

An efficient method for in vitro gene delivery via regulation of cellular endocytosis pathway

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Abstract: Transfection efficiency was the primary goal for in vitro gene delivery mediated by nonviral gene carriers. Here, we report a modified gene transfection method that could greatly increase the efficiency of, and accelerate the process mediated by, 25 kDa branched polyethyleneimine and Lipofectamine™ 2000 in a broad range of cell strains, including tumor, normal, primary, and embryonic stem cells. In this method, the combination of transfection procedure with optimized complexation volume had a determinant effect on gene delivery result. The superiorities of the method were found to be related to the change of cellular endocytosis pathway and decrease of particle size. The efficient and simple method established in this study can be widely used for in vitro gene delivery into cultured cells. We think it may also be applicable for many more nonviral gene delivery materials than polyethyleneimine and liposome.

Keywords: gene delivery, gene expression, endocytosis, polyethyleneimine, Lipofectamine™ 2000

Introduction

Delivery of exogenous gene into cells has great importance in gene function research, genetic modification of cells, and gene therapy.¹⁻⁴ Viral vector owns the highest gene transfection efficiency, but its wide application has been limited by safety issues.^{5,6} Non-viral gene carriers such as cationic lipids and polymers have been potential alternatives for safe and efficient gene delivery due to their good biocompatibility, versatility, and controllable molecule size.⁷⁻⁹ However, their gene transfection efficiencies are much lower than those of viral vectors.¹⁰ Therefore, increasing gene delivery efficiency has been the primary challenge for nonviral material-mediated gene transfer trials.¹¹

The gene delivery efficiency of nonviral materials is usually influenced by many factors, such as the quantity and quality of deoxyribonucleic acid (DNA), ratio of DNA to material, serum, antibiotics, solvent for DNA and material, incubation time, and mixing order.¹²⁻¹⁸ Therefore, the readily available off-the-shelf gene carriers usually provide standard transfection procedures that have been considered to generate the highest gene transfection efficiency for cultured cells.^{19,20} Branched polyethyleneimine (PEI, Mw ≈ 25 kDa), a typical cationic polymer, has been commonly used as the 'gold standard' for in vitro gene delivery.²¹ Lipofectamine™ 2000 (Lipo2000) is one of the most widely used commercial cationic lipids with high efficiency for in vitro gene transfer.²² Both materials have their recommended optimal transfection procedures that have been used for many years.

In this study, we will present a modified method applicable for improving the efficiency of PEI- and Lipo2000-mediated gene delivery into a wide range of cell strains. Exploration of the mechanism underlying this method may also give us innovative insights into transgene delivery procedure.

Materials and methods

Materials

The plasmids pEGFP-C1 (Clontech, Takara Biotechnology Co. Ltd, Dalian, People's Republic of China) encoding enhanced green fluorescent protein (EGFP) and pGL3-control (Promega, Madison, WI, USA) containing the modified coding region for firefly (*Photinus pyralis*) luciferase (Luc) were amplified in *Escherichia coli* DH5 α and extracted using the PureLink[®] HiPure Plasmid Filter Maxiprep Kit (Invitrogen, Carlsbad, CA, USA). Branched PEI (25 kDa), methyl- β -cyclodextrin (M β CD), amiloride hydrochloride, and chlorpromazine hydrochloride were purchased from Sigma-Aldrich (St Louis, MO, USA) and used without further treatment. Lipo2000, fetal bovine serum (FBS), Dulbecco's Modified Eagle's Medium (DMEM), RPMI-1640, Minimum Essential Medium (MEM with Earle's Balanced Salt Solution), and Opti-MEM[®] I Medium were purchased from Invitrogen.

Cell culture

Hep G2 (human hepatocellular carcinoma; American Type Culture Collection [ATCC] No HB-8065TM), SMMC-7721 (human hepatocellular carcinoma), MCF-7 (human breast adenocarcinoma; ATCC No HTB-22TM), AGS (human gastric adenocarcinoma; ATCC No CRL-1739TM), Hela (human cervix adenocarcinoma; ATCC No CCL-2TM), 4T1 (mouse breast cancer; ATCC No CRL-2539TM), NIH/3T3 (mouse embryo fibroblast; ATCC No CRL-1568TM), C2C12 (mouse muscle myoblast; ATCC No CRL-1772TM), R1 (mouse embryonic stem cell; ATCC No CRL-11114TM), and L6 (rat skeletal muscle myoblast; ATCC No CRL-1458TM) cell lines were purchased from the cell bank of the Chinese Academy of Science (Shanghai, People's Republic of China) and the China Center for Type Culture Collection (Wuhan, People's Republic of China). Cells were cultured according to the recommended procedures by ATCC. For primary fibroblast isolation, a piece of sterile neonatal foreskin was separated, washed with phosphate-buffered saline (PBS), minced thoroughly, and transferred into a 0.1 mM trypsin/1 mM ethylenediaminetetraacetic acid solution. After trypsinization at 37°C for ~30 minutes, an equal volume of DMEM/FBS (10%) was added and the tissue/medium mixture was pipetted up and down a few times for tissue dissociation. After 5-minutes' incubation at room temperature, the supernatant was transferred into a new tube and centrifuged at 2,000 g for 5 minutes at 4°C. Cells were resuspended in fresh DMEM containing 15% FBS and 1% penicillin-streptomycin. All cells were cultured as adherent monolayer in a CO₂ incubator at 37°C under saturated humidity.

Cell transfection

One day before transfection, cells were seeded in six-well or 24-well plates and cultured in growth medium without antibiotics. For PEI-mediated transfection cases, the suitable cell confluency was 70%–80% at the time of transfection, while for Lipo2000, it was 90%–95%. Plasmids 4 μ g/well were used for transfection in six-well and 0.8 μ g/well for 24-well format. PEI, Lipo2000, and pDNA were freshly diluted to equal volumes with Opti-MEM[®] I medium prior to use. The polyplexes were prepared by adding PEI into pDNA using an N/P ratio of 10. The lipoplexes were prepared by adding pDNA into Lipo2000 using a DNA (μ g) to Lipo2000 (μ L) ratio of 1:2.5. The polyplexes and lipoplexes were incubated for 20 minutes at room temperature. In standard PEI and Lipo2000 transfection cases, the growth medium was replaced by fresh FBS⁻/antibiotics⁻ medium (2 mL for six-well and 0.5 mL for 24-well plate) and then the complexes (500 μ L for six-well and 100 μ L for 24-well plate) were added. In the modified transfection procedures for PEI or Lipo2000, the growth medium was completely removed and different volumes of polyplexes or lipoplexes were gently added to the corresponding cells. After incubation at 37°C for a certain time (4 hours for standard or 0.5–4 hours for modified method), the growth medium was replaced with fresh FBS⁺/antibiotics⁺ media and cultured for 1–3 days for detections.

Gene transfection efficiency assays

To qualitatively evaluate the transfection efficiency, cells transfected with pEGFP-C1 were observed and imaged using an inverted fluorescent microscope (Leica DMI 4000B; Wetzlar, Germany) for EGFP expression at days 1 and 2 after transfection.

The transfection efficiency was quantitatively analyzed by detecting and comparing the luciferase activity between the standard and modified transfection groups. Two days after transfection, cells treated with pGL3-control were washed with PBS, digested with 0.25% (w/v) trypsin, harvested in 1 mL PBS, and lysed by adding 200 μ L 1 \times lysis reporter buffer (Promega). The cell lysates were centrifuged at 12,000 g for 5 minutes at 4°C. Total protein concentrations in the supernatants were determined using the BCA Protein Assay Kit (Pierce, Rockford, IL, USA) with bovine serum albumin as the standard. To detect the luciferase activity, 20 μ L of supernatant was added in a 96-well plate and mixed with 100 μ L luciferase substrate (Promega), and the light emission was measured for 10 seconds in a multifunctional microplate reader (Varioskan Flash; Thermo Fisher). The gene transfection efficiency was represented by luciferase

activity, which was normalized to total protein concentration and calculated as relative light units per milligram of total protein.

The transfection efficiency was also quantitatively detected at cellular level. In brief, by using fluorescence-activated cell sorting (FACS; Calibur; Becton Dickinson, Franklin Lakes, NJ, USA), the proportion of EGFP-expressing cells in 1×10^4 cells was analyzed. Data were treated by CellQuest software (Becton Dickinson).

Cytotoxicity assay

Cytotoxicity was presented as the percentage of cell viability, which was evaluated by a cell counting kit-8 (CCK-8; Dojindo Laboratories, Kumamoto, Japan) method. In 300 μ L antibiotics-free medium, 5.0×10^3 or 1.0×10^4 cells/well were seeded in 24-well plates and cultured to 70%–80% confluence for PEI or ~90% for Lipo2000 at the time of transfection. The PEI/pDNA or Lipo2000/pDNA complexes containing 0.8 μ g pEGFP-C1 were added to corresponding wells according to the transfection procedures. To each well 30 μ L of CCK-8 was added and the plates were incubated at 37°C for an additional 2 hours. Absorbance values were measured at a wavelength of 490 nm and a reference wavelength of 630 nm using a multifunctional microplate reader (Varioskan Flash; Thermo Fisher). The cell viability was calculated according to the formula: cell viability (%) = $[(A_{\text{sample}} - A_{\text{blank}}) / (A_{\text{control}} - A_{\text{blank}})] \times 100\%$. Here, A_{sample} , A_{control} , and A_{blank} are the absorbance values of transfected cells, untransfected cells, and the medium, respectively.

DNA labeling and intracellular traffic analysis

The plasmid pGL3-control was labeled with TOTO[®]-3 iodide (Molecular Probes, Inc. Eugene, OR, USA) according to previously described protocol.²³ Cells were treated with PEI/TOTO[®]-3-pDNA polyplexes using the standard and modified methods for 4 hours and 1.5 hours, respectively. To observe the intracellular traffic of labeled pDNA, cells were washed with PBS twice and analyzed using confocal laser scanning microscopy (Leica TCP SP5) with the excitation wavelength at 642 nm and emission wavelength at 660 nm.

Particle size and zeta potential analysis

For the standard method, 4 μ g pDNA (pGL3-control) and 5 μ L PEI (1 mg/mL) or 4 μ g pDNA and 10 μ L Lipo2000 were diluted, respectively, in 250 μ L H₂O, mixed, incubated for 20 minutes at room temperature, and then diluted to 2 mL with H₂O. For the modified method, 6 μ g pDNA and

7.5 μ L PEI or 6 μ g pDNA and 15 μ L Lipo2000 were diluted, respectively, in 525 μ L H₂O, mixed, and incubated for 20 minutes at room temperature. One milliliter of the complexes was used for size and zeta potential measurements using a Malvern Zetasizer Nano ZS90 (Malvern Instruments Ltd, Malvern, UK) at 25°C.

Endocytosis inhibition assay

Hep G2 cells were pretreated with 5 mM m β CD, 1 mM amiloride hydrochloride, or 10 μ g/mL chlorpromazine hydrochloride for 1 hour at 37°C, respectively. The subsequent transfection of pEGFP-C1 was performed as described and cells were analyzed 48 hours later for EGFP expression by FACS.

Statistical analysis

Each experiment was performed in triplicate and repeated three times unless stated otherwise. Data were statistically analyzed using the software SPSS 11.5 and presented as mean \pm standard deviation. The value of $P < 0.05$ was considered as a significant difference, while $P < 0.01$ was highly significant.

Results

The influence of polyplex volume on transfection efficiency

The transfection efficiencies were qualitatively evaluated by observing the expression of EGFP in Hep G2 cells. In the transfection cases using the modified method, the efficiencies were higher than those of the PEI standard method when the volumes of PEI/pEGFP-C1 polyplexes were above 400 μ L (Figure 1A). The results were further quantitatively analyzed by FACS (Figure 1B). The efficiencies increased gradually in the cases from 300 μ L to 700 μ L of polyplexes and then decreased. At the volume of 700 μ L, the peak value was significantly higher than others and approximately 2.8-fold that of PEI standard transfection (Figure 1C). The results indicated that the modified transfection method significantly improved the gene transfection efficiency in a volume-dependent manner.

The influence of transfection procedure on transfection efficiency

For the modified transfection procedure, the optimal volume was 700 μ L in the six-well plate; for standard transfection, it was 2,500 μ L (500 μ L complex + 2,000 μ L medium). Therefore, we wondered whether the increase of transfection efficiency was due only to the decrease of complexation volume. To address this question, we compared the

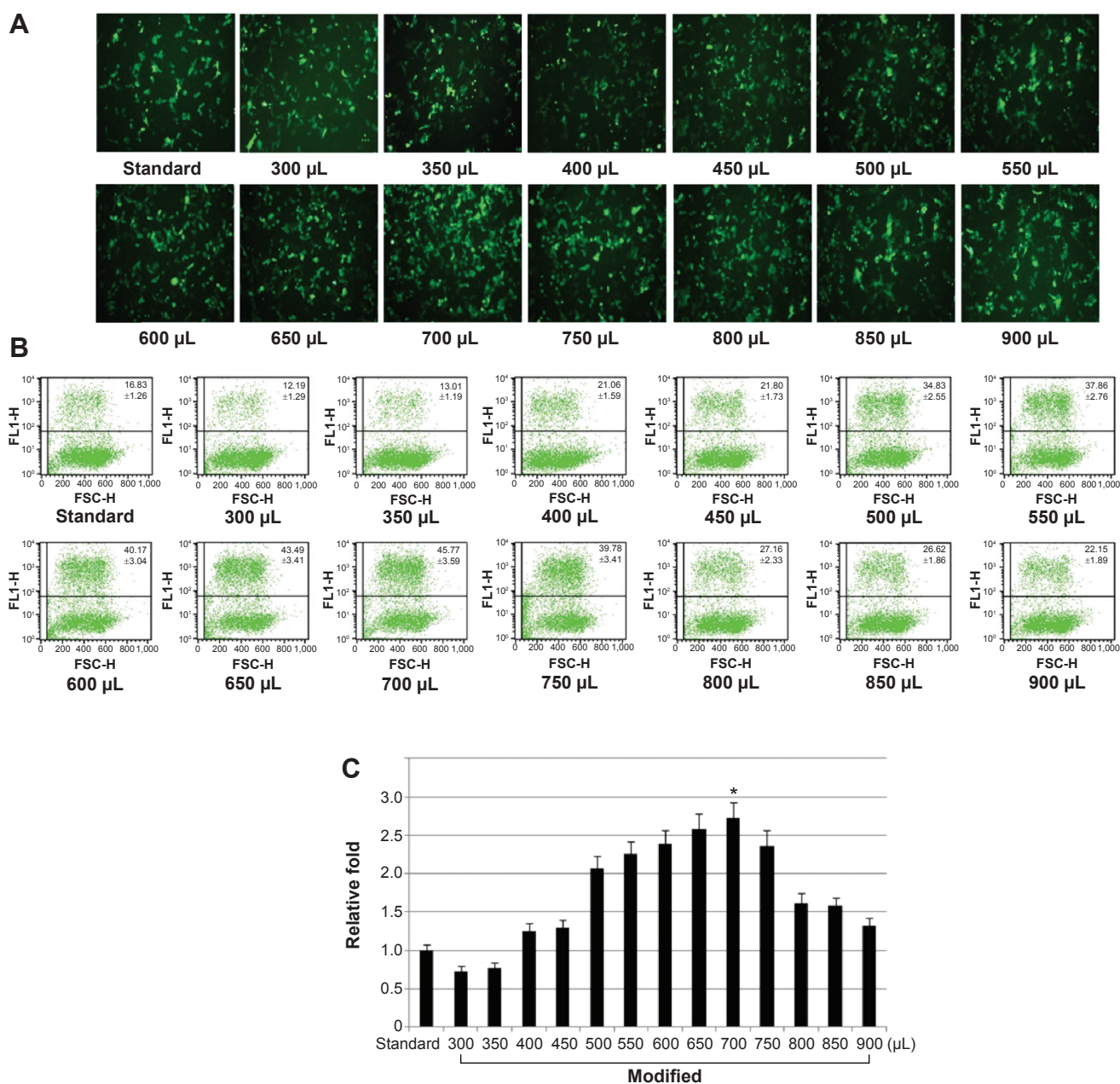


Figure 1 Qualitative and quantitative evaluation of the relationship between transfection efficiency and polyplexation volume. Cell strain: Hep G2. Plate: six-well format. Complex: polyethylenimine (PEI)/pEGFP-C1 polyplex. Standard: PEI standard transfection. 300–900 μ L: different volumes of the polyplexes in the modified transfection cases (complexes were added to cells without medium and incubated for 1.5 hours). Time: 48 hours after transfection. **(A)** Cells were evaluated with an inverted fluorescent microscope. Magnification: 100 \times . **(B)** Cells were analyzed by fluorescence-activated cell sorting. **(C)** Transfection efficiencies were analyzed by normalizing the values to those of the standard.

Note: * $P < 0.05$ versus others.

efficiencies of modified and traditional procedures at a fixed volume of 700 μ L using three different cell strains. In the modified transfection, the medium was discarded and 700 μ L polyplex was added to the cells; in traditional trials that mimic the standard procedure, the medium was exchanged with 200 μ L of fresh medium and then the 500 μ L polyplex was added in. The data showed that the modified method presented significantly higher transfection efficiencies than

those of the traditional procedure in Hep G2, NIH/3T3, and SMMC-7721 cell strains (Figure 2A; $P < 0.01$). To further investigate the results, we systematically compared the efficiencies of the two methods among a series of complexation volumes from 200 μ L to 500 μ L in a 24-well format using both Lipo2000/pEGFP-C1 lipoplexes (Figure 2B) and PEI/pEGFP-C1 polyplexes (Figure 2C). The results indicated that 1) at each volume, the modified method presented obviously

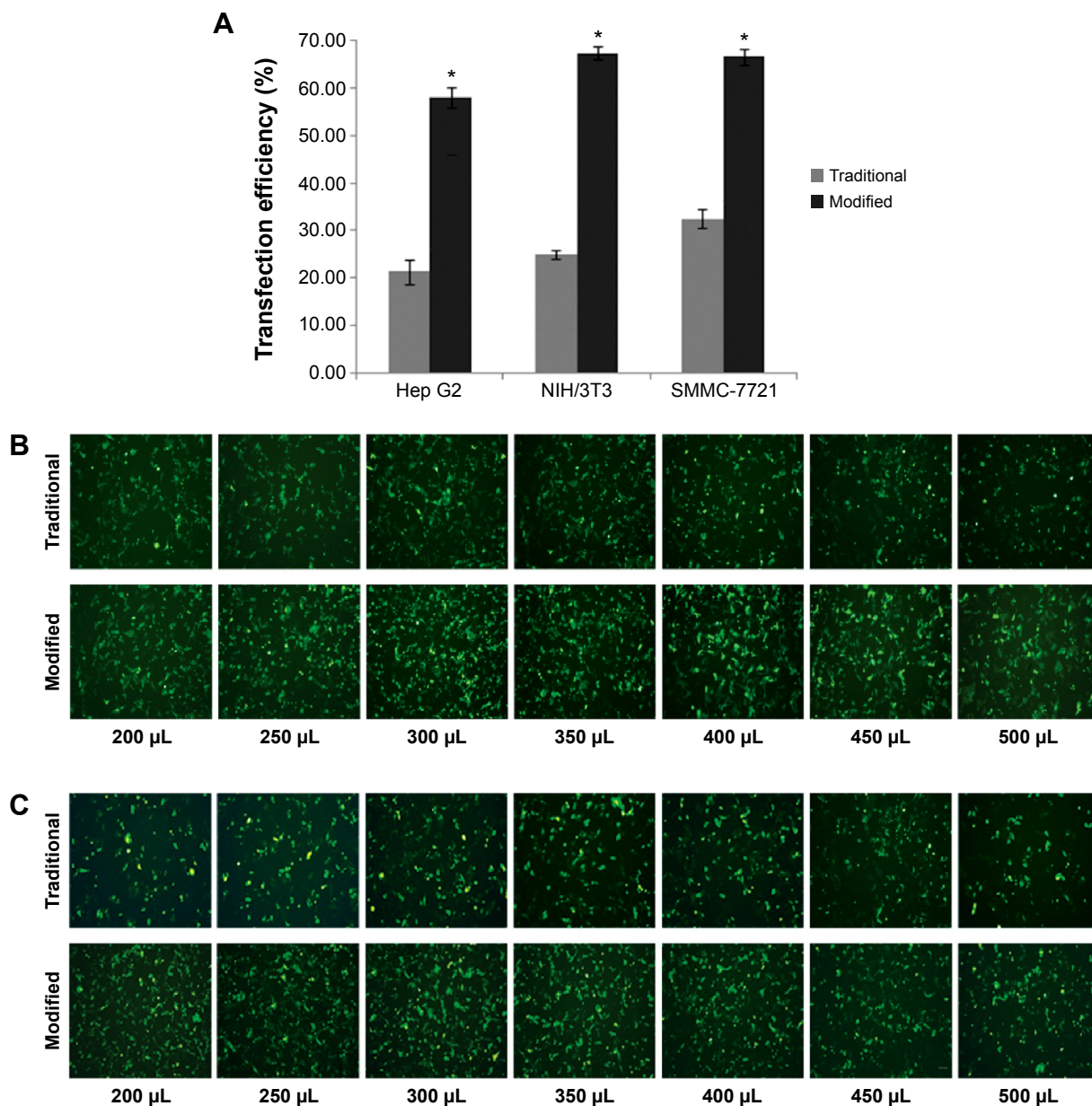


Figure 2 Systematic comparisons of the transfection efficiencies between modified and traditional procedures. (A) Quantitative analysis of the transfection efficiency in different cell strains. Plate: six-well format. Complex: polyethyleneimine (PEI)/pEGFP-C1. Modified: 700 µL of polyplex was added to cells without medium. Traditional: 500 µL polyplex was added to cells cultured in 200 µL fresh medium. Transfection efficiencies were analyzed by fluorescence-activated cell sorting 48 hours after transfection. (B) and (C) qualitative evaluation of the transfection efficiencies in different complexation volumes using Lipofectamine™ 2000/pEGFP-C1 lipoplexes (B) and PEI/pEGFP-C1 polyplexes (C). Cell line: Hep G2. Plate: 24-well format. Magnification: 100×. Time: 48 hours after transfection. Modified: the complexation complexes with indicated volumes were added to cells without medium and incubated for 1.5 hours. Traditional: 100 µL complex was added to cells with medium to get indicated final volumes from 200 µL to 500 µL.

Note: * $P < 0.01$ versus traditional.

higher transfection efficiency than the traditional method; 2) in the modified cases, the transfection efficiencies varied in a volume-dependent manner and 300 µL was the optimal volume; and 3) changing the complexation volume only had a slight influence on transfection efficiency in traditional cases. Altogether, the results confirmed that the improved efficiency was due to the combination of decreased complexation volume with transfection procedure. The best modified

transfection method was to add a suitable volume (700 µL for the six-well plate and 300 µL for the 24-well plate) of transfection complex to cells without medium.

Optimization of the incubation time of polyplex with cells

In the modified procedure, a shorter incubation time of polyplex with cells can also result in high transgene

expression level when compared with that in the standard procedure (Figure 3A). The peak values of transfection were acquired after 1.5–2.5 hours' incubation, which were significantly higher than those of others except in the case of 3.0 hours ($P < 0.05$) (Figure 3B). The results indicated that the modified procedure presented a faster transfection process and the suitable incubation time was reduced to 1.5–2.5 hours.

The influence of transfection methods on cell viability

In traditional transfection cases, a longer incubation time usually generates higher transfection efficiency but stronger cytotoxicity.²⁴ In the modified method, removal of cell medium followed with direct addition of transfection complex onto cells might also exert certain influence on cellular physiological

state. Therefore, a suitable incubation time should be identified to balance transfection efficiency and cytotoxicity. In both PEI- (Figure 4A) and Lipo2000-mediated (Figure 4B) transfection trials, the 0.5 hours' incubation had the minimum cytotoxicity, but its transfection efficiency was too low (Figure 3B). In comparison, the 1 hour and 1.5 hours of incubation showed similar cytotoxicity as that of the standard method (Figure 4A, B; $P > 0.05$). Considering that the 1.5 hours of incubation generated high transfection efficiency (Figure 3B), we selected it as the optimal parameter. It was further verified by transfecting other cell lines. Using the modified method with 1.5 hours' incubation time in both PEI- and Lipo2000-mediated transfection, Hep G2 and NIH/3T3 cells showed equal viability to those in the standard method; Hela cells in modified trials even had higher cell viability than those in the standard cases (Figure 4C).

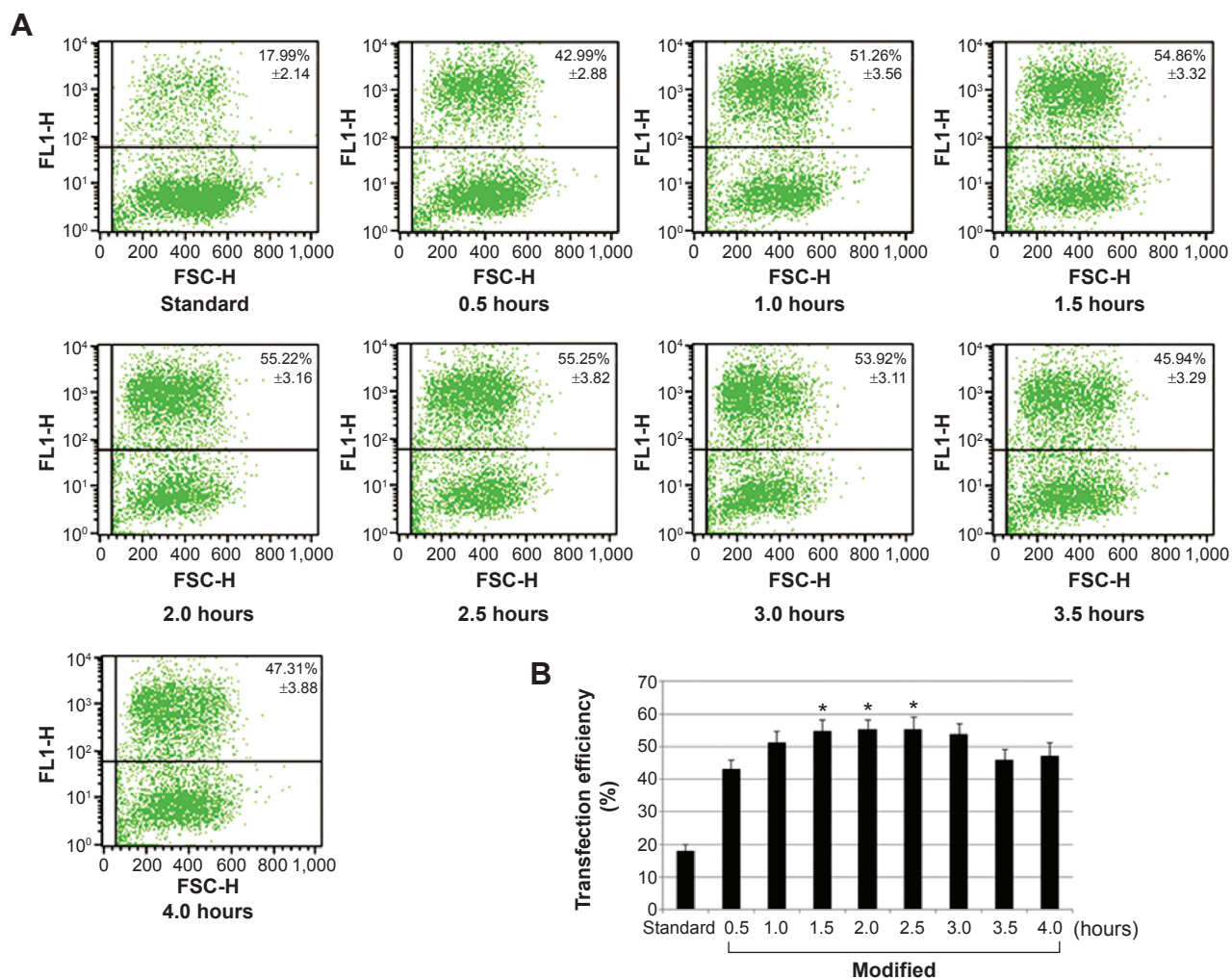


Figure 3 Optimization of the incubation time for transfection. Cell line: Hep G2. Plate: six-well format. Complex: polyethyleneimine (PEI)/pEGFP-C1. Standard: standard transfection. 0.5–4.0 hours: different incubation time in the modified transfection cases with 700 μ L transfection complex. (A) Transfection efficiency analyzed by fluorescence-activated cell sorting 48 hours after transfection. (B) Comparison of the transfection efficiency.

Note: * $P < 0.05$ versus others except 3.0 hours (1.5 hours vs 3.0 hours: $P > 0.05$).

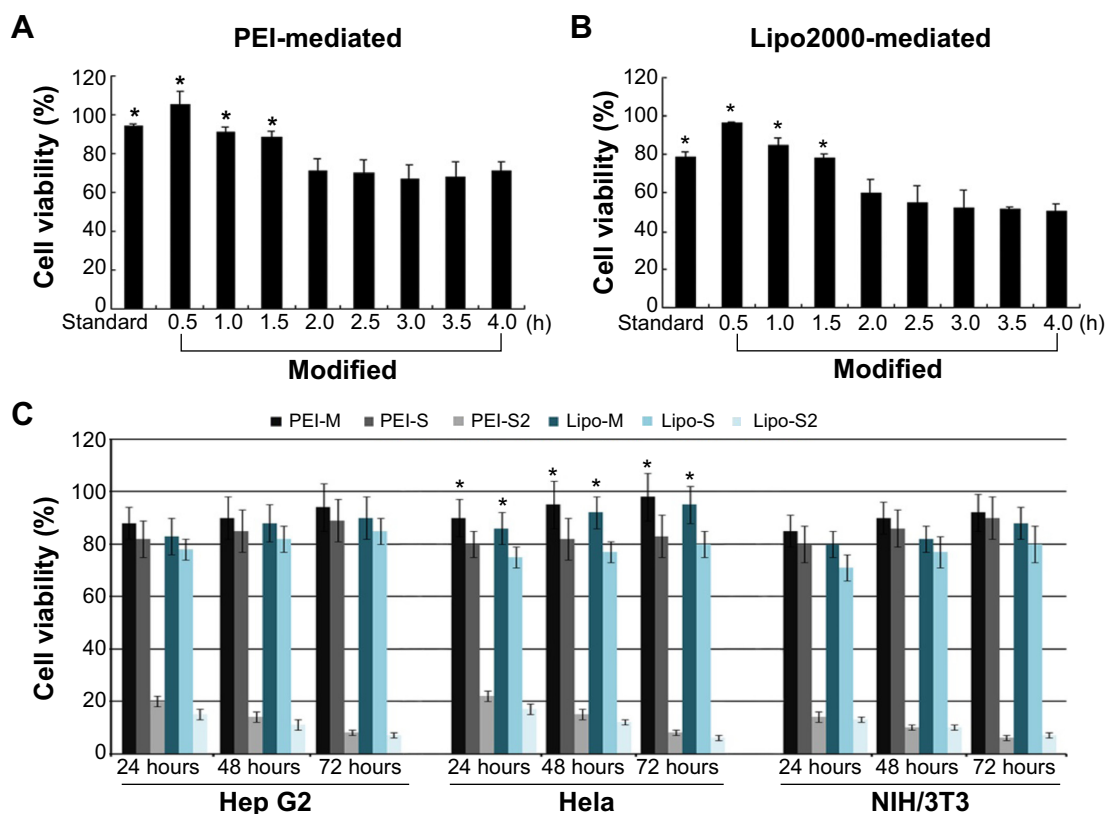


Figure 4 Cell viability assay in different transfection cases. (A) and (B) The influence of incubation time on cell viability in polyethyleneimine (PEI)- (A) and Lipofectamine™ 2000 (Lipo2000)-mediated (B) transfection cases. Cell line: Hep G2. Plate: 24-well format. Time: 48 hours after transfection. Standard: standard transfection. Modified: modified transfection (300 μ L/well transfection complex and 1.5 hours' incubation). 0.5–4.0 hours: different incubation time in modified transfection cases with 300 μ L/well transfection complex ($P < 0.05$ vs others). (C) Comparison of the cell viability in different transfection methods. PEI- and Lipo-M: PEI- and Lipo2000-modified transfection; PEI- and Lipo-S: PEI and Lipo2000 standard transfection; PEI- and Lipo-S2: PEI and Lipo2000 standard transfection with increased complex concentration. 24, 48, and 72 hours: cell viability was analyzed 24, 48, and 72 hours after transfection ($P < 0.05$ vs standard transfection).

The concentration of PEI/pDNA or Lipo2000/pDNA complex in the modified method was about 3.57 times that in the standard method. We wondered whether the standard transfection with the same complex concentration as that in modified method can also provide higher transfection efficiency. However, most of the cells died in these cases (Figure 4C; PEI-S2 and Lipo-S2). It demonstrated that the simple increase of complex concentration brought higher cytotoxicity instead of higher transfection efficiency.

Application of the modified method on the transfection of different cell strains

Based on the aforementioned outcomes, we concluded that the optimal procedure for the modified transfection method contained four steps (Table 1).

Further, we wanted to know whether the modified method can be widely used to transfect different cell strains. As confirmed by both FACS (Figure 5A and B) and luciferase activity assay (Figure 5C and D), for each cell strain, the modified method greatly improved the transfection efficiency mediated

by PEI (Figure 5A and C) and Lipo2000 (Figure 5B and D). The cell strains included different cancer cells (AGS, HeLa, Hep G2, SMMC-7721, MCF-7, and 4T1), normal cells (C2C12, L6, and NIH/3T3), embryonic stem cells (R1), and even primary skin fibroblasts (primary). The cell strains can also be divided into human (AGS, HeLa, Hep G2, MCF-7, and primary), rat (L6), and mice (C2C12, NIH/3T3, and R1) cells. The results indicated that the modified method was suitable, at least, for PEI and Lipo2000 in gene transfer into a wide range of cell strains.

Internalization course of PEI/pDNA polyplex

By monitoring the internalization course of PEI/pDNA polyplex, we hope to find some clues to explore the reason

Table 1 Optimal procedure for the modified transfection method

Removal of culture medium
Addition of the transfection complex (700 μ L for six-well and 300 μ L for 24-well format) to cells
Incubation of the complex with cells for 1.5 hours
Replacement of the transfection complex with fresh medium followed with additional culture for transgene expression

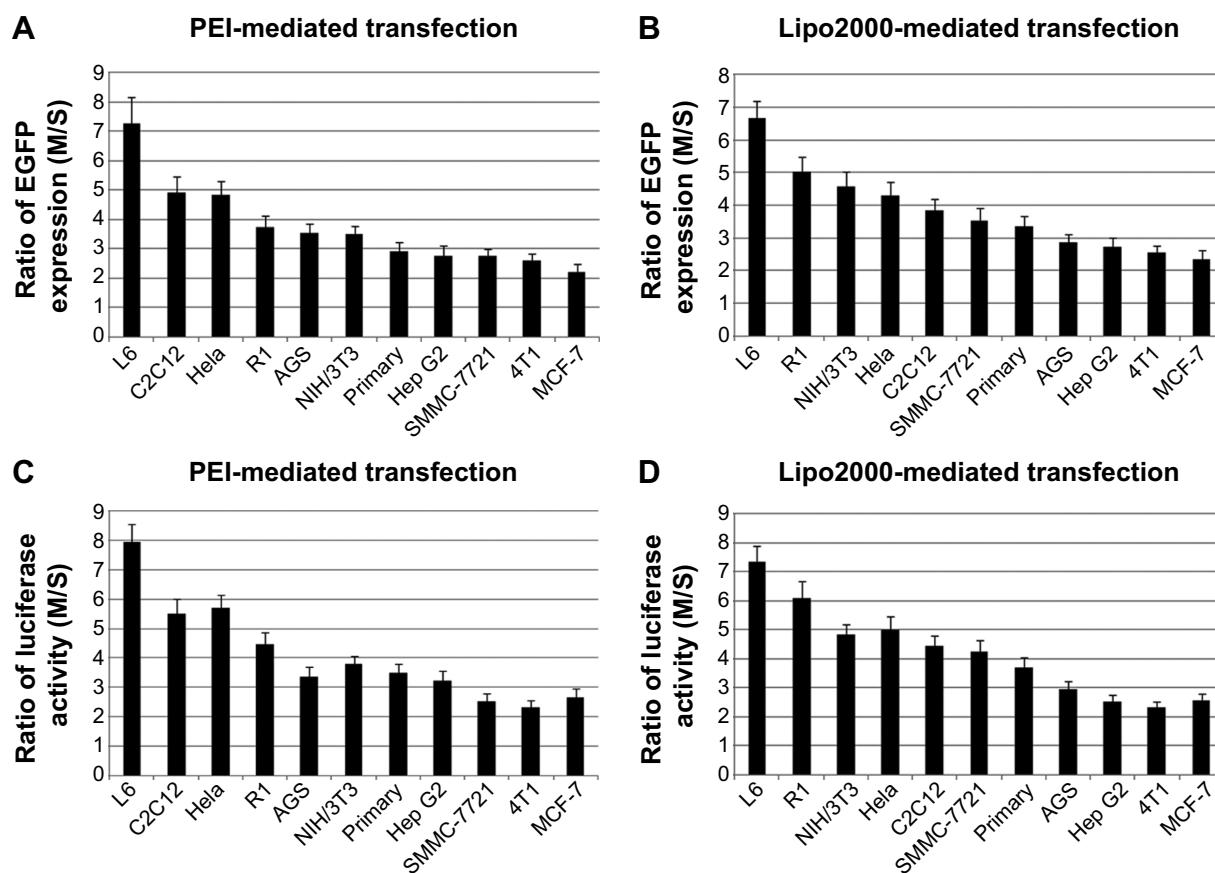


Figure 5 Comparison of the transfection efficiency between the standard and modified methods in different cell strains. The transfection efficiencies were measured by fluorescence-activated cell sorting (A) and (B) and luciferase activity assay (C) and (D), and presented as the relative folds by comparing the data between modified (M) and standard (S) methods. Plate: six-well format. Time: 48 hours after transfection.

Abbreviations: PEI, polyethyleneimine; Lipo2000, Lipofectamine™ 2000.

why the modified method had better performance on gene delivery. As shown in Figure 6, in the modified procedure, the polyplexes attached to cell membranes from 0.5 hour penetrated into the cells from 1.0 hour to 1.5 hours and dispersed within the cytoplasm/nucleus from 1.5 hours to 2.5 hours after transfection. In the standard procedure, the attachment/penetration was a slower kinetics process within 2.5 hours, and the dispersion of the polyplexes in the cells took place 2 hours after transfection. The results indicated that the modified method provided a more active kinetics process for the polyplex internalization than that in the standard method. Meanwhile, the proportion of fluorescent cells in the modified transfection was higher than that in the standard method. This may be the reason why the modified method provided higher transgene expression than the standard.

Endocytosis pathways in different transfection methods

The difference of internalization kinetics of the polyplex between the two methods may be related to their different

endocytosis pathways. So we investigated the cellular endocytosis by excluding specific pathways with endocytosis inhibitors. When Hep G2 cells were, respectively, treated with chlorpromazine, M β CD, and amiloride, the cellular transfection efficiencies mediated by PEI and Lipo2000 standard methods were all significantly decreased, indicating that the uptake of polyplexes and lipoplexes were, simultaneously, through clathrin-, caveolae-, and macropinocytosis-mediated endocytosis pathways (Figure 7). However, when macropinocytosis-mediated endocytosis in the PEI-modified method (Figure 7A) and clathrin-mediated endocytosis in the Lipo2000-modified method (Figure 7B) were inhibited, respectively, the gene transfection efficiencies were not decreased. The results revealed that macropinocytosis-mediated endocytosis in the PEI-modified method and clathrin-mediated endocytosis in the Lipo2000-modified method were excluded. We further found that for both PEI and Lipo2000 transfection, the particle size prepared in the modified method was much smaller than that in the standard method, while the zeta potentials were not changed (Table 2).

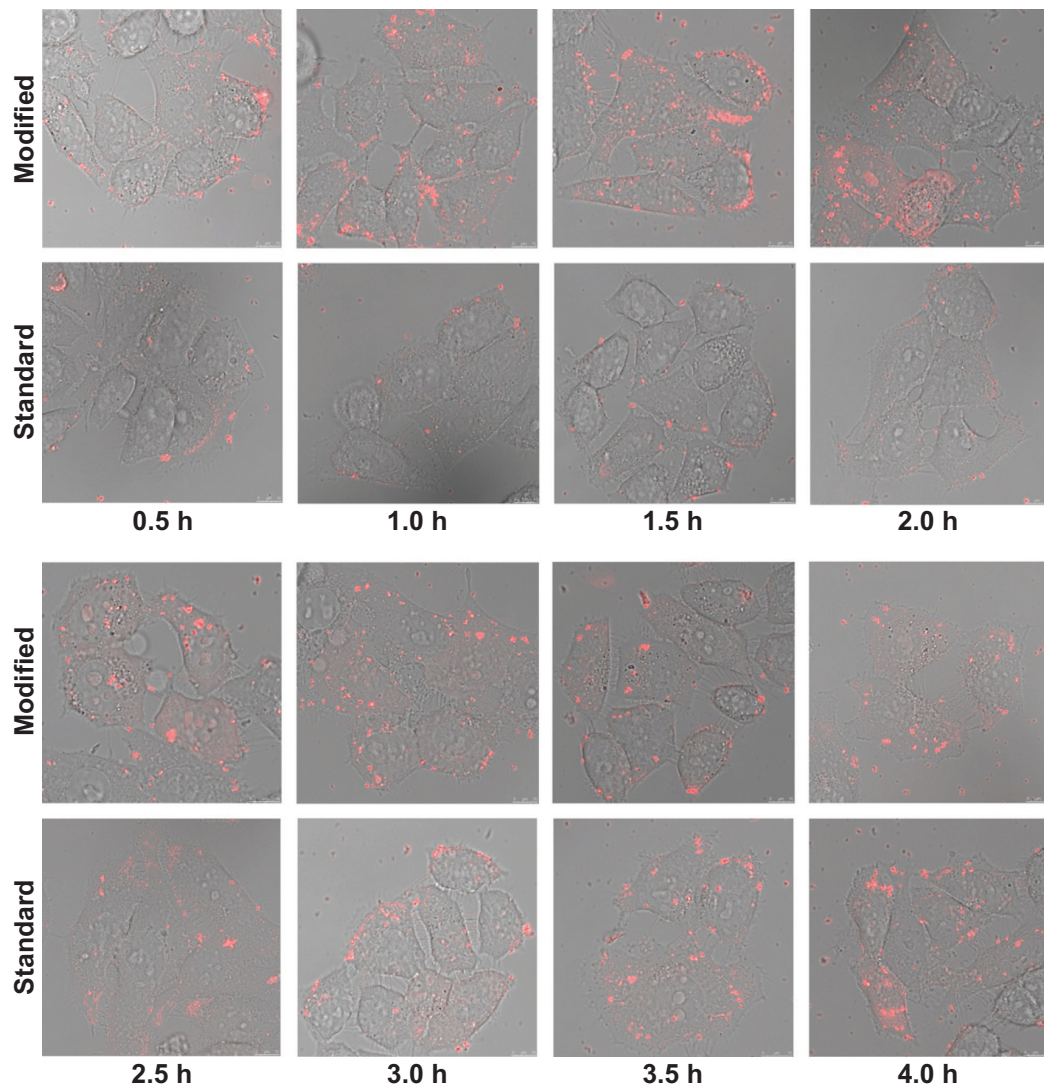


Figure 6 Time lapse images of the cellular uptake of polyplex. Hep G2 cells cultured in a 24-well plate were transfected with the polyplex containing 0.8 μg TOTO[®]-3-labeled pGL3-control and polyethyleneimine (N/P=10) using both methods. Time lapse images of the internalization of the polyplex were taken at every 0.5 hour using confocal laser scanning microscopy. Scale bar: 10 μm .

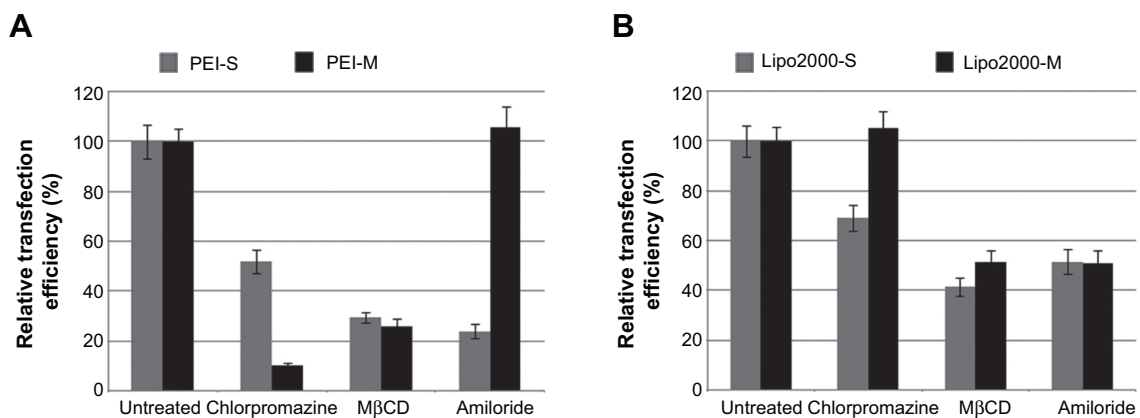


Figure 7 Analysis of relative transfection efficiency after the inhibition of specific pathways by endocytosis inhibitors. Cell line: Hep G2. Plate: six-well format. S: standard transfection. M: modified transfection. Cells were treated with inhibitors for 1 hour followed with pEGFP-C1 transfection. Proportion of enhanced green fluorescent protein-expressing cells in each group was analyzed by fluorescence-activated cell sorting 48 hours later and the data were normalized to those of the untreated group. **Abbreviations:** PEI, polyethyleneimine; Lipo2000, Lipofectamine™ 2000; MβCD, methyl-β-cyclodextrin.

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Discussion

For nonviral gene delivery material, the optimal transfection procedure represented the balance between high efficiency and low cytotoxicity, which was commonly acquired by optimizing a lot of parameters.¹⁰ For this reason, the commercially available gene carriers such as PEI, cationic lipids, and polymers usually have their specific standard protocols, which have been considered as the optimal procedures in gene delivery cases. However, the standard procedure can also be empirically improved in some cases, and compared with the complicated and time-consuming modifications on gene delivery carriers, optimization of transfection method sometimes seems to be more simple and efficient. In a reported case, the authors showed an optimized protocol for more efficient transfection of adherent primary mammalian cells using 25 kDa branched PEI.¹² In this study, the modified method was applicable for, at least, PEI and liposome to deliver genes into a wider range of cell lines.

The determinant point in the modified method is the combination of transfection procedure with optimized complexation volume. The complexation volume can greatly affect the transfection efficiency: the smaller the volume, the higher the efficiency. Although the smaller volume possesses a higher complex concentration, a simple increase of complex concentration only brought higher cytotoxicity instead of higher transfection efficiency (Figure 4C; PEI-S2 and Lipo-S2). Therefore, the decrease of complexation volume should be combined with the new transfection procedure in which the DNA/material complex was added to the cells immediately after the removal of cell culture medium. In fact, when the volume was reduced to 300 μ L, cells located at the margin of the well had the highest transfection efficiency, but cells in the middle of the well were rarely transfected because this area was only slightly covered by the transfection mixture due to the surface tension (data not shown). So the global efficiency in the 300 μ L system was less than that in the 700 μ L system (Figure 1), because cells in the latter were well covered by the transfection mixture. If the transfection procedure was similar to the standard procedure, the reduction of complexation volume only had a very limited increase on the transfection efficiency. More interestingly,

in the equal volume, the modified procedure had higher transfection efficiency than that of the standard procedure (Figure 2). The results implied that the transfection procedure had a more important role than complexation volume in gene delivery.

The additional superiority of the modified method was the faster transfection process (Figure 3). Usually, shorter incubation time results in lower cytotoxicity, but higher transfection efficiency needs longer incubation time.²⁵ To resolve this paradox, an optimal incubation time should be experimentally determined to balance transfection efficiency and cytotoxicity. If the biocompatibility of a material was not good enough but its gene transfection ability was excellent, an alternative solution was to reduce the incubation time to control its cytotoxicity using the modified method. In addition, it is well known that some cell lines, such as stem cells and primarily cultured cells, are difficult to transfect and sensitive to toxic reagent.²⁶ The modified method offered significantly increased efficiency for all the tested cell strains (Figure 5).

There are multiple pathways for cellular uptake of particles and solutes.^{27,28} The endocytosis pathways of DNA/material particles were commonly classified as clathrin-dependent, caveolae-mediated, caveolae- and clathrin-independent, and macropinocytosis.^{29–31} Endocytosis inhibitors were usually applied to analyze the endocytosis pathways of cells, but their concentrations and treatment time should be well determined to minimize cytotoxicity.^{32,33} Meanwhile, inhibition of certain endocytosis pathways may artificially upregulate other internalization routes that are not originally involved in the uptake of transfection particles because many chemical inhibitors are not specific to a single internalization pathway.²⁷ Therefore, inhibition experiments should be rationally designed to observe effects at a suitable time in referable parallel treatments.³⁴ In the standard transfection, both PEI- and Lipo2000-mediated endocytosis involved clathrin-dependent, caveolae-mediated, and macropinocytosis pathways. In the modified transfection, however, the endocytosis pathways were changed: clathrin-dependent and caveolae-mediated for PEI while caveolae-mediated and macropinocytosis for Lipo2000 (Figure 7). We think that the regulation is mainly due to the change of transfection procedure. When the transfection mixture was added, cells suffered a sudden stimulation. In the modified transfection, the culture medium was already removed and the concentration of complexes was increased, making the stimulation so great that the cellular endocytosis pathways were changed. In the standard or traditional transfection, the stimulation was buffered by the culture medium. The decreased size and increased concentration of pDNA/

Table 2 Size and zeta potential of lipoplexes and polyplexes

	Size (nm)		Zeta potential (mV)	
Lipo-S	314.47±6.82	<i>P</i> <0.01	17.87±0.85	<i>P</i> >0.05
Lipo-M	146.7±1.31		18.7±0.6	
PEI-S	88.19±1.3	<i>P</i> <0.05	22.88±1.99	<i>P</i> >0.05
PEI-M	65.09±2.65		23.43±3.26	

Abbreviations: Lipo, Lipofectamine™ 2000; S, standard transfection; M, modified transfection; PEI, polyethyleneimine.

material particles in the modified method may also influence the endocytosis pathway and kinetics of internalization (Figure 6). In general, complexes may undergo physicochemical changes when mixed at different concentrations, and the aggregation of particles may also be influenced by the different ionic strength in the medium.^{12,35} The complexation of pDNA/material particle was formed in 700 μ L solution for the modified transfection while in 500 μ L and then diluted to 2,500 μ L system for standard transfection, leading to a different particle concentration and size. Altogether, the change of cellular endocytosis pathway, decrease of particle size, and increase of particle concentration made the modified transfection method a potent tool for nonviral material-mediated gene delivery into a wide range of cell strains.

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Disclosure

The authors declare no conflict of interest.

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