

RESEARCH ARTICLE

Rational design, fabrication, characterization and *in vitro* testing of biodegradable microparticles that generate targeted and sustained transgene expression in HepG2 liver cells

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Abstract

Poly(lactide-co-glycolide) (PLGA) microparticles have significant potential for sustained delivery of plasmid DNA (pDNA). However, unmodified PLGA microparticles have poor transfection efficiencies. In this study, we use several approaches to enhance the transfection efficiencies of PLGA microparticles in a HepG2 liver cell line. Polyethylenimine (PEI) is used to condense the pDNA prior to loading into the PLGA microparticles. This provides enhanced loading efficiencies and greater protection to the pDNA during the entrapment process. In addition, the pDNA used (ApoE) incorporates a hybrid liver-specific murine albumin enhancer/ α 1 antitrypsin promoter (AlbE/haAT) to enhance transgene expression in human liver (HepG2) cells. The percentage of surfactant used in the preparation of the microparticles, the polymer composition of the PLGA, the ratio of the PEI to pDNA (N/P), the structure of the PEI and the potential utility of a galactose targeting ligand were then investigated to further optimize the efficacy of the cationic microparticle non-viral delivery system in transfecting HepG2 cells. For each PLGA PEI-pDNA microparticle formulation prepared, we evaluated particle size, ζ -potential, loading of pDNA, cytotoxicity, and transgene expression in HepG2 cells and control human embryonic kidney (HEK293) and monkey African green kidney fibroblast-like (COS7) cells. Loading PLGA particles with PEI-ApoE pDNA complexes resulted in a significant reduction in particle size when compared to PLGA microparticles loaded with ApoE pDNA alone. Scanning electron microscopy images showed that all the particle formulations were smooth and spherical in appearance. Incorporation of the cationic PEI in the PLGA particles changed the ζ -potential from negative to positive. Complexing PEI with ApoE pDNA increased the loading efficiency of the ApoE pDNA into the PLGA microparticles. The cytotoxicity of PLGA particles loaded with PEI-ApoE pDNA complexes was similar to PLGA particles loaded with ApoE pDNA alone. The transfection efficiency of all particle formulations prepared with ApoE pDNA was significantly higher in HepG2 cells when compared to HEK293 and COS7 cell lines. The release of PEI-pDNA complexes from particles prepared with different PLGA polymer compositions including PLGA 50–50, PLGA 75–25, and PLGA 85–15 was sustained in all cases but the release profile was dependent on the polymer composition. Transmission electron microscopy images showed that PEI-pDNA complexes remained structurally intact after release. The optimum formulation for PLGA particles loaded with PEI-ApoE pDNA complexes was prepared using 2% polyvinyl alcohol, 50–50 PLGA compositions and N/P ratios of 5–10. Strong sustained transgene expression in HepG2 cells was generated by PLGA PEI-ApoE pDNA particles up to the full 13 days tested.

Keywords: Polyethylenimine (PEI), liver, targeting ligands, Poly(lactide-co-glycolide) (PLGA), HepG2, PVA, microparticle, polyplexes, complexes, non-viral gene delivery, galactose, formulation, luciferase, plasmid DNA

Introduction

Gene therapy has the potential to provide new treatments for a variety of inherited and acquired diseases (Salem et al., 2003; Abbas et al., 2008; Intra and Salem, 2010). Liver is an attractive target tissue for gene de-

livery because of its metabolic capacity and size (Sun et al., 2005; Wang et al., 2008). Liver tissue has a strong vascular network that allows for delivery of target genes to the liver and the distribution of protein production from liver to the systemic circulation (Kim et al., 2005).

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In order to achieve this goal, it requires a delivery system that generates high transfection efficiency without causing cytotoxicity. Viral vectors are widely used for many gene therapies because of their strong transfection efficiency, but they have several drawbacks (Wiethoff and Middaugh, 2003; Gao et al., 2005). These include immunogenicity of adenoviruses, packaging constraints of adeno-associated virus and the necessity for cell mitosis for most retroviruses. In contrast, non-viral vectors are not immunogenic. Advantages to non-viral vectors include ease of scale-up, storage stability, and quality control. Drawbacks to non-viral gene delivery are poor transfection and transient gene expression, necessitating regular administration to increase efficacy. A number of hepatocyte-targeting gene delivery vectors have been reported (Midoux et al., 1993; Hashida et al., 1997; Zanta et al., 1997; Sagara and Kim, 2002; Kunath et al., 2003a; Cook et al., 2005; Sun et al., 2005; Zhang, et al., 2005; Hohokabe et al., 2007; Chen et al., 2008; Wang et al., 2008).

Polymeric gene carriers have been considered important for enhancing stability and improving the transport properties of gene materials. The polymeric drug carriers have been used as a reservoir in which drug molecules can be incorporated by means of chemical, physical, or electrostatic interactions, depending on their physicochemical properties. PLGA is biodegradable, biocompatible and has been approved by the Food and Drug Administration for human use in a variety of applications (Zhang et al., 2007a,b,c; Abbas et al., 2008; Goforth, 2009; Krishnamachari and Salem, 2009). Encapsulation of plasmid DNA (pDNA) in biodegradable polymer potentially offers a way to protect pDNA from degradation. In addition, PLGA particles can facilitate sustained release of drug and pDNA (Shive and Anderson, 1997; Zhang et al., 2007a,b; Abbas et al., 2008; Intra and Salem, 2010). Hence, it can mediate longer-term gene expression.

The hybrid liver-specific promoter murine albumin enhancer/ $\alpha 1$ antitrypsin promoter was first cloned by Kramer *et al.* in 2003. This liver-specific promoter is a chimeric construction of promoter and enhancer regions of albumin and $\alpha 1$ antitrypsin. Albumin, an abundant protein in adult serum, is mostly expressed in liver. The human $\alpha 1$ antitrypsin (hAAT) is one of the predominant proteinase inhibitors in serum and is primarily synthesized in hepatocytes. Previous studies have shown that the inclusion of this hybrid liver-specific promoter (AlbE/hAAT) increased and sustained transgene expression in hepatocytes (Kang et al., 2005).

PLGA particles loaded with polyethylenimine (PEI)-pDNA complexes is an efficient approach for delivering pDNA to target cells because of increased entrapment efficiencies, protection of pDNA from endonuclease degradation, enhancing transgene expression and low cytotoxicity (Zhang et al., 2008). In this study, PLGA PEI particles were used to deliver pDNA encoding firefly luciferase gene driven by a hybrid liver-specific

promoter (AlbE/hAAT) which is denoted as ApoE pDNA for increased transgene expression in hepatocytes. Different formulation parameters were investigated in order to optimize the preparation of PLGA PEI-ApoE pDNA particles. PLGA PEI-ApoE pDNA particles were evaluated for physicochemical characteristics including size, morphology, surface charge, and ApoE pDNA loading efficiency. The effect of polymer composition on *in vitro* release of PEI-ApoE pDNA complexes was determined. In addition, the conformational stability of released PEI-ApoE pDNA complexes was examined. *In vitro* cytotoxicity, transgene expression and sustained transgene expression of PLGA particles loaded with PEI-ApoE pDNA were also evaluated in HepG2, HEK293, and COS7 cell lines.

Materials and methods

Materials

Poly(D,L-lactide-co-glycolide) (PLGA) (Figure 1, molar ratio: 50-50, inherent viscosity: 0.41 dL/g; molar ratio: 75-25, inherent viscosity: 0.47 dL/g and molar ratio 85-15, inherent viscosity: 0.99 dL/g) were purchased from Absorbable Polymers International (Pelham, AL). Branched PEI (BPEI) (MW 25 kDa, Figure 1), polyvinyl alcohol (PVA) (Mw 30-70 kDa) 88% hydrolysed and dichloromethane (DCM) were obtained from Sigma-Aldrich Chemical Company (Milwaukee, WI). The bicinchoninic acid (BCA) protein assay kit was obtained from Pierce Biotechnology Inc. (Rockford, IL). Dulbecco's modified Eagle's medium (DMEM) was supplied from Gibco BRL (Grand Island, NY). The luciferase assay system was obtained from Promega (Madison, WI). The MTT (3-[4,5-dimethylthiazol-2-yl]-2, 5-diphenyl tetrazolium bromide) assay system was purchased from Sigma-Aldrich Chemical Company.

Preparation of GPEI

Galactosylated PEI (GPEI) was prepared as previously described (Zanta et al., 1997). Briefly, 1 g of BPEI (25 kDa) and 477 mg of lactose were dissolved in 45 mL of sodium borate buffer (200 mM, pH 8.5). The reaction mixture was incubated at 40°C for 48 h after the addition of sodium cyanoborohydride at 5 molar excess. The GPEI was purified by ultrafiltration using Pierce dialysis membrane (MWCO 10,000 Da, Pierce Biotechnology Inc.) and washing with distilled water. Finally, GPEI was lyophilized (Labconco FreeZone 4.5, Kansas City, MO) and stored at -20°C. The degree of galactose conjugated was determined by the resorcinol test which showed 5% conjugation of galactose to PEI. The ratio of PEI nitrogen to DNA phosphate (N/P) calculation for GPEI was based on a mean M_w of 61 kDa.

Cell culture

HepG2 human hepatocellular carcinoma cells, COS7 monkey African green kidney and HEK293 human embryonic kidney cells were obtained from American

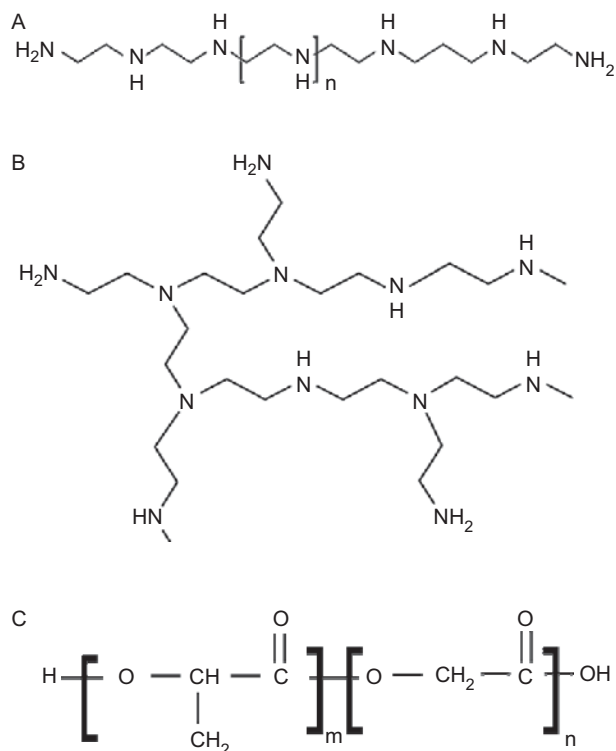


Figure 1. Chemical structure of (A) linear polyethylenimine, (B) branched polyethylenimine, and (C) poly(lactide-co-glycolide).

Type Culture Collection (Rockville, MD). The cells were maintained in DMEM supplemented with 10% fetal bovine serum, streptomycin at 100 $\mu\text{g}/\text{mL}$, penicillin at 100 U/mL, and 4 mM L-glutamine at 37°C in a humidified 5% CO_2 -containing atmosphere.

Amplification and purification of pDNA

ApoE pDNA encoding firefly luciferase driven by the murine albumin enhancer/ $\alpha 1$ -antitrypsin promoter (AlbE/hAAT) was kindly provided from Dr. Paul McCray (Carver College of Medicine, University of Iowa). The plasmid was transformed in *Escherichia coli* DH5 α and amplified in Terrific Broth media at 37°C overnight on a plate shaker set at 300 rpm. The plasmid was purified by an endotoxin-free QIAGEN Giga plasmid purification kit (Qiagen, Valencia, CA) according to the manufacturer's protocol. Purified DNA was dissolved in saline, and its purity and concentration were determined by UV absorbance at 260 and 280 nm using a SpectraMax Plus³⁸⁴ Microplate Spectrophotometer (Molecular Devices, Sunnyvale, CA).

Preparation of PEI-ApoE pDNA complexes

PEI-ApoE pDNA complexes were prepared in double distilled water by addition of PEI to ApoE pDNA solution at the desired N/P ratio and immediately vortexed for 20 sec. The solutions were incubated for 30 min at room temperature for complex formation. All experiments were repeated at least twice.

PLGA particles loaded with ApoE pDNA

PLGA particles loaded with ApoE pDNA alone were prepared using a water in oil in water (w/o/w) double-emulsion solvent-evaporation technique. One hundred milligram of 50–50 PLGA was dissolved in 5 mL of DCM. ApoE pDNA in 0.5% (w/v) PVA solution was prepared at a concentration of 4 mg/mL. Using a microtip probe sonicator set at level 2 (Sonic Dismembrator Model 100, Fisher Scientific, Pittsburgh, PA), 500 μL of the PVA solution containing 2 mg of ApoE pDNA was mixed with the PLGA/DCM solution for 20 sec to form the first emulsion. This emulsion was then rapidly added to 50 mL of 0.5% (w/v) PVA solution with stirring at 13,500 rpm for 30 sec using an IKA Ultra-Turrax T25 basic homogenizer (IKA, Wilmington, NC). The mixture was stirred overnight during which time the DCM solvent was evaporated. The particles were then washed three times with deionized water and lyophilized. The supernatant was collected and analyzed spectrophotometrically at 260 nm using a SpectraMax Plus³⁸⁴ Microplate Spectrophotometer for pDNA content. The amount of pDNA loaded in the PLGA particles was calculated by subtracting the pDNA content in the supernatant from the initial concentration of pDNA added. Particles were stored at -20°C until use.

PLGA particles loaded with PEI-ApoE pDNA complexes

PEI-ApoE pDNA complexes at N/P ratio of 2, 5, or 10 were prepared by mixing 2 mg of ApoE pDNA with 0.52, 1.3, or 2.6 mg of PEI in 500 μL of PVA solution (various concentrations), respectively. The mixture was vortexed for 20 sec, and incubated for 30 min at room temperature. Then 500 μL of PEI-ApoE pDNA complexes solution with N/P ratio of 2, 5, or 10 was mixed with 5 mL of DCM containing 100 mg of PLGA using the microtip probe sonicator set at level 2 for 30 sec to form the first emulsion. This emulsion was then rapidly added to 50 mL of PVA solution (various concentrations) and homogenized at 13,500 rpm for 30 sec. The mixture was stirred overnight for evaporation of the DCM solvent. The particles were then washed three times with deionized water and lyophilized. Particles were stored at -20°C until use.

Particle characterization

Microparticle size and ζ -potentials were measured using the Zetasizer Nano ZS (Malvern, Southborough, MA). Polystyrene of known particle size was used to calibrate the Zetasizer Nano ZS. Briefly, the particles were suspended in deionized water at a concentration of 1 mg/mL. The size measurements were performed at 25°C at a 173° scattering angle. The mean hydrodynamic diameter was determined by cumulant analysis (Zetasizer Nano ZS). The ζ -potential determinations were based on electrophoretic mobility of the PEI-pDNA complexes in deionized water (pH 4.7), which were performed using folded capillary cells (Malvern). Particle with known ζ -potential was used as the standard to calibrate the Zetasizer Nano ZS.

Surface morphology analysis

Particle morphology was assessed by scanning electron microscopy (SEM, Hitachi S-4000, Schaumburg, IL). Air-dried particles were placed on adhesive carbon tabs mounted on SEM specimen stubs. The specimen stubs were coated with approximately 5 nm of gold by ion beam evaporation before examination in the SEM operated at 5 kV accelerating voltage.

pDNA loading estimation

pDNA loading in PLGA particles was determined by dissolving 200 mg of lyophilized particles in 2 mL of chloroform. The pDNA was extracted in TE buffer (10 mM, pH 8.3). The amount of pDNA was evaluated based on absorbance at 260 nm. Empty PLGA particles in chloroform were spiked with a known amount of pDNA. After the procedure, recovery of the extracted pDNA was found to be complete. pDNA concentrations in PEI-pDNA complexes were measured at 480 nm/520 nm (Ex/Em) using the PicoGreen reagent (Invitrogen, Carlsbad, CA).

TEM analysis of PEI-ApoE pDNA complexes

PEI-ApoE pDNA complexes, before encapsulation in PLGA particles and following their release (at day 5) were visualized by transmission electron microscopy (TEM). PEI-ApoE pDNA complexes (20 μ L) were adsorbed onto carbon-coated grids for 1 min and negatively stained for 1 min with 2% uranyl acetate. After drying, samples were examined with a JEOL JEM-1230 TEM.

In vitro release of PEI-ApoE pDNA complexes from PLGA particles loaded with PEI-ApoE pDNA complexes

PEI-ApoE pDNA complexes release was determined by suspending triplicate samples of 5–10 mg of particles exactly weighted in 1.5 mL of phosphate-buffered saline (PBS) buffer, pH 7.4. The samples were incubated at 37°C with 100 rpm agitation. The sample triplicates were centrifuged at 5000 rpm for 8 min. One milliliter of supernatant was taken at each time point for complexes analysis and 1 mL of fresh PBS was added into the particle suspension. For quantification of the released PEI-ApoE pDNA complexes, a standard curve of PEI-ApoE pDNA complexes with known concentrations of pDNA was constructed with different concentrations of PEI-ApoE pDNA complexes in PBS (pH 7.4). The pDNA concentrations were determined from the unknown sample by extrapolating from a the standard curve. The pDNA concentration in the sample was quantified spectrophotometrically at 480 nm/520 nm (Ex/Em) using the PicoGreen reagent.

Cytotoxicity evaluation using the MTT assay

Cytotoxicity of the PLGA particles loaded with ApoE alone and PLGA particles loaded with PEI-ApoE pDNA complexes was evaluated using the MTT assay. PEI-ApoE pDNA complexes alone were used as a control. HepG2, COS7, and HEK293 cells were seeded in a 96-well plate at a density of 1×10^4 cells/well. Twenty-four hours later, cells were incubated with 200 μ L of complete DMEM

containing PLGA particles loaded with ApoE alone, PLGA particles loaded with PEI-ApoE pDNA complexes or PEI-ApoE pDNA complexes at various concentrations. After 4 h of incubation, the medium in each well was replaced with 100 μ L of fresh complete medium. MTT solution in PBS was added to each well and incubated with cells for an additional 2 h. Cells were lysed with 100 μ L of the extraction buffer (20% sodium dodecyl sulfate in 50% dimethylformamide, pH 4.7) overnight. The optical density of the lysate was measured at 550 nm using a Spectramax plus³⁸⁴ Microplate Spectrophotometer. Values are expressed as a percentage of the control to which no particles are added.

Evaluation of luciferase expression in HepG2, COS7, and HEK293 cells

Cells were seeded into a 24-well plate at a density of 1.5×10^5 /well of HepG2, 8×10^4 /well of COS7 and HEK293 cells 24 h before transfection. pDNA and PLGA PEI-pDNA particles (0.2 mg/well) were added to the cells in the transfection medium (serum-free) and incubated for 4 h at 37°C, followed by further incubation in serum containing medium for 44 h. The concentration of the particles was chosen from an estimated pDNA loading and a target pDNA dose of 1 μ g/well. After 44 h of incubation, cells were treated with 200 μ L of lysis buffer (Promega). The lysate was subjected to two cycles of freezing and thawing, then transferred into tubes and centrifuged at 13,200 rpm for 5 min. Twenty microliters of supernatant was added to 100 μ L of luciferase assay reagent (Promega) and samples were measured on a luminometer for 10 sec (Lumat LB 9507, EG&G Berthold, Oak Ridge, TN). The relative light units (RLU) were normalized against protein concentration in the cell extracts, measured by a BCA protein assay kit. Luciferase activity was expressed as (RLU/mg protein in the cell lysate). The data are reported as mean \pm standard deviation for triplicate samples. Every transfection experiment was repeated at least twice.

Measuring sustained *in vitro* transfection activity

The experiment was performed as described above except the incubation time of particles varied. The cells were incubated with particles for 4 h, 1, 3, 5, 7, and 11 days. After the pre-determined incubation time, the medium containing particles was removed and washed with PBS. Then completed medium was added to cells which were incubated for another 44 h for protein expression. Measurement of luciferase activity and protein assay were carried out as described above.

Statistical analysis

Group data are reported as mean \pm SD. Differences between groups were analyzed by one-way analysis of variance with a Tukey post-test analysis. Levels of significance were accepted at the $P < 0.05$ level. Statistical analyses were performed using Prism 3.02 software (Graphpad Software, Inc., San Diego, CA).

Results

Parameters evaluated in preparation and characterization of PLGA PEI-ApoE pDNA microparticles

PLGA particles loaded with PEI-ApoE pDNA complexes were prepared by encapsulating PEI-ApoE pDNA complexes into PLGA particles using the double-emulsion solvent-evaporation technique. This method had been selected based on the results of our previous work evaluating and comparing different cationic microparticle formulation methodologies (Zhang et al., 2008). Figure 2 shows a schematic diagram of the preparation method for PLGA PEI-ApoE pDNA particles.

The concentration of surfactant used in the preparation of PLGA pDNA particles can have a significant impact on particle size and the transgene expression generated by biodegradable particles (Prabha and Labhassetwar, 2004). To characterize the impact of surfactant concentration on transgene expression, five different concentrations of PVA including 0.5, 1.0, 1.5, 2.0, and 5% were used to prepare PLGA particles loaded with PEI-ApoE pDNA complexes (indicated as formulations 1, 2, 3, 4, and 5, respectively, in Table 1).

Previous studies have shown that the polymer composition can significantly impact the degradation rate

of PLGA particles (Wu and Wang, 2001). This can result in different release rates of PEI-ApoE pDNA complexes. Therefore, in this study we hypothesized that using PLGA with different polymer compositions might also result in changes in transgene expression. We compared three different types of PLGA in this study including PLGA 50-50, PLGA 75-25, and PLGA 85-15.

To characterize whether changing the structure of the PEI or conjugating a ligand to PEI that binds specific receptors could modulate transgene expression, three different types of PEI including BPEI, linear PEI (LPEI) and GPEI were used to complex with ApoE pDNA followed by loading into PLGA particles. GPEI has a galactose group that is known to be one of the most effective binding ligands for hepatocytes. The galactose ligand binds to the asialoglycoproteins receptor on the surface on hepatocytes (Zanta et al., 1997). We hypothesized that the galactose moiety in combination with the hybrid liver-specific promoter could enhance transfection in hepatocytes even further.

A correlation between N/P ratio and transfection efficiency has been explored by several researchers (Oh et al., 2002; Intra and Salem, 2008). However, the effects of N/P ratio on transfection efficiency after the PEI-pDNA complexes have been loaded into PLGA particles has yet to be determined. Therefore, PLGA particles loaded with PEI-ApoE pDNA complexes at three different N/P ratios were also prepared and evaluated.

Particle size, ζ -potential, and surface morphology of PLGA PEI-ApoE particles

The presence of PEI within the internal aqueous phase affected particle size, ζ -potential and surface morphology of PLGA particles. The mean diameter of all PLGA PEI-ApoE pDNA particles significantly decreased upon addition of PEI ($P < 0.001$). The mean particle size was found to dependent on the PVA concentration used in the particle fabrication ($P < 0.001$). Particle sizes of PLGA particles loaded with PEI-ApoE pDNA complexes prepared with 0.5–1.5% PVA range from 1000 to 1200 nm (Table 2). Increasing the PVA concentration above 1.5% resulted

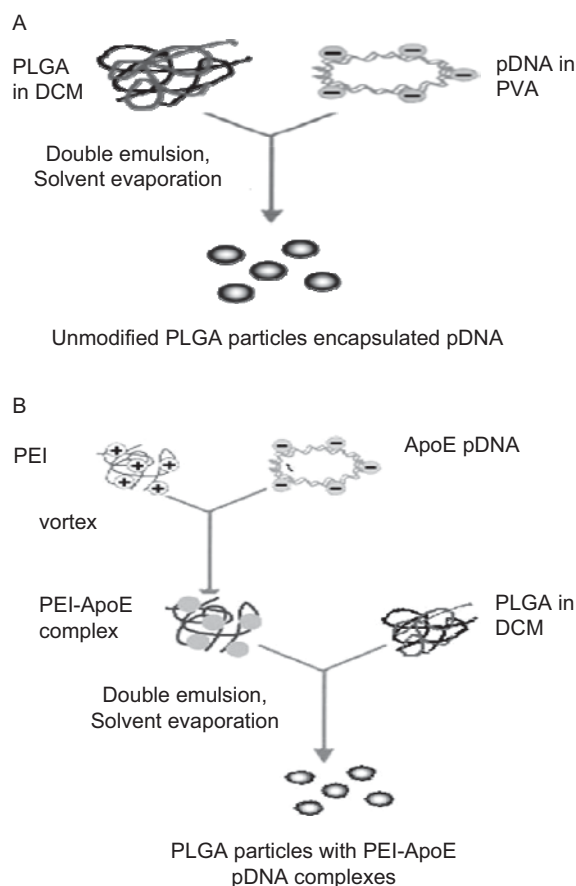


Figure 2. Schematic of the preparation of (A) PLGA particles loaded with ApoE pDNA and (B) PLGA particles loaded with PEI-ApoE pDNA complexes.

Table 1. Formulation parameters for PLGA particles loaded with PEI-ApoE pDNA complexes (formulation 1-11) prepared by the double-emulsion solvent-evaporation technique.

Formulation	PLGA	PEI structure	PVA (%)	N/P ratio
1	50-50	Branched	0.5	5
2	50-50	Branched	1.0	5
3	50-50	Branched	1.5	5
4	50-50	Branched	2.0	5
5	50-50	Branched	5.0	5
6	50-50	Linear	0.5	5
7	50-50	Galactosylated	0.5	5
8	50-50	Branched	0.5	2
9	50-50	Branched	0.5	10
10	75-25	Branched	0.5	5
11	85-15	Branched	0.5	5
PLGA	50-50	—	0.5	5

in a further decrease in particle size which ranged from 650 to 850 nm (Table 2). The mean particle diameter of particles prepared by BPEI and GPEI was similar (900–1000 nm) whereas that of particles prepared by LPEI was lower (752 nm). Furthermore, increasing the content of PEI in the PLGA particles does not significantly change particle size. This result is consistent with our previous observations of PLGA PEI particle preparations (Zhang et al., 2008).

Unmodified PLGA particles loaded ApoE pDNA displayed a net negative surface charge of -26.15 mV. After the introduction of PEI, all formulations except formulation 8 (PLGA particles loaded with PEI-ApoE pDNA complexes at N/P ratio of 2) show a net positive surface charge with values ranging from $+2.40$ to $+9.88$ mV. PLGA particles loaded with PEI-ApoE pDNA complexes at N/P ratio of 10 displayed the highest ζ -potential among all formulations (Tables 2–5). As shown in Table 2, the ζ -potential of PLGA particles loaded with PEI-pDNA complexes

decreased from $+3.90$ to $+2.40$ mV as the concentration of PVA increased from 0.5% to 5%. There is no significant difference in ζ -potential for PLGA PEI-ApoE particles prepared with different PLGA polymer compositions (Table 3). In addition, for PLGA PEI-ApoE particles prepared with different types of PEI, the surface charge of particles formed with LPEI displayed the highest ζ -potential followed by those prepared by BPEI and GPEI, respectively (Table 4). The surface charge of PLGA PEI-ApoE particles was more positive in the group prepared with higher N/P ratios (Table 5). Figure 3 displays SEM images of PLGA particles. SEM was used to study surface morphology of PLGA particles. All PLGA PEI-ApoE particles are smooth and spherical in appearance.

pDNA loading in PLGA particles

The pDNA loading efficiency was determined by dissolving PLGA particles in chloroform and PEI-ApoE pDNA complexes were extracted in TE buffer (10 mM,

Table 2. Physical characterization of the PLGA particles loaded with PEI-ApoE pDNA complexes formulated using different PVA concentrations.

Formulation	PVA (%)	PLGA	Mean particle size (nm)	Zeta potential (mV)	pDNA loading (μg pDNA/mg particle)
1	0.5	50-50	1067 ± 14	3.90 ± 0.01	8.28 ± 0.28
2	1.0	50-50	1037 ± 40	3.13 ± 0.03	9.50 ± 0.25
3	1.5	50-50	1151 ± 64	2.93 ± 0.01	9.98 ± 0.14
4	2.0	50-50	682 ± 10	2.64 ± 0.04	10.26 ± 0.25
5	5.0	50-50	812 ± 52	2.40 ± 0.01	10.54 ± 0.71

The results are the mean of three experiments \pm SD.

Table 3. Physical characterization of the PLGA particles loaded with PEI-ApoE pDNA complexes formulated using different PLGA polymer compositions.

Formulation	PVA (%)	PLGA	Mean particle size (nm)	Zeta potential (mV)	pDNA loading (μg pDNA/mg particle)
1	0.5	50-50	1067 ± 14	3.90 ± 0.01	8.28 ± 0.28
10	0.5	75-25	1360 ± 27	4.15 ± 0.02	8.39 ± 0.74
11	0.5	85-15	1335 ± 81	3.91 ± 0.02	10.09 ± 0.42

The results are the mean of three experiments \pm SD.

Table 4. Physical characterization of the PLGA particles loaded with PEI-ApoE pDNA complexes formulated using different forms of PEI.

Formulation	PVA (%)	PLGA	Mean particle size (nan)	Zeta potential (mV)	pDNA loading (μg pDNA/mg particle)
1	0.5	BPEI	1067 ± 14	3.90 ± 0.01	8.28 ± 0.28
6	0.5	LPEI	752 ± 114	4.23 ± 0.16	10.33 ± 0.38
7	0.5	GPEI	909 ± 81	3.15 ± 0.04	8.76 ± 0.28

The results are the mean of three experiments \pm SD.

Table 5. Physical characterization of the PLGA particles loaded with PEI-ApoE pDNA complexes formulated using PEI-ApoE pDNA complexes at different N/P ratios.

Formulation	PVA (%)	PLGA	Mean particle size (nm)	Zeta potential (mV)	pDNA loading (μg pDNA/mg particle)
8	0.5	2	1316 ± 8	-10.62 ± 0.05	7.48 ± 0.08
1	0.5	5	1067 ± 14	3.90 ± 0.01	8.28 ± 0.28
9	0.5	10	1218 ± 13	9.88 ± 0.01	10.24 ± 0.23

The results are the mean of three experiments \pm SD.

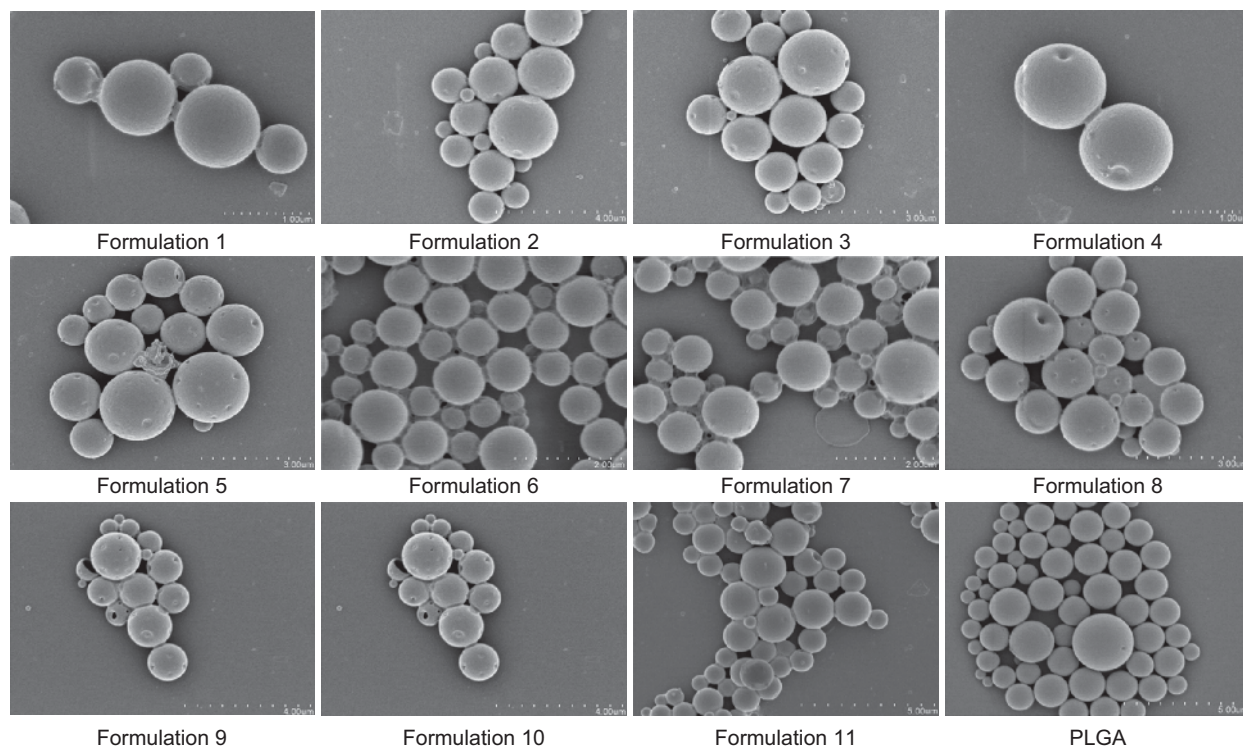


Figure 3. Scanning electron micrographs of PLGA particles loaded with PEI-ApoE pDNA complexes (formulation 1-11) and PLGA particles loaded with ApoE pDNA.

pH 8.3). Then the amount of PEI-ApoE pDNA was evaluated by UV spectroscopy and the picogreen reagent using standard calibration curves in order to determine the amount of pDNA (Intra and Salem, 2010). In this study, all PLGA particles loaded with PEI-ApoE pDNA complexes displayed significantly higher loading of ApoE pDNA in comparison to unmodified PLGA particles loaded with ApoE pDNA ($P < 0.001$). The loading efficiency of ApoE pDNA in PLGA PEI particles increased from 8.28 to 10.54 μg pDNA/mg particle as the PVA concentration was increased from 0.5 to 5% (Table 2). Amongst the particles prepared with different PLGA polymer compositions, the pDNA loading efficiency of 50-50 PLGA was found to be the lowest (8.28 μg pDNA/mg particles) and there was no significant increase ($P > 0.05$) in the ApoE pDNA loading of 75-25 PLGA (8.39 μg pDNA/mg particles) when compared with 50-50 PLGA. There was a significant increase ($P < 0.001$) in the ApoE pDNA loading of 85-15 PLGA (10.09 μg pDNA/mg particles) in comparison to 50-50 PLGA and 75-25 PLGA. Complex formation of PEI and ApoE with different types of PEI prior to encapsulation into PLGA particles resulted in differences in ApoE pDNA loading efficiency. PLGA PEI-pDNA particles prepared from LPEI resulted in the highest pDNA loading (10.33 μg pDNA/mg particles) followed by those prepared from GPEI (8.76 μg pDNA/mg particles) and BPEI (8.28 μg pDNA/mg particles), respectively (Table 4). As shown in Table 5, increasing the N/P ratio of PEI-ApoE pDNA complexes leads to increases in ApoE pDNA loading efficiency. For exam-

ple, the ApoE pDNA loading efficiency increased from 7.50 to 10.28 μg pDNA/mg particles as the N/P ratio increased from 2 to 10.

***In vitro* release of PEI-ApoE pDNA complexes**

The release profiles of PEI-ApoE pDNA complexes from all the formulations are reported in Figure 4. The experiments measuring release of PEI-ApoE pDNA complexes from PLGA particles were carried out in PBS (pH 7.4) at 37°C. PLGA particles loaded with PEI-ApoE pDNA complexes prepared from different copolymer compositions; PLGA 50-50, PLGA 75-25, and PLGA 85-15 showed a decrease in PEI-ApoE pDNA complex release rate as the lactide content of the polymer increases. Particles prepared by PLGA 50-50 were characterized by a monophasic profile. The initial burst release corresponded to about 17% of the total PEI-ApoE pDNA complexes loaded. This was followed by a slower phase that lasted approximately 7 days. PLGA particles loaded with PEI-ApoE pDNA complexes prepared from 50-50 PLGA displayed 100% release of total entrapped PEI-ApoE pDNA complexes in 7 days. PLGA particles loaded with PEI-ApoE pDNA complexes prepared from 75-25 PLGA demonstrated a biphasic release profile with an initial burst release of 15% followed by slower controlled release up to 22 days. Total complexes released from PLGA particles prepared from 75-25 PLGA were observed at 22 days. The release of PEI-ApoE pDNA complexes from PLGA particles prepared with 85-15 PLGA was characterized by a triphasic release profile. It started with a small initial burst release of 8% followed

by a slower controlled release phase. No release is observed from day 9 to day 18. Beyond day 19, the release rate increases and is followed by a lag phase (no release) up to day 30. PLGA particles prepared from 85–15 PLGA released 70% of the total pDNA-ApoE pDNA complexes in 30 days.

Physical characterization of released PEI-ApoE pDNA complexes

Characteristics of PEI-ApoE pDNA complexes before and after their release from PLGA particles were evaluated by TEM in order to determine whether the formulation process altered the morphology of PEI-ApoE pDNA complexes. PEI-ApoE pDNA complexes released from PLGA particles had a spherical shape, similar to their shape before encapsulation (Figure 5). After encapsulation, some PEI-ApoE pDNA complexes appear to have a less distinct area of uranyl acetate staining, which may be due to more pDNA exposed on the surface of the complexes.

Effect of PVA concentration on transfection efficiencies of PLGA particles loaded with PEI-ApoE pDNA complexes

The concentration of PVA used to prepare PLGA particles affects transfection efficiency (Figure 6). As shown in Figure 6a and 6b, particles prepared from 2% PVA showed the highest transfection efficiency in HepG2 and COS7 cells (7.86×10^7 and 3.34×10^7 RLU/mg protein, respectively) with no significant difference in comparison to BPEI-ApoE pDNA complexes at the same N/P ratio ($P > 0.05$). The transfection efficiency was PVA concentration dependent in HEK293 cells. As the PVA concentration used in the PLGA particle preparation increased, the transfection efficiency was enhanced from 4.06×10^4 to 1.3×10^5 RLU/mg protein (Figure 6c).

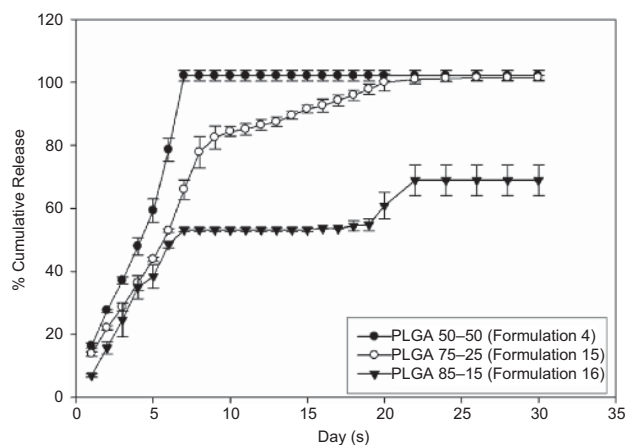


Figure 4. Cumulative (%) release of PEI-ApoE pDNA complexes from PLGA particles loaded with PEI-ApoE pDNA complexes prepared using PLGA 50–50, PLGA 75–25, and PLGA 85–15. Experiments were carried out in PBS buffer pH 7.4 at 37°C. Data is presented as means \pm SD and are representative of three independent experiments.

Effect of PLGA composition on transfection efficiencies of PLGA-PEI-ApoE particles

In HepG2 cells, PLGA particles made with 50–50 PLGA displayed the highest transgene expression at 3.11×10^6 RLU/mg protein in comparison with those made with 75–25 PLGA at 9.12×10^5 RLU/mg protein and 85–15 PLGA at 7.86×10^5 RLU/mg protein (Figure 7). In contrast, for transgene expression in COS7 cells, PLGA particles prepared from 50–50 PLGA showed similar transfection efficiency as PLGA particles prepared from 75–25 PLGA (3.19×10^5 and 3.12×10^5 RLU/mg protein for PLGA particles prepared from 50–50 PLGA and 75–25 PLGA, respectively). The transgene expression in COS7 cells was the lowest in the cells treated with PLGA particles prepared from 85–15 PLGA (1.92×10^5 RLU/mg protein).

Effect of PEI structure and galactose ligand conjugation to PEI on transfection efficiencies generated by PLGA PEI-ApoE pDNA particles

Two forms of PEI were used to complex with ApoE pDNA prior to encapsulation in PLGA particles. As shown in Figure 8, PLGA particles loaded with BPEI and GPEI displayed similar transgene expression profiles in HepG2 cells (3.11×10^6 and 2.33×10^6 RLU/mg protein respectively) whilst PLGA particles loaded with LPEI resulted in the lowest transgene expression (3.16×10^4 RLU/mg protein). In COS7 cells, LPEI showed the highest transfection efficiency followed by BPEI and GPEI, respectively.

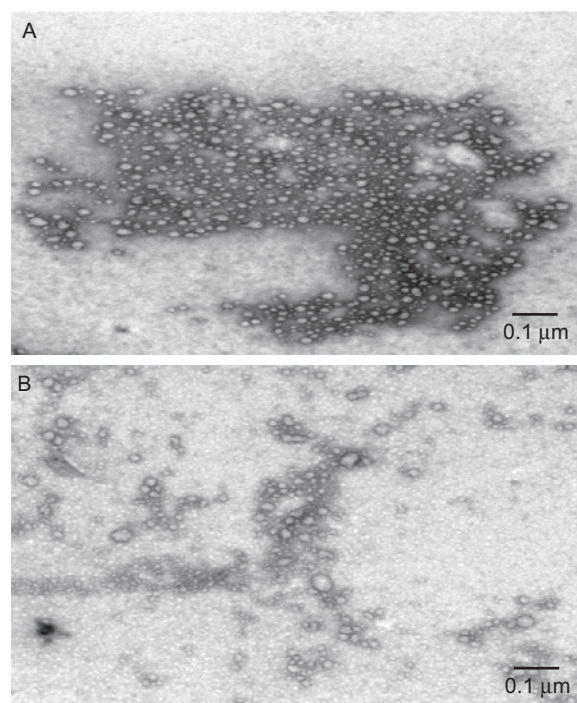


Figure 5. Transmission electron micrographs of PEI-ApoE pDNA complexes (A) before encapsulation in PLGA particles and (B) 5 days after release from PLGA particles. The bar represents 100 nm.

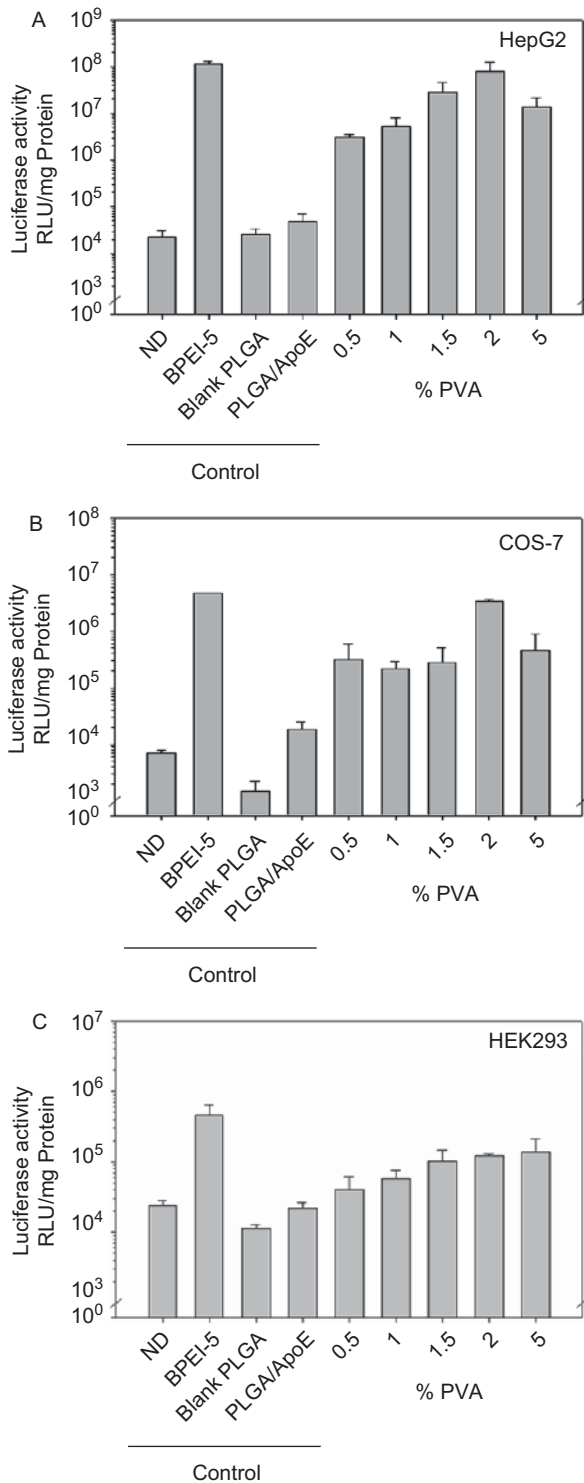


Figure 6. Luciferase activity of (A) HepG2, (B) HEK293, (C) COS7 cells that have been treated with unmodified PLGA particles loaded with ApoE pDNA (PLGA/ApoE), PLGA particles loaded with PEI-ApoE pDNA complexes prepared using varying concentrations of PVA solution (0.5, 1.0, 1.5, 2.0, and 5% (w/v)). Transfection was performed by incubating these formulations with cells for 4h (reporter gene: ApoE pDNA; pDNA: 1 μ g/well). Data is presented as the mean \pm standard deviation ($n=3$). ND represents naked DNA.

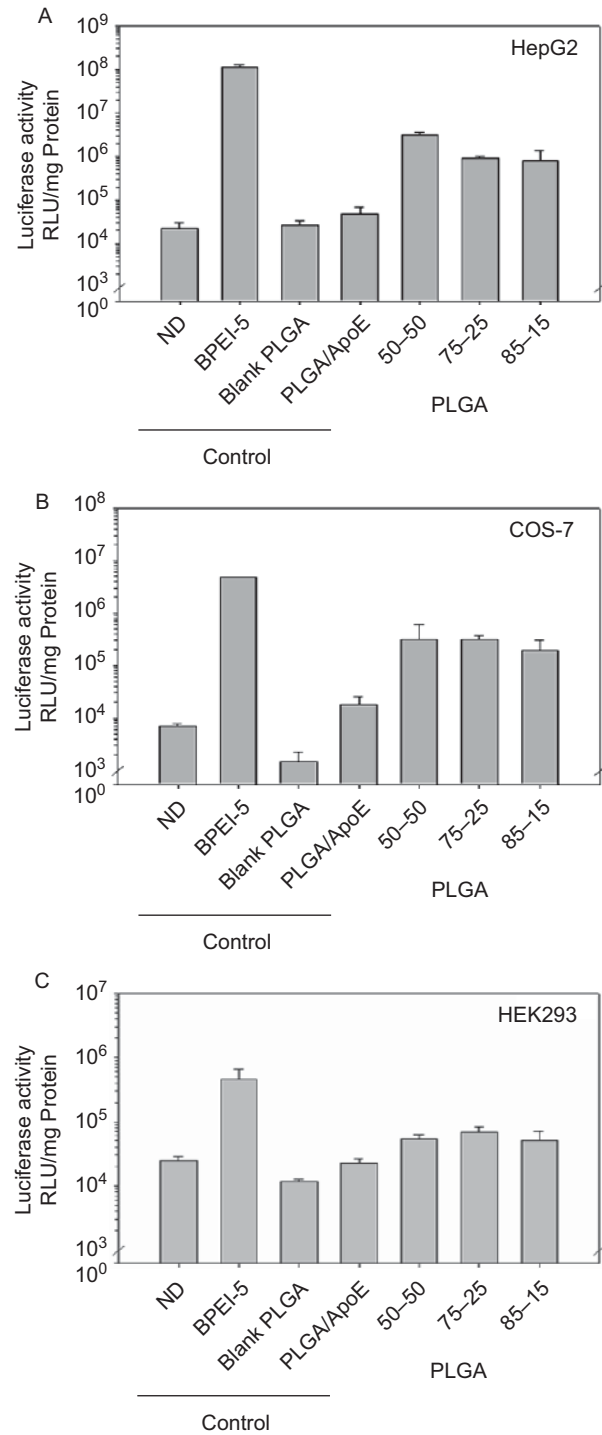


Figure 7. Luciferase activity from (A) HepG2, (B) HEK293, (C) COS7 cells that have been treated with unmodified PLGA particles loaded with ApoE pDNA (PLGA/ApoE), PLGA particles loaded with PEI-ApoE pDNA complexes prepared with varying polymer compositions of PLGA polymer (PLGA 50-50, PLGA 75-25, and PLGA 85-15). Transfection experiments were performed by incubating these formulations with cells for 4h (reporter gene: ApoE pDNA; pDNA: 1 μ g/well). Data is presented as the mean \pm standard deviation ($n=3$). ND represents naked DNA.

Effect of N/P ratio of PEI to ApoE pDNA on transfection efficiency of PLGA particles loaded with PEI-ApoE pDNA complexes

The N/P ratio used for PEI-ApoE pDNA complexes formation prior to encapsulation in PLGA particles also affects the transfection efficiency of PLGA particles in HepG2, COS7, and HEK293 cells (Figure 9). The transfection efficiency of PLGA particles increases with increasing N/P ratio used for PEI-ApoE pDNA complexes formation prior to encapsulation in PLGA particles in all cells. PEI-ApoE pDNA complexes made at N/P = 10 yield the highest transgene expression in all cell lines. In HepG2 and HEK293 cells, there is no significant difference between PLGA particles containing PEI-ApoE pDNA complexes prepared at N/P = 5 and N/P = 10 ($P > 0.05$).

PLGA particles loaded with PEI-ApoE pDNA complexes generate sustained transgene expression in HepG2 cells

In vitro transgene expression generated from all PLGA particles loaded with PEI-ApoE complexes were evaluated and compared with PEI-ApoE complexes and PLGA particles loaded with ApoE pDNA in HepG2 cells. Two different factors; PVA concentrations and PLGA polymer compositions were also evaluated. All PLGA PEI-ApoE particles prepared with different PVA concentrations and different polymer compositions displayed sustained transgene expression in HepG2 cells (Figures 10 and 11). For the first 4 h, transgene expression was higher in the PEI group than that of all groups with PLGA particles loaded with PEI-ApoE complexes. The transgene expression in the cationic PLGA particle groups was maintained for 13 days, whereas the transgene expression resulting from BPEI-ApoE pDNA complexes significantly decreased after day 1 ($P < 0.01$). The transgene expression generated by cationic PLGA particles groups was 449- and 390-fold higher than that of the BPEI-ApoE pDNA complexes at day 13 for particles prepared with 5% PVA and PLGA 85-15, respectively.

Evaluation of cytotoxicity of PLGA PEI-ApoE microparticles

Cytotoxicity is a major hurdle for use of polycationic gene delivery systems. *In vitro* cytotoxicity was evaluated in HepG2, HEK293, and COS7 cells with increasing doses of PLGA particles loaded with PEI-ApoE pDNA complexes (from 7.8 to 1000 μg of particles per milliliter of DMEM). BPEI 25 kDa, LPEI 25 kDa, and GPEI 25 kDa were used as a control. The LD_{50} value which represents 50% cell viability is reported. For BPEI 25 kDa, the LD_{50} value of HepG2 cells was reached at 150 $\mu\text{g}/\text{mL}$ of the polymer concentration (Table 6). The LD_{50} value of HepG2 cells treated with GPEI was 170 $\mu\text{g}/\text{mL}$ of the polymer concentration. PLGA particles loaded with PEI-ApoE pDNA complexes showed similar cytotoxicity profile in HepG2 cells in comparison with unmodified PLGA particles

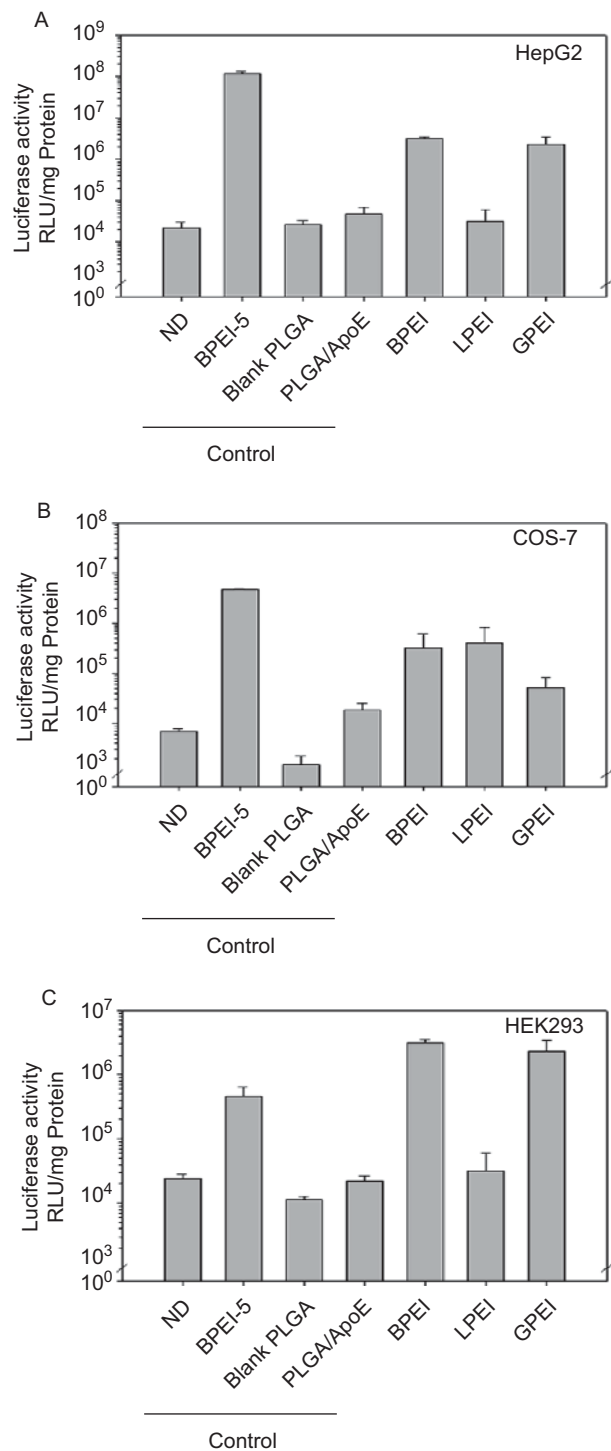


Figure 8. Luciferase activity from (A) HepG2, (B) HEK293, (C) COS7 cells that have been treated with unmodified PLGA particles loaded with ApoE pDNA (PLGA/ApoE), PLGA particles loaded with PEI-ApoE pDNA complexes prepared with varying types of PEI polymer (branched PEI (BPEI), linear PEI (LPEI), and galactosylated PEI (GPEI)). Transfection experiments were performed by incubating these formulations with cells for 4 h (reporter gene: ApoE pDNA; pDNA: 1 $\mu\text{g}/\text{well}$). Data is presented as the mean \pm standard deviation ($n=3$). ND represents naked DNA.

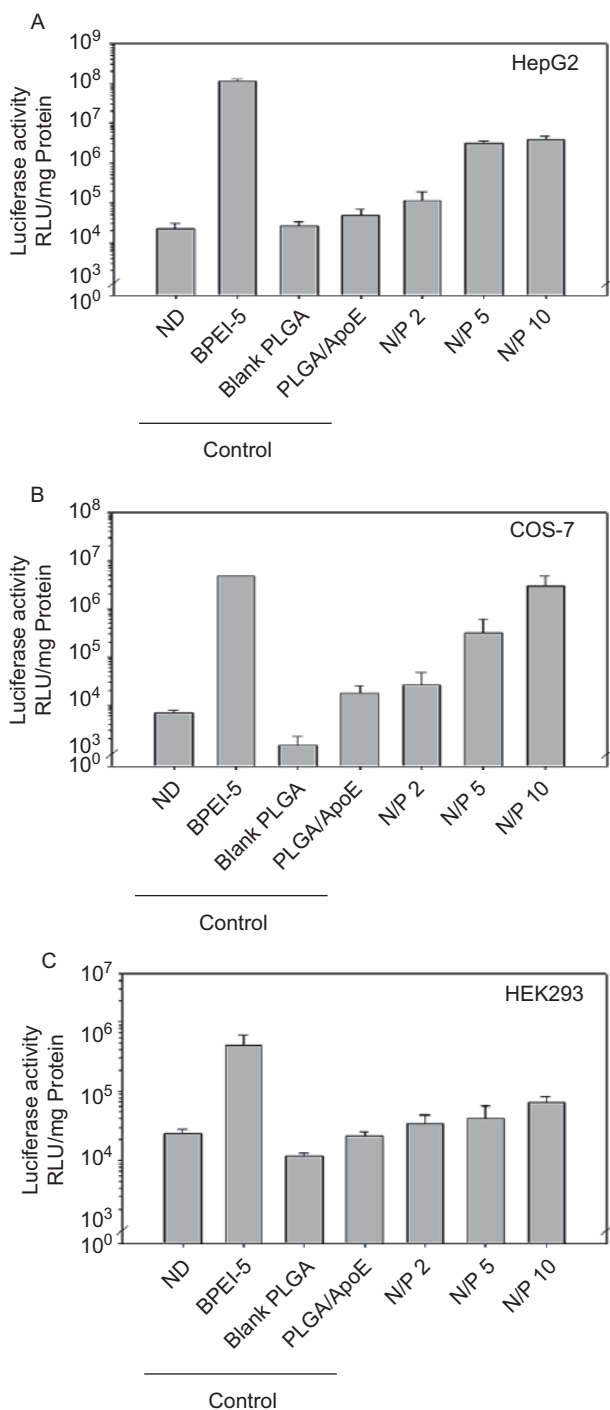


Figure 9. Luciferase activity from (A) HepG2, (B) HEK293, (C) COS7 cells that have been treated with unmodified PLGA particles loaded with ApoE pDNA (PLGA/ApoE), PLGA particles loaded with PEI-ApoE pDNA complexes prepared with varying N/P ratio of PEI-ApoE pDNA complexes (2, 5, and 10). Transfection experiments were performed by incubating these particles with cells for 4 h (reporter gene: ApoE pDNA; pDNA: 1 μ g/well). Data is presented as the mean \pm standard deviation ($n=3$). ND represents naked DNA.

with LD₅₀ value greater than 1000 μ g/mL of the polymer concentration. The LD₅₀ value of BPEI, LPEI, and GPEI was 150, 170, and 27 μ g particles/mL of the DMEM medium, respectively (Table 7). As shown in Tables 6 and 7,

the cytotoxicity of all PLGA PEI-ApoE particles was significantly lower than that of BPEI, LPEI, and GPEI. The LD₅₀ value of PLGA particles loaded with PEI-ApoE pDNA complexes was similar to unmodified PLGA particles with values greater than 1000 μ g particles/mL of the DMEM medium. In addition, the concentration dependent effect on the cell viability was observed in all the cell lines (data not shown).

Discussion

Gene delivery to liver cells is of interest for systemic delivery of transgene products and for the treatment of liver diseases. There are many barriers to efficient delivery of pDNA to the hepatocytes. An important step in the advancement of gene therapy is the development of an efficient, targeted gene delivery system. This delivery system should provide pDNA stability, protection from nuclease and endosomal degradation, intracellular protection from endosomal degradation, release of pDNA from gene carrier and uptake into the liver cells. Moreover, the safety of the delivery system is major concern. In addition, long-term expression of the transgene is desirable. In this study, we complex ApoE pDNA encoding firefly luciferase driven by a hybrid liver-specific promoter (AlbE/hAAT) to a cationic polymer, PEI, and load the complexes into biodegradable PLGA particles. PEI complexes with pDNA by electrostatic interactions with positively charged amine groups in PEI and negatively charged phosphate groups in pDNA. The N/P ratio is charge balance between positively charged amine groups in PEI and negatively charged phosphate groups in pDNA (Dunlap et al., 1997). In all PEI-ApoE complexes, increasing the N/P ratio resulted in particle size reduction. The decrease in particle size may be explained by (i) condensation of PEI and pDNA and (ii) repulsion force of charged particles that prevent particles from aggregation. The condensation property of PEI is a very important factor in its use in non-viral gene delivery (Kunath et al., 2003b).

Particle size is one of the critical factors in determining the transfection efficiency of non-viral gene vectors. Complex formation between PEI and pDNA before encapsulation into the PLGA particles leads to a decrease in the mean particle size of PLGA particles (Zhang et al., 2008; Intra and Salem, 2010). In this study, the mean particle size of PLGA PEI particles was found to be dependent on the PVA concentrations used in the particle formation. Higher PVA concentrations resulted in the formation of smaller PLGA particles. Significant increases in the net shear stress can be induced by increases in PVA concentration. As the concentration of PVA increases, more PVA can be oriented at the water and organic solvent interface to reduce the interfacial tension (Nandi et al., 2001). In addition, PLGA particles prepared with PLGA 50-50 had lower mean particle sizes than particles prepared with PLGA 75-25 and PLGA 85-15. An increase in the lactide

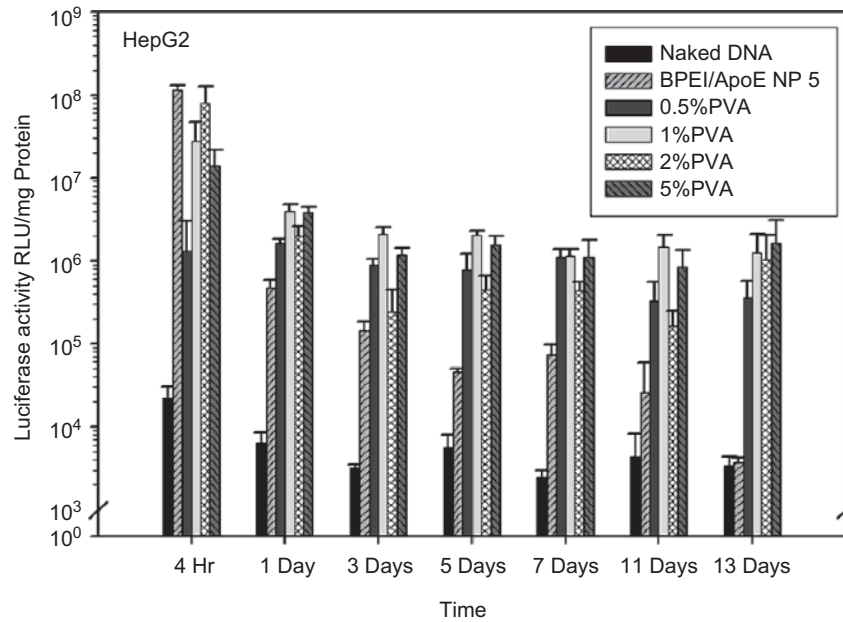


Figure 10. Luciferase activity from HepG2 cells that have been treated with naked ApoE pDNA, BPEI-ApoE pDNA complexes at N/P ratio 5 and PLGA particles loaded with PEI-ApoE pDNA complexes prepared with varying concentrations of PVA solution (0.5, 1.5, 2.0, and 5% (w/v)). Transfection was performed by incubating these particles with cells for 4h, 1 day, 3 days, 5 days, 7 days, 11 days, and 13 days (reporter gene: ApoE pDNA; pDNA: 1 μ g/well). Data is presented as the mean \pm standard deviation ($n=3$).

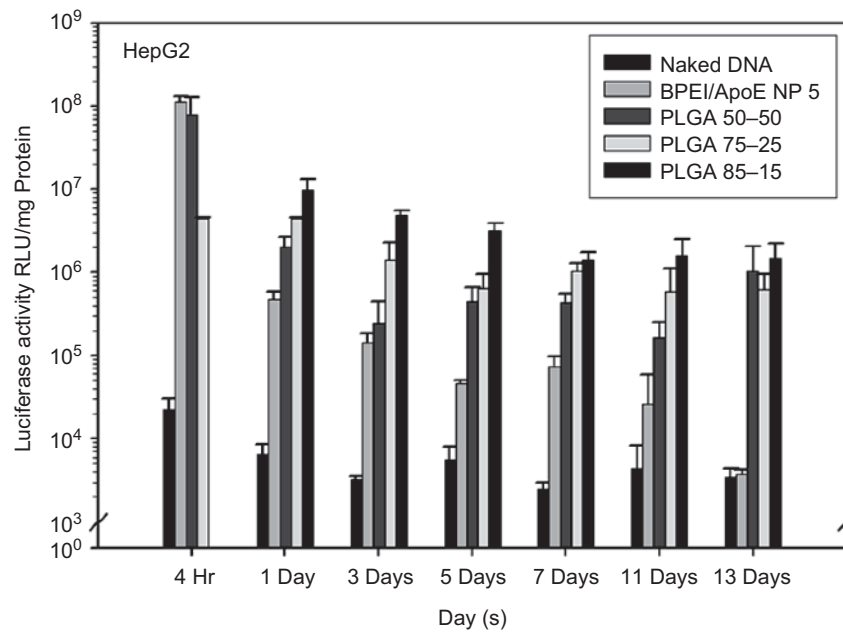


Figure 11. Luciferase activity of HepG2 cells that have been treated with naked ApoE pDNA, BPEI-ApoE pDNA complexes at N/P ratio 5 and PLGA particles loaded with PEI-ApoE pDNA complexes prepared with varying polymer compositions of PLGA polymer (PLGA 50-50, PLGA 75-25, and PLGA 85-15). Transfection was performed by incubating these particles with cells for 4h, 1 day, 3 days, 5 days, 7 days, 11 days, and 13 days (reporter gene: ApoE pDNA; pDNA: 1 μ g/well). Data is presented as the mean \pm standard deviation ($n=3$).

composition of PLGA led to an increase in the mean particle size of PLGA PEI-ApoE particles.

The surface charge of the non-viral delivery system is expected to influence its interactions with various biological components as well as its cellular interaction and cellular uptake. PLGA particles loaded with ApoE pDNA had a net negative ζ -potential. After the introduction

of PEI, the ζ -potential of all formulation turned positive except formulation 8 (PLGA 50:50, BPEI, 0.5% PVA, and N/P 2). Increasing the concentration of PVA used in PLGA particles formation resulted in a decrease in the positive surface charge of PLGA particles. This may be explained by PVA surfactant residue left on the surface of PLGA particles. After the removal of the organic solvent,

Table 6. Cytotoxicity of PEI-ApoE pDNA complexes in free form or loaded in PLGA particles and PLGA particles loaded with ApoE pDNA in HepG2 cell lines.

Sample	LD ₅₀ (µg particles/mL of DMEM medium)
Branched PEI	150 ± 19
Linear PEI	>1000
GPEI	170 ± 15
PLGA	>1000
Formulation 1	>1000
Formulation 2	>1000
Formulation 3	>1000
Formulation 4	>1000
Formulation 5	>1000
Formulation 6	>1000
Formulation 7	>1000
Formulation 8	>1000
Formulation 9	>1000
Formulation 10	>1000
Formulation 11	>1000

The results are reported as 50% of cell viability (LD₅₀). The results are the mean of three experiments ± SD.

Table 7. Cytotoxicity of PEI-ApoE pDNA complexes in free form or loaded in PLGA particles and PLGA particles loaded with ApoE pDNA in HEK293 cell lines.

Sample	LD ₅₀ (µg particles/mL of DMEM medium)
Branched PEI	100 ± 17
Linear PEI	170 ± 21
GPEI	80 ± 8
PLGA	>1000
Formulation 1	>1000
Formulation 2	>1000
Formulation 3	>1000
Formulation 4	>1000
Formulation 5	>1000
Formulation 6	>1000
Formulation 7	>1000
Formulation 8	>1000
Formulation 9	>1000
Formulation 10	>1000
Formulation 11	>1000

The results are reported as 50% of cell viability (LD₅₀). The results are the mean of three experiments ± SD.

PVA residues can physically adsorb onto the surface of particles (Galindo-Rodriguez et al., 2004). All of the PLGA particles loaded with PEI-ApoE pDNA had smooth surfaces and spherical shape.

pDNA complexation with PEI significantly enhanced the loading of pDNA from 3.52 µg pDNA/mg particles to 10.54 µg pDNA/mg particles. This is consistent with ours and other previous studies on the use of cationic excipients to increase loading efficiency of pDNA in PLGA particles (Abbas et al., 2008). Other factors that can also affect the loading of macromolecules include drug/polymer interactions and emulsion stability. For example,

Table 8. Cytotoxicity of PEI-ApoE pDNA complexes in free form or loaded in PLGA particles and PLGA particles loaded with ApoE pDNA particles in COS-7 cell lines.

Sample	LD ₅₀ (µg particles/mL of DMEM medium)
Branched PEI	150 ± 24
Linear PEI	170 ± 15
GPEI	27 ± 5
PLGA	>1000
Formulation 1	>1000
Formulation 2	>1000
Formulation 3	>1000
Formulation 4	>1000
Formulation 5	>1000
Formulation 6	>1000
Formulation 7	>1000
Formulation 8	>1000
Formulation 9	>1000
Formulation 10	>1000
Formulation 11	>1000

The results are reported as 50% of cell viability (LD₅₀). The results are the mean of three experiments ± SD.

one study showed that the interaction of gene materials such as oligonucleotides with PEI and PLGA occurs at the interface and lead to a decrease in concentration of gene materials in the aqueous phase (De Rosa et al., 2002). This can be used to increase loading in the PLGA PEI-ApoE particle delivery system. The pDNA loading efficiency of PLGA particles prepared from PLGA 50–50 was found to be the lowest. There was no significant increase ($P > 0.05$) in the pDNA loading efficiency of PLGA particles prepared from PLGA 75–25 in comparison with PLGA particles prepared from PLGA 50–50. However, there was a significant increase ($P < 0.05$) in the pDNA loading efficiency of PLGA particles prepared from PLGA 85–15 in comparison with PLGA particles prepared from PLGA 50–50. This can be attributed to the fact that as lactide content of PLGA copolymer increases (from PLGA 50–50 to PLGA 85–15), the hydrophobicity of the copolymer also increases because of the hydrophobic nature of the lactide content in the copolymer.

In general, the release of drugs from PLGA particles can be described by two mechanisms. In the first, the drugs can diffuse through channels formed during particle fabrication. Second, the release of the drugs is dependent on polymer degradation. Therefore, the degradation rate plays an important role in drug release. *In vitro* release of PEI-ApoE complexes from PLGA particles using three different PLGA polymers including PLGA 50–50, PLGA 75–25, and PLGA 85–15 were studied. One hundred percent release of PEI-pDNA complexes from PLGA particles prepared from PLGA 50–50 was reached within 6 days. In contrast, PLGA particles made by PLGA 85–15 resulted in complexes release beyond 30 days. PEI-pDNA complexes released from PLGA particles prepared from PLGA 85–15 reached 70% cumulative release at day 30. PLGA particles made by PLGA 75–25 showed a biphasic release

profile. A small burst release was observed from all PLGA batches. This phase was attributed to the complexes on or close to the particle surface. The different amounts of complexes released in the initial phase could be from (i) different amounts of complexes adsorbed onto the wall of particles that lead to immediate release causing burst effect and (ii) different sizes of channels on the surface of PLGA particles. After the initial phase, the sustained release of complexes from PLGA particles could result from diffusion through PLGA channels as well as the erosion of PLGA polymer. The degradation rate of PLGA 50–50 is fastest followed by PLGA 75–25 and PLGA 85–15, respectively. In our study, the release of complexes from PLGA particle reaches 100% within 6 days and 21 days for PLGA 50–50 and PLGA 75–25, respectively. The increase in the release rate from the particles could also result from increased in water uptake by the PEI moiety which leads to increased in hydrolysis of the PLGA particles.

It has been reported that PLGA particle preparation using the double-emulsion solvent-evaporation method can cause pDNA damage during the fabrication process. Shear force generated by ultrasonication and homogenization can cause nicking of naked pDNA (Walter et al., 1999; Abbas et al., 2008). In our study, PEI-ApoE pDNA complexes that release from PLGA particles maintained their physical characteristics including size and morphology. This result suggests that the double-emulsion solvent-evaporation technique does not damage PEI-ApoE pDNA complexes appreciably. This may be explained by the protection effect of PEI. For example, Chumakova et al. (2008) have reported that by forming complexes of pDNA with PEI, the polymer can protect pDNA from ultrasound-induced damage. In comparison with freshly prepared complexes on the TEM micrograph, the negative stain of released complexes was less discrete (Figure 5). This difference may be explained by the partial dissociation of PEI-ApoE complexes over time.

Transgene expression mediated by PLGA PEI-ApoE particles were evaluated in three different cell lines including HepG2, COS7, and HEK293 (Figures 6–9). ApoE pDNA is driven by the murine albumin enhancer/ α 1-antitrypsin promoter (AlbE/hAAT) which is specific liver promoter. In all formulations, the transfection efficiency of PLGA PEI-ApoE particles was highest in HepG2 cells. This system is therefore attractive for hepatocyte specific gene delivery. All PLGA particles loaded with PEI-ApoE pDNA complexes showed significantly higher transfection efficiencies than unmodified PLGA particles loaded with ApoE pDNA ($P < 0.05$). PLGA particles loaded with PEI-pDNA complexes were prepared with solution of PVA concentrations of 0.5, 1.0, 1.5, 2.0, and 5.0% (w/v). The optimal transfection efficiency was observed in all cell lines transfected with particles prepared with 2.0% (w/v) PVA solution. The transgene expression of PLGA particles loaded with BPEI-ApoE pDNA complexes showed no significant difference in comparison to PLGA particles loaded with GPEI-ApoE pDNA complexes. Therefore, in

this study, the use of a galactose ligand *in combination* with a hybrid liver promoter (AlbE/hAAT) did not show any additive enhancement in transgene expression generated in HepG2 cells.

In this study, the cells transfected with PLGA particles loaded with PEI-ApoE pDNA complexes demonstrated significant and sustained luciferase expression (Figures 10–11). The slow intracellular release of the entrapped PEI-ApoE complexes from PLGA particles is hypothesized to be a convincing explanation for this observation. Sustained delivery of PEI-ApoE complexes in the extracellular matrix of target tissue should prolong the duration of ApoE pDNA expression due to a continuous PEI-ApoE pDNA complexes uptake by the target cells. In addition, this study also shows that released complexes maintain their biological activity.

PEI has been shown to significantly promote pDNA transfection efficiency but also has high cytotoxicity. Higher molecular weight PEI showed higher cytotoxicity (Fischer et al., 1999). Cell viability of all PLGA particles loaded with PEI-ApoE pDNA complexes formulations was evaluated and compared to BPEI-ApoE pDNA complexes. PLGA particles loaded with PEI-ApoE pDNA complexes show similar cytotoxicity as the unmodified PLGA particles in HEK293, COS7, and HepG2 cells (Table 8). PLGA particles loaded with PEI-ApoE pDNA complexes displayed significantly lower cytotoxicity than BPEI-ApoE pDNA, LPEI-ApoE pDNA, and GPEI-ApoE pDNA complexes *in vitro* ($P < 0.001$) (Table 6). For BPEI, concentrations above 150 μ g/mL medium resulted in lower than 50% cell viability in HepG2 and COS7 cells. In comparison, for PLGA particles and all PLGA particles loaded with PEI-ApoE pDNA complexes, more than 50% of the cells are viable at concentrations as high as 1000 μ g/mL medium in both cell lines. This shows that PLGA particles loaded with PEI-ApoE pDNA complexes retain the low cytotoxicity properties of PLGA particles. No significant difference in the toxicity of the PLGA particles loaded with PEI-ApoE pDNA complexes is observed among the different formulations. Low cytotoxicity of PLGA particles loaded with PEI-ApoE pDNA complexes could also be attributed to the slow release of PEI-ApoE pDNA complexes from PLGA particles, which reduces the direct interaction of cationic polymer with cells.

Conclusions

In this study, we rationally designed and developed a non-viral vector for targeted and sustained delivery of genes to liver cells. Cationic PEI polymers were used to condense pDNA and improve transfection efficiencies. These complexes were loaded into biodegradable PLGA particles to provide sustained release and improve toxicity profiles of the delivery system. A liver-specific promoter and the use of a galactose ligand were evaluated to improve the transgene expression generated in HepG2 cells. Several parameters including the composition of the PLGA polymer, the PVA

surfactant concentration, the structure of the PEI and the N/P ratio of the PEI-pDNA complexes were investigated to optimize the transgene expression generated by the PLGA PEI-ApoE pDNA delivery system. Encapsulation of PEI-ApoE pDNA complexes in PLGA particles resulted in particle size reduction, ζ -potential increases and pDNA loading enhancement. Additionally, the PLGA PEI-ApoE pDNA particles showed sustained transgene expression in HepG2 cells for at least 13 days. The PLGA PEI-pDNA particles had cytotoxicity profiles that were significantly lower than PEI and similar to unmodified PLGA as demonstrated by MTT assays. The optimum PLGA PEI-ApoE pDNA particles for transfection of HepG2 cells was achieved by preparing particles using a 2% PVA surfactant concentration and a 50-50 PLGA composition.

Declaration of interest

We gratefully acknowledge support from the American Cancer Society (RSG-09-015-01-CDD), the National Cancer Institute at the National Institutes of Health (1R21CA13345-01/ 1R21CA128414-01A2), and the Pharmaceutical Research and Manufacturers of America Foundation. J. Intra acknowledges support from the Parenteral Drug Association for a predoctoral fellowship. We have no declarations of conflicts of interest.

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