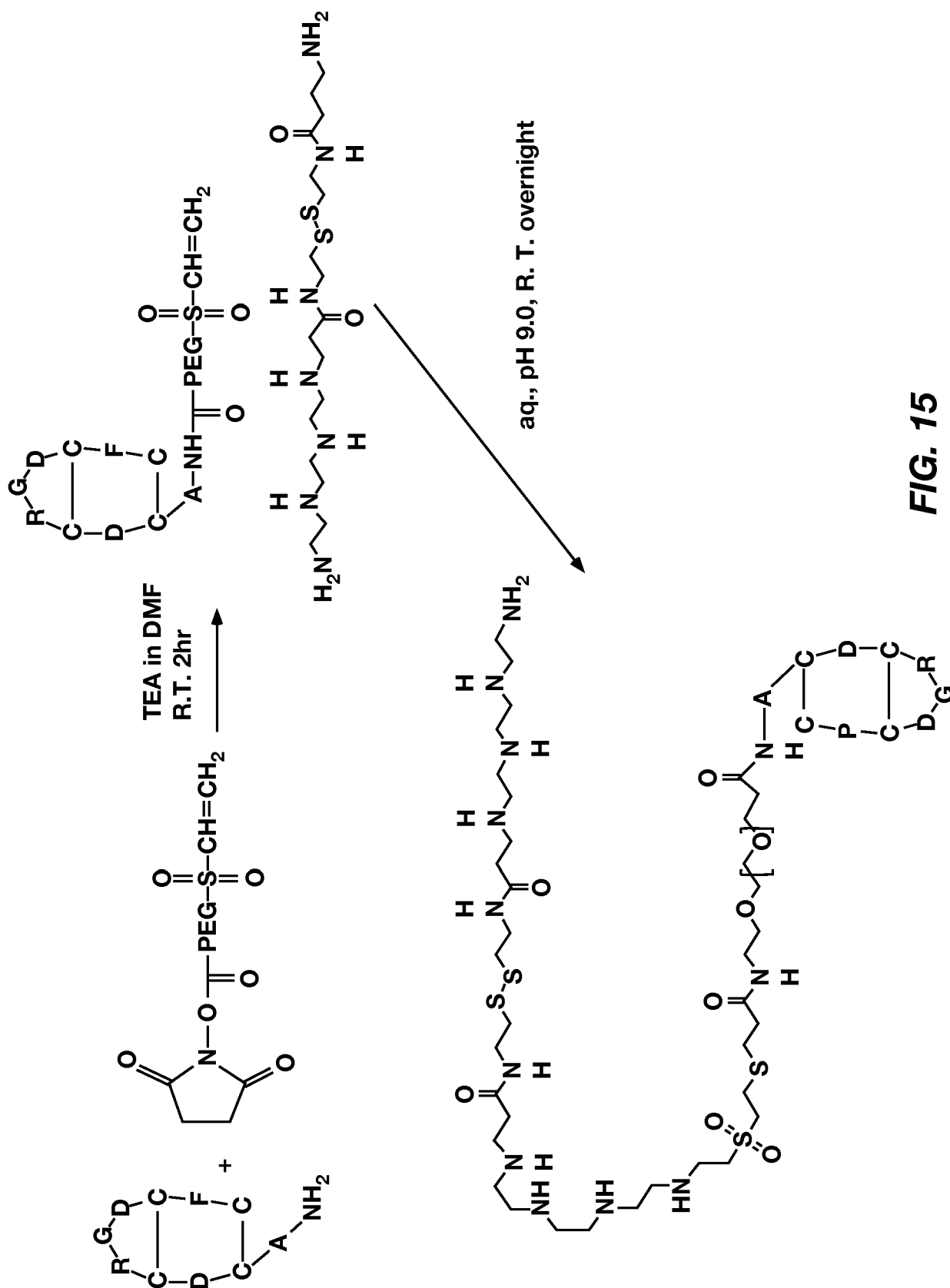


FIG. 15



FIG. 16

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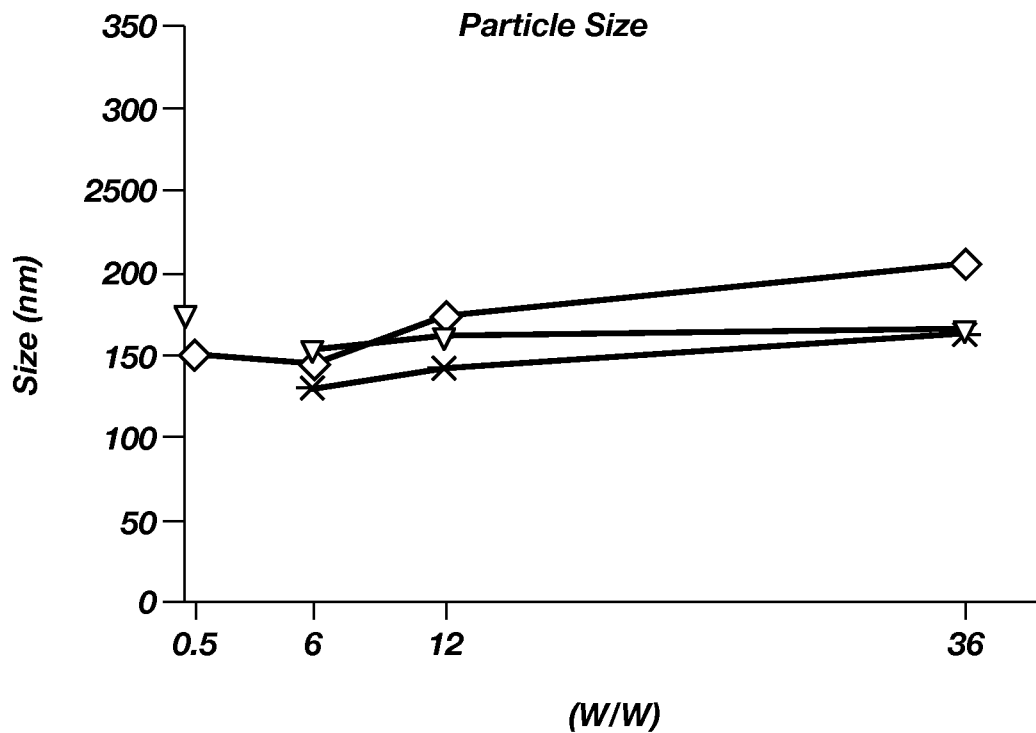


FIG. 17

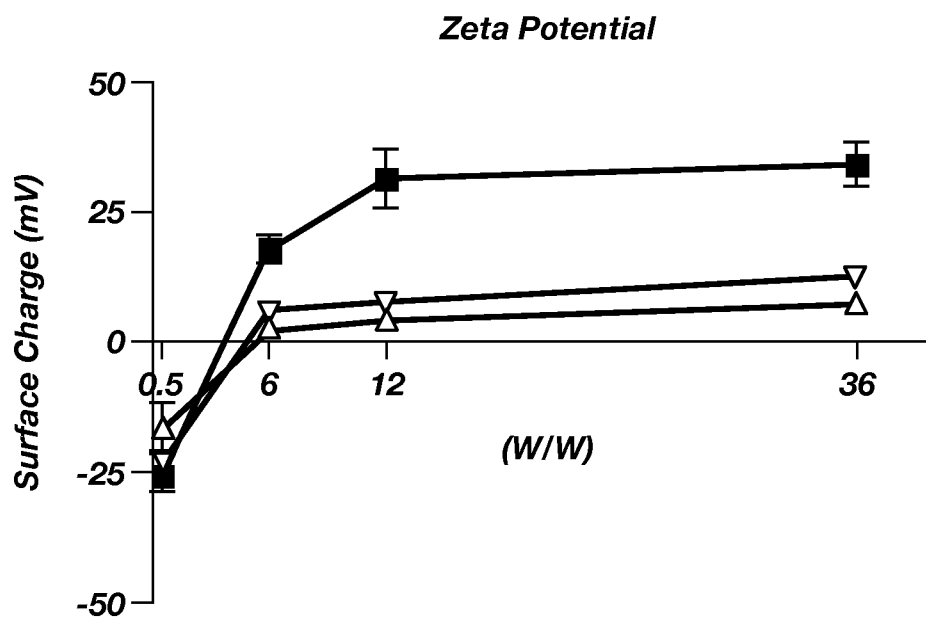
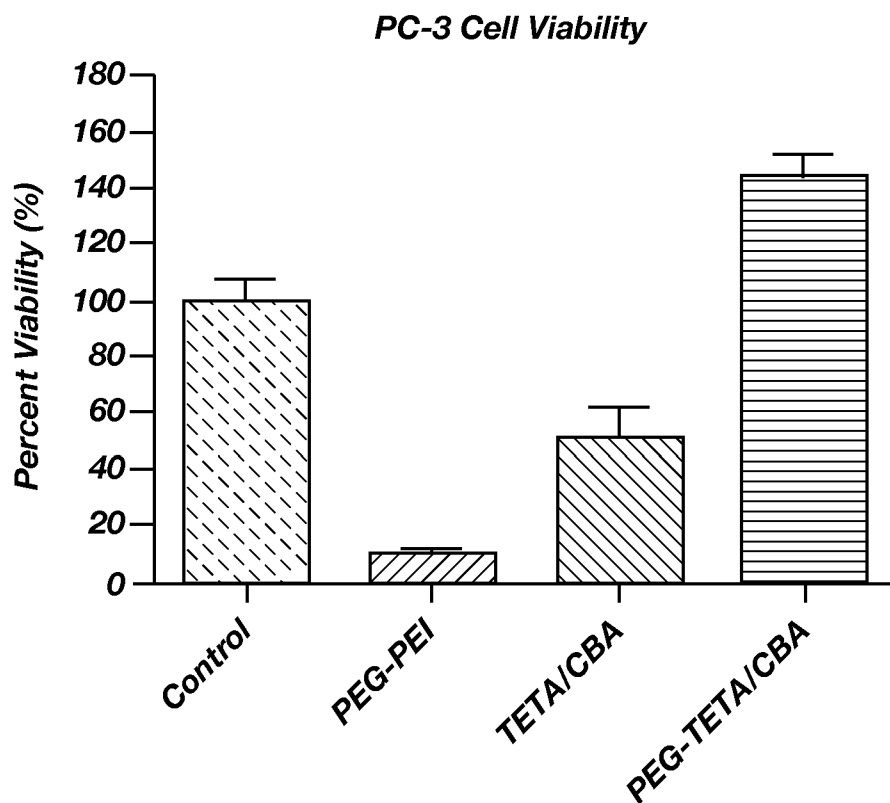
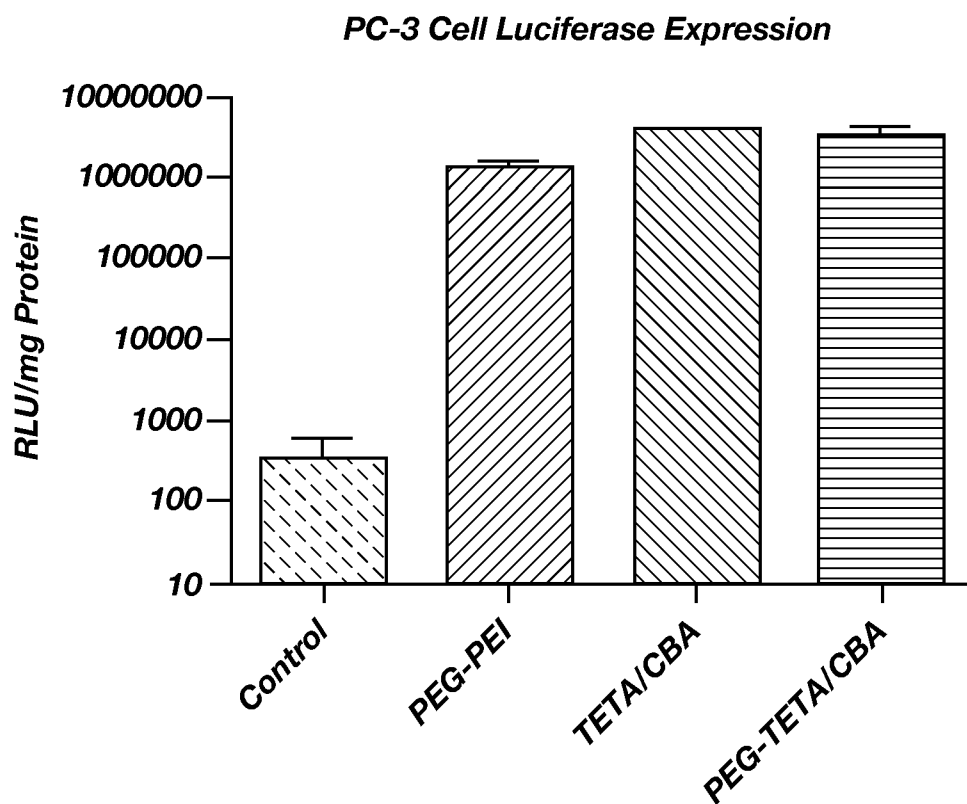


FIG. 18

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**FIG. 19****FIG. 20**