Preparation of ultrasound microbubbles crosslinked to albumin nanoparticles packaged with tissue-type plasminogen activator gene plasmid and method of in vivo transfection

Ji Jun
Ji Shang-Yi
He Xia
Ling Wen-Ping

Department of Pathology, Shenzhen Sun Yat-Sen Cardiovascular Hospital, Shenzhen, GuangDong Province, People’s Republic of China

Aims: To observe the effect of constructed ultrasound microbubble crosslinked to albumin nanoparticles packaged with tissue-type plasminogen activator (tPA) gene plasmid on the in vivo transfection.

Methods: The rabbits were chosen for all experiments. A highly expressive gene plasmid for tPA was constructed and packaged into a prepared nanoparticle with bovine serum albumin (BSA). This albumin nanoparticle packaged with tPA gene plasmid was crosslinked to an ultrasound microbubble prepared with BSA and sucrose to form a nano-targeting vector system for tPA gene transfection. The transfection and effective expression of tPA in heart, liver, leg skeletal muscle and the cervical rib were detected with polyclonal antibodies to tPA using immunohistochemical method; the tPA level and D-dimer content of blood were also tested.

Results: The expression of tPA could be seen in the tissues mentioned above, with the increase in blood tPA level and D-dimer content from 0.20 ± 0.05 µg/L and 81.76 ± 9.84 µg/L before the operation, to the higher levels of 0.44 ± 0.05 µg/L and 669.28 ± 97.74 µg/L after transfection.

Conclusion: The nano-targeting vector system for tPA gene was constructed successfully. This provides a new theory and experimental method for the nano-targeted transgene.

Keywords: tissue-type plasminogen activator, albumin nanoparticle, ultrasound microbubble, targeting transfection

Introduction
Long-term or lifelong anticoagulant therapy is involved with transluminal balloon angioplasty and stenting, coronary bypass, and heart valve replacement, which may induce bleeding complications. Tissue-type plasminogen activator (tPA) is the main activator of the fibrinolytic system and is synthesized by the vessel endothelial cells and released into the blood circulation under the stimulation of fibrin. The function of tPA is to make the plasminogen transfer to plasmin, which promotes fibrinolysis. Because the cDNA of tPA has been cloned successfully, tPA reverse-transcript virus vector can be constructed and transfected in vitro to the epithelial cells with a high expression of tPA protein. It has been demonstrated experimentally that thrombosis and restenosis after coronary stenting or bypass can be prevented with a long-term outcome when transfected epithelial cells are spread on the surface of stents or cover vascular anastomosis.
Human recombinant tPa has also been used clinically in thrombolytic treatments for acute myocardial infarction, brain embolism, and pulmonary infarction with very good results, but it is expensive for patients. We aim to construct the tPa gene plasmid and to transf ect it to the cells of a tissue or an organ, followed by long-term anticoagulation and thrombolysis. In our early study, we constructed a tPa gene plasmid and transf ected it to pig cardiocytes using the surgery dacron suture as a gene vector, and successfully prevented thrombosis after the valve replacement and anastomotic restenosis after coronary bypass. Although the surgery dacron suture-carrying gene has been proved to be safe, convenient, and effective, it is traumatic and unfavorable for patients when used as a drug carrier.

Using polymer-constructed nanoparticles as a carrier for gene transf ection is a new method that has been used for the past few years. Albumin has been used as a material for nanoparticles because of its high biocompatibility and biodegradability with no immunogenicity or cellular toxicity. With a positive charge on their surface when pH value in the reactive system is adjusted to acidity, the albumin nanoparticles can not only absorb the gene DNA with a negative charge by a form of static electricity but also carry them in a form of package. The main method for in vivo transf ection is to inject an objective gene, such as a plasmid DNA or an adenovirus vector, directly to the targeted tissue or cells. This method is difficult clinically because of traumatic occlusion.

Ultrasound contrast agents have been used in diagnostic ultrasound imaging in the past few years. It has been demonstrated that ultrasound microbubbles can take the drugs or genes to the targeted tissue and cells for the purpose of treatment or transfection safely and effectively. The fundamental principles of ultrasound for targeted treatment are: i) cavitation and machinery effects produced by the therapeutic ultrasound, which is another kind of ultrasound heating, injure the cell membrane slightly and are reversibly followed by an increase in its permeability; ii) the capillary vessels (≤7 mm) in the ultrasound field are injured and the interspaces between endothelial cells become wide, through which drugs or specific genes can get to the targeted tissue and cells; and iii) when the ultrasound microbubbles in the blood circulation pass through the tissue or organs treated with the ultrasound, they are quickly destroyed by a force produced in the ultrasound field, and the drugs or genes carried are released to the tissue or organs for the purpose of treatment or transfection. The systemic toxicity and side effects produced by the drugs and genes significantly decrease because of targeted localization.

In this study, a highly expressive tPa gene plasmid was constructed and a bovine serum albumin (BSA) nanoultrasound microbubble targeting vector for the tPa gene was made. Rabbit heart, liver, and skeleton muscle in the hind leg and rib were chosen as the targeted tissue for the gene transf ection.

**Materials and methods**

**Animals**

Twenty-five healthy male New Zealand rabbits, 2.0–2.5 kilogram in weight, were used in an experimental study and provided from Southern Medical University Animal Center, GuangZhou, GuangDong Province, China.

**Main reagents and instruments**

Chinese hamster ovary (CHO) cell lines, pSecTag2B plasmid, E. Coli JM109, rabbit antihuman tPa antibody, FITC-coupled sheep anti rabbit immunoglobulin G antibody, rabbit antihuman polyclonal antibody, immunohistochemical reagents, and BSA were purchased from JingMei Biotech, ShenZhen, GuangDong Province, China. Restriction enzymes HindIII, KpnI, BamHI and XhoI, Vent DNA polymerase, T4 DNA ligase, QIA prep spin miniprep kit, QIA quick gel extraction kit, QIAGEN polymerase chain reaction (PCR) product purification kit, and DNA marker DL 2000 were purchased from New England Biolabs, Hong Kong, China. Perfluoropropane (Halocarbon-218) was supplied from JieRui Co. Ltd, FuShan, GuangDong Province, China. The diagnostic ultrasonic generator is a product of NingBo Scientz Biotech Co. Ltd. The therapeutic ultrasound unit (US-700) was made by ITO Co. Ltd, Tokyo, Japan.

**Construction and expression of the pSecTag2B tPa gene**

Three EST sequences were obtained from Internet Blast according to the tPa gene sequence. The ID numbers were 6251209, 4861268, and 5190656, respectively. The primers were synthesized as follows: tPa-1F: 5′-CCC AAG CTT ATG GAT GCA ATG AAG AGA GGG-3′, tPa-1R: 5′-GGG GTA CCA CGG TAG TCT GAC CCA TTC-3′, tPa-2F: 5′-GGG GTA CCC ACA GCC TCA CCG AGT CG-3′, tPa-2R: 5′-CGG GAT CCA GCA GGA GCT GAG TAT GCC-3′, tPa-3F: 5′-CGG GAT CCT CTC TGC CGC CCA CTG CT-3′, and tPa-3R: 5′-CCC TCG AGG CGG TCG CAT GTT GTC AC-3′. As the PCR amplification template,
three EST clone strains were abstracted and the three tPa fragments amplified. The pSecTag 2B and three tPa fragments tPa-1, tPa-2, and tPa-3 were digested by HindIII and Xhol, HindIII and KpnI, KpnI and BamHI, and BamHI and Xhol, respectively. These enzymatic products were purified with QIAGEN PCR product purification kit and were linked by T4 DNA ligase at 14°C overnight. The linked products were transfected to E. Coli JM109, and the resistance colony in the aminobenzyl penicillin LB plate culture was chosen. This tPa plasmid was sequenced and transfected to CHO cells by calcium phosphate coprecipitation. The expression of tPa was detected using a rabbit antihuman tPa antibody by indirect immunofluorescence method.

Preparation of the BSA nanoparticles loaded with tPa gene plasmid

The preparation of BSA nanoparticles loaded with tPa gene plasmid was carried out according to the methods published by Arnedo et al10 and Zhang et al18 with some improvements. Briefly, 2 mg tPa plasmid DNA was firstly incubated with 10 mL albumin aqueous solution (1% w/v; pH 5.5). Then, the aqueous phase was desolvated with ethanol dropwise (ethanol:water = 2:1). The coacervates were hardened with 30 µL glutaraldehyde (concentration: 0.5%, w/v) for 2 hours. After ethanol was eliminated by evaporation, the nanoparticles were purified by centrifugation at 17,000 rpm for 30 minutes to eliminate free albumin and excess crosslinking agent. The purified nanoparticles by centrifugation were resuspended in pure water and dispersed with ultrasound generator (180 W, 20 kHz, for 30 sec) and stored at 4°C for further use. Some of the nanoparticles were taken for determination of particle size, morphological observation, surface Zeta electric potential, envelopment rate, and electrophoresis for gel retardation.

Preparation of the ultrasonic microbubbles

The preparation of albumin ultrasonic microbubbles was carried out according to the method reported by Du et al9 with some improvements. Ten milliliters of BSA (5%, w/v) with sucrose (final concentration: 10%, w/v) was prepared in 50 mL plastic centrifuge tubes and saturated with oxygen and perfluoropropane (flow rate: 6 mL/min) by turning for 10 minutes and dispersed with ultrasonic generator (180 W, 20 kHz, for 1 min). All procedures were carried out under sterile conditions. The prepared microbubbles were stored at 4°C for further use.

The linkage of the nanoparticles to the microbubbles

The nanoparticles packaged with tPa gene plasmid (containing 1 mg plasmid DNA) were mixed with 5 mL microbubbles (containing 1.2 × 10⁴/mL) at room temperature. Ten microliters of 50% glutaraldehyde (final concentration: 0.1%, w/v) was added to the mixture prepared as stated previously for incubation for 2 hours at 4°C. This linked product was centrifuged at 200 rpm for 1 minute, the floatage was taken and washed with 0.9% sodium chloride under the centrifugation at 200 rpm three times, and the supernatant was stored at 4°C for ultrasound targeted transfection.

Targeted transfection

New Zealand rabbits were used in all experiments. All surgical procedures were carried out on the animals under general anesthesia (25 mg/kg sodium pentobarbital). The heart and liver were observed with two-dimensional diagnostic ultrasound. Soon after injecting 5 mL, the linked ultrasonic microbubbles from the rabbit ear margin vein and a strong resonance of the ultrasound signal were clearly seen. All experimental animals were randomly divided into two groups, the experimental group and the control group. The experimental group (n = 1) was divided into three subgroups, heart (n = 6), liver (n = 6), and skeleton (n = 4). Those organs and the tissue were treated with therapeutic ultrasound (1 MHz, 1.5 w/cm², 30 min, according to Ling et al20 with some improvements) after injection of the ultrasound microbubbles. The control groups (n = 9) were also divided into three subgroups, signal microbubbles (n = 3) (no ultrasound treatment was operated after the microbubble injection), nanogene (n = 3) (the nanoparticles with tPa gene plasmid without the microbubble linkage were injected into the animals with the treatment of the ultrasound), and blank control (n = 3) (only physiologic saline injection and ultrasound treatment). The whole experiment was observed for 4 weeks after the operation.

Pathology and immunohistochemical stain

All animals were sacrificed after the observation finished. The hearts, livers, targeted muscles, and ribs with ultrasound exposure were taken for pathology and immunohistochemical staining in which a polyclonal antibody to tPa (dilution: 1:100) was used with the indirect immunohistochemical method for tPa targeted expression. Lungs and kidneys were taken for study on targeting specificity and systemic toxicity of the transfected tPa gene. The blood was taken for D-dimer with the immunoturbidimetric method and tPa
concentration with the enzyme-linked immunosorbent assay method following the manufacturer’s instruction.

Statistical analysis
The results are expressed in x ± s. The Student’s t-test was used for statistics, with a statistically significant difference of P < 0.05.

Results
Construction of pSecTag2B-tPa and gene expression
The results of the indirect immunofluorescence in the transferred CHO cells are shown in Figure 1. The strong positive reaction of the fluorescence in the gene-transferred CHO cell is present (Figure 1: a1). It is negative in the control group (Figure 1: a2).

Analytic results for albumin nano-tPa gene plasmid
Particle size analysis showed that the grain size ranged from 49.10 nm to 152.40 nm and was 132 nm, on average, with very even distribution. The polydispersity was 0.33. The surface Zeta potential ranged from +31.32 to +41.42 mV. The enveloping rate was calculated as 73.58% according to the following formula: (Wg − Wf)/Wg × 100%, where Wg is the initial plasmid DNA added, and Wf is the amount of plasmid DNA determined in the supernatants obtained during the purification steps with analysis by ultraviolet spectrophotometer. All of the analytic results met the needs of the experimental study.

Gel retardation and DNase I protection
Figure 1b shows the result of 0.9% agarose gel electrophoresis, from which we can see that the plasmid DNA packaged by the nanoparticles with or without DNase I digestion could not move in the electric field and was retarded in the initial well (DNA:albumin was 1:100). The plasmid DNA nonpackaged with the nanoparticles was digested by the DNase I, and the moving strap in the gel electrophoresis could not be seen, indicating that the nanoparticles could protect the plasmid DNA from DNase I digestion.

Morphological and physical features of ultrasound microbubbles
The size of the prepared microbubbles was 2–5 micrometer in diameter and cystic with very good dispersity, which could fit the vascular ultrasound visualization (Figure 1c). The 10% sucrose–albumin linked to the nanoparticles was heat resistant (40°C, 30 min) and could be stored at 4°C for 30 days without morphological change.

Ultrasound imaging and targeted transfection
Figure 2 shows the changes to the ultrasound images before and after the ultrasound treatment. After venous injection of the microbubbles with tPa gene plasmid, the ultrasound visualization of the heart (Figure 2: a1, a2, a3) and liver (Figure 2: b1, b2, b3) obviously increased compared with preinjection and decreased to the normal image following the ultrasound treatment. There was no change in ultrasound visualization in the nanogene and blank control groups.

Histopathology and tPa gene expression
Routine pathology showed that myocardial cells, liver cells, and skeletal muscle cells exposed to the ultrasound treatment were larger than those normal cells with the rich cytoplasm. Immunohistochemical analysis was performed on all tissue samples. TPA-positive cells were scattered in the tissue with exposure to the ultrasound treatment. These positive cells included myocardial cells (Figure 3: a1, a2), skeletal muscle cells (Figure 3: b), and liver cells (Figure 3: c). In the liver, these positive cells were mainly distributed near the portal area with a radiating arrangement. In skeleton and cartilage tissue, some chondrocytes in the germinal layer (Figure 3: d), myeloid cells, and some interstitial cells were tPA positive (Figure 3: e).
Table 1 shows the changes in the blood content of D-dimer and tPA before and after the targeted transgene. We could see that the blood contents of D-dimer and tPA increased significantly 4 weeks after the targeted transgene, which indicates a higher fibrinolysis.

**Discussion**

We have demonstrated in several experimental studies that the constructed tPA plasmid used in the present study carried by surgery dacron suture or chitosan nanospheres could be transfected in vivo to cardio-ocytes and skeleton muscle cells with a high expression of tPA protein. But all these methods were traumatic and might be suitable only for experimental or surgical use. Albumin chosen as nanomaterial not only has some benefits as mentioned previously but also was easy to link to the protein ultrasound microbubbles for the purpose of targeting. There are two different methods for nanoparticles to carry genes. One of them is the single-step method, in which a gene is added to the albumin solution before the nanoparticles are prepared, and the gene DNA is encapsulated in the matrix of the nanoparticles. This method has some advantages because of the higher concentration of the drug, delayed and controllable release, enzyme resistance, and a longer time to transfect the gene. Another method is the two-step method, in which the albumin nanoparticles are prepared before the gene is added, and the pH value of the solution is adjusted to be acidic, which makes a more positive charge on the surface of the nanoparticles and links easily to the gene DNA with the negative charge. The nanoparticles made this way are equal in size, and the particle diameter is changed easily but without the advantages mentioned previously. In this study, the one-step method was carried out to prepare the albumin nanoparticles packaged with the tPA gene plasmid, which met the needs of the study because of its suitable size (132.0 nm in diameter), good dispersity, high encapsulation rate (73.58%), and effective DNase I protection according to the results of gel electrophoresis.

Determining how to get the albumin nano-tPA gene to the targeted tissue after being injected into the blood is the main purpose in this study. Albumin can be crosslinked to another protein by glutaraldehyde. Thus, when albumin nanoparticles
Some researchers found that the prepared microbubbles are crosslinked to the fragment of an antibody, they can connect to the specific ligand on the surface of the cells in a tissue for the purpose of targeting. Albunin has been used as a material for preparation of ultrasonic microbubbles because of its innocuousness and easy preparation. We prepared the albumin ultrasound microbubbles and linked them to the constructed albumin nano-tPA plasmid successfully using glutaraldehyde at a final concentration of 0.1% (w/v) because the microbubbles could be denaturated and destroyed easily at a higher concentration (1%, w/v) of glutaraldehyde and could not be crosslinked with the nanoparticles in a lower concentration (0.05%, w/v) of glutaraldehyde.

Table 1 Blood content of D-dimer and tissue-type plasminogen activator (tPA) before and after the targeting transgene (µg/L, x ± s)

<table>
<thead>
<tr>
<th>Group</th>
<th>N</th>
<th>D-dimer</th>
<th>tPA</th>
</tr>
</thead>
<tbody>
<tr>
<td>Before transgene</td>
<td>10</td>
<td>81.76 ± 9.84</td>
<td>0.20 ± 0.05</td>
</tr>
<tr>
<td>After transgene</td>
<td>10</td>
<td>669.28 ± 97.74</td>
<td>0.44 ± 0.05*</td>
</tr>
</tbody>
</table>

Notes: *P<0.05, **P<0.01.
such as effectiveness, stability, and manipulability, are not yet solved.

**Acknowledgment**

This work was supported by National High Technology Research and Development Program 863 (Project: No. 2007AA021809, No. 2007AA021803, No. 2007AA021904).

**Disclosure**

The authors report no conflicts of interest in this work.

**References**