Preventive Effect of Albumin Nano TPA Gene Plasmid Ultrasound Microbubble Carrier System on Thrombosis after Cardiac Valve Replacement

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After cardiac valve replacement, most patients will have different degrees of thrombosis, different parts of the thrombus, and even more frequent occurrence of postoperative thrombosis; therefore, the prevention of postoperative thrombosis is particularly important. The purpose of this study was to investigate the preventive effect of albumin nano tPA gene plasmid ultrasound microbubble carrier system on thrombosis, especially in cardiac valve. The experimental control method was used. Firstly, 11 dogs meeting the experimental requirements were selected. Secondly, the data of albumin nanoparticles and microbubbles were analyzed. The average size of albumin particles was 132.0 nm, the average size of microbubbles was 3.1 ± 1.6 μm, and the zeta potential was 13.70 ± 1.95 MV. The concentration of microbubbles was 4.2 ± 1.3 × 10^11/ml. Finally, 11 dogs were divided into two groups. The experimental group was treated with albumin nanoparticles ultrasound microbubble, and the others were the control group. It was found that the levels of tPA and D-dimer in the experimental group were significantly increased and maintained at a high level at 1 week after operation, and the prothrombin time was detected and the international normalized ratio was calculated at the same time. No significant changes were found in the experimental group.

1. Introduction

Valvular heart disease is a kind of serious heart disease, which could lead to heart failure, dyspnea, and edema; valve replacement should be carried out in time. Congenital heart valve disease has life-threatening consequences, which requires early valve replacement. In addition, after cardiac valve replacement, hemorheology changes abnormally and produces emboli, which leads to the possibility of cerebral embolism, reaching 50%. Albumin nanoparticles were initially used only as diagnostic reagents. Now, albumin nanoparticles are a relatively mature drug delivery system, and related drugs and diagnostic agents have been put on the market [1, 2]. Albumin nanoparticles can be administered through a variety of ways, such as intravenous injection, intramuscular injection, oral administration, and intra-arterial injection [3, 4]. Tissue-type plasminogen activator (tPA) is a tissue plasminogen activator that selectively activates plasmin [5, 6]. However, the way in which plasmin activator gene can be effectively transferred into terminal cells has not been solved. Therefore, whether the plasma tissue-type activator can be effectively expressed for a long time and play a local anticoagulant effect is still a key problem to be solved [7, 8]. Oschatz pointed out in the study that clinical transcatheter heart valve thrombosis is more common than previously thought, which is characterized by imaging abnormalities, gradient increase, and N-terminal pro brain natriuretic peptide level, which mostly occurs after balloon dilated transcatheter aortic valve replacement and valve replacement [9, 10]. To test this idea, he studied 87 patients [11, 12]. Among the 87 patients, 46 (53%) had mitral valve replacement, 27 (31%) had aortic valve replacement, and 14 (16%) had double valve replacement. The research of Oschatz is accidental and its reliability is not high [13, 14]. In
order to explain the influence of element measurement on bubble size distribution, Sauter's average diameter, and volume mass transfer coefficient as performance indicators of pressure dissolved microbubble generator, Dasi studied the influence of swirl chamber, crushing plate, and other elements on the performance of pressurized dissolved microbubble generator [15, 16]. In the process of the experiment, Dasi used a crushing disk and B-type cyclone chamber to improve the efficiency of the pressurized dissolved microbubble generator. At the same time, the experimental results were quantitatively summarized [17, 18]. Dasi's research results show that in terms of influencing bubble size, the crushing disk has the best efficiency, and its contribution rate to D50 and Sauter's average diameter is 27.84% and 45.16%, respectively. In other cases, the influence of swirl chamber on mass transfer coefficient is better than that in other cases [19, 20]. Dasi's research has great reference value, but the economy of its research method needs to be improved [21, 22]. In order to prepare ultrasound microbubbles carrying herpes simplex virus thymidine kinase (HSV-TK) gene and target HepG2 cells, Carlyle observed its targeting ability in vitro and its inhibitory effect on proliferation of HepG2 cells. Carlyle accumulated ultrasonic microbubbles carrying HSV-TK on the surface of HepG2 cells and detected proliferating cell nuclear antigen (PCNA) and thiazolyl blue (MTT) [23, 24]. He found that the proliferation ability of targeted microbubbles was significantly decreased, and the apoptosis was significantly increased. The cell invasion test of Carlyle showed that the gene targeted microbubbles group (22.18 ± 2.01) was significantly less than that of the control group and the non-targeted microbubbles group, which had a good inhibitory effect on the proliferation and invasion of HepG2 cells [25]. Therefore, gene targeted microbubbles have a good inhibitory effect on HepG2 cells. Ultrasound targeted microbubble destruction (UTMD) is a noninvasive microRNA delivery method. Carlyle's research method is relatively advanced, but its practicability still needs to be further verified. Based on the analysis of the basic state and characteristics of albumin nanoparticles and ultrasonic microbubble system, 11 dogs were selected and divided into two groups. In the experimental group, albumin nanoparticles ultrasound microbubble carrier was used to treat the thrombus after cardiac valve replacement preventive effect.

2. Materials and Methods

2.1. Relevant Materials

(1) Eleven healthy domestic dogs weighing 28–36 kg were male. They were divided into control group (n = 5) and treated with cardiac valve replacement and nonanticoagulant therapy. In the experimental group, 6 rats underwent cardiac valve replacement and targeted gene therapy.

(2) The reagents used included plasmid pSec Tag 2B, competent cells E. coli JM1109, hamster ovary cells, dh5a competent bacteria, RNAiso Plus, PrimeScript, RT Master Mix (perfect real time), SYBR Premix Ex Taq™ II (Tli RNaseH Plus), HindIII, KpnI, BamHI, and Xhol, and E.Z.N.A. Endo-Free Plasmid Maxi Kit. Human umbilical vein endothelial cell line EA.hy926 Cells. DMEM/F12 basic medium, dipalmitoylphosphatidylcholine (DSPC). Methods: Vent DNA polymerase and T4 DNA ligase, fetal serum, 0.25% Trypsin-EDTA, and Opti-MEM, respectively. Sheep anti-human tPA polyclonal antibody, rabbit anti-sheep polyclonal antibody, and AssayMax human tissue-type plasma activator ELISA kit. Bovine serum albumin (BSA), tPA ELISA kit, and halo-carbon-218 were used.

(3) The related instruments include rotary evaporator, rotary evaporator, vacuum drying oven, one hundred thousandth electronic balance, ultrasonic cleaner, ultrasonic gene transfection system, laser particle size detection, and zeta potential analyzer, probe diameter 0.8 cm², probe area, probe frequency 3 MHz, energy range 0–6 w/cm, built-in working cycle pulse setting and digital timer, optical microscope, ultrasonic vibrometer, CO₂ incubator, ultra clean table, precision pH meter, fluorescence microscope, desktop high-speed low-temperature centrifuge, circulating water vacuum pump, and multifunctional enzyme marker.

3. Methods

After operation, a thromboembolism model was established in dogs to replace heart valves and plasma activator gene (TPA) encapsulated in nanoparticles. Albumin nanoparticles were prepared and transported by albumin ultrasound microbubble channel. The effects of albumin nanoparticles on thrombosis and local vascular smooth muscle cell proliferation were observed. The plasmids with high expression of tPA gene were prepared, and plasma nanoparticles carrying tPA were prepared from albumin. Albumin was used to prepare glycosidic acid ultrasound microbubble channel and combined with tPA-loaded plasma nanoparticles to form targeted diagnostic ultrasound microbubble vector. After mechanical tricuspid valve replacement in dogs, targeted ultrasound diagnosis was performed to observe the preventive effect of postoperative thrombosis. The expression of tPA antigen in heart tissue was detected by immunohistochemistry, and the content of tPA and D-dimer in blood was detected by ELISA. At the end of the experiment, local thrombosis was observed in each group. It should be noted that the heart valve should be taken as the center, about 1 cm of coronary artery and vein bridge should be taken, and the heart valve tissue from the anterior wall of the heart should be taken for routine, immunohistochemistry, and in situ hybridization. Immunohistochemistry was used to detect the expression of tPA in myocardium. The first antibody was sheep anti-human tPA polyclonal antibody (dilution 1/100). The expression of proliferating cell nuclear antigen (PCNA) was detected by LSAB method. Anti-PCNA mouse monoclonal antibody (diluted 1/400) was performed according to the kit. Human
lung cancer was positive for PCNA and PBS was used as negative control instead of antibody. A PDGF-B mRNA oligonucleotide detector was designed. The sequence was position 91 5-gat CGC ACC AAT GCC AAC tccgt GTG TGG CCC CCC TGC GTG gag GTG cag-3141. The effectiveness and specificity of digoxin were detected by dot blot hybridization. The concentration of digoxin (1.5 ng/UL) was determined. The expression of PDGF-B mRNA was detected by conventional in situ hybridization. Then, the internal thickness and area of coronary artery were measured by pathological image analyzer, and the ratio of corneal thickness was calculated accurately. The formula was as follows: lumen stenosis rate = (1 – existing lumen area/area below the elastic membrane) × 100%. The number of PCNA positive cells in 200 coronary artery cells was measured as the number of in situ hybridization positive cells. After statistical calculation, the internal area is 0.25 cm^2 to × 4 (i.e., 1 cm^2), which is lower than the high-power field (×400) of the microscope.

3.1. Construction and Expression of Gene Plasmid. In this experiment, three pairs of primers were designed according to three EST sequences and tPA gene sequences to amplify the three fragments of tPA from three EST cloning plasmids. Three EST clones were cultured on LB containing ampicillin and chloramphenicol. Three EST clone plasmids were extracted and three tPA amplified fragments were used as PCR vectors. The enhancement products were recovered and identified in sequence analysis. HindIII and XhoI, HindIII and KpnI, KpnI and BamHI, and BamHI and XhoI were digested by pSec Tag 2B and three tPA—tPA-1, tPA-2, and tPA-3, respectively. The products were cleaned with Qiagen PCR product purification kit. Meanwhile, T4 DNA ligase was used. The decontamination products were transformed into E. coli JM109 cells by 14°C overnight. The resistant colonies were screened from LB tablets containing ampicillin. The recombinant plasmids were detected by PCR, and the recombinant plasmids were extracted and identified. The recombinant plasmid was transfected into CHO cells by calcium phosphate coprecipitation method. The expression of tPA was detected by indirect immunofluorescence.

The primers for tPA and the three EST sequences are as follows: t-PA-1F: 5′-CCCaagcttATGGATGCATGAAAGA-GAGGG-3′, t-PA-1R: 5′-GGggtacccACGTTAGGCT-GACCCATTCC-3′, t-PA-2F: 5′-GGggtacccACAGCGCT-CACCGATGC-3′, t-PA-2R: 5′-GGggtacccACGAG-AAGGTATGAGTGC-3′, t-PA-3F: 5′-GGggtacccTCTGCGCCAGCTTGCT-3′, t-PA-3R: 5′-CCtccgagGCG GTGCCATGTTGTCAC-3′.

The primer sequence of the primer sequence is t-PA-1F: 5′-GCCaagcttATGGATGCGATGAAAGA-GAGGG-3′, t-PA-1R: 5′-GGggtacccACGTTAGGCT-GACCCATTCC-3′, t-PA-2F: 5′-GGggtacccACAGCGCT-CACCGATGC-3′, t-PA-2R: 5′-GGggtacccACGAG-AAGGTATGAGTGC-3′, t-PA-3F: 5′-GGggtacccTCTGCGCCAGCTTGCT-3′, t-PA-3R: 5′-CCtccgagGCG GTGCCATGTTGTCAC-3′.

3.2. Preparation of Ultrasound Microbubbles. First, a sterile 10 ml bovine serum albumin liquid (5% (g/M)) was prepared, in which 10% sucrose was prepared and placed in a 50 ml covered plastic centrifuge tube. Oxygen and charcoal gas were used to corrode the liquid (flow rate: 6 ml/min), about 10 min, ultrasonic treatment was carried out for 1 minute in turn (condition: 180 W, stable frequency: 20 kHz), and the prepared microspheres were stored at 4°C for future use. The morphology, size, and number of microvessels were observed under microscope. The size and surface zeta potential of the microvessels were measured by Zetasizer 3000 analyzer.

Secondly, on the one hand, the albumin nanoparticles (containing 1mg plasmid) were added into 5ml albumin microbubbles and 10 ml of 50% glutaraldehyde solution and then incubated at 4°C for 2 hours for interconnection (the final concentration of glutaraldehyde solution was 0.1%). On the other hand, during the preparation of nanoparticles, the supernatant was collected after centrifugation, and the amount of plasmid remaining in the nanoparticles was detected. The encapsulation ratio shall be calculated according to the ratio of total amount to balance. The amount of plasmids carrying albumin nanoparticles ultrasound microbubble complex was calculated.

With normal serum, centrifugation, stratification, centrifugation speed of 200 r/min, for 1 minute, suspension foam was obtained to obtain nano microbubbles containing targeted ultrasound contrast factor (carrier) gene 0.88 × 10^7 ~ 1.8 × 10^9/ml microbubbles were found in 10 ml targeted contrast medium, and the gene plasmid content was 2 mg. It is then stored at 4 degrees Celsius for later use.

Finally, Zetasizer 3000 was fully used to determine the size and zeta potential of albumin nanoparticles. Albumin nanoparticles need gel retardation test and encapsulation efficiency test. The prepared albumin nanoparticles (containing 2mg plasmid) were added into 10 ml albumin microbubbles, added 50% glutaraldehyde solution 10 ml, and incubated for 2 h at 4 degrees Celsius. After cross-linking, centrifugation (rotating speed 200 r/min, 1 minute), suspension foam was obtained, and then the carrier was obtained. The microbubble concentration was adjusted to (0.8–1.8) × 10^9 cells/ml.

3.3. Targeted Ultrasound Therapy of TPA Gene-Loaded Albumin Nanobubbles. The distribution of albumin nanoparticles in vivo mainly depends on its size and surface...
properties. Nanoparticles have a solid particle skeleton. Only nanoparticles with a diameter less than 100 nm can pass through the hepatic sinusoidal endothelial cell window and accumulate in the liver without entering the human body. Nanoparticles modified with hydrophilic groups in the range of 100–200 nm can avoid phagocytosis of reticuloendothelial system and become a long circulating system. To obtain colloidal system with good physical and chemical properties is a prerequisite for the use of protein nanoparticles as drug carriers and surfactants. Therefore, the preparation process has an important impact on the in vivo activity and targeting of albumin nanoparticles after intravenous injection. After tricuspid valve replacement in the experimental group, the heart rate of chest wall surface was detected by two-dimensional ultrasound. Imaging. After intravenous injection of 10 ml albumin nanovesicles containing tPA gene, cardiac imaging was significantly improved. The chest wall was directly connected to the right ventricular tube of the anterior wall of the heart for 30 minutes. The chest wall was examined by two-dimensional ultrasound to observe the description of the heart. The ultrasonic frequency was 1 MHz and the intensity was 1.5 W/cm². In the control group, the tricuspid valves were replaced by conventional cardiopulmonary bypass (CPB), and then they were intravenously injected with 10 ml normal saline and irradiated with ultrasound (as before). Intravenous injection of tPA 10 ml gene (containing tPA 2 mg) of nano albumin microvascular can significantly improve the development of related organs. Postoperative antibiotics were used to prevent traumatic infection. There was no anticoagulant therapy in each group. Before and after the operation (four weeks), the content of D-dimer and tPA were detected by ELISA. A diphosphorylase is used to remove the phosphorus group at the fifth end of DNA. Dephosphorylation of 5-limb vectors can reduce the occurrence of self-cyclization of vectors, thus reducing the production of false positive clones.

In order to improve the therapeutic potential of UTMD, microbubble gene vectors should have sufficient capacity to concentrate DNA and/or specific cell or tissue selectivity. Therefore, after 70–80% digestion of EA.Y926 cells the day before the experiment, 5000 wells were divided into 96 vertical plates and cultured overnight at 70–80% fusion rate. Albumin nanoparticles with plasmid content of 0.05 μg, 0.1 μg, 1 μg, and 10 μg were added into each axis, which was composed of 0.5 μg/ml plasmid, 1 μg/ml, 10 μg/ml, and 100 μg/ml, and added into the conventional culture medium, and incubated at room temperature for 20 hours. Then, add 10 u1CCK-8 solution to each well. The plates were incubated in incubator for 4 hours. The absorbance at 450 nm was determined by enzyme-linked immunosorbent assay, and the number of units was calculated according to the standard curve.

4. Results and Discussion

On the one hand, with the development of molecular biology, tissue-type plasminogen activator (tPA) has been used as a target for gene therapy in thrombosis-related diseases, so as to achieve the purpose of anticoagulation. This not only avoids the systemic bleeding complications caused by drug thrombolysis, but also plays a role in the local long term to prevent thrombosis. Intravenous tissue plasminogen activator (tPA) was first introduced as a safe and effective thrombolytic agent, and then new thrombolytic agents, anticoagulants, and antiplatelet drugs were introduced. These drugs were considered as potential safety drugs with more favorable interactions. In addition to chemical thrombolysis, other techniques, including transcranial ultrasound thrombolysis and microbubble cavitation, have been introduced.

4.1. Analysis of Albumin Nanoparticles and Microbubbles. The average diameter of nanoparticles is 132.0 nm and the average surface area of zeta is +31.88 ± 40.03 mV. The average entrapment efficiency was 73.97%. Gel retardation analysis and DNase I protection experiments showed that nanoparticles had protective effect on DNA plasma. The ultrasonic microsphere was round and vesicular, with uniform size and good dispersion, and the diameter was 2–5 μm. The albumin microbubbles containing 10% sucrose were stored at 4 °C for 30 days with the same morphology and good thermal stability (40°C for 30 minutes). After interaction with albumin nanoparticles, the above properties did not change significantly. The particle size and zeta potential are shown in Table 1.

Under the microscope, the cationic microbubble suspension was observed. The microbubbles were spherical in shape, and the size distribution was uneven. There was no obvious fusion and rupture. It is not difficult to see from Table 1 that the average size of microbubbles is 3.1 ± 1.6 μm and zeta potential is 13.70 ± 1.95 MV. The microbubble concentration was (4.2 ± 1.3) × 10/ml. The particle size was 218.1 ± 1.1 nm and zeta potential was −15.30 ± 0.91 MV. The average size of nanoparticles ultrasound microbubble complex was 4.5 ± 1.8 μm. The zeta potential was 2.79 ± 1.40 MV. The concentration was (3.0 ± 1.3) × 10/ml.

The results of particle size, morphology, and entrapment efficiency of tPA-loaded albumin nanoparticles are shown in Figure 1.

According to Figure 1 and SEM results, albumin nanoparticles are spherical, uniform in size, and well dispersed. The particle size analysis of Zetasizer 3000 showed that the average particle size was 132.0 nm, the maximum particle size was 153.1 nm, and the minimum particle size was 48.9 nm. The average multiple scattering index is 0.34. The average surface area of zeta was 31.32 ± 41.42 MV. The blocking rate of the sample was detected by UV spectrophotometer, and the total DNA content of plasmid was determined. The results showed that the average absorption efficiency was 73.58%, which met the experimental requirements.

4.2. Survival Rate of Albumin Nanoparticle Ultrasound Microbubble Complex. In vitro and in vivo ultrasound imaging experiments show that complex of albumin nanoparticle microbubbles has a significant impact on ultrasound imaging, indicating that the microbubbles still
have good acoustic performance even after inhalation of nanoparticles. We treated the complex of albumin nanoparticle ultrasound microbubbles with endothelial cells EA.hy926. Albumin nanoparticles were detected in the acute toxic cells of ultrasound microbubble complex for 20 hours. The cell survival rate of albumin nanoparticles ultrasound microbubble complex is shown in Figure 2.

According to the data in Figure 2, for most concentration groups, there was no obvious cytotoxicity effect; only slight decrease of cell activity was observed in the highest concentration group, and the average cell activity was between 80%–120% compared with the control group.

4.3. Comparison of tPA Content before and after Operation. Thrombolysis or microthrombosis in situ may be the cause of poor clinical improvement after recanalization in acute ischemic stroke. Therefore, the thickness and area of heart valve are one of the important factors. The comparison of intimal thickness and area between the 5 normal dogs in the control group who underwent cardiac valve replacement and nonanticoagulant therapy alone and the 6 dogs in the control group receiving targeted gene transfer therapy at the same time with cardiac valve replacement is shown in Figure 3.

It can be seen from Figure 3 that the average thickness of the heart valve in the control group is 67.22 nm, and the average thickness of the six ordinary dogs in the experimental group is 17.14 nm. In the comparison of area size, the average thickness of the control group is 24.95 mm, and the data of the experimental group is 21.13 mm. The detection results of blood D-dimer and tPA before and 4 weeks after targeted gene transfer in experimental group and control group are shown in Table 2.

It can be seen from Table 2 that the D-dimer and tPA content of the control group before operation were 78.36 ± 6.12 and 0.19 ± 0.08, respectively, and the two values after operation were 80.50 ± 3.35 and 0.18 ± 0.03, respectively. At the same time, the levels of D-dimer and tPA were 79.22 ± 4.56, 0.17 ± 0.05, 856.78 ± 100.20, and 0.67 ± 0.11 in the control group before and after the operation. The two indexes of fibrinolytic activity in the experimental group were significantly higher than those before the gene transfer, the control group before and after the operation. There was no significant difference in the ratio. The changes of tPA and D-dimer in venous blood of experimental group at different periods after operation are shown in Figure 4.

It can be seen from Figure 4 that the average value of tPA in the experimental group before operation is 0.19, and it changes to 0.50 one week after operation. At this time, D-dimer also changes from 89.56 to 835.28. In the following weeks, the content of tPA increased steadily until the eighth week, and then slightly decreased to 0.79 at the 12th week after operation, but the tPA content was still very high, and the change trend of D-dimer was roughly the same as that of tPA. In the experimental group, the two indexes were significantly increased and maintained at a high level 1 week after operation until the end of the experimental observation at 12 weeks after operation. Prothrombin time was detected and the international normalized ratio was calculated, and no significant change was found.

### Table 1: Particle size and zeta potential.

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<th>Particle size</th>
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<td>Plasmid albumin nanoparticles</td>
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<td>−15.30 ± 0.91 mV</td>
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<td>Cationic microbubbles</td>
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<td>Albumin nanoparticles ultrasound microbubble complex</td>
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<td>2.7 ± 1.40 mV</td>
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![Figure 1: Analysis of particle size of Zetasizer 3000 instrument.](image)
5. Conclusions

Most methods of gene transfer into human body are to inject the target gene into target tissue directly. For cardiovascular diseases, direct injection of myocardial or plasmid DNA or adenovector into cardiac catheters is carried out. In view of the application of ultrasound contrast agent, microbubbles have been confirmed to have the function of directional delivery while enhancing ultrasound imaging by using microbubbles. It can target gene or drug delivery to achieve the purpose of treatment of diseases and can create a safe and effective drug ultrasound-mediated targeted delivery system.

At present, there are many studies on albumin as liquid membrane microbubbles, which are nontoxic and easy to prepare. The disadvantage is its poor stability. In this experiment, 10% sucrose was selected as glycoprotein, which not only avoided the above conditions, but also had good microbubble stability. Because the inert gas fluorocarbon is the most commonly used gas in the preparation of ultrasound microbubbles, in this experiment, we used the physical inhalation (one-step method) to prepare albumin nanoparticles loaded with tPA gene. The particle size was about 132.0 nm, the particle size was relatively uniform, the dispersion was good, and the entrapment efficiency was 73.58%. Gel electrophoresis analysis showed that it had a good protective effect on DNA enzyme. The slight damage of ultrasound on microbubbles may be reversible damage and permeability of target cell membrane caused by ultrasound or mechanical factors, resulting in the rupture of microbubbles with diameter of 7 mm and the expansion of endothelial cell space. The target gene reaches tissue cells...
through the gap between microbubble rupture and endothelial cells. In this study, tPA gene expression plasmid was constructed, and the appropriate vector was selected to transfer tPA into the body cells to obtain effective and sustained tPA production, so as to achieve the purpose of long-term anticoagulation and prevention of thrombosis. Using three EST cloning plasmids as matrix, three tPA fragments were amplified and cloned into pSec Tag 2B plasmid to prepare recombinant pSec Tag 2B tPA. In the experimental group, although the normal endothelial cells attached to the surface of the biological valve, the degree of valve leaf degeneration has been slowed down, but the incidence of valve thrombosis is still high due to the incomplete and partial shedding of endothelium and no anticoagulant treatment after operation. The results showed that the targeted release and gene expression of albumin nanoparticles loaded with gene plasmids could be enhanced by ultrasound destruction of microbubbles. This method is a simple and effective new technology for targeted gene transfer. With the development of molecular biology and ultrasound medicine technology, it will provide a new way to study the prevention of human diseases.

Data Availability

The data used to support the findings of this study are included within the article.

Disclosure

Feng Qi and Linlin Zhao are considered as the co-first authors.

Conflicts of Interest

The authors declare no conflicts of interest regarding the publication of this paper.

Authors’ Contributions

Feng Qi and Linlin Zhao contributed equally to this work.

References


