

Quaternized Polyamidoamine Dendrimers as Novel Gene Delivery System: Relationship between Degree of Quaternization and Their Influences

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Quaternary ammonium groups were introduced to Starburst polyamidoamine (PAMAM) dendrimers for a gene carrier. These quaternary dendritic carriers exhibited reduced cytotoxicity on 293T cells compared to parent dendrimers examined and their transfection efficiency were similar with parent dendrimers. Quaternization could be a promising tool to improve properties of dendrimers as a gene delivery carrier.

Key Words : Gene delivery, PAMAM, Quaternary amine, Cytotoxicity

Introduction

Dendrimers are highly branched polymers with well-defined and three-dimensional structure. Starburst polyamidoamine (PAMAM) dendrimers consist of tertiary amines in core, primary amines on a surface and amide backbone.¹ They have the captured interest of researchers because of successful use for nonviral gene delivery^{2,3} and commercially available polymers as whole (amine-terminated groups) or half (carboxylate-terminated groups) generation. Basic dendrimers *e.g.*, poly(propylene imine) and PAMAM, were modified to quaternary amine groups for a pH-sensitive controlled release system,⁴ a dendritic anion conductor,⁵ effective antimicrobials⁶ and a photosensitizer for photodynamic therapy.⁷ Here, we report starburst PAMAM dendrimers (generation 4, containing 62 interior tertiary amines and 64 exterior primary amines) possessing quaternary amine groups as novel nonviral gene delivery system. An important feature of quaternized PAMAM (QPAM) dendrimers was that primary amines which had significant toxic effect to cell membrane⁸⁻¹¹ could be decreased. Exchange of primary amines and tertiary amines to quaternary ammonium salts could be expected to reduce the cytotoxicity giving hydrophilic nature. Moreover, their charge density could be increased by methylation of interior tertiary amine groups so that the amount for complexation with DNA could be reduced.

Materials and Methods

Materials. PAMAM G4 (Starburst), Methyl iodide, anhydrous N,N-dimethylformamide (DMF) and PEI (average molecular weight 25 kDa) were purchased from Aldrich (Milwaukee, WI). PGL3-control vector (plasmid DNA) was purchased from Promega (Madison, WI). Fetal bovine serum (FBS) and Dubecco's modified Eagle's medium (DMEM) were purchased from GIBCO (Gaithersburg, MD).

Synthesis of partially quaternized PAMAM (QPAM).

The solvent (methanol) dissolved PAMAM was evaporated and dried prior to reaction. PAMAM (0.1 g, 0.7 mmol) dissolved in DMF (0.5 mL) were added to appropriate methyl iodide (0.2 equiv., 0.5 equiv. and 0.8 equiv. per mole of the possible number of tertiary amino groups and primary amino groups) diluted in DMF (0.5 mL) and stirred for 24 h at room temperature or 37 °C. The resulting clear solution was precipitated in diethyl ether and the residues were vacuum-dried for overnight. After dried residues were redissolved in 1 mL of water, the solution was placed into a dialysis membrane (SpectraPor, MwCO 6000-8000), and dialyzed against 2 M NaCl and pure water in succession. Purified QPAMs were dried and obtained as white powder. ¹H NMR (300 MHz, D₂O) δ 2.46 (br m, CH₂CO), 2.65-3.1 (br m, NCH₂CH₂ and CH₂CH₂CONH₂), 3.17-3.22 (br s, CH₃), 3.47-3.32 (br m, CH₂N⁺), 3.71 CONHCH₂). The level of quaternization was determined by ratio intergration of peak at 3.2 ppm versus theoretically calculated integration of full quaternization.

Atomic force microscopy (AFM). Atomic force microscopy (Nanoscope IIIa system, Digital Instruments, Inc., Santa Barbara, CA) was used for imaging the morphology of complexes at 4 : 1 (N/P or +/-) ratio. Complex were formed at 1 µg/mL plasmid DNA concentration in water for 30 min. Samples were applied to freshly cleaved mica and absorbed on the mica for 5 min. After excess fluid on the mica was wicked off using filter paper, samples on mica were dried at room temperature prior to imaging. The image mode was set to tapping mode and average scan speed was 1.5 Hz.

Dynamic light scattering (DLS) measurement. The size of complexes was determined using a BI-200SM Goniometer (Brookhaven instruments corporation, Holtsville, NY, USA) with a Lxel laser model 95 argon laser (100 mW output power at a wavelength of 514.5 nm). Correlator, PD2000 (Precision Detectors) was used and the scattering angle was 90 °C.

Cytotoxicity assay. MTT assay was performed for the cytotoxicity assay. 293T cells (1 × 10⁴ cells/well) were seeded in 96-well plate and grown in 95 µL DMEM containing 10% FBS, supplying 5% CO₂ at 37 °C for overnight. Cells were exposed QPAMs, PAMAM G4 and PEI 25 kDa as controls

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with various concentration for 1 day. After that, they were added 25 μL of MTT stock solution (5 mg/mL) and incubated for 2 h, then added 100 μL of extraction buffer (20% w/v of SDS in 50% DMF, pH 4.7). Absorbance was measured at 570 nm after overnight incubation.

Cell transfection. 293T cells were seeded in 24-well plate at a density of 5×10^4 cells/well and grown in DMEM containing 10% FBS for 1 day. Two μg of DNA per well was complexed dropping into QPAMs in FBS free DMEM for 30 min at room temperature. Complexes in FBS free DMEM were transfected to the cells with replacing old medium. Following 4 h incubating cells, culture medium was exchanged to fresh DMEM containing 10% FBS. After incubating for 2 days, cells were washed with PBS with removing the growth medium and lysed for 30 min at room temperature adding reporter lysis buffer. Luciferase activity was measured in a LB 9507 luminometer (Berthold, Germany) and protein content was measured by BCA assay (Pierce, Rockford, IL).

Results and Discussion

Synthesis of QPAMs. QPAMs (Figure 1) with various degrees of quaternization were synthesized by partial methylation of tertiary amine groups and primary amine groups. The level of quaternization was determined by the method proton integration values at 3.2 ppm in ^1H NMR (300 MHz, D_2O) that were divided by theoretically calculated integration values of fully quaternized dendrimers. Approximately 10% (0.1-QPAM), 50% (0.5-QPAM) and 70% (0.7-QPAM) of quaternary ammonium salts were produced at PAMAM G4.

Characterization of QPAMs-complexed polyplexes. To visualize their morphology of the polyplexes between QPAMs and DNA, the particles were observed by atomic force microscopy (AFM) and the images are shown in Figure 2. The single population particles were formed respectively, and their shapes were assumed to be several tens nano-sized spherical particles.

Wolfert *et al.* reported that cationic polymers containing quaternary ammonium groups showed efficient complex formation with smallest diameters, in according to studying polymers possessing among primary, tertiary and quaternary amino groups on the properties of complexes formed with DNA.¹² To verify that QPAM/DNA complexes are strong and efficient enough, the particle size of these complexes were performed using dynamic light scattering (DLS). Figure 3 shows the particle size of QPAM dendrimers prepared in water. Interestingly, relatively large particles were detected at N/P ratios of 0.5 (0.1-QPAM), 2 (0.5-QPAM) and 1 (0.7-QPAM). Probably at these ratios, QPAM molecules functioned as a crosslinker for plamid molecules yielding large aggregates. This is seen from the neutralization between DNA and QPAMs by canceling the net charge¹³ and the resulting minimized the charge-to-charge repulsion between complex particles might cause the large particles.¹⁴ Increase of dendrimers/DNA values up to 10

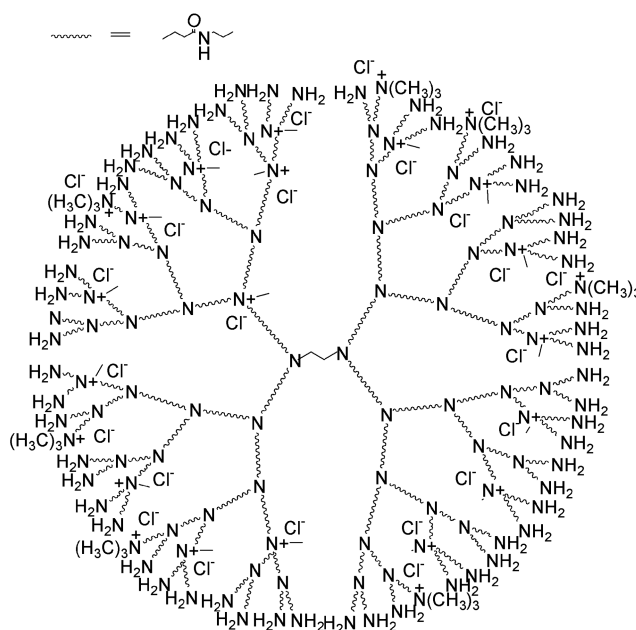


Figure 1. The structure of QPAM. Primary amines and tertiary amines of PAMAM were converted into quaternary amine groups at a degree of 10% (0.1Q-PAM), 50% (0.5Q-PAM) and 70% (0.7Q-PAM).

resulted in recharging of the polyplexes and decreasing in their size as small as 56 nm.

Cytotoxicity issue depends on the degree of quaternization. Cell viability studies were performed on 293T cells using an MTT assay.¹⁵ Relative cell viability (%) versus dendrimer concentration is shown in Figure 4. The minimum viability of cells exposed to the solution of QPAMs at various concentration was 75% relative to controls, indicating low cytotoxicity of QPAMs compared to PAMAM and PEI. The cells exposed all the QPAMs were more viable than those of parent dendrimers, and a positive correlation between cell viability and degree of quaternization was detected; PAMAM < 0.1-QPAM < 0.5-QPAM < 0.7-QPAM with cell viability in according to the degree of quaternization. In this study, PEI was found to be polymer with the highest cytotoxicity. The polycations characterized rigid and global structure (e.g. PAMAM dendrimers) were found to be low toxicity because of the difficulties to attach to membranes or to uptake to cells causing cytotoxicity, whereas polymers with linear or branched and flexible structure (e.g. PEI) showed higher cell damaging effects.⁹ It was also reported that the quaternization of the PEI polymer resulted in a decrease of toxicity because transformation of an water soluble polymer resulted in an improved safety profile.¹⁶ Quaternary ammonium groups are always charged and strongly hydrophilic thus, unmodified PAMAM dendrimers composed of primary amine groups and tertiary amines groups may not be as hydrophilic as modified PAMAM with quaternary ammonium groups. These quaternary ammonium groups modified dendrimer-based delivery systems are supposed to be useful for reduction of the toxicity of PAMAM dendrimers.

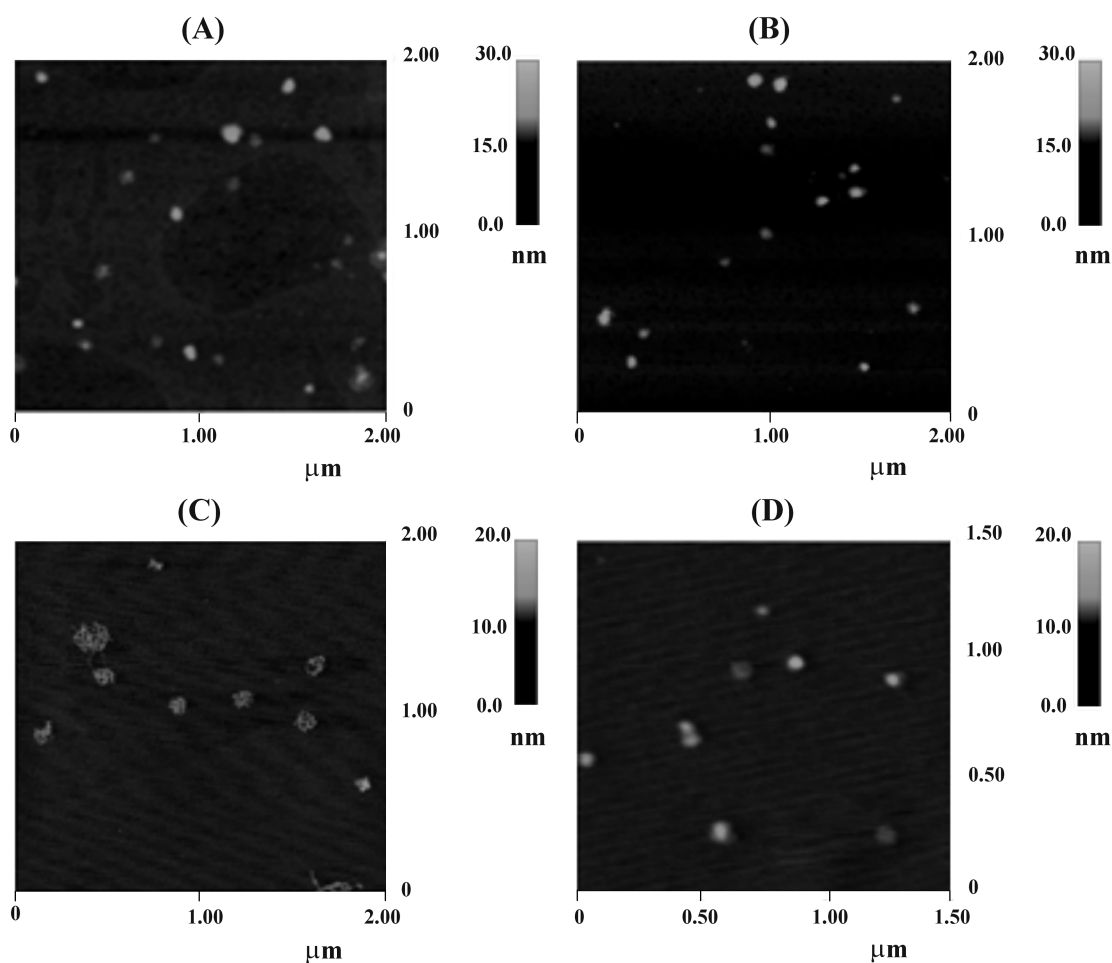


Figure 2. Morphology of particles imaged by atomic force microscopy (AFM) at charge ratio (+/-) = 4.0 or N/P ratio = 4.0. (A) PAMAM G4/plasmid DNA particles. (B) 0.1-QPAM/plasmid DNA particles. (C) 0.5-QPAM/plasmid DNA particles (E) 0.7-QPAM/plasmid DNA particles. The N/P ratio was calculated from the number of amines and of QPAMs and the number of phosphate groups of DNA.

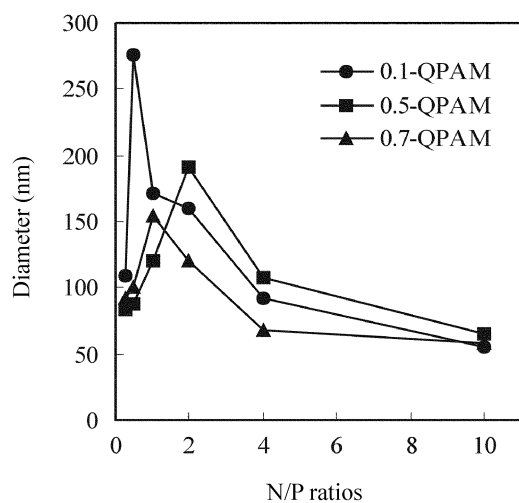


Figure 3. Particle sizes of quaternized PAMAM derivatives determined by dynamic light scattering (DLS). Diameters indicate the sizes which dendrimers formed in 5 μg/mL of plasmid DNA concentration.

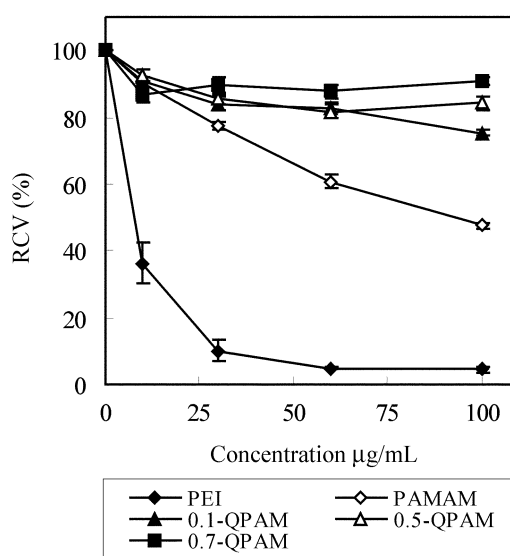


Figure 4. Cytotoxicity assay on 293T cell line for the study of the effect of quaternization to PAMAM G4 and PEI 25 kDa as a control. Relative cell viability (RCV, %) is expressed as a percent of viable cells divided by untreated cells.

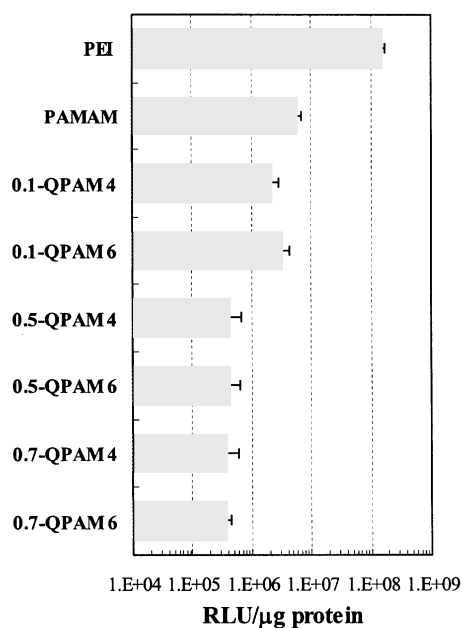


Figure 5. Transfection efficiency on 293T cell line. Cells were transfected with DNA of 2 μ g in the absence of serum for 4 h. Transgene expression was measured 48 h posttransfection. Numbers indicate N/P ratios of dendrimers/DNA. Data are determined in triplicate and expressed relative light unit (RLU) per μ g protein.

Transfection efficiency in vitro. The quaternization effect of PAMAM on transfection is shown in Figure 5. We have transfected 293T cells with plasmid DNA, observing luciferase activities. The levels of expression in the case of 0.1-QPAM were in the same order of magnitude as those obtained with nonmodified PAMAM. Using 0.5-QPAM and 0.7-QPAM, the result of quaternization exhibited an observed approximately 10-fold reduction of transfection levels comparing to PAMAM. Quaternized polycations are so completely charged at any pH as a strong polyelectrolyte that they lack pH responsiveness on the endosome, perhaps via the so-called proton sponge mechanism.¹⁷ Therefore, the reduced transfection activity associated with quaternized polycation may involve relatively poor access to nucleus.

Conclusion

In summary, we have introduced a series of quaternized PAMAM by simple and efficient methylation of PAMAM G4. The complexes were highly efficient in condensation of

plasmid DNA. Although the transfection efficiency of QPAM derivatives except to 0.1-QPAM is lower by an order of magnitude than PAMAM G4, our systems have the merit of much lower cytotoxicity comparing to parent dendrimers by producing hydrophilic nature. The influence of quaternization on the polycations is currently under investigation as our further studies.

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