Combination of doxorubicin liposomes with left atrial appendage radiofrequency catheter ablation to reduce post-ablation recovery of electrical conduction

Ying Zhuge1,2,*
Huan-Er Ni1,2,*
Yu-Jie Wang3
Mu-Ye He4
Jun-Shan Wang3
Feng Gao3,4
Fang Wang1

1Department of Cardiology, Shanghai General Hospital, Shanghai Jiao Tong University School of Medicine, Shanghai 200800, People’s Republic of China; 2Department of Cardiomyocytes in the ablated area and the proximal transition zone of the LAA.

Aims: To determine whether use of radiofrequency catheter ablation (RFCA) combined with intravenously administered liposomal doxorubicin (L-DOX) facilitates a reduction in the recovery of post-ablation electrical conduction.

Methods: Circumferential ablation was performed on the epicardial surface of the left atrial appendage (LAA) in New Zealand White rabbits, and L-DOX was then administered intravenously. Fluorescence spectrophotometry was used to assess reagent bio-distribution, while Western blots and immunohistochemistry were used to assess the localization of the apoptotic markers Bel-2, Bax, and cleaved CASP3 in the LAA. Liver, kidney, and cardiac functions were also measured to evaluate the safety of this approach.

Results: At 1 week and 1 month after RFCA, a pacing electrocardiogram could not be detected in most of the rabbits that had received the combined RFCA and L-DOX therapy. L-DOX began to target the LAA on the second day after RFCA. L-DOX treatment increased the apoptosis of cardiomyocytes in the regions peripheral to the necrotic area induced by RFCA. Doxorubicin had some effect on liver and kidney function, but these effects were reversible and did not affect survival.

Conclusion: The present results provide evidence that L-DOX treatment can reduce the recovery of electrical conduction after RFCA therapy owing to L-DOX-induced apoptosis of cardiomyocytes in the ablated area and the proximal transition zone of the LAA.

Keywords: doxorubicin, liposome, radiofrequency catheter ablation, atrial fibrillation, recovered electrical conduction

Introduction

Atrial fibrillation (AF) is the most common form of arrhythmia and is associated with high morbidity and mortality.1 However, current pharmaceutical approaches are generally not sufficient to treat this condition, owing to recurrent AF episodes and side effects associated with long-term treatment. Over the last decade, radiofrequency catheter ablation (RFCA) has been established as the preferred treatment for a subset of patients with symptomatic AF, offering these individuals clear improvements in their quality of life.2 RFCA approaches such as pulmonary vein isolation are effective treatments for drug-refractory AF.1,3–5 However, while RFCA treatment induces acute coagulation and necrosis of myocardial cells at the localized site where high temperature is initially applied, peripheral myocardial cells can undergo sub-lethal and reversible damage owing to lower temperatures, enabling some of these cells to recover.

Correspondence: Fang Wang
Department of Cardiology, Shanghai General Hospital, Shanghai Jiao Tong University School of Medicine, 100 Haining Road, Hongkou District, Shanghai 200080, People’s Republic of China
Tel +86 21 6324 0090
Fax +86 21 6324 0090
Email onlyfang1@hotmail.com

*These authors contributed equally to this work

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over time. As a result, incomplete ablation coupled with the recovery of acutely damaged myocardial cells can contribute to a recovery of electrical conduction at the ablated site.6–9

RFCA is known to induce an inflammatory reaction.10 According to a previous study, doxorubicin (DOX) is one of the most widely used clinical anthracycline anti-tumor drugs11 and liposomes have inflammation-targeting properties.12 In our project, liposomal doxorubicin (L-DOX) were injected into the auricular veins of rabbits after circumferential ablation of the anterior and posterior wall of the left atrial appendage (LAA), which is a source of localized re-entrant tachycardia and an important non-pulmonary vein trigger of AF.13–15 We found that L-DOX could effectively reach the LAA ablation site owing to its inflammation-targeting capabilities, and once there, it could completely kill acutely damaged myocardial cells that would otherwise be able to recover, thus maximizing the therapeutic efficacy of RFCA (Figure 1).

Methods

RFCA operation procedures

The operation was performed using aseptic techniques with sterile instruments. Before surgery, rabbits were injected with 1% pentobarbital sodium that was administered into the auricular vein at a dose of 3 mL/kg. After the corneal reflex was no longer apparent, rabbits were placed on a warming operative platform (37°C) in the supine position. The third and fourth left ribs were cut, the chest was opened, and the whole heart was then exposed using a chest expander. An incision was made in the pericardium to expose the myocardial wall, and the LAA was gently exteriorized using smooth forceps and a medical suture line was threaded through the distal end of the LAA.

Before ablation, the following parameters were set on the Cardiac Ablation Generator (IBI-1500T8, Irvine Biomedical Inc., Irvine, CA, USA): power =15 W, temperature =55°C, time =15 seconds. The LAA was then slightly raised using the medical suture line, and a 4 mm quadrupole temperature-controlled ablation catheter (Celsius, Biosense Webster Inc., Irwindale, CA, USA) was used to perform circumferential ablation around the anterior and posterior wall of the LAA (Figure S1). The number of ablation lesions or the distance between lesions was not pre-defined. RFCA was conducted until a bidirectional conduction block of the LAA was achieved. As such, the empirical electrical isolation of the LAA was the endpoint of this ablation procedure.

Fluorescence imaging

IR-775 chloride, a near-infrared fluorescent dye with an excitation wavelength of 740 nm and an emission wavelength of 790 nm, was used to reveal in vivo agent distributions.

Figure 1 Illustration of the use of L-DOX in combination with LAA RFCA to reduce the recovery of electrical conduction.

Abbreviations: L-DOX, liposomal doxorubicin; LAA, left atrial appendage; RFCA, radiofrequency catheter ablation.
The IR-775 liposome (L-IR-775) preparation protocols are available in the online Supplementary material. Twenty-four adult New Zealand White rabbits were randomly divided into four groups: group A, blank; group B, intravenous administration of L-IR-775 (0.5 mg IR-775/kg) after thoracotomy without RFCA (sham-operated [SO] + L-IR-775 group); group C, intravenous administration of free IR-775 (0.5 mg/kg) after RFCA (RFCA+IR-775 group); group D, intravenous administration of L-IR-775 (0.5 mg IR-775/kg) after RFCA (RFCA+L-IR-775 group). Group D was subdivided into two groups in which the rabbits were sacrificed after 24 hours and 48 hours respectively. Free IR-775 and L-IR-775 were administered as a single dose through intravenous auricular vein administration.

Organs of interest (including heart, liver, spleen, lung, and kidney) were harvested, rinsed with saline, and placed in a culture dish. This culture dish was placed in an In-vivo Imaging System (Lumina III, PerkinElmer Inc., Waltham, MA, USA) equipped with a fluorescent filter ($\lambda_{em}=740$ nm, $\lambda_{ex}=790$ nm). The settings were as follows: subject height =1.5 cm, near infrared excitation light intensity =8 mW/cm$^2$, pixel resolution =1,024×1,024, exposure time =1 second, binning =2×2.

Pacing and mapping maneuver

The L-DOX preparation protocols are available in the online Supplementary material. Thirty adult New Zealand White rabbits were randomly divided into three groups: group A, RFCA; group B, intravenous administration of free DOX (4 mg/kg) after RFCA (RFCA+DOX (4 mg/kg) group); group C, intravenous administration of L-DOX (4 mg DOX/kg) after RFCA (RFCA+L-DOX (4 mg DOX/kg) group). Free DOX and L-DOX were administered as a single intravenous dose in the auricular vein.

Electrical conduction between the interior and exterior circumferential ablation lesions was assessed in all animals before ablation, as well as 5 minutes, 1 week, and 1 month after RFCA. The rabbits’ chests were closed, and wounds were sutured and disinfected 5 minutes after RFCA. After waking up, the animals were sent to the laboratory animal center for feeding. All animals received a dose of 400,000 U/day of penicillin intramuscularly for 3 consecutive days. One week and 1 month after the surgery, each rabbit’s chest was reopened and a pacing electrode was placed on the epicardium to detect the pacing electrocardiogram.

The electrocardiogram was monitored and stored on a BL420 Biological Signal Acquisition and Analysis System (BL420S, Taimeng Software Co. Ltd., Chengdu, People’s Republic of China). The tip of the stimulating electrode was attached to the anterior or posterior wall of the LAA inside the circumferential ablation lesion (Figure S2) and it was connected to the Cardiac Electrophysiological Stimulator (CES, DF-5A, Suzhou Dongfang Electronic Instrument Factory, Suzhou, People’s Republic of China). Rapid S1S1 (150 ms cycle length) procedural stimuli were adapted to pace.

The myocardial electrogram was used to reflect the muscle excitatory activity of the LAA muscle to verify whether it was captured by the pacing signal (Figures S3 and S4). The electrical conduction block is available in the online Supplementary material (Figure S5). Successful LAA conduction block was defined as no detection of pacing electrocardiogram while pacing myocardial tissues inside the circumferential ablation lesions.

L-DOX tissue distribution

Twenty-five adult New Zealand White rabbits were randomly divided into two groups: group A (n=5), RFCA+DOX (4 mg/kg); group B (n=20), RFCA+L-DOX (4 mg DOX/kg).

Five rabbits in group A were sacrificed on the second day after intravenous injection. Twenty rabbits in group B were randomly divided into four smaller groups, which were sacrificed on the first, second, fifth, and seventh day after intravenous injection. Key organ samples (including heart, liver, spleen, lung, and kidney) were harvested and stored at −80°C. The concentration of DOX in tissues was measured by fluorescence spectrophotometry with a fluorescence analysis spectrometer (Lumina, Thermo Fisher Scientific, Waltham, MA, USA), following previously published protocols16,17 (the specific detection steps are available in the online Supplementary material).

Detection of LAA cardiomyocyte apoptosis

Fifteen adult New Zealand White rabbits were randomly divided into three groups: group A, SO; group B, RFCA; group C, RFCA+L-DOX (4 mg DOX/kg).

After 1 week, rabbits were sacrificed, hearts were collected and the LAA was carefully cut out. The LAA was stored in normal saline in order to remove blood, and was then fixed in 4% paraformaldehyde and frozen at −80°C until further analysis. TUNEL staining was used to assess cardiomyocyte apoptosis in the ablated area and in the transition zone of the LAA (the specific detection steps are available in the online Supplementary material).
Western blot analysis of LAA proteins
The expression of Bcl-2, Bax, and cleaved CASP3 in the ablated area and transition zone of the LAA were detected by Western blotting. Fifteen adult New Zealand White rabbits were randomly divided into three groups: group A, SO; group B, RFCA; group C, RFCA+L-DOX (4 mg DOX/kg).

After 1 week, rabbits were sacrificed and the LAA was collected as previously described. We then used SDS-PAGE to analyze proteins. Those proteins were electro-transferred to PVDF membranes after electrophoresis. These membranes were used for immunoblotting with appropriate antibodies. The protein bands were scanned and quantified by using ImageJ software (specific detection steps are available in the online Supplementary material).

Semi-quantitative immunohistochemical (IHC) examination
A total of 15 adult New Zealand White rabbits were randomly divided into three groups: group A, SO; group B, RFCA; group C, RFCA+L-DOX (4 mg DOX/kg).

After 1 week, rabbits were sacrificed and LAA specimens were collected. The expression of Bcl-2, Bax, and cleaved CASP3 was assessed by IHC staining using appropriate antibodies (specific detection steps are available in the online Supplementary material).

Evaluation of kidney and liver function
Twenty adult New Zealand White rabbits were randomly divided into four groups: group A, SO; group B, RFCA; group C, RFCA+L-DOX (3 mg DOX/kg); group D, RFCA+L-DOX (4 mg DOX/kg).

On the first, third, fifth, and seventh day after intravenous injection, a blood sample (1 mL) was collected from the auricular vein. These blood samples were centrifuged at 5,000 rpm/min for 15 minutes, and serum was collected and stored at −20°C until further analysis.

Using respective colorimetric diagnostic kits (Nanjing Jiancheng Bioengineering Institute, Nanjing, People’s Republic of China), the levels of serum ALT and AST were determined to evaluate liver function and hepatotoxicity, while blood urea nitrogen (BUN) and serum creatinine were measured to evaluate renal function and nephrotoxicity. The results were calculated as instructed by the manufacturers.

Echocardiography determination
Fifteen adult New Zealand White rabbits were randomly divided into three groups: group A, SO; group B, RFCA; group C, RFCA+L-DOX (4 mg DOX/kg).

After 1 week, rabbits were subjected to induction of anesthesia at a concentration of 4% and were maintained in a deeply anesthetized state with 2% isoflurane.

We invited two experienced ultrasonographic doctors who were blinded to the grouping of rabbits to perform transthoracic echocardiography. The left ventricular ejection fraction (LVEF), left ventricular fraction of shorting (LVFS), left ventricular end diastolic diameter (LVEDD), stroke volume, left ventricular posterior wall depth (LVPWD), and left ventricular end diastolic volume (LVEDV) of each rabbit were assessed using a 12 MHz transducer connected to a commercially available echocardiographic system (Phillips Sonos 7500, Amsterdam, Holland). All measurements are averages from three consecutive cardiac cycles.

Statistical analysis
IBM SPSS Statistics 22.0 software was used for all statistical analyses, and data were expressed as means ± SD based on three to six independent experiments. Paired Student’s t-tests were used to evaluate statistical significance. A P-value of less than 0.05 was considered statistically significant.

Ethics statement
All animal experiments were approved by the Laboratory Animal Center, Shanghai Tenth People’s Hospital, and all animal procedures were in accordance with the guidelines from Directive 2010/63/EU of the European Parliament on the protection of animals used for scientific purposes. For all in vivo experiments, adult New Zealand White rabbits (4–6 months old, 2.2–2.8 kg body weight) purchased from Shanghai Shengwang Experimental Animal Breeding Co. Ltd. (Shanghai, People’s Republic of China) were used. Male and female adult New Zealand White rabbits were maintained in a temperature-controlled (25°C) facility with a 12-hour light/dark cycle. Figure S6 shows the number and group assignment of the animals.

Results
The circumferential RFCA lesions
Upon completion of the ablation procedure, the circumferential ablation lesions were visible. There were approximately 5–6 ablation points in the anterior wall of the LAA, and approximately 4–5 ablation points in the posterior wall. The lesions associated with each ablation point were continuous and without gaps. The ablation points were coupled to each other to form a complete closed annular ablation ring, which passed through the anterior and posterior walls and
surrounded the root of the LAA to achieve empirical LAA electrical isolation.

Fluorescence imaging
L-IR-775 was not detectable in the LAA at 24 hours after surgery, but at 48 hours after surgery it was detectable in the LAA, and it was more concentrated at the ablation line than in other parts of the heart (Figure S7). The fluorescein intensity of LAA IR-775 was ranked in the following order: RFCA+L-IR-775 > RFCA+IR-775 > SO+L-IR-775 > blank (Figure 2A and B). The fluorescein intensity of IR-775 in the LAA was higher than in any other organs in the RFCA+L-IR-775 group (Figure 2C and D).

Cardiac electrophysiological examination
Representative electrophysiological recordings and the rates of electrical conduction block in four different phases are shown in Figure 3 and Table S1. Before ablation (Phase 1), pacing electrocardiogram can be detected in all groups. At 5 minutes after ablation (Phase 2), pacing electrocardiogram could not be detected in all groups. At 1 week (Phase 3) and 1 month after ablation (Phase 4), pacing electrocardiogram could not be detected in most rabbits in the RFCA+L-DOX group, while it was detectable in most rabbits in the RFCA group that did not receive L-DOX. Since all of the rabbits injected with free DOX only survived for a maximum of 48 hours due to acute toxicity, a pacing electrocardiogram could not be performed at later times in these animals.

L-DOX tissue distribution
The concentration of L-DOX in the LAA and in other parts of the heart was roughly the same on the first day after surgery. However, on the second day after surgery, the concentration of L-DOX in the LAA was higher than that in the other parts of heart. More interestingly, the...
Figure 3 The representative electrophysiological recordings of two groups in four different phases.

Notes: Group A, RFCA (A); Group C, RFCA+L-DOX (B). The dotted line indicates the start of pacing stimulation. n=10 per group.

Abbreviations: L-DOX, liposomal doxorubicin; RFCA, radiofrequency catheter ablation.
concentration of L-DOX at the LAA ablation site on the fifth and the seventh day after surgery was significantly higher than in other parts of the heart or in other organs. Liposomes remained in the liver and spleen for a long time, likely due to the ability of liposomes to passively target the liver and spleen. However, concentrations of L-DOX in lung and kidney were significantly lower than that in other organs (Figure 4A). On the second day after surgery, compared with free DOX, liposomes increased the content of DOX in the LAA ablation area and significantly reduced the DOX content in other parts of the heart and kidney (Figure 4B). The drug targeting index (DTI) of L-DOX for each organ is shown in Table S2. On the second day after surgery, the DTI of the LAA was the highest, and was significantly higher than that of the heart.

### Detection of LAA cardiomyocyte apoptosis

Figure S8 shows the TUNEL assay findings. The percentage of TUNEL-positive cells in the LAA slides was more significantly increased in the RFCA+L-DOX group (85.25%±4.31%) than in the RFCA group (63.03%±5.51%, P<0.01) or the SO group (63.63%±1.61%, P<0.001).

### Western blot analysis of LAA proteins

Western blot assays were utilized to detect protein levels of Bcl-2, Bax, and cleaved CASP3. Bcl-2 protein content in the RFCA+L-DOX group (0.32±0.10) was lower than that in the SO group (0.69±0.18, P<0.01). Bax protein content in the RFCA+L-DOX group (1.87±0.11) was significantly higher than that in the RFCA group (1.22±0.19, P<0.01). Cleaved CASP3 protein content in the RFCA+L-DOX group (1.76±0.14) was significantly higher than that in the RFCA group (1.20±0.15, P<0.001). However, the difference between the RFCA group and the SO group was substantially less apparent than the difference between the RFCA+L-DOX group and the SO group (Figure 5).

### Semi-quantitative IHC examinations

IHC localization of the apoptotic markers Bcl-2, Bax, and cleaved CASP3 was assessed in the LAA. The levels of Bcl-2 in the RFCA+L-DOX group were significantly decreased relative to the SO group and RFCA group, whereas the Bax and cleaved CASP3 protein levels were significantly increased (P<0.001). L-DOX treatment significantly down-regulated Bcl-2 expression while up-regulating Bax and cleaved CASP3 expression (P<0.001). Nevertheless, no significant differences were detected in the expression of these three proteins after treatment between the RFCA group and the SO group (Figure 6).

### Evaluation of kidney and liver function

Compared with the SO group, hepatic enzyme activities (ALT, AST) and serum renal product biomarkers (BUN, serum creatinine) were not significantly altered in RFCA group. Significant increases in levels of ALT, AST, BUN, and serum creatinine were observed in rabbits on the first 3 days in the RFCA+L-DOX groups in a dose-dependent manner. However, after the third day, the levels of those indicators began to decline (Figure 7).

### Echocardiography examination

The results of echocardiographic examination demonstrated that RFCA+L-DOX treatment significantly reduced LVEF
Figure 5 Expression level of apoptosis markers.
Notes: (A) Western blotting results for markers of apoptosis in different groups. Quantitative measurements of the band intensities of Bcl-2 (B), Bax (C), and cleaved CASP3 (D) are shown, following normalization to β-actin. *P<0.05, ***P<0.001 vs SO treatment; **P<0.01, ###P<0.001 vs RFCA treatment. n=5 per group, data are presented as means ± SD.
Abbreviations: L-DOX, liposomal doxorubicin; NS, not significant; RFCA, radiofrequency catheter ablation; SO, sham-operated.

Figure 6 The expression of Bcl-2, Bax, and cleaved CASP3 in different groups (10×40).
Notes: (A) Representative Bcl-2, Bax, and cleaved CASP3 protein levels in different treatment groups were measured via immunohistochemistry analysis. (B–D) Bcl-2, Bax, and cleaved CASP3 protein expression in each group. ***P<0.001 vs SO treatment; ###P<0.001 vs RFCA treatment. n=5 per group, data are presented as means ± SD.
Abbreviations: L-DOX, liposomal doxorubicin; NS, not significant; RFCA, radiofrequency catheter ablation; SO, sham-operated.
and LVFS compared with RFCA treatment on the seventh day. In contrast, LVEDD, stroke volume, LVPWD, and LVEDV were increased in RFCA+L-DOX group (Figure S9).

**Discussion**

We made the following observations in this study. First, we demonstrated that fluorescent liposomes (L-IR-775) are targeted to the LAA after RFCA. Second, we demonstrated that similar liposomes were able to target DOX to the LAA ablation sites. Third, L-DOX was shown to reduce the recovery of electrical conduction after RFCA. Fourth, L-DOX induced cardiomyocyte apoptosis at the LAA ablation line. Fifth, L-DOX in combination with RFCA of LAA almost had no serious adverse effects in rabbits.

At 48 hours after surgery, L-IR-775 was targeted to the LAA. We postulated that RFCA of the LAA induced an inflammatory reaction, enabling the inflammation-targeting of the fluorescently-labeled liposomes which were then phagocytosed by local macrophages. In the SO+L-IR-775 group, the IR-775 fluorescent signal in the LAA was very weak. This may be a result of the thoracotomy, which may have induced pericardial inflammation and limited L-IR-775 targeting of the inflammatory site. The IR-775 fluorescence intensity in the LAA in the RFCA+L-IR-775 group was higher than that in the RFCA+IR-775 group because L-IR-775 had superior inflammation-targeting compared to the IR-775 solution. In summary, these results showed that the LAA-targeting ability of L-IR-775 was closely associated with ablation injury.

We found that electrical conduction failed to recover at 1 week or 1 month after ablation in the majority of rabbits in the RFCA+L-DOX group, whereas it did recover in most rabbits in RFCA group. RFCA injury is thermally mediated, resulting in coagulative tissue necrosis. As the heating of tissue is reduced with increasing distance from the catheter tip, cells in regions peripheral to the necrotic area may undergo heating insufficient to cause necrosis, but sufficient to provoke acute cell damage. This may explain the electrical
conduction recovery after RFCA and long-term recurrence after AF ablation. The combination of L-DOX and RFCA in our study appeared to be more effective in mediating a permanent conduction block. This targeted therapeutic effect may be useful in reducing recurrence of AF after ablation.

The concentration of L-DOX in the LAA on the second day after surgery was higher than that in other regions of the heart. This suggests that the liposomes began to target the LAA and changed the distribution characteristics of DOX in vivo. In addition, L-DOX has a strong affinity for the liver, spleen, and especially LAA ablation sites where many macrophages are present. The accumulation of L-DOX in the ablation site of LAA also gradually increased over time. This may be as a result of liposomes increasing DOX circulation in vivo, and being able to subsequently target the inflamed sites created by RFCA.19

Our positive TUNEL testing results in the RFCA group are consistent with the data reported by Winston et al,20 who investigated the periphery of the RF lesion in the myocardial tissue of pigs. However, some investigators have observed no detectable induction of apoptosis at the periphery of RF lesions in the rat myocardium.14 The TUNEL method aims to detect DNA strand breaks that occur during apoptosis. Nonetheless, necrosis induced by RFCA can also produce DNA breaks and thus may allow for TUNEL staining. Although the TUNEL test was clearly positive at the ablation site, additional testing including Western blotting and immunohistochemistry were consistently negative in the RFCA group. Our study did find, however, that L-DOX administration markedly aggravated cardiomyocyte apoptosis after RFCA. L-DOX could thus kill the remaining cardiomyocytes in the ablation area and transition zone of the LAA, thus overcoming the previously incomplete RFCA.

L-DOX produced a transient, reversible injury to the liver and kidney in rabbits in a dose-dependent manner. We observed decreased LVEF in both the RFCA+L-DOX group and the RFCA group compared with the SO group, but a more pronounced decrease in LVEF was observed in the former compared to the latter. In addition to cardiac “depressive” effects associated with anesthesia and thoracotomy, the cardiotoxicity of DOX might have contributed to the decreased LVEF. The optimal dose of DOX must therefore be investigated in our future studies.

Since L-DOX such as DOXIL® and MYOCET® have been marketed and used clinically, this paper may yield lots of clinical applications. For instance, after RFCA surgery, the commercially available L-DOX can be administered to AF patients, and it is expected that L-DOX can increase the apoptosis of cardiomyocytes in the ablated area and transition zone in order to reduce long-term recurrence after RFCA. Nevertheless, this clinical application will have to be tested first because DOX is an agent with cardiotoxicity and its adverse effects on cardiac function cannot be avoided; thus, the study of effective dose may face challenges when ensuring safety.

We acknowledge several limitations. First, DOX is a potent anti-tumor agent with cardiotoxicity. Although liposomes are used as carriers for DOX, the adverse effects of DOX on cardiac function cannot be avoided, especially when using a large dose, as we did in this study. We hope that in our further studies, the minimum effective dose can be obtained and the side effects on the heart can be avoided as much as possible. Second, radiofrequency ablation combined with another, safer nano-formulation requires further investigation. Third, we call for further study in additional models before translating to clinical trials.

Conclusion
In this study, an L-DOX formulation with potential inflammation-targeting properties was successfully developed, and its potential use for LAA-targeting after circumferential ablation around the anterior and posterior wall of the LAA was investigated. The present results provide evidence that L-DOX reduces the recovery of electrical conduction after RFCA due to L-DOX-induced apoptosis of cardiomyocytes in the ablated area and transition zone of the LAA.

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Disclosure
The authors report no conflicts of interest in this work.

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