ABSTRACT
Background: Cardiac resynchronisation therapy (CRT) improves outcomes in selected patients with heart failure and left ventricular dysfunction. One mechanism of benefit is believed to be favourable ventricular remodelling. Whether CRT also decreases the frequency of ventricular arrhythmias and risk of sudden death is unknown.

Objective: To determine the effect of CRT on frequency of ventricular arrhythmias and appropriate ICD therapies.

Design: Retrospective cohort study.

Setting: Single-centre, tertiary care facility (Mayo Clinic).

Patients: 52 patients (46 male), aged 70 (SD 10) years, who underwent upgrade from an implantable cardioverter defibrillator (ICD) to a CRT-defibrillator were included.

Interventions: Upgrade of ICD to CRT-defibrillator.

Main Outcome Measures: Frequency of ventricular arrhythmias prior to and following upgrade to CRT device.

Results: Ejection fraction increased from 22% (SD 8%) to 27% (SD 11%) following CRT. However, the frequency of non-sustained ventricular arrhythmias, sustained ventricular arrhythmias, and ventricular fibrillation was not significantly changed prior to and following CRT (2.38 (SD 9.78) vs 58.51 (SD 412.73) per patient per month, p = 0.66; 0.07 (SD 0.17) vs 0.16 (SD 0.52), p = 0.70; 0.05 (SD 0.12) vs 0.25 (SD 1.40), p = 0.12).

Conclusions: CRT is not associated with a decrease in the frequency of ventricular arrhythmias or appropriate device therapy. Thus, use of CRT alone is not beneficial in decreasing the frequency of ventricular arrhythmias or the risk of appropriate ICD therapies.

Cardiac resynchronisation therapy (CRT) is associated with fewer hospitalisations and decreased mortality in patients with congestive heart failure. These beneficial effects are thought to result from favourable left ventricular reverse remodelling, which may occur following CRT. Since adverse remodelling (ventricular dilatation), which is a consequence of worsening congestive heart failure, may be associated with increased likelihood of ventricular arrhythmias, CRT alone might be expected to decrease the incidence of ventricular arrhythmias and need for implantable cardioverter defibrillator (ICD) therapy. Whether CRT alone is associated with decreased ventricular arrhythmias or risk of sudden cardiac death is unclear since previous studies have yielded conflicting results.

Since advanced or worsening heart failure may increase the risk of ventricular arrhythmias and appropriate ICD therapies, patients with ICDs and worsening heart failure represent an especially high-risk group who may suffer significant morbidity due to frequent ICD therapies. Since CRT may promote favourable ventricular remodelling and reduce the severity of heart failure, use of CRT might provide an additional therapeutic option to decrease the frequency of appropriate ICD therapies in patients at risk. To address the impact of CRT on the occurrence of ventricular arrhythmias we analysed the frequency of ventricular arrhythmias in the absence and presence of CRT.

PATIENTS AND METHODS
The protocol was reviewed and approved by the Mayo Clinic Institutional Review Board.

Patients
All consecutive patients who underwent upgrade from a single or dual chamber ICD to CRT-defibrillator at Mayo Clinic between January 1999 and November 2005 who had ≥1 month of follow-up prior to and following device upgrade were included. Clinical and follow-up data were prospectively entered into a centralised clinical record that includes all clinical evaluations, ICD interrogations and electrogams, echocardiographic data, digital echocardiographic images, operative notes, and digital radiographic records including chest x rays, at Mayo Clinic and affiliated hospitals, and retrospectively analysed. Left ventricular dimensions and ejection fraction prior to and following CRT were measured by echocardiogram in all patients.

CRT-defibrillator upgrade
All patients were considered eligible for CRT based on accepted clinical criteria. In all cases, left ventricular pacing was initially attempted via the coronary sinus. If a coronary sinus lead could not be placed, an epicardial left ventricular lead was placed via a thoracotomy. The choice of device, coronary sinus lead, and decision to abandon coronary sinus lead placement were at the discretion of the operator. All patients were observed for 24 h after upgrade, and coronary sinus lead location and thresholds confirmed with a chest x ray and device interrogation.
Appropriate device function and pacing thresholds were assessed at 24 h and at each follow-up. Device reprogramming or lead revision, if necessary, was performed.

Ventricular arrhythmias
Ventricular arrhythmias were classified as non-sustained (terminated spontaneously), sustained (requiring antitachycardia pacing or shock therapy, excluding ventricular fibrillation), or ventricular fibrillation (cycle length <250 ms). Frequency of ventricular arrhythmias was determined by calculating the mean number of episodes per patient per month. All ventricular arrhythmia events were retrieved at the time of device interrogation during follow-up and entered into a centralised database. Each episode of ventricular arrhythmia was reviewed and classified as non-sustained, sustained ventricular tachycardia or ventricular fibrillation according to our definitions. The frequency (mean number of ventricular arrhythmia episodes per patient per month) of ventricular arrhythmias and appropriate ICD therapies was determined. Since all patients according to the study design underwent upgrade from an ICD to CRT device, episode data from the ICD were obtained retrospectively (although from the same centralised database).

Comparison of frequency of ventricular arrhythmias
All patients in the study underwent upgrade from an ICD to CRT device, such that each patient served as his or her own control for comparison of frequency of ventricular arrhythmias prior to and following CRT. However, to account for differences in the frequency of ventricular arrhythmias that may have resulted from progression of congestive heart failure or left ventricular dysfunction, we also compared our study group with a cohort of control subjects matched for age, gender, ejection fraction, and ICD device, who had not undergone upgrade of an ICD to a CRT device. To ensure an equivalent length of follow-up to our study patients, we also matched the implant date of the control subjects to the initial ICD implant date of the study patients.

CRT-defibrillator follow-up
All CRT devices were interrogated at 24 h and once again at 1 month after implantation, and subsequently every 3 months thereafter. Stored electrograms were obtained during each device interrogation, entered into a centralised database and reviewed and classified according to the same definitions.

Statistical analysis
Differences in the frequency of ventricular arrhythmias (mean number of episodes per patient per month) prior to and following CRT were compared using a Wilcoxon signed rank test. A p value of <0.05 was considered significant. All-cause mortality or need for heart transplantation following CRT was determined (Kaplan–Meier estimates).

RESULTS
Baseline characteristics
A total of 53 patients underwent upgrade from ICD to CRT-defibrillator between January 1999 and November 2005 and were included. Baseline characteristics are summarised in table 1. There were 46 men (88%) and six women (12%). Mean age was 70 (SD 10) years. Underlying cardiac disease was ischaemic in 41 patients (79%) and non-ischaemic in 11 patients (21%). All patients were New York Heart Association functional class II–III at the time of device upgrade. Initial ICD placement was for secondary prevention of sudden cardiac death in 50 patients (96%) and primary prevention in two patients (4%). Indication for ICD implant in the secondary prevention patients was sustained ventricular arrhythmias or cardiac arrest in 33 patients (65%), and non-sustained ventricular arrhythmias in association with syncope and inducible ventricular arrhythmias (sustained monomorphic ventricular tachycardia in all cases, induction of non-sustained ventricular arrhythmias (<30 beats) or polymorphic ventricular tachycardia was not considered an indication for ICD implantation) at electrophysiology study in 17 patients (33%). Atrial fibrillation was present in 55 patients (65%).

Left ventricular end-diastolic and end-systolic dimensions were 68.11 (SD 9.66) mm and 60.45 (SD 10.50) mm, respectively, and mean ejection fraction was 22% (SD 8%) prior to CRT.

Medications prior to upgrade to a CRT device are summarised in table 2. All patients were on maximal tolerated medical therapy for congestive heart failure. Thirty-six patients (69%) were taking ACE inhibitors or angiotensin receptor blockers, 35 (67%) were taking β-blockers, and 35 (65%) were on digoxin. A total of 24 patients (46%) were on antiarrhythmic drug therapy, with 20 (38%) on Amiodarone, five (10%) on Mexilitene, and one (2%) on Sotalol. Non-antiarrhythmic cardiac medications are shown in table 2.

CRT device upgrade
All patients met accepted clinical criteria for upgrade to a CRT device. In 12 patients (23%) who also met criteria for CRT, upgrade occurred at the time of planned device generator change. No patients were upgraded specifically for the purpose of reducing ventricular arrhythmia burden. Initial implantation success and need for coronary sinus lead revisions are shown in table 3. A coronary sinus lead was successfully placed in 49 patients (94%). In three patients (6%), an epicardial left ventricular lead was required. In one of these patients, the epicardial lead failed and a coronary sinus lead was subsequently placed successfully. The epicardial left ventricular lead remained stable in the other two patients. Mean fluoroscopy time was 41.57 (SD 30.26) minutes, which was not significantly different when compared with 46.30 (SD 28.97) minutes in our entire CRT implant population (p = 0.30).

Acute lead revision (<24 h) was required in three patients; this was for increased pacing thresholds in two, and macro-dislodgement of the coronary sinus lead in one patient. Chronic
lead revision (>24 h) was required in two patients at 1 and 19 months following device upgrade. In one of these, right atrial capture was present due to dislodgement of the left ventricular lead into the right atrium. The other had increased pacing thresholds requiring multiple lead revisions. In the remaining 45 patients, the coronary sinus lead remained stable over a mean follow-up time of 14 (SD 12.75) months without the need for lead revision.

Following CRT

Patient characteristics following CRT are summarised in table 3. Mean ejection fraction increased to 27% (SD 11%), an increase of 5% (SD 9%) (p<0.01) from baseline. No significant change was observed in left ventricular end diastolic and end systolic dimensions (+0.12 (SD 5.83) mm, p = 0.64 and −0.60 (SD 5.34) mm, p = 0.90, respectively).

Medication use prior to and following upgrade to a CRT-defibrillator is summarised in table 2. Although use of Digoxin and β-blockers did increase following upgrade (33 versus 36 patients, p = 0.53, 33 versus 38 patients, p = 0.52, respectively), while use of ACE inhibitors and angiotensin receptor blockers decreased (36 versus 31 patients, p = 0.31), these differences were not significant. Use of antiarrhythmic drugs was also unchanged following CRT-defibrillator upgrade.

Survival

Survival of patients following upgrade to CRT device was 88%, 77%, 66%, 61% at 6, 12, 24, and 36 months, respectively.

Frequency of ventricular arrhythmias

The frequency of ventricular arrhythmias and appropriate ICD therapies were determined 38.40 (SD 33.02) and 15.63 (SD 33.02) months prior to and following CRT, respectively.

Sustained ventricular arrhythmias

The frequency of sustained ventricular arrhythmias (requiring either antitachycardia pacing or shock therapy, excluding ventricular fibrillation) was 0.07 (SD 0.17) prior to and 0.16 (SD 0.52) following CRT (mean per patient per month) and was not significantly different (p = 0.70; fig 2).

Ventricular fibrillation

The frequency of ventricular fibrillation was 0.05 (SD 0.12) prior to and 0.25 (SD 1.40) following CRT (mean per patient per month) and was also not significantly different (p = 0.12; fig 3).

Non-sustained ventricular arrhythmias

The frequency of non-sustained ventricular arrhythmias was also unchanged after CRT and was 2.38 (SD 9.78) prior to and 58.51 (SD 412.73) (mean per patient per month) following CRT (p = 0.66; fig 1A). The large standard deviation was due to one patient with >40 000 asymptomatic episodes of non-sustained ventricular arrhythmias following CRT. After exclusion of that patient from analysis, no significant difference in the frequency of non-sustained ventricular arrhythmias prior to and following CRT was seen (1.08 (SD 2.68) versus 1.28 (SD 3.58), mean per patient per month, prior to and following CRT (p = 0.65; fig 1B)).

Comparison with control group

To determine whether the potential beneficial effect of CRT on the frequency of ventricular arrhythmias was offset by worsening congestive heart failure, we identified 52 patients matched for age, gender, ejection fraction, device type, and implant date to serve as a control group. Of these, 34 had >1 month of follow-up and were used for comparison of frequency of ventricular arrhythmias prior to and following CRT. During a mean follow-up time of 48.47 (SD 52.03) months, the frequency of sustained ventricular arrhythmias, ventricular fibrillation, and non-sustained ventricular arrhythmias was 2.49 (SD 12.69), 0.04 (SD 0.12), and 2.00 (SD 6.46), respectively. No significant differences in the frequency of ventricular arrhythmias prior to or following CRT (p = 0.27, 0.73, 0.82 prior to CRT, p = 0.29, 0.29, 0.53 following CRT) were observed in comparison with the study patients.

DISCUSSION

The major finding of this study is that, in patients with severe left ventricular dysfunction, use of CRT to decrease the frequency of ventricular arrhythmias or appropriate device therapy is not warranted.

Our findings contrast with those of previous studies that have reported both an increase and a decrease in the frequency of ventricular arrhythmias following CRT.6 8 11–13 The mechanism
by which CRT is thought to improve outcome is via decreased wall stress due to reversed remodelling. Although we did not observe a significant decrease in left ventricular size, we did observe a similar increase in ejection fraction following CRT to that of two previous studies that found decreased frequency of ventricular arrhythmias following upgrade from an ICD to a CRT-defibrillator, suggesting that our patients had a similar haemodynamic benefit from CRT, yet no change in the frequency of ventricular arrhythmias.\(^{48}\) These studies, however, were limited by small sample size and may have overestimated the effect of CRT on ventricular arrhythmia burden. In contrast, our study represents the largest single-centre series of patients undergoing upgrade from an ICD to a CRT-defibrillator, with the longest follow-up time. Moreover, by comparison of our study patients with a matched control group, we have demonstrated that the frequency of ventricular arrhythmias is similar in patients with and without CRT devices, which further supports the notion that CRT does not impact on the occurrence of ventricular arrhythmias.

It has been reported that CRT may also have a pro-arrhythmic potential. Although several mechanisms may underlie this, animal studies using epicardial pacing via the coronary sinus have demonstrated prolongation of the QT interval and torsades des pointes.\(^{12}\) A pro-arrhythmic effect of CRT has also been reported in humans.\(^{11,13,14}\) Although we did in fact observe an increase in the number of ventricular arrhythmia episodes after CRT, this difference was not statistically significant.

Our data demonstrating a neutral effect of CRT on ventricular arrhythmias is consistent with the findings of the CONTAK-CD and Multisite InSync Randomized Clinical Evaluation-ICD (MIRACLE-ICD) trials, which also found no effect of CRT on ventricular arrhythmias or appropriate ICD therapies.\(^{1,5}\) The importance of our findings is that all data are derived from a single-centre experience with uniform and complete follow-up, thereby avoiding possible confounding due to differences in device follow-up and programming. The impact of CRT alone on mortality is less clear. A recent meta-analysis including data from the extension phase of Cardiac Resynchronization-Heart Failure (CARE-HF) provided evidence against a benefit of CRT alone on risk of sudden cardiac death.\(^{15,16}\) These findings are consistent with ours and suggest that the mortality benefit of CRT is not related to a reduction in the frequency of ventricular arrhythmias but is more likely related to improved haemodynamic status as a consequence of ventricular reverse remodelling.

In conclusion, in patients with usual indications for CRT undergoing upgrade from ICD therapy, CRT alone is not associated with a reduction in the frequency of ventricular arrhythmias or ventricular fibrillation. Therefore, these data do not support the use of CRT alone as an effective therapy for frequent ventricular arrhythmias and appropriate ICD shocks in patients with worsening heart failure at risk of sudden cardiac death.

**STUDY LIMITATIONS**

Due to the retrospective study design, differences in device programming and/or initiation of antiarrhythmic drug therapy occurring as a result of a documented ventricular arrhythmia
CONCLUSIONS
Based upon our findings, CRT does not decrease the frequency of ventricular arrhythmias, and therefore upgrade of an ICD to a device capable of CRT to decrease the frequency of appropriate ICD therapies or to reduce symptoms from ventricular arrhythmias is unlikely to be of benefit.

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REFERENCES
Effect of cardiac resynchronisation therapy on occurrence of ventricular arrhythmia in patients with implantable cardioverter defibrillators undergoing upgrade to cardiac resynchronisation therapy devices

G Lin, R F Rea, S C Hammill, D L Hayes and P A Brady

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