

Research Paper

Zwitterionic Poly(carboxybetaine)-based Cationic Liposomes for Effective Delivery of Small Interfering RNA Therapeutics without Accelerated Blood Clearance Phenomenon

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Abstract

For efficient delivery of small interfering RNA (siRNA) to the target diseased site *in vivo*, it is important to design suitable vehicles to control the blood circulation of siRNA. It has been shown that surface modification of cationic liposome/siRNA complexes (lipoplexes) with polyethylene glycol (PEG) could enhance the circulation time of lipoplexes. However, the first injection of PEGylated lipoplexes *in vivo* induces accelerated blood clearance and enhances hepatic accumulation of the following injected PEGylated lipoplexes, which is known as the accelerated blood clearance (ABC) phenomenon. Herein, we developed zwitterionic poly(carboxybetaine) (PCB) modified lipoplexes for the delivery of siRNA therapeutics, which could avoid protein adsorption and enhance the stability of lipoplexes as that for PEG. Quite different from the PEGylation, the PCBylated lipoplexes could avoid ABC phenomenon, which extended the blood circulation time and enhanced the tumor accumulation of lipoplexes *in vivo*. After accumulation in tumor site, the PCBylation could promote the cellular uptake and endosomal/lysosomal escape of lipoplexes due to its unique chemical structure and pH-sensitive ability. With excellent tumor accumulation, cellular uptake and endosomal/lysosomal escape abilities, the PCBylated lipoplexes significantly inhibited tumor growth and induced tumor cell apoptosis.

Key words: Cationic liposomes, siRNA therapeutics, Poly(carboxybetaine), Polyethylene glycol, Accelerated blood clearance (ABC) phenomenon.

Introduction

Despite small interfering RNA (siRNA) has garnered much interest as a potential drug for cancer therapy, the clinical application of siRNA still faces considerable obstacles due to its inherent problems such as instability in physiological environment, poor

membrane permeability and short serum half-life [1-4]. Therefore, toward the success in siRNA therapeutics, it is significant to develop efficient and safe vectors to extend the circulation time of siRNA and deliver siRNA into cytoplasm of the target cells [5, 6].

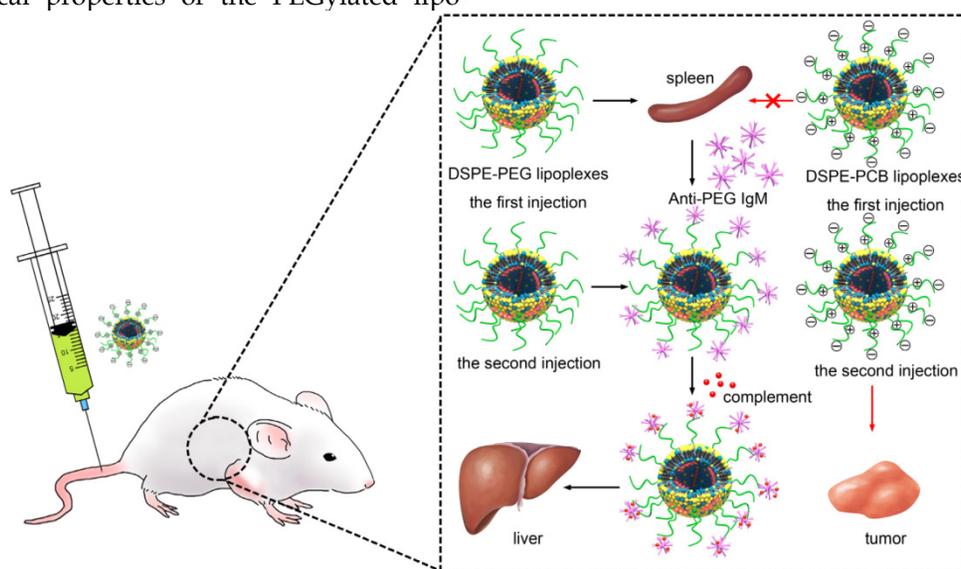
Among them, cationic liposomes have been reported to display a strong ability to protect siRNA from RNase degradation and to accumulate in tumor *via* the enhanced permeability and retention (EPR) effects [7-9]. However, cationic liposome/siRNA complexes (lipoplexes) are prone to have nonspecific interaction with plasma protein *in vivo* and are rapidly cleared by the reticuloendothelial system (RES) before reaching the tumor site, resulting in low gene silencing efficiency [10, 11].

One strategy to circumvent this problem is the use of polyethylene glycol (PEG)-lipids to shield the positive charge on the lipoplexes surface to increase the serum stability and blood circulation time of lipoplexes [12-14]. Unfortunately, the steric hindrance due to PEGylation reduces the electrostatic interactions required for siRNA encapsulation, cellular uptake and endosomal/lysosomal escape of the lipoplexes [15]. Most importantly, it has been shown that upon subsequent administrations of PEGylated liposomes, an immune response could be elicited, leading to a rapid blood clearance, which further compromised the lipoplexes in clinical application [16]. PEGylated lipoplexes are intended to stimulate the spleen to produce anti-PEG IgM after the first administration, which selectively binds to PEG on the surface of the second administered lipoplexes to cause rapid blood elimination and enhanced hepatic uptake (Scheme 1) [17-19]. This effect is known as the "accelerated blood clearance (ABC) phenomenon" [20-22]. The ABC phenomenon represents a tremendous challenge to the clinical application of PEGylated lipoplexes due to the accelerated blood clearance and decreased target diseased site accumulation [23].

There have been many strategies to avoid the induction of ABC phenomenon through changing the physicochemical properties of the PEGylated lipo-

somes. Semple and co-workers used PEG-lipids (PEG-CerC14) with a smaller C14 lipid anchor to modify liposomes to abrogate ABC phenomenon, which could exchange out of the carrier immediately after injection, with greater than 50% loss of PEG-lipid in approximately 3 min. However, the lipid exchange would result in the defects in the membranes of the liposomes [24]. It has also been shown that liposomes modified with cleavable PEG-lipid derivatives (PEG-CHEMS and PEG-CHMC) could lessen or eliminate the ABC phenomenon. The cleavable PEG-lipid derivatives were constructed by linking PEG to lipids through ester bonds, and the chemical bond would gradually be cleaved during circulation. However, the effect on long circulation of the cleavable PEGylated liposomes was worse than that of non-cleavable PEGylated liposomes [25].

Therefore, it is urgently needed to develop a strategy to avoid the immunogenicity of PEGylation without significantly limiting the *in vivo* performance of liposomes. Zwitterionic polymers such as poly (sulfobetaine) (PSB) and poly (carboxybetaine) (PCB) have been demonstrated to be promising alternatives to PEG due to their ability to resist nonspecific protein adsorption [26, 27]. In our previous study, it has been shown that zwitterionic polymer PCB based lipid DSPE-PCB₂₀ had the same ability to enhance the serum stability of lipoplexes. However, quite different from the PEGylation, zwitterionic DSPE-PCB₂₀ could offer stability without interfering with the siRNA encapsulation efficiency and endosomal/lysosomal escape ability of lipoplexes, which was favorable for the systemic delivery of siRNA [28]. However, whether PCBylation could avoid ABC phenomenon and enhance the target tumor accumulation of lipoplexes *in vivo* has not been verified.



Scheme 1. Schematic illustration of the ABC process of the PEGylated and PCBylated lipoplexes.

In this study, cationic liposomes based siRNA delivery system with the modification of PCB was developed. The performances of PCBylated lipoplexes in pharmacokinetics, biodistribution and tumor therapy were investigated. Our findings demonstrated that PCBylation could avoid ABC phenomenon, which facilitated the tumor accumulation and therefore enhanced the antitumor activity of lipoplexes *in vivo* (Scheme 1).

Materials and Methods

Materials

Dimethyldioctadecylammonium bromide (DDAB), cholesterol (95%) and 4',6-diamidino-2-phenylindole dihydrochloride (DAPI) were purchased from J&K Scientific Ltd. 1,2-Distearoyl-sn-glycerol-3-phosphoethanolamine-N-[methoxy (polyethylene glycol)-2000] (DSPE-PEG 2000) was from Shanghai Advanced Vehicle Technology Ltd. Co. LysoTracker Red was obtained from Invitrogen. Glo lysis buffer (×1) was purchased from Promega. Fluorescence labeled Cy5-labeled siRNA (antisense strand, 5'-UUUGAAGUAUGCCUCAAGG dTdT-3'), FAM-labeled siRNA (antisense strand, 5'-UUUGAAGUAUGCCUCAAGGdTdT-3'), as well as siRNA targeting Plk1 mRNA (*siPlk1*, antisense strand, 5'-UAAGGAGGGUGAUCUUCUUCAdTdT-3') were synthesized by Suzhou Ribo Life Science Co. (Kunshan, China). All other reagents used were obtained commercially at analytical grade.

Animals

Male SD rats weighting 150-200 g and female BALB/c nu/nu mice weighting 20-25 g were purchased from the Academy of Military Medical Sciences of China. The animal had free access to water and animal chow. All care and handling of animals was performed in accordance with the protocols approved by the Institutional Animals Care and Use Committee of Peking University.

Preparation and characterization of cationic liposomes and lipoplexes

Distearoyl phosphoethanolamine-poly(carboxybetaine)₂₀ (DSPE-PCB₂₀) was used to make modification of cationic liposomes as it had comparable ability in enhancing the serum stability of lipoplexes with that of DSPE-PEG 2000 in our previous work [28]. Cationic liposomes with composition of DDAB/cholesterol/DSPE-PCB₂₀ or DDAB/cholesterol/DSPE-PEG 2000 (1/1/0.2, molar ratio) were prepared by thin lipid film method. Briefly, all lipids were dissolved in chloroform and were subsequently evaporated to form dry film using a rotary evaporator under vacuum at 45 °C. The resulting thin lipid film

was then hydrated with phosphate buffered saline (PBS, pH=7.4) and sonicated at 37 °C for 30 min. Finally, the cationic liposomes were sized by repeated extrusion through EmulsiFlex-C5 high-pressure homogenizer (Avestin, Canada). The lipoplexes were prepared by mixing cationic liposomes with siRNA together at designed N/P ratios at room temperature for 30 min. The mean diameter and zeta potential of the resulting cationic liposomes and lipoplexes in DMEM were determined using a Malvern Nano ZS instrument (Malvern Instruments). The morphological analysis was carried out by cryogenic transmission electron microscopy (Cryo-TEM, FEI Tecnai 20, The Netherlands).

siRNA encapsulation efficiency

The encapsulation efficiency of siRNA in the lipoplexes was evaluated by Quant-iT™ RiboGreen® RNA Reagent (Invitrogen). The amount of un-encapsulated siRNA in supernatant was measured after centrifugation at 5000×g for 5 min. For low-range assay, 100 µL of the reagent was added to microplate wells and the samples were quantified using microplate reader (SpectraMax M5, Molecular Devices, CA, USA) with the excitation wavelength of 485 nm and emission wavelength of 530 nm. The encapsulation efficiency of siRNA was calculated using the formula:

$$\text{Encapsulation efficiency (\%)} = (F_0 - F_1) / F_0 \times 100\%$$

Where F_0 was the fluorescence of siRNA used for encapsulation, F_1 was the fluorescence of siRNA in supernatant.

Serum stability of lipoplexes

The serum stability of lipoplexes was examined by incubating lipoplexes in DMEM containing 10% FBS at 37 °C under gentle stirring. At predetermined time points, the mean diameter of the lipoplexes was monitored by DLS.

The protonation of DSPE-PCB₂₀ lipoplexes in acidic environment

The ability of protonation in acidic environment for DSPE-PCB₂₀ lipoplexes was determined by incubating lipoplexes in PBs solution with different pH from 7.4 to 3.5 at 37 °C for 30 min. The zeta potential of the lipoplexes was assessed with a Zetasizer Nano ZS instrument.

Cytotoxicity of lipoplexes

The cytotoxicity of lipoplexes was assessed with a MTT viability assay against Hela cells. Hela cells were seeded at 1×10^4 cells per well on a 96-well plate and allowed to adhere overnight at 37 °C for 24 h. Afterwards, 100 µL of various concentration of DSPE-PEG and DSPE-PCB₂₀ lipoplexes at N/P ratio of

5/1 with DMEM containing 10% FBS were added to each well and incubated with the cells for 24 h. 20 μ L of MTT stock solution (5 mg/mL) was then added to each well and incubated for an additional 2 h. After incubation, the medium was replaced with 100 μ L of DMSO and the absorbance was read at 490 nm using a microplate reader (Tecan, Switzerland). Cell viability was normalized to that of HeLa cells cultured in the culture medium with PBS treatment.

Flow cytometry measurement

The cellular uptake of the lipoplexes was evaluated by flow cytometry. Briefly, HeLa cells were seeded in 12-well plates at 1×10^5 cells per well in 500 μ L of culture medium and allowed to adhere for 24 h. FAM-labeled siRNA was complexed with cationic liposomes at N/P ratio of 5/1 for 30 min at room temperature and then added to each well with the DMEM containing 10% FBS with the final concentration of 2 μ g siRNA/mL. The lipoplexes were incubated with cells for 0.25, 0.5, 1, 1.5 and 2.5 h, respectively. Afterwards, the medium was removed and cells were washed three times with cold PBS, trypsinized and harvested in PBS for flow cytometry analysis using BD Calibur flow cytometry (BD Co., USA).

To further elucidate the mechanisms underlying the cellular internalization of DSPE-PEG and DSPE-PCB₂₀ lipoplexes, the cellular uptake study was performed at 4 °C or in the presence of various endocytic inhibitors. Briefly, HeLa cells were pre-incubated with the endocytic inhibitors for 30 min, and the lipoplexes with FAM-labeled siRNA at N/P ratio of 5/1 were added for 2 h uptake experiment at 37 °C. Afterwards, the medium was removed and cells were washed three times with cold PBS, trypsinized and harvested in PBS for flow cytometry analysis using BD Calibur flow cytometry (BD Co., USA). Results were expressed as percentage uptake level of control cells that were incubated with lipoplexes at 37 °C for 2 h in the absence of endocytic inhibitors.

Confocal microscopy

Cellular internalization and location of cationic liposomes with FAM-labeled siRNA were tested using confocal microscopy. Briefly, HeLa cells were seeded in culture dish and incubated for 24 h. Afterwards, DSPE-PEG and DSPE-PCB₂₀ lipoplexes at N/P ratio of 5/1 including 1 μ g FAM-labeled siRNA were added to each well and incubated with the cells for 2 h and 8 h, respectively. Subsequently, the cells were washed three times with PBS followed by staining with LysoTracker Red for 30 min at 37 °C. After another three rinses with PBS, the cells were fixed by 4%

paraformaldehyde for 10 min and the nuclei were stained using DAPI for 10 min. The cells were then detected using a Zeiss LSM780 confocal microscopy (Zeiss Co., Germany).

Quantitative evaluation of intracellular siRNA release

To determine the amount of siRNA release from DSPE-PEG and DSPE-PCB₂₀ lipoplexes inside the cells, HeLa cells were seeded in 12-well plates (1×10^5) for 24 h. The cationic liposome/FAM-labeled siRNA complexes were added to each well with the concentration of 2 μ g/mL and then incubated for 8 h. At designed time point, the cells were washed with PBS twice and lysed in 100 μ L of lysis buffer for 30 min. After the centrifugation of lysates at $15000 \times g$ for 15 min at 4 °C, 50 μ L of the supernatant was transferred to a black 96-well plate for fluorescence measurement using microplate reader with the excitation wavelength of 488 nm and emission wavelength of 525 nm. The percentage of released siRNA of all internalized siRNA was calculated using the following equation:

$$\text{siRNA released (\%)} = (F_1 - F_2) / (F_0 - F_2) \times 100\%$$

Where F_0 was the fluorescence of all internalized siRNA (lysis buffer is Glo lysis buffer and 2% SDS), F_1 was the fluorescence of siRNA in supernatant (lysis buffer is only Glo lysis buffer), F_2 was the fluorescence induced by untreated cells.

Blood clearance and tissue accumulation of lipoplexes

To determine the blood clearance and tissue accumulation of lipoplexes, SD rats were randomly divided into 5 groups ($n=3$). For the first injection, rats were intravenously injected with DSPE-PEG and DSPE-PCB₂₀ lipoplexes at a dose of 1 mg Cy5-labeled siRNA/kg. Control animals received PBS instead of lipoplexes. Five days later, the same lipoplexes were injected intravenously to the same rats at a dose of 1 mg Cy5-labeled siRNA/kg. At selected post-injection time points after two injections, blood (200 μ L) was sampled *via* eye puncture. The liver and spleen were removed 1 h after the injection. Plasma was collected after centrifugation at $903 \times g$ for 10 min at 4 °C. The concentration of the Cy5-labeled siRNA in plasma was detected using microplate reader (SpectraMax M5, Molecular Devices, CA, USA) for fluorescence measurement after demulsification with 2% Tween 80. The distribution of Cy5-labeled siRNA in liver and spleen was detected using a Kodak *in vivo* imaging system (Kodak In-Vivo Imaging System FX Pro, Carestream Health, USA).

Detection of IgM and IgG protein level

SD Rats were pre-treated with DSPE-PEG and

DSPE-PCB₂₀ lipoplexes (N/P=5/1) at a dose of 1 mg Cy5-labeled siRNA/kg (n=5). Five days later, blood samples were collected *via* eye puncture. Plasma was collected after centrifugation at 903×g for 10 min at 4 °C. The IgM level in plasma was determined using ELISA method according to the manufacturer's instructions (Rat IgM Ready-SET-Go!®, eBioscience). The absorbance was measured at 450 nm using a microplate reader (SpectraMax M5, Molecular Devices, CA, USA).

200 µL of plasma samples collected on day 5 after the first injection of DSPE-PEG lipoplexes were incubated with 50 µL of PBS, DSPE-PEG and DSPE-PCB₂₀ lipoplexes at 37 °C for 15 min. After incubation, lipoplexes were separated out by centrifugation at 14462×g for 30 min. The IgM in supernatant was detected using ELISA method according to the manufacturer's instructions (Rat IgM Ready-SET-Go!®, eBioscience). The absorbance was measured at 450 nm using a microplate reader.

To detect the IgG protein level, other SD rats were pre-treated with two injections of DSPE-PEG and DSPE-PCB₂₀ lipoplexes (N/P=5/1) at a dose of 1 mg Cy5-labeled siRNA/kg (n=5), respectively. The time interval between the two injections was 7 days. Two weeks after the second injection, blood samples were collected *via* eye puncture. Plasma was collected after centrifugation at 903×g for 10 min at 4 °C. The IgG level in plasma was determined using ELISA method (Rat IgG total Ready-SET-Go!®, eBioscience) according to the manufacturer's instruction. The absorbance was measured at 450 nm using a microplate reader (SpectraMax M5, Molecular Devices, CA, USA).

The standard curves of IgM and IgG level were shown in Supplementary Material: Figure S1.

Biodistribution of lipoplexes in tumor-bearing nude mice

To detect the biodistribution of DSPE-PEG and DSPE-PCB₂₀ lipoplexes in Hela tumor-bearing nude mice, mice were randomly divided into 4 groups (n=10). For the first injection, mice were intravenously injected with PBS, Cy5-labeled siRNA, DSPE-PEG and DSPE-PCB₂₀/Cy5-labeled siRNA lipoplexes (N/P=5/1) at a dose of 0.5 mg Cy5-labeled siRNA/kg, respectively. 24 h after the first injection, five mice of each group were sacrificed and the main organs and tumor were harvested. Five days later, the same samples were injected intravenously to the same mice at a dose of 0.5 mg Cy5-labeled siRNA/kg. At 24 h post-injection, the mice were sacrificed and the main organs and tumor were harvested. The biodistribution of Cy5-labeled siRNA *in vivo* in organs and tumor was detected using a Kodak *in vivo* imaging system (Ko-

dak In-Vivo Imaging System FX Pro. Carestream Health, USA).

To further quantify the accumulated dose of Cy5-labeled siRNA in liver and tumor, the excised tissues were homogenized in 300 or 1000 µL lysis buffer (300 µL for tumors and 1000 µL for livers) and incubated at 65 °C for 10 min. 100 µL of supernatant was collected after centrifugation at 16162×g for 10 min. Cy5-labeled siRNA in supernatant was extracted and quantified by microplate reader (SpectraMax M5, Molecular Devices, CA, USA) with the excitation wavelength of 649 nm and emission wavelength of 670 nm. The accumulated dose in each tissue was calculated from a standard curve obtained by extracting known amounts of Cy5-labeled siRNA in tissues obtained from un-injected animals [29].

Tumor suppression study

The anti-tumor efficacy was investigated using Hela tumor-bearing nude mice. When the tumor volume reached approximately 50 mm³, mice were randomly divided into four groups (n=5) and treated with PBS, siPlk1, DSPE-PEG and DSPE-PCB₂₀ lipoplexes with siPlk1 at a dose of 2.5 mg/kg every other day, respectively. Body weight and tumor volume were monitored before each injection. The tumor volume was measured using the formula:

$$V (\text{mm}^3) = (a \times b^2) / 2$$

Where a and b are the major and minor axes of the tumor, respectively.

At 24 h post-injection, the mice were sacrificed and the main organs and tumor were harvested. The level of Plk1 mRNA and protein in tumor was analyzed by qRT-PCR and WB. The apoptosis of tumor sections was observed after staining with terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling (TUNEL) assay.

Statistical analysis

Quantitative data were expressed as mean ± SD. Means were compared using Student's *t* test. *p* values < 0.05 were considered statistically significant.

Results and Discussion

The characteristics of cationic liposomes

Zwitterionic polymer based lipid distearoyl phosphoethanolamine-poly(carboxybetaine)₂₀ (DSPE-PCB₂₀) was used to make modification of cationic liposomes as it had comparable ability in enhancing the serum stability of cationic liposomes with that of DSPE-PEG 2000 in our previous work (Figure 1A) [28]. As shown in Figure 1B, DSPE-PEG and DSPE-PCB₂₀ cationic liposomes had similar diameters with narrow distributions, about 119.2±16.2 nm and

113.2±20.9 nm, respectively. The morphology of cationic liposomes was further observed by Cryo-TEM. The Cryo-TEM images in Figure 1C showed that both DSPE-PEG and DSPE-PCB₂₀ cationic liposomes had spherical shape with an average size of 80 nm, which is slightly smaller than that measured by DLS due to the dehydration during sample preparation. The surface zeta potentials of both PEGylated and PCBylated cationic liposomes in DMEM medium were comparable, about 10.2±2.7 mV and 14.2±3.0 mV, which is significantly lower than that of unmodified cationic liposomes (33.5±4.5 mV) with the same molar ratio of DDAB and cholesterol (Figure 1B). The results indicated that PCB had the same ability to reduce the surface charge of cationic liposomes as PEG did.

The positive charge of cationic liposomes allowed for binding of negatively charged siRNA to form lipoplexes. To assess the siRNA binding ability of DSPE-PEG and DSPE-PCB₂₀ cationic liposomes, percentage of siRNA encapsulation was further determined by Quant-iT™ RiboGreen® RNA Reagent. The encapsulation efficiency was only 14.6±5.4% for DSPE-PEG cationic liposomes at N/P ratio of 1/1, while the encapsulation efficiency could reach to 49.0±5.3% for DSPE-PCB₂₀ cationic liposomes (Figure 1D). The excellent siRNA binding ability of DSPE-PCB₂₀ cationic liposomes might due to the unique structure of zwitterionic polymers that the cationic quaternary amine groups assisted the cationic liposomes in retaining the siRNA [30].

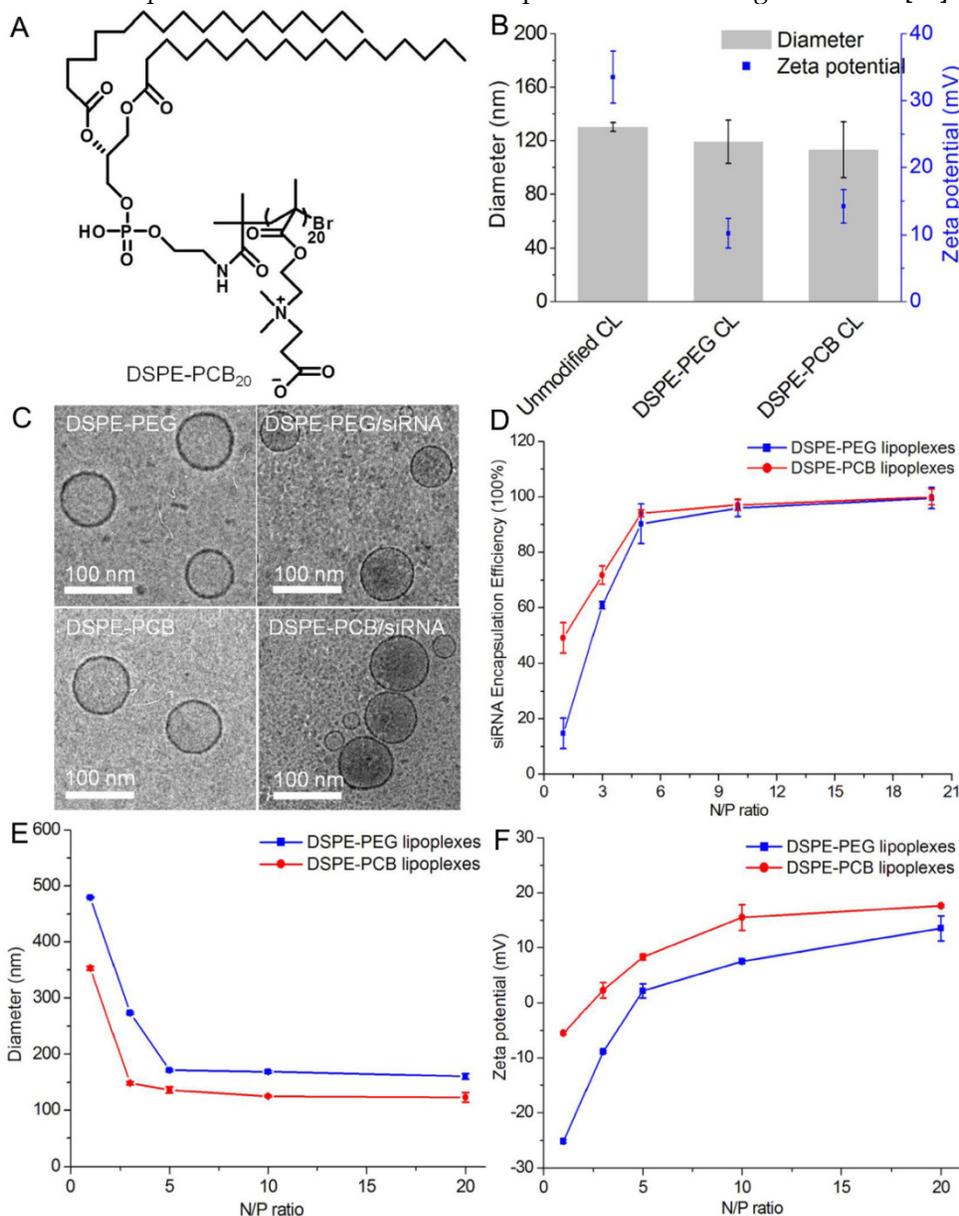


Figure 1. Characterization of cationic liposomes and lipoplexes. (A) The chemical structure of DSPE-PCB₂₀. (B) Mean particle diameter and zeta potential of cationic liposomes in DMEM. (C) Cryo-TEM images of cationic liposomes and lipoplexes at N/P ratio of 5/1. (D) The siRNA encapsulation efficiency of lipoplexes evaluated by Quant-iT™ RiboGreen® RNA Reagent. (E) The mean diameter of lipoplexes at various N/P ratios in DMEM. (F) The zeta potential of lipoplexes at various N/P ratios in DMEM. Data are shown as the mean ± S.D. of three independent experiments.

The size and surface property of lipoplexes are important physiochemical parameters in designing the cancer-targeting delivery system [31]. As shown in Figure 1E, the diameter of both lipoplexes was decreased with the increase of N/P ratio. It was about 100-200 nm when the N/P ratio ranging from 3/1 to 20/1, which was suitable for the prolonged blood circulation to perform the EPR effect [32]. On the contrary, the corresponding zeta potential values of lipoplexes were increased with the increase of N/P ratio. As shown in Figure 1D, the siRNA encapsulation efficiency of lipoplexes was low at N/P ratio of 1/1 or 3/1, which was not favorable for siRNA delivery. When the N/P ratio reached 5/1 or larger than 5/1, the siRNA encapsulation efficiency was nearly 100%. However, extra cationic liposomes could induce cytotoxicity [33, 34]. Therefore, lipoplexes with N/P ratio of 5/1 were chosen for the following experiments as they had appropriate siRNA encapsulation efficiency, size and zeta potentials characteristics. Additionally, both DSPE-PEG and DSPE-PCB₂₀ lipoplexes at N/P ratio of 5/1 had spherical shape (Figure 1C).

Serum stability of lipoplexes

The instability of unmodified lipoplexes when exposed to plasma proteins impeded the clinical use of cationic liposomes as systemic delivery vehicles of siRNA [35]. Therefore, the serum stability of DSPE-PCB₂₀ lipoplexes was investigated whether the PCB modification could overcome this drawback. DSPE-PCB₂₀ lipoplexes at N/P ratio of 5/1 were incubated with DMEM containing 10% FBS, and the changes in the sizes of the lipoplexes were analyzed by DLS measurements. As shown in Figure 2 and Supplementary Material: Figure S2, both DSPE-PCB₂₀ and DSPE-PEG lipoplexes exhibited excellent serum stability as demonstrated by the consistent diameter after incubation for 84 h at 37 °C. The results indicated that the presence of PCB on the surface of lipoplexes prevented nonspecific protein adsorption and aggregation of the lipoplexes as PEG did.

In vitro cytotoxicity of lipoplexes

The biocompatibility of lipoplexes was crucial for its clinical application. Therefore, the cytotoxicity of DSPE-PEG and DSPE-PCB₂₀ lipoplexes to Hela cells was evaluated using the MTT assay. As shown in Figure 3, cell viability of DSPE-PEG and DSPE-PCB₂₀ lipoplexes at N/P ratio of 5/1 remained nearly 90% when the siRNA concentration reached 8 µg/mL. The results suggested that DSPE-PCB₂₀ lipoplexes had good biocompatibility as that for DSPE-PEG lipoplexes, which were suitable for *in vivo* application.

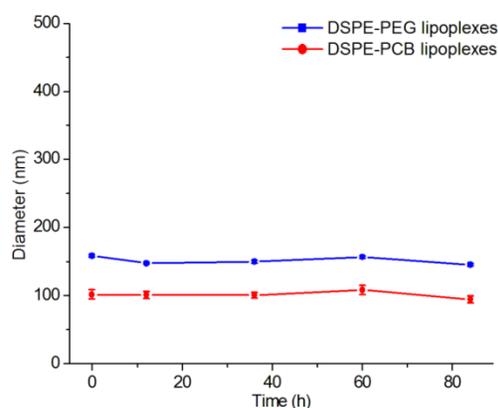


Figure 2. The changes of lipoplexes sizes in the presence of 10% FBS with the extension of time. Data are shown as the mean \pm S.D. of three independent experiments.

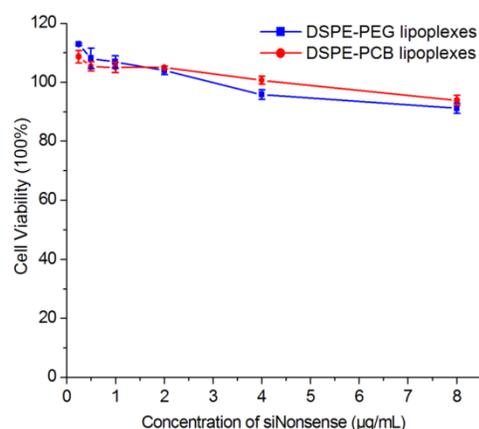


Figure 3. Cytotoxicity of DSPE-PEG and DSPE-PCB₂₀ cationic liposomes with siNonsense to Hela cells.

ABC phenomenon of PCBylated lipoplexes

PEGylated cationic liposome/siRNA complexes could induce ABC phenomenon after repeated administration, which resulted in enhanced liver accumulation and decreased tumor accumulation of lipoplexes [36, 37]. The ABC phenomenon significantly impeded the clinical use of PEGylated lipoplexes [38]. In our previous work, it has been shown that PCBylation could overcome the barriers associated with the use of PEGylation in cellular level [28]. However, whether PCBylated lipoplexes would avoid ABC phenomenon *in vivo* has not been verified. To answer the question, two doses of DSPE-PCB₂₀ lipoplexes were injected into rats with a time interval of 5 days. As shown in Figure 4A, pretreatment rats with DSPE-PEG lipoplexes triggered a rapid clearance of the second dose from the circulatory system, in which siRNA concentration in plasma dropped to 48.74% ID within 5 min. Additionally, 1 hour's biodistribution after the second injection showed a highly increased liver uptake, about 2 times of that for the first injection (Figure 4C). However, the blood clearance trend and

hepatic uptake of DSPE-PCB₂₀ lipoplexes were basically the same for the two injections (Figure 4B and 4D). The half-life of the two injections for DSPE-PEG lipoplexes and DSPE-PCB₂₀ lipoplexes was calculated according to the siRNA concentration profiles, respectively. The half-life of DSPE-PEG lipoplexes for the first injection was 18.72 h, and only 6.45 h for the second injection, while it was about 20.05 h for both injections of DSPE-PCB₂₀ lipoplexes. The results indicated that PCBylation could avoid ABC phenomenon and enhance the blood circulation time of lipoplexes.

It has been confirmed that there was a positive correlation between the anti-PEG IgM production and the accelerated clearance of the PEGylated liposomes [39, 40]. Whether the PCBylated lipoplexes would initiate the production of anti-PCB IgM was evaluated by administration of DSPE-PCB₂₀ lipoplexes on rats. The plasma IgM level was detected using ELISA 5 days later. DSPE-PEG lipoplexes obviously induced the production of IgM, which was about 4 times of that for PBS groups (Figure 5A). In contrast, the IgM level has no significant difference compared with the PBS groups for the DSPE-PCB₂₀ lipoplexes. The results demonstrated that PCBylation would not stimulate the production of IgM, and therefore would avoid the ABC phenomenon against the following administration.

The IgM level reached peaked after 5 days of the first injection and it has also been shown that the IgG

protein level continued to increase with additional injections. The production of IgG is dependent on IgM stimulation and it is a secondary and more effective immune response to foreign materials in the body [41]. The IgG level was tested two weeks after the second injection and the second injection was 7 days after the first injection. As shown in Fig. 5B, DSPE-PEG groups induced the production of IgG, which was about 2.5 times of that for PBS group. In contrast, there was no significant difference between the PBS groups and the DSPE-PCB₂₀ lipoplexes groups. The result further confirmed that the DSPE-PCB could avoid the stimulation of ABC phenomenon.

To further determine whether the generated IgM specifically bound to PEGylated lipoplexes, the PEGylated and PCBylated lipoplexes were incubated with the plasma obtained from rats received DSPE-PEG lipoplexes 5 days later. As shown in Figure 6, the IgM level was significantly decreased after incubation with the DSPE-PEG lipoplexes, while there was no difference of IgM level after incubation with DSPE-PCB₂₀ lipoplexes. The results supported the idea that the elevated IgM selectively bound to DSPE-PEG lipoplexes and activated the ABC phenomenon. In contrast, PCBylation could avoid the ABC phenomenon and extend the blood circulation time of lipoplexes.

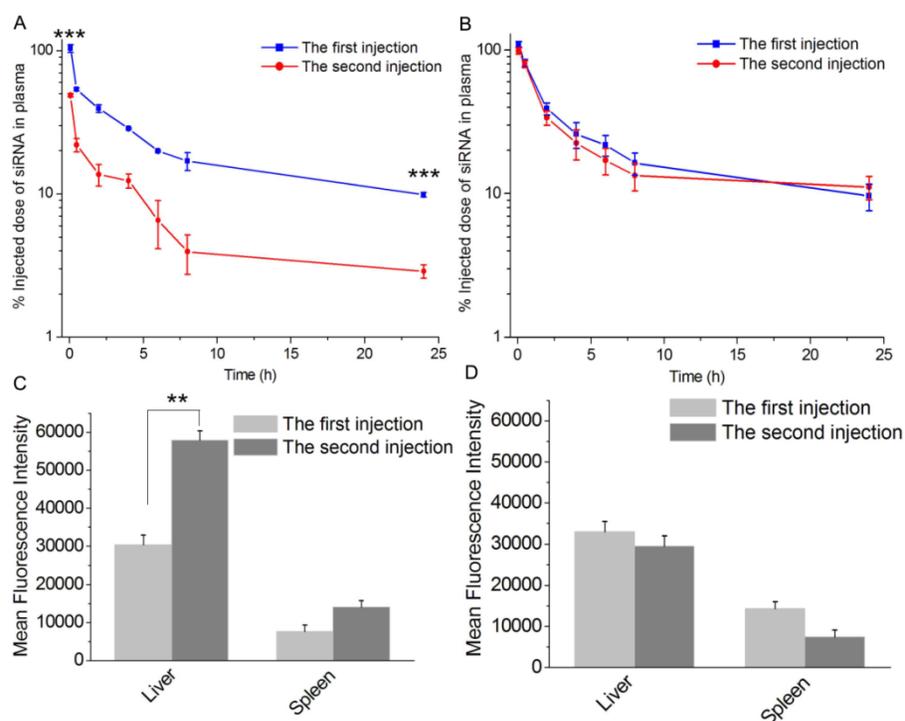


Figure 4. Pharmacokinetic and biodistribution study of DSPE-PEG and DSPE-PCB₂₀ lipoplexes in rats after two doses of injection with time interval of 5 days. (A) Blood clearance profile of DSPE-PEG lipoplexes. (B) Blood clearance profile of DSPE-PCB₂₀ lipoplexes. (C) Distribution of DSPE-PEG lipoplexes in liver and spleen. (D) Distribution of DSPE-PCB₂₀ lipoplexes in liver and spleen. Lipoplexes at N/P ratio of 5/1 were administered to SD rats with Cy5-labeled siRNA concentration of 1 mg/kg via the tail vein. Data are shown as the mean \pm S.D. of three independent experiments. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.005$ ($n = 3$).

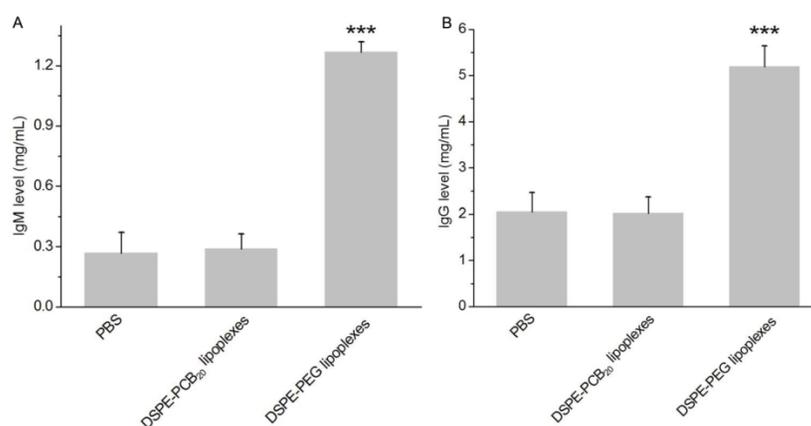


Figure 5. (A) IgM level in plasma of SD rats following administrated with DSPE-PEG and DSPE-PCB₂₀ lipoplexes 5 days later. (B) IgG level in plasma of SD rats following administrated with two injections of DSPE-PEG and DSPE-PCB₂₀ lipoplexes. The time interval between the two injections was 7 days and the plasma was obtained two weeks after the second injection. Lipoplexes at N/P ratio of 5/1 were administered to SD rats with Cy5-labeled siRNA concentration of 1 mg/kg via the tail vein. Data are shown as the mean \pm S.D. of three independent experiments. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.005$ (n=5).

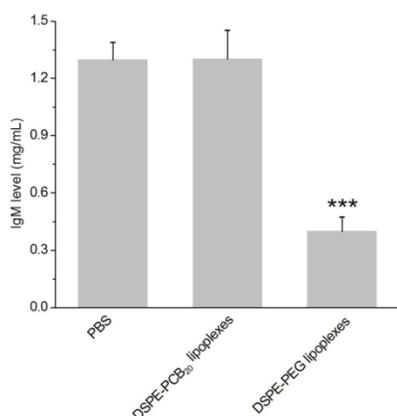


Figure 6. IgM level in plasma after incubation with DSPE-PEG and DSPE-PCB₂₀ lipoplexes at 37 °C for 15 min. The plasma was obtained from SD rats administrated with DSPE-PEG lipoplexes with Cy5-labeled siRNA concentration of 1 mg/kg. Data are shown as the mean \pm S.D. of three independent experiments. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.005$ (n=3).

Biodistribution of lipoplexes in tumor-bearing nude mice

The ABC phenomenon against the following dose of DSPE-PEG lipoplexes might interfere with its accumulation in tumor site. To demonstrate this, the biodistribution of lipoplexes with Cy5-labeled siRNA at N/P ratio of 5/1 was analyzed after two injections with a time interval of 5 days on Hela tumor-bearing nude mice. In comparison with naked Cy5-labeled siRNA, all lipoplexes formulations significantly enhanced the siRNA accumulation in tumor site due to the EPR effect. As shown in Figure 7A and 7C, the mean fluorescence intensity of Cy5-labeled siRNA in liver and tumor for DSPE-PEG and DSPE-PCB₂₀ was basic the same after the first injection. As expectedly, the mean fluorescence intensity of Cy5-labeled siRNA in liver for DSPE-PEG lipoplexes was about 1.5 times of that for DSPE-PCB₂₀ lipoplexes after the second injection, while the mean fluorescence intensity of Cy5-labeled siRNA in tumor for DSPE-PCB₂₀ lipoplexes was about 1.3 times of that for DSPE-PEG lip-

oplexes (Figure 7B and 7D). As shown in Figure 7E and 7F, the quantitative results further confirmed the results. The results demonstrated that DSPE-PCB₂₀ avoided the ABC phenomenon, which resulted in higher accumulation of Cy5-labeled siRNA in tumor for the following injection.

Cellular uptake and intracellular distribution of lipoplexes

After the lipoplexes accumulated in tumor site, the siRNA molecules must be transported into tumor cells, and more siRNA in the cytoplasm would induce more significant gene silencing effect [42]. The cellular uptake of DSPE-PCB₂₀ lipoplexes at N/P ratio of 5/1 was evaluated after incubation with Hela cells by flow cytometric analysis. As shown in Figure 8A, the fluorescence intensity of FAM-labeled siRNA in Hela cells was increased with the extension of culturing time from 0.25 to 2.5 h. In addition, the fluorescence intensity of FAM-labeled siRNA for DSPE-PCB₂₀ lipoplexes was about 2 times of that for DSPE-PEG lipoplexes after 2.5 h incubation, indicating that DSPE-PCB₂₀ lipoplexes had better cellular uptake ability than that of DSPE-PEG lipoplexes. This might be attributed to the unique chemical structure of DSPE-PCB₂₀ that the cationic quaternary amine groups assisted the cationic liposomes in retaining the siRNA and cellular uptake.

To further probe the mechanisms underlying the cellular internalization of lipoplexes, the cellular uptake study was performed at lower temperature of 4 °C or in the presence of various endocytic inhibitors. As shown in Figure 8B, DSPE-PCB₂₀ and DSPE-PEG lipoplexes were internalized *via* energy dependent endocytosis, as cellular uptake of both lipoplexes was decreased about 70% at 4 °C. In addition, chlorpromazine and m β CD significantly inhibited the cellular uptake level of both lipoplexes, indicating that the lipoplexes were endocytosed *via* both the caveolae- and clathrin-mediated endocytosis pathways [43].

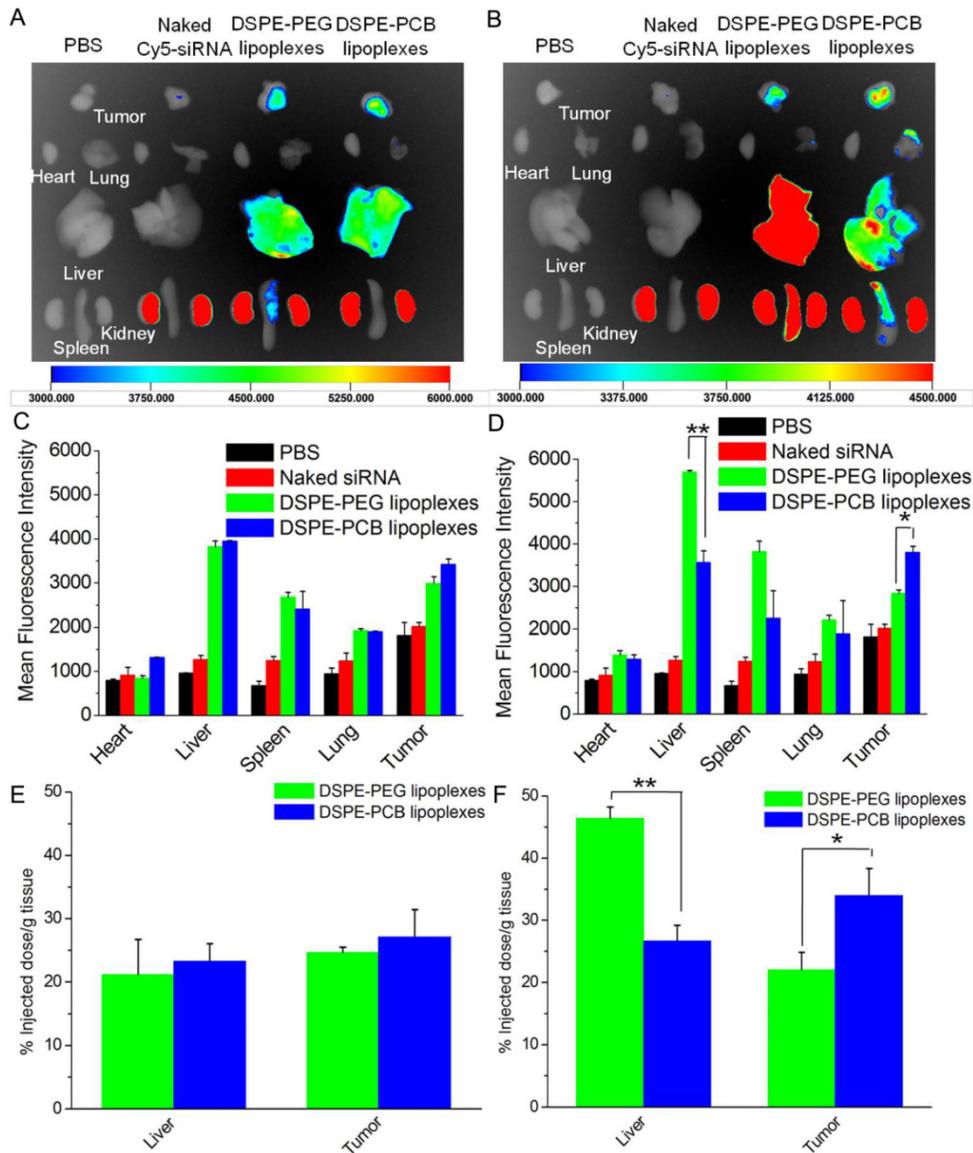


Figure 7. *In vivo* distribution of lipoplexes in Hela tumor-bearing nude mice after 24 h of two injections. (A) The fluorescence images of major organs and tumor after 24 h of the first injection. (B) The fluorescence images of major organs and tumor after 24 h of the second injection. (C) The mean fluorescence intensity of Cy5-labeled siRNA of major organs and tumor of Figure A. (D) The mean fluorescence intensity of Cy5-labeled siRNA of major organs and tumor of Figure B. (E) Quantitative results of Cy5-labeled siRNA in liver and tumor of the first injection. (F) Quantitative results of Cy5-labeled siRNA in liver and tumor of the second injection. Lipoplexes at N/P ratio of 5/1 were administered to mice at a dose of 0.5 mg Cy5-labeled siRNA/kg via the tail vein. Data are shown as the mean \pm S.D. of three independent experiments. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.005$ ($n = 3$).

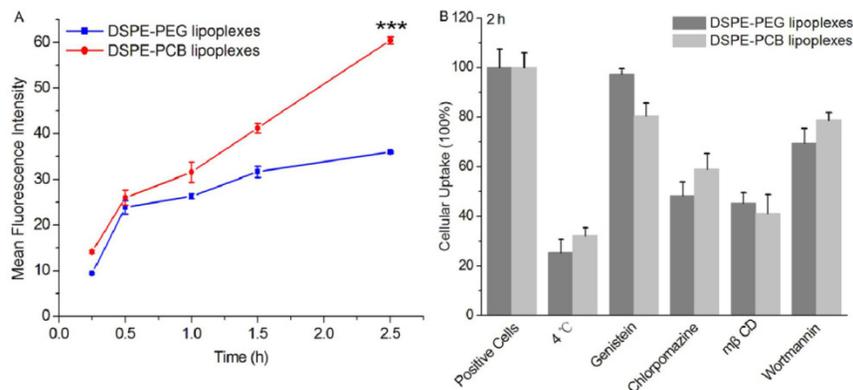


Figure 8. Flow cytometric analyses of cellular internalization of cationic liposome/FAM-labeled siRNA lipoplexes in Hela cells after incubation for different times. (A) Comparison of cellular internalization between DSPE-PEG and DSPE-PCB₂₀ lipoplexes with the extension of time. (B) Mechanistic probes of the intracellular kinetics of the DSPE-PEG and DSPE-PCB₂₀ lipoplexes by monitoring the cellular uptake level at 4 °C or in the presence of various endocytic inhibitors. Data are shown as the mean \pm S.D. of three independent experiments.

As lipoplexes enter cells by an endocytosis pathway, the endosomal/lysosomal escape of siRNA is important for the subsequent post-transcriptional gene silencing in the cytoplasm [44]. To investigate whether siRNA loaded lipoplexes could efficiently escape from endosomes/lysosomes, Hela cells were treated with DSPE-PEG and DSPE-PCB₂₀ cationic liposome/FAM-labeled siRNA complexes at N/P ratio of 5/1 for different periods of time, and the localization of lipoplexes in cells was observed by confocal laser scanning microscopy (CLSM). The endosomes/lysosomes were stained with LysoTracker Red, and the cell nuclei were counterstained with DAPI. As shown in Figure 9A, both DSPE-PEG and DSPE-PCB₂₀ lipoplexes were mainly colocalized with the LysoTracker Red stained organelles after 2 h of incubation, suggesting that lipoplexes resided in endosomes/lysosomes. However, after 8 h of incubation, the separation of the green and red fluorescence was more significant when the cells were incubated with DSPE-PCB₂₀ lipoplexes, suggesting that FAM-labeled siRNA more efficiently escaped from the endosomes/lysosomes following the delivery of DSPE-PCB₂₀ lipoplexes. In contrast, the FAM-labeled

siRNA still resided in endosomes/lysosomes for DSPE-PEG lipoplexes after 8 h of incubation. The quantitative results of co-localization ratio for DSPE-PCB₂₀ lipoplexes decreased about 41.6%, while only 22.8% for DSPE-PEG lipoplexes after 8 h incubation (Figure 9B). The percentage of released FAM-labeled siRNA from cationic liposomes of all internalized siRNA at various time points also confirmed the expedited endosomal/lysosomal escape of DSPE-PCB₂₀ lipoplexes. As shown in Figure 9C, the cumulative release of siRNA at 8 h was 53.6±2.0% for DSPE-PCB₂₀ lipoplexes, while only 12.9±1.4% for DSPE-PEG lipoplexes. As demonstrated in our previous work, the excellent endosomal/lysosomal escape ability of DSPE-PCB₂₀ lipoplexes was due to the pH sensitive ability of PCB₂₀ that the carboxyl acid groups of PCB₂₀ was protonation under acidic condition, resulting in increased zeta potential from 4.1±0.1 mV at pH 7.4 to 20.2±0.2 mV at pH 3.5 (Figure 9D). The increase in positive charge of DSPE-PCB₂₀ lipoplexes promoted the fusion of lipoplexes with anionic endosomal/lysosomal membrane, and releasing siRNA into the cytoplasm [28].

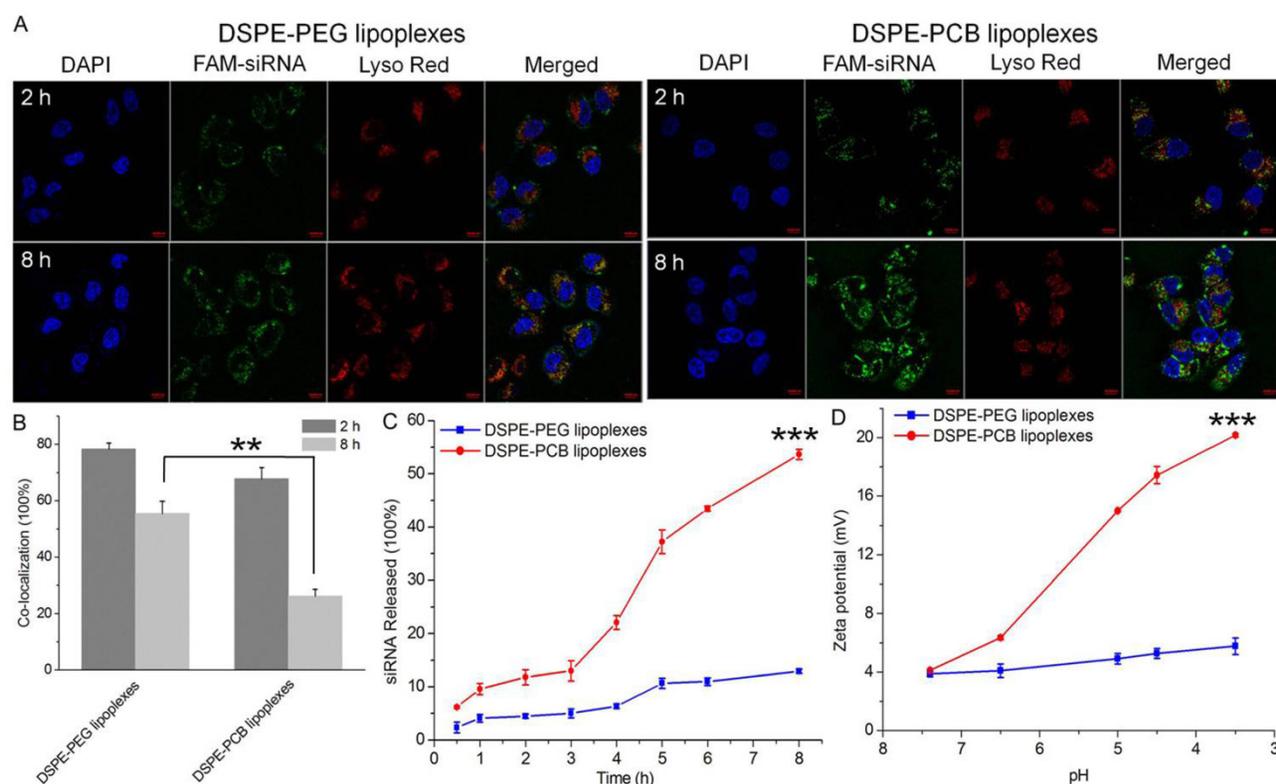


Figure 9. Assessment of cellular internalization and endosomal/lysosomal escape of lipoplexes in Hela cells after incubation for different time points. (A) Images detected by confocal laser scanning microscopy (CLSM). Cell nuclei and endosomes/lysosomes were stained with DAPI (blue) and LysoTracker Red (red). Scale bar is 10 μ m. (B) Co-localization ratio of FAM-labeled siRNA and LysoTracker Red after 2 h and 8 h incubation. (C) The percentage of siRNA released from lipoplexes after incubation with the extension of time. (D) The changes of zeta potential of lipoplexes in different pH values PBs. Data are shown as the mean \pm S.D. of three independent experiments.

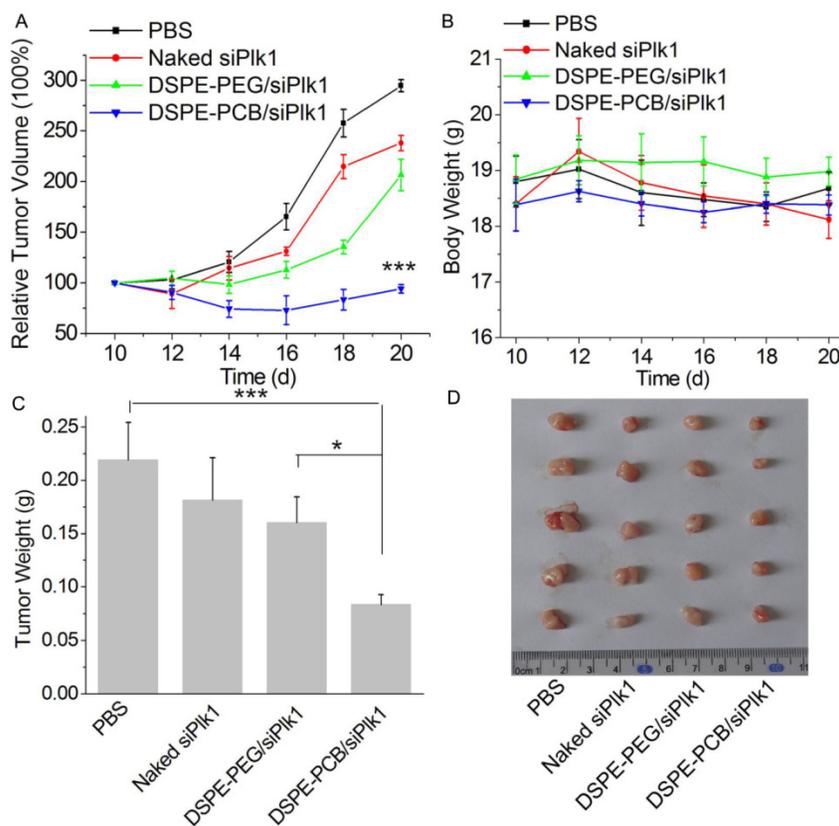


Figure 10. *In vivo* anti-tumor study of lipoplexes in Hela tumor-bearing nude mice after intravenous injection with PBS, siPlk1, DSPE-PEG and DSPE-PCB₂₀ lipoplexes with siPlk1 at N/P ratio of 5/1 at siPlk1 dose of 2.5 mg/kg (n=5). (A) Relative tumor volume-time curve. (B) Body weight-time curve. (C) The weights of the removed tumors. (D) The image of the solid tumors removed from different treatment groups. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.005$.

Tumor therapeutic activity of lipoplexes

It has been demonstrated that PCBylation could avoid ABC phenomenon and enhance the tumor accumulation of lipoplexes after repeated injections *in vivo*. After the lipoplexes accumulating in tumor site, PCBylation could promote the siRNA molecules be transported into tumor cells, and accelerate its release into cytoplasm. To reveal the antitumor growth effect of DSPE-PCB₂₀ lipoplexes *in vivo*, Hela tumor-bearing nude mice were received intravenous injection of DSPE-PCB₂₀ lipoplexes at siRNA dose of 2.5 mg/kg. As illustrated in Figure 10A, treatment with siPlk1 alone did not show significantly tumor growth inhibition in comparison with PBS groups. The DSPE-PCB₂₀ lipoplexes with siPlk1 significantly inhibited tumor growth with the final relative tumor volume (RTV) of 94.2 ± 4.2 , which was much lower than that of DSPE-PEG lipoplexes groups (206.5 ± 15.6). At the same time, both the DSPE-PEG and DSPE-PCB₂₀ lipoplexes exhibited neglectable toxicity, which indicated the good biocompatibility of PEGylated and PCBylated lipoplexes (Figure 10B). The excised tumor weight also exhibited the corresponding weight (Figure 10C and 10D). The results indicated that DSPE-PCB₂₀ lipoplexes with siPlk1 exhibited excellent antitumor activity.

To demonstrate that retarded tumor growth by DSPE-PCB₂₀ lipoplexes was related to Plk1 down-regulation in tumor cells, the tumors were excised 24 h after the last injection. Tumor Plk1 mRNA and protein expression were analyzed by qRT-PCR and WB analyses, respectively. As shown in Figure 11A, treatment with siPlk1 alone did not show reductions in Plk1 mRNA level due to its lower tumor accumulation and cellular uptake. In contrast, Plk1 mRNA level showed a $80.4 \pm 2.5\%$ reduction after treatment with DSPE-PCB₂₀ lipoplexes with siPlk1 compared to the level in tumors following treatment with PBS, which is much higher than that of DSPE-PEG lipoplexes ($53.2 \pm 3.8\%$). WB analyses of Plk1 protein levels in each tumor mass (Figure 11B and 11C) also revealed a significant reduction in Plk1 protein levels when the mice were treated with DSPE-PCB₂₀ lipoplexes. These results demonstrated that DSPE-PCB₂₀ lipoplexes could significantly down-regulate the Plk1 gene expression *in vivo*, which resulted in higher efficiency tumor growth suppression. Plk1 inhibition has been shown to be associated with apoptosis induction [45, 46]. Cell apoptosis in the tumors after treatment with various formulations were analyzed by TUNEL (Figure 11D), that the DSPE-PCB₂₀ lipoplexes with siPlk1 induced more significant cell apoptosis in tumors.

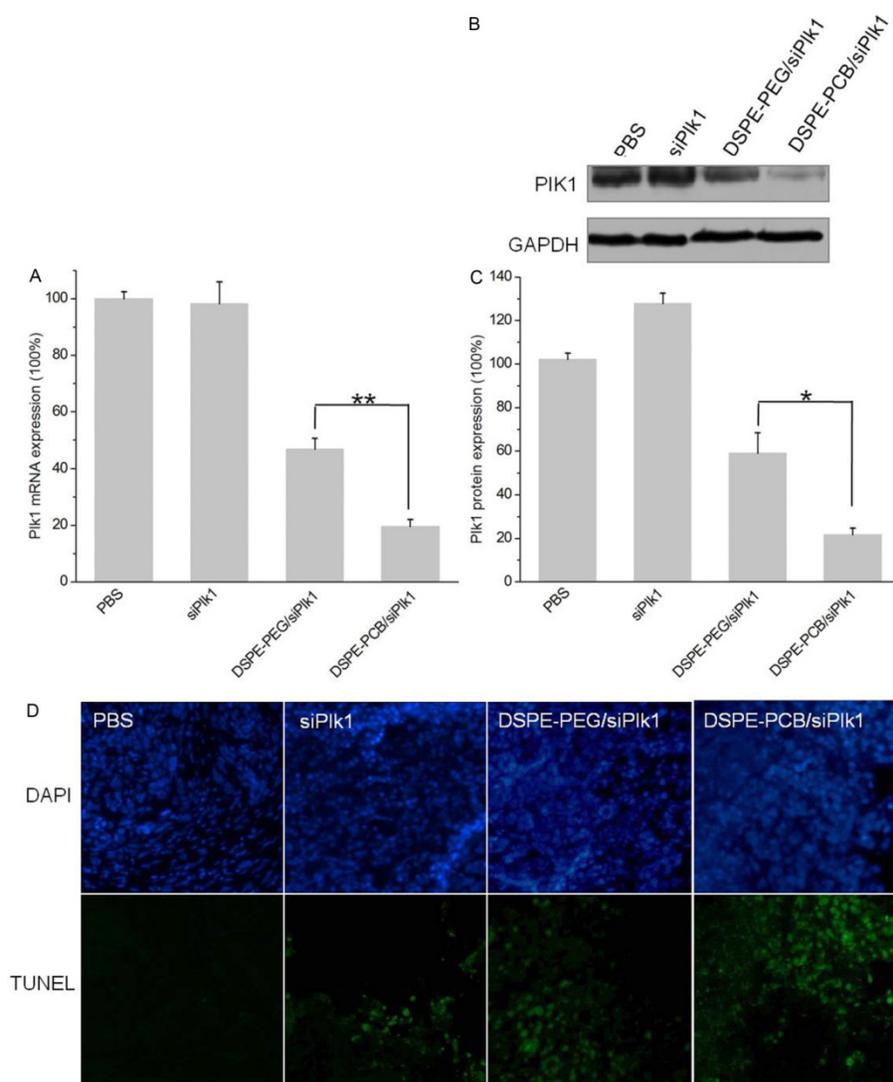


Figure 11. *In vivo* gene silencing efficiency of DSPE-PEG and DSPE-PCB₂₀ lipoplexes. (A) Expression of Plk1 mRNA. (B, C) Expression of Plk1 protein. (D) TUNEL analyses of tumor tissue after treatments. The tumor tissues were collected 24 h after the last injection. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.005$.

Conclusions

We have developed zwitterionic polymer PCB modified cationic liposomes for the systemic delivery of siRNA therapeutics, which could overcome the barriers associated with the use of PEGylation. The PCBylated cationic liposomes could avoid ABC phenomenon, which extended the blood circulation time and enhanced the tumor accumulation of siRNA following injection. In addition, the DSPE-PCB₂₀ cationic liposomes exhibited higher efficiency of siRNA encapsulation, cellular uptake and endosomal/lysosomal escape abilities, which in turn lead to remarkable release siRNA into cytoplasm. With excellent *in vitro* and *in vivo* behavior, the PCBylated cationic liposomes significantly inhibited tumor growth following systemic administration, suggesting its potential application for the delivery of siRNA therapeutics.

Supplementary Material

Figures S1 – S2.

<http://www.thno.org/v05p0583s1.pdf>

Acknowledgments

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Competing Interests

The authors have declared that no competing interest exists.

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