Enhancement of cardiac contractility by refractory period stimulation in conjunction with cardiac resynchronization therapy

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Abstract

Background: Cardiac resynchronization therapy (CRT) improves cardiac function in patients with heart failure by synchronizing contraction of the right and left ventricles (RV, LV). This study was to determine if electrical stimulation during the refractory period of bi-ventricular (BiV) pacing would provide additional LV mechanical improvement.

Methods: 16 patients receiving CRT-ICD devices were enrolled. After the CRT leads were in place, the leads were connected to an EP stimulator for acute delivery of electrical stimuli during the absolute refractory period of each cardiac beat for a 30-sec interval. Four types of refractory period stimulation were delivered: stimulation at the local LV pacing site during intrinsic rhythm, at the local RV pacing site during intrinsic rhythm, at the local LV pacing site during BiV pacing, and simultaneously at LV and RV during BiV pacing. The LV pressure was recorded for dp/dtmax measurement to assess changes of cardiac contractility.

Results: Overall LV dp/dtmax increased to 1453 ± 522 mmHg/Sec during refractory period stimulation from a baseline of 1383 ± 476 mmHg/Sec (P=0.013). The LV dp/dtmax was significantly increased by an average 12.8% when refractory period stimulation was simultaneously delivered at the LV free wall and the RV apex during BiV pacing. There was no significant improvement in LV dp/dtmax by other three types of stimulation.

Conclusion: Refractory period stimulation, when delivered to both RV and LV simultaneously, can improve LV mechanical function on top of BiV pacing, which may provide additional benefits for CRT patients.

Introduction

Congestive heart failure (CHF) is a syndrome caused by impaired ventricular function. Despite improvements in pharmacological treatment, the prognosis for patients with heart failure remains poor - the risk of death annually is 5-10% in patients with mild symptoms and 30-40% in those with advanced disease [1,2]. More recently, cardiac resynchronization therapy (CRT) that synchronizes cardiac contraction of left and right ventricles (LV, RV) has been shown to provide further morbidity and mortality benefits in addition to pharmacologic therapy in patients with ventricular dyssynchrony (wide QRS) and a reduced left ventricular ejection fraction ≤35% [3,4]. However, approximately 30% of patients who receive CRT are non-responders [5-7].

Today’s stimulation system of cardiac contractility modulation used in patients with CHF utilizes two right ventricular leads that attach to the septum of the right ventricle and apply electrical stimulation during the absolute refractory period [16,18]. The localized improvement, however, might not be synchronized with global activity of ventricles, especially in patients with impaired synchronization of activation conduction in ventricles. Thus, the purpose of the present study was to test the hypothesis that the magnitude of the myocardial contractility improvement by refractory period stimulation would be greater if the stimuli were applied through the CRT lead system.

Methods

The present study was a single-center, non-randomized acute study in CRT-indicated patients who underwent the implantation of a CRT-ICD. The study protocol was approved by the hospital Investigational

Key words: cardiac resynchronization therapy, refractory period stimulation, cardiac contractility, heart failure

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Research Board and in compliance with the Declaration of Helsinki and with the laws and regulations in China. All patients completed written informed consent.

Patient selection

The study enrolled sixteen patients between October 2011 and December 2012. Enrolled patients met the following criteria: (1) patients were indicated for receiving CRT-ICD, (2) patients underwent the placement of intracardiac leads for CRT-ICD, (3) patients were greater than 18 years of age, and (4) patients were willing and able to give informed consent.

Patients were excluded if (1) patients had experienced angina in the last three months, (2) patients’ measurement of left ventricular (LV) pressure was not technically feasible (e.g. atrial fibrillation, frequent premature ventricular contractions), (3) patients were enrolled in other clinical studies that would confound the results of this study, or (4) patients were pregnant. Demographic and medical history was collected at enrollment.

Study procedures

Lead placement: Each patient underwent standard procedures of CRT-ICD implantation under local anesthesia. Briefly, the atrial lead (model 4574, Medtronic, Inc., Minnesota, USA) was placed in the right atrial appendage, the RV lead (Model 3076, Medtronic, Inc., Minnesota, USA) was placed in the RV with the tip attaching to the apex for RV apical pacing, and the LV lead (Models 4193, 4194, or 4195, Medtronic, Inc., Minnesota, USA) was placed in a tributary of the coronary sinus for stimulating the LV free wall. The LV lead was targeted to a lateral or a posterolateral cardiac vein. The procedure of lead placement was performed under fluoroscopy. After all leads were positioned, the leads were connected to an EP stimulator (Model Transtar MX9505T, MedLab-U, Nanjing Medease Science and Technology Co., Ltd, Nanjing, China) for delivering refractory period stimulation as described below. Once the tests were completed, all leads were connected to a CRT-ICD device that was then implanted inferior to the left clavicle.

Hemodynamic monitoring: An LV pressure catheter (Model Transtar MX9505T, Smith Medical ASD, Dublin, OH 43016 USA) was inserted via the left radial artery into the LV chamber. The pressure transducer was then connected with a data acquisition system (Model Transtar MX9505T, MedLab-U, Nanjing Medease Science and Technology co., Ltd, Nanjing, China) that recorded the LV pressure at a sampling rate of 1 KHz along with simultaneous ECG. Pressure data were analyzed offline by a custom Matlab program. The LV pressure parameters for each beat consisted of LV systolic pressure, LV end-diastolic pressure, dp/dt max, and heart rate. Pressure parameters measured during premature cardiac beats were excluded in the analysis.

Test of refractory period stimulation: The absolute refractory period (ARP) of ventricles was measured in each patient to ensure no refractory period stimuli would be delivered during the later phase of the ARP. Ten S1 stimuli were delivered with the amplitude at twice pacing threshold and S1-S1 interval of 90% of the intrinsic R-R intervals based on the ECG or 10 beats faster than the sinus rate. The last S1 was followed by an S2 with S1-S2 starting at approximately 300 msec at 10% pacing threshold and S1-S1 interval of 90% of the intrinsic R-R intervals having amplitude of 8 volts. Four types of refractory period stimulation were tested. Test 1: refractory period stimulation was locally delivered via the tip electrode of the LV lead and the RV coil (return electrode) during intrinsic sinus rhythm. Test 2: stimulation was locally delivered via the tip electrode of the RV lead at the RV apex and the RV coil during intrinsic sinus rhythm. Test 3: stimulation was locally delivered via the tip electrode of the LV lead and the RV coil during BiV pacing. Test 4: stimulation was simultaneously delivered at the LV tip electrode and the RV tip electrode during BiV pacing and the return electrode was the RV coil electrode. Each test was performed for 30 sec with a 2-minute recovery interval between adjacent tests.

Data analysis

The baseline data was defined as the 30-sec data that was collected before corresponding test, e.g., the 30-sec data of the intrinsic rhythm before tests 1 and 2 and the 30-sec data of BiV pacing before tests 3 and test 4. The test data was defined as the data in which refractory period stimulation was delivered. The beat-to-beat LV pressure data was analyzed for each of 30-sec data segments and an averaged value from each 30-sec data segment was then obtained for further comparisons. Absolute values and percent changes (measured variables during test over the baseline) were compared between baseline and during tests described above. Data were presented as mean ± 5D or percentage wherever applicable. Student paired or unpaired t test for two means was used for comparisons. Statistically significant results were defined as a probability of p<0.05.

Results

All patients (age: 65 ± 11 years old, 11 males) were indicated for CRT-ICD and received optimal medical therapy prior to CRT-ICD implantation. The average left ventricular ejection fraction was 29 ± 6%, QRS duration was 156 ± 29 msec, and the NYHA classification was 3 ± 0.7. At the CRT-ICD implantation and tests of refractory period stimulation, all 16 patients were under beta-blocker therapy, 13 (81%) under angiotensin-converting enzyme inhibitor, and 10 patients (63%) under diuretics. Most patients (14, 86%) had dilated cardiomyopathy and 2 patients had a history of ventricular tachyarrhythmia. There were a total of 55 tests available for the data analysis in 16 patients.

Refractory period stimulation produced a positive effect on cardiac mechanical function as shown in LV pressure recording in Figure 1. When the refractory period stimulation was delivered via both RV and LV electrodes during BiV pacing (Figure 1B), the LV pressure and its dp/dt max were greater than during BiV pacing only (Figure 1A), suggesting a cardiac mechanical improvement. Overall, LV dp/dt max increased to 1453 ± 522 mmHg/Sec during refractory period stimulation from the baseline 1383 ± 476 mmHg/Sec without refractory period stimulation (P=0.013 vs. stimulation tests), representing an overall increase of 5.2 ± 10.4% in LV dp/dt max by refractory period stimulation.

The response to refractory period stimulation varied depending

significant responses and 31 tests (56%) yielded insignificant responses. The averaged percent change in LV dp/dt\textsubscript{max} from the baseline was 14.1 ± 8.8% for significant responses and -0.9 ± 4.7% for insignificant responses (P<0.0001 between two responses). When the maximum response (one value) was selected from multiple tests of refractory period stimulation in each of 16 patients, the averaged maximum percent change in LV dp/dt\textsubscript{max} produced by refractory period stimulation was 12.3 ± 9.1% over the baseline value. Of 16 patients, 12 (75%) patients experienced at least one significant response out of multiple tests and an averaged maximum percent change in the LV dp/dt\textsubscript{max} by refractory period stimulation was 15.8 ± 7.5% from the baseline; 4 (25%) patients had only insignificant responses during multiple tests of refractory period stimulation (the averaged maximum percent change in the LV dp/dt\textsubscript{max} of 1.8 ± 3.0% from the baseline).

Left ventricular systolic pressure during the intrinsic sinus rhythm was 138.0 ± 11.7 mmHg at the baseline and 136.1 ± 11.4 mmHg during refractory period stimulation (P=0.103 vs. during sinus rhythm). The LV systolic pressure was 127.8 ± 19.8 mmHg during the baseline BiV pacing and 132.8 ± 19.6 mmHg during the BiV pacing plus refractory period stimulation (P=0.004 vs. baseline BiV pacing). The LV diastolic pressure during the intrinsic sinus rhythm was 15.7 ± 11.6 mmHg at the baseline and 13.2 ± 11.3 mmHg during refractory period stimulation (P=0.293 vs. sinus rhythm without refractory period stimulation). The LV diastolic pressure was 14.1 ± 8.8 mmHg during baseline BiV pacing and 12.6 ± 7.6 mmHg during BiV pacing plus refractory period stimulation (P=0.094).

**Discussion**

**Major findings**

This study demonstrated for the first time that refractory period stimuli delivered through the existing CRT lead system could improve left ventricular mechanical function in terms of LV dp/dt\textsubscript{max} in CRT-indicated patients. The overall maximum percent change during refractory period stimulation compared to the baseline could reach 12% improvement in LV dp/dt\textsubscript{max}. The response to refractory period stimuli was significant when the stimuli were delivered simultaneously to both ventricles during bi-ventricular pacing.

**Prior studies**

Studies in both animals and humans have demonstrated the enhancement of ventricular contractile performance by electrical stimuli applied to cardiac tissue during the absolute refractory period [8-19]. In mechanism studies of isolated ventricular muscle, monophasic pulses delivered during the refractory period can modulate not only the cardiac contractility but also the calcium (Ca\textsuperscript{2+}) flux across the sarcolemma [8-10]. The administration of beta-blockers in that study cannot diminish the contractility modulation by the electrical stimulation during the cardiac refractory period [8,11,14]. The studies in canine heart failure model demonstrated that in addition to the acute and chronic effects of electrical modulation on contractility, changes in gene expression (genetic remodeling) might contribute to a therapeutic effect of electrical modulation on the failing heart [13-15].

Clinical evaluations of electrical cardiac contractility modulation for heart failure therapy were carried out in a cross-over study in 120 patients (Fix-HF4) [16] and a randomized trial in 420 heart failure patients (Fix-FH5) [18]. Both studies used an implantable device (Optimizer) that delivers an approximately 8V dual biphasic waveform 40 ms after a ventricular depolarization through the conventional lead.
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The parameters of refractory period stimulation. It is unknown whether the response can be further improved by modulating non-responder rate to the refractory period stimulation delivered via study, 75% of 16 tested patients met the response criterion. Thus, the define the response to refractory period stimulation in the present contractility modulation [16-19]. When this criterion was used to criterion for the positive response to an appropriate lead placement in study, the area affected by stimuli might be increased for a greater role in achieving contractility enhancement.

An acute increase in LV dp/dt\textsubscript{max} greater than 5% was used as a criterion for the positive response to an appropriate lead placement in the RV septum and acceptance of device implant for electrical cardiac contractility modulation [16-19]. When this criterion was used to define the response to refractory period stimulation in the present study, 75% of 16 tested patients met the response criterion. Thus, the non-responder rate to the refractory period stimulation delivered via the CRT lead system could be 25% based on the present study. It is unknown whether the response can be further improved by modulating the parameters of refractory period stimulation.

**Limitations**

It has long been recognized that the LV dp/dt\textsubscript{max} represents the contractility of ventricular myocardium. However, the association of the LV dp/dt\textsubscript{max} with clinical assessments and patient performance is unknown. It is also unclear whether a 5-15% increase of the LV dp/dt\textsubscript{max} would contribute to a significant improvement in clinical performance in heart failure patients.

The present study was an acute investigation without follow up. Thus, it is unknown whether a long-term application of refractory period stimulation through a CRT lead system would lead to any long-term clinical benefits such as improvements in echocardiographic and cardiac functional assessments. Long-term clinical trials are thus needed.

BiV pacing used in the present study was not equivalent to true CRT pacing due to the limitation of the EP stimulator that did not have a full capability for true CRT pacing.

The present study was performed in 16 patients. This small sample size may limit the interpretation of the findings in the present study to the CRT-indicated population. A large population study will be required to confirm the findings in the present study.

**Clinical implications**

CRT restores the synchronization of the left and right ventricles from the dys synchrony caused by inter/intra-ventricular conduction delay in selected patients (systolic dysfunction and wide ECG QRS duration). However, the CRT non-responder rate remains high, accounting for one third of CRT recipients [5-7]. Cardiac contractility modulation has been shown to improve ventricular contractility, but the current cardiac contractility modulation device only delivers stimuli at the septum of the right ventricle without helping in synchronization. Moreover, today’s device of cardiac contractility modulation must be used in patients with implantable cardioverter-defibrillator [16,18], which means the cardiac electrical therapy with two independent implantable devices. There is the possibility that ventricular mechanical function can be further improved if CRT works in conjunction with cardiac contractility modulation [21], e.g., synchronization plus contractility enhancement by one single implantable device. The present study demonstrates for the first time the feasibility that cardiac contractility modulation in terms of refractory period stimulation, when delivered simultaneously to both the left and right ventricles through an existing CRT-ICD lead system, can provide additional improvement of LV dp/ dt\textsubscript{max}. This finding may add clinical benefits on top of CRT in CRT-ICD patients, especially for those who are CRT non-responders. Another applicable possibility is to deliver refractory period stimulation only in the event when an acute weakening of myocardial contractility is detected. By doing so, refractory period stimulation is delivered on demand with battery energy saving.

**References**


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