Telemonitoring in heart failure patients treated by cardiac resynchronisation therapy with defibrillator (CRT-D): the TELECART Study

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SUMMARY
Aim: Telemonitoring (TM) is a safe and efficient monitoring system for internal cardioverter defibrillator device (ICD) recipients. TM has been used to track info on the clinical status of heart failure patients treated by ICD and/or cardiac resynchronisation therapy defibrillator (CRT-D). The aim of this study was to investigate the impact of TM on clinical outcomes in a population of CRT-D patients with heart failure. Methods: In a multicentre, randomised study, patients with chronic heart failure, New York Heart Association (NYHA) functional class II or III, left bundle branch block, severe left ventricle ejection fraction reduction (LVEF < 35%) have been identified and screened. Results: One hundred and ninety-one patients have been randomised to receive either a CRT-D with TM or a CRT-D with traditional ambulatory monitoring (control group) and completed the 12-month study follow-up. Primary endpoints were all cause death, cardiac death and hospital admission for heart failure. Secondary endpoints were atrial fibrillation, sustained episodes, non-sustained and self terminated ventricular tachycardia, ventricular fibrillation, ICD shocks and percentage of CRT-D responder patients. Univariate analysis identified the following factors predicting hospitalisation: TM, age, chronic kidney disease, hypercholesterolaemia, LVEF and NYHA class. At multivariate analysis, TM was the only factor predicting heart failure hospitalisation (hazard ratio 0.6, 0.42–0.79, 95% CI, p = 0.002), without affecting overall mortality and cardiac deaths events. Conclusions: Taken together, our data indicate the importance of TM in predicting heart failure hospitalisation in patients treated with CRT-D.

What’s known
Telemonitoring is an efficient monitoring system for internal cardioverter defibrillator device (ICD) recipients. It has been used to track info on the clinical status of heart failure patients treated by ICD and/or cardiac resynchronisation therapy.

What’s new
This study demonstrates that telemonitoring is an independent prognostic factor predicting heart failure hospitalisation in patients treated with cardiac resynchronisation therapy defibrillator (CRT-D).

Introduction
Since its introduction in the clinical scenario, telemonitoring (TM) has rapidly become a diffused system to monitor patient’s clinical course and device function (1–6). Large clinical trials have studied the TM impact on clinical or device-related events, medical care and resource consumption and follow-up visits costs (1,7–12). Previous trials have investigated the TM impact in a population of heart failure patients treated by ICD, and/or cardiac resynchronisation therapy defibrillator (CRT-D). In heart failure patients, CRT-D is the choice treatment to improve symptoms, quality of life, NYHA class and clinical outcomes, and to prevent heart failure episodes and progression (13–15). CRT-D induces a reverse cardiac remodelling in a percentage of 65% of treated patients, the so-called ‘CRT-D responders’ (13,16,17). On the other hand, CRT-D non-responders display poor outcomes, mainly related to a progressive ventricular dysfunction with an increased risk of worse clinical outcomes (16,17).

In this scenario, TM, which has just been successfully utilised in heart failure patients treated by ICD and/or CRT-D, may represent a helpful tool to improve clinical outcomes and CRT-D response. To our knowledge there are no clinical studies focused on TM impact in a homogenous population of heart failure patients (chronic heart failure patients with left bundle branch block, and left ventricle ejection fraction reduction (LVEF) < 35%, in NYHA class II/III) treated by CRT-D. Herein, we have investigated in a randomised, multicentre, prospective study conducted on a population of heart failure...
patients treated by CRT-D, the impact of CRT on primary and secondary clinical study endpoints.

Methods

In a randomised, multicentre study conducted in different Italian centres (Catholic University of Sacred Heart, Campobasso; ‘John Paul II’ Research and Care Foundation, Campobasso; Second University of Naples, Naples) between September 2010 and September 2014 (follow-up has been closed in June 2015), we have enrolled patients with standard indications (18) for a CRT-D implant, enabled with [Biotronik (Berlin, Germany) CRT-D models, Lumax 640HF, Iforia 3HF, Iforia 5HF] or without TM technology [Inogen CRT-D, Incepta CRT-D; Boston Scientific (Marlborough, MA, USA), Unify Assura CRT-D; St Jude Medical (St. Paul, MN, USA), Brava CRT-D; Medtronic (Minneapolis, MN, USA)].

Informed consent was obtained from all participants included in the study. All procedures were in accordance with the ethical standards of the institutional and/or national research committee and with the Helsinki declaration. Enrolled patients had chronic heart failure lasting for at least 3 months, New York Heart Association (NYHA) functional class II or III, left bundle branch block, severe left ventricle ejection fraction reduction (LVEF < 35%) and an indication for CRT-D treatment according to the American College of Cardiology/American Heart Association guidelines (18). Exclusion criteria were as follows: age < 18 or > 75 years, ejection fraction > 35%, previous internal ICD, CRT-D and/or pacemaker implant, prior cardiac surgery, absence of informed written consent and any condition that would make survival for 1 year unlikely. All patients were informed of the nature of the study and provided written consent. Screened patients to receive a CRT-D were defined according to the American College of Cardiology/American Heart Association guidelines for the management of patients with heart failure refractory to a maximal medical therapy (18). After screening phase, 196 patients have been enrolled in the study (these patients met criteria reported above). This population of patients receiving CRT-D has been randomly divided in TM group and traditional monitoring (control) group. Baseline parameters have been determined before interventions and the follow-up has been concluded 12 months after CRT-D implant. Responders to CRT-D treatment have been defined as previously described (16).

Interventions

All patients identified to be treated with CRT-D (heart failure, NYHA class 2–3, LVEF < 35% and left bundle branch block) were randomly assigned (1 : 1) to receive TM (TM group) in addition to standard care (traditional ambulatory clinical and instrumental assessment), or to standard care without TM (control group) for 12 months. The random allocation sequence with variable and randomised block size (sizes four) was computer-generated and concealed from the sites. A small portable patient device receives the data and relays them automatically over mobile phone links to the Home Monitoring Service Center. Data were processed automatically and posted on a server. In the TM group, transmitted data were reviewed by independent investigators according to their clinical routine. In parallel, transmitted data were reviewed by a central monitoring unit composed of trained study nurses and supporting physicians, located at the Giovanni Paolo II Research and Care Foundation, Campobasso (Italy).

The role of this unit was to ensure the awareness of investigational sites to predefined medical events including ventricular and atrial tachyarrhythmia episodes, low percentage of biventricular pacing, increase in the frequency of ventricular extrasystoles, decreased patient activity and abnormal intracardiac electrogram, as described in previous reports (19). On working days, the central monitoring unit redundantly forwarded these events and standard technical safety notifications issued by the TM system to investigational sites. The investigational site had to confirm receipt of the reports within 48 h. A clinical response to TM observations was done at the discretion of investigators. When contacting patients on the basis of TM data, the investigators did a standard (prespecified in the protocol) telephone interview to establish whether the patient’s overall condition or dyspnoea had worsened, whether the patient was regularly taking prescribed drugs, and whether the patient’s weight had increased by more than 2 kg over the preceding 3 days, followed in any case by a clinical examination. The investigators reported the additional clinical follow-up and whether a visit to the family doctor was recommended. In the control group, no study participant had access to TM data, but followed regularly in outpatient clinic ambulatory follow-up visits. All patients were treated according to international guidelines (18).

Surgical procedure

The left ventricle lead was inserted transvenously via the subclavian route. A coronary sinus venogram was obtained using a balloon catheter, and the left ventricle pacing lead was inserted through the coronary sinus with the help of an 8-F or 9-F guiding catheter and positioned as far as possible in the venous system, preferably in the lateral or posterolateral vein.
The atrial and right ventricular leads were placed in the right atrial appendage and the right ventricular apex, respectively. All leads were connected to a dual-chamber biventricular implantable cardiac device, with defibrillator function (CRT-D). The atrioventricular interval was optimised by Ritter's method with transthoracic echocardiography, as previously described (16).

Patients monitoring
Patients were scheduled for in office follow-up visits 10 days after clinical discharge and after 1, 3, 6 and 12 months by the treating physician (TM and control group), and every patient was under continuous, automatic remote monitoring during the entire study (TM group). The frequency of TM data analysis and the response to TM alerts was left to the investigator’s discretion.

The study has been conducted in accordance with the Declaration of Helsinki. The protocol has been approved by the Ethics Committees of participating Institutions. All the patients gave their written informed consent to participate in the trial.

Data collection and use
All data have been collected at admission visits, follow-up visits and clinical database and during TM (TM group) and clinical examination (control and TM group) follow-up. Clinical evaluation included physical examination, vital signs, review of adverse events, fasting venous blood withdrawal (at least 12 h from last meal) have been performed for glycaemia and lipid profile at every visit. Follow-up visits have been scheduled 10 days after hospital discharge and at 1, 3, 6 and 12 months by the treating physician (the 12th month visit was conducted at the end of follow-up). At each clinical follow-up, NYHA classification was re-assessed and patients graded their overall condition as unchanged or slightly, moderately, or markedly worsened, or improved since randomisation by global self-assessment. All patients have been instructed to regularly assess body weight, occurrence of dyspnoea and any clinical symptom. At each visit, patients have been asked whether medical events or symptoms suggestive of cardiac arrhythmias occurred; moreover, both ECG and ECG Holter-monitoring have been performed to detect the presence of asymptomatic arrhythmias (16).

Endpoints

Primary endpoints
The primary endpoints were all cause death, cardiac death and hospital admission for heart failure (hospitalisation for objective worsening evidence of change in clinical status, NYHA functional class changing, patient’s symptoms and quality of life, or moderately to markedly worse self-reported overall condition compared with at randomisation). Heart failure worsening has been also reported as unplanned overnight admission to hospital, worse NYHA functional class, or had moderately to markedly worse self-reported overall condition compared with at randomisation, as described in previous studies (7,8,20).

Secondary endpoints
Atrial fibrillation (AF) sustained episodes, non-sustained and self terminated ventricular tachyarrhythmia (VTI), sustained VT and ventricular fibrillation (VF), ICD shocks, percentage of CRT-D responder patients. The determination of endpoints was adjudicated by an independent clinical committee, according to criteria prespecified in the protocol.

Statistical analyses
Continuous data, non-normally distributed, has been compared with the Mann–Whitney–Wilcoxon rank sum test. We compared categorical data, including the primary endpoint with the exact Pearson’s χ² test. Cox regression models were used to calculate hazard ratios. We considered a two-sided p value of less than 0.05 as statistically significant. Sample size was calculated using a power of 80% and confidence of 95%. The analysis was performed by using SPSS version 21 (Chicago, IL, USA).

Results
We included in the study 196 eligible patients; 191 received a CRT – subdivided in TM treatment and traditional CRT-D ambulatory monitoring (Figure 1); 183 patients terminated the study follow-up (94 patients in control group, no TM, and 89 in TM group).

Mean population age was 72.2 ± 7.2. The clinical characteristics at enrolment were similar and balanced between two groups, as shown in detail in Table 1. The study population was represented by chronic heart failure patients, in maximal pharmacological treatment, receiving CRT-D. No significant difference was observed when comparing pharmacological treatment between the two groups (Table 2). At 1-year follow-up primary and secondary study endpoints have been examined, comparing TM to control group (Table 3). We evaluated all different parameters revealed by TM or by traditional visits, to differentiate CRT-D responders from non-responders and to study the primary and the secondary study endpoints. The patients have been then divided
in CRT-D responders and CRT-D non-responders, as indicated by clinical characteristics and response during follow-up to the CRT-D using criteria previously described (16). At 1-year follow-up 26 patients were in persistent AF (7 patients in TM group vs. 19 patients in control group, p = 0.048). There was a significant difference in hospitalisation events (15.7 vs. 28.7, p = 0.02) comparing TM patients to control group. There was no significant difference when considering all cause mortality (7.9 vs. 8.5, p = 0.54) or cardiac death events (3.4 vs. 5.3, p = 0.39), comparing TM to non-TM patients. We also detected no

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**Table 1** Baseline parameters of the study population

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<thead>
<tr>
<th></th>
<th>Total</th>
<th>TM</th>
<th>non-TM</th>
<th>p</th>
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<tbody>
<tr>
<td>Number of patients</td>
<td>183</td>
<td>89</td>
<td>94</td>
<td></td>
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<tr>
<td>Age</td>
<td>72.2 ± 7.2</td>
<td>71.8 ± 8.5</td>
<td>72.6 ± 5.7</td>
<td>0.43</td>
</tr>
<tr>
<td>Male gender, n (%)</td>
<td>139 (75.9)</td>
<td>64 (71.9)</td>
<td>75 (79.8)</td>
<td>0.23</td>
</tr>
<tr>
<td>Hypertension, n (%)</td>
<td>109 (59.5)</td>
<td>52 (58.4)</td>
<td>57 (60.6)</td>
<td>0.46</td>
</tr>
<tr>
<td>Hypercholesterolaemia, n (%)</td>
<td>106 (57.9)</td>
<td>61 (68.5)</td>
<td>45 (47.9)</td>
<td>0.07</td>
</tr>
<tr>
<td>Diabetes, n (%)</td>
<td>110 (60.1)</td>
<td>53 (59.6)</td>
<td>57 (60.6)</td>
<td>0.51</td>
</tr>
<tr>
<td>Glucose (mg/dl)</td>
<td>136.4 ± 44.8</td>
<td>141.1 ± 48</td>
<td>132.6 ± 41</td>
<td>0.17</td>
</tr>
<tr>
<td>Creatinine</td>
<td>1.23 ± 0.48</td>
<td>1.23 ± 0.48</td>
<td>1.22 ± 0.36</td>
<td>0.46</td>
</tr>
<tr>
<td>NYHA (II/III)</td>
<td>83/100</td>
<td>37/52</td>
<td>46/48</td>
<td>0.19</td>
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Data are expressed as mean ± standard deviation. HF, heart failure; NYHA, New York Heart Association; TM, telemonitoring.
significant differences when examining responder percentage, stroke events and number of sustained VT/VF episodes or ICD shocks events. Notably, at 1-year follow-up seven patients in TM group vs. 17 patients in control group reported ventricular non-sustained tachyarrhythmias events (VTt) (p = 0.04).

When considering secondary endpoints, we did not observe a significant difference in ICD shock events (event numbers 10 vs. 16, p = 0.208), CRT-D responder patients percentage [n = 60 (67.4%) vs. 59 (62.8%), p = 0.31], stroke events (n = 3 vs. 4, p = 0.549) and VT/VF events (n = 18 vs. 23, p = 0.35) comparing TM to non-TM patients. A significant difference has been found when examining sustained AF episodes (events 7 vs. 19, p = 0.048) and non-sustained VT episodes (VTt episodes 7 vs. 17, p value 0.04), comparing TM to non-TM patients.

Then, we evaluated the relative benefits of TM in CRT-D responders and non-responders by univariate analysis of factors predicting heart failure hospitalisation (Table 4). Strikingly, at multivariate analysis of factors predicting heart failure hospitalisation (Table 5), TM is the only factor predicting heart failure hospitalisation [hazard ratio (HR) 0.6, 0.42–0.79, 95% CI, p value 0.002].

### Discussion

To our knowledge, this is the first study reporting TM as a powerful diagnostic tool that can independently predict heart failure hospitalisation in patients
Home Monitoring in Heart Failure: the TELECART study

The results of the IN-TIME (19) study, with a 1-year cardiovascular mortality of ~2.7% in the TM group vs. 6.8% in the control group (log-rank p = 0.012; HR 0.37, 95% CI 0.16–0.83), the Authors concluded that TM may reduce the percentage of cardiovascular mortality as compared with non-TM patients, treated by ICD/CRT-D (19). This discrepancy might be because of the differences in CRT-D non-responders percentage between TM and non-TM groups, which may impact heart failure progression (26). In our population we have similar CRT-D response percentages comparing TM vs. non-TM, and TM does not ameliorate the CRT-D response percentage.

A significant difference between TM and non-TM patients (15.7% vs. 28.7%) has been observed when examining the hospital admission for heart failure disease progression (objective worsening evidence of change in clinical status, NYHA functional class changing, patient’s symptoms and quality of life, or moderately to markedly worse self-reported overall condition) (16). Thus, TM does not reduce overall cause mortality and cardiac death, in CRT-D recipients but may change the clinical course of disease progression. Continuous monitoring and data collection, interpretation and alarm settings may help the clinicians in immediate therapy management and treatment.

<table>
<thead>
<tr>
<th>Table 5 Multivariate analysis of factors predicting hospitalisation</th>
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<td><strong>HR</strong></td>
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<td>---------------------------------</td>
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<tr>
<td>TM vs non-TM</td>
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<tr>
<td>Age (year)</td>
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<tr>
<td>Hypercholesterolaemia (&lt; vs &gt;)</td>
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<tr>
<td>Chronic kidney disease (&lt; vs &gt;)</td>
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<tr>
<td>NYHA class (class)</td>
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<td>LV ejection fraction (%)</td>
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</table>

After testing for collinearity, variables with p < 0.10 on univariate analysis were included in the multivariate analysis. Data are expressed as mean ± standard deviation. NYHA, New York Heart Association; LV, left ventricle; TM, telemonitoring; HR, hazard ratio. CI, confidence interval. * p < 0.05.
adjustments to have the better CRT-D response. We could consider to use TM to achieve a better clinical control of CRT-D recipients exploiting common monitoring mechanisms, including low percentage of biventricular pacing, early detection of the onset or progression of arrhythmias, number of device interventions, early recognition of leads and device dysfunction. Continuous monitoring may be reflected in a better patient therapeutic management and lower hospital admission, as compared with periodic outpatient follow-up.

We did not detect significant differences when examining stroke events, number of sustained VT/VF episodes or ICD shocks events. A significant difference comparing TM to non-TM patients has been found when examining sustained AF and non-sustained VT episodes. This effect may be in part attributable to a higher prevalence of immediate anti-arrhythmic treatment to restore sinus rhythm in TM patients as compared with non-TM (27).

In our study, LVEF and NYHA functional class are predictive factors of hospitalisation. In line with this observation, LVEF has been classified as an independent factor for heart failure worse prognosis in the general population and in CRT-D recipients (26). Indeed, LVEF improvement is an index to define CRT-D-positive response, and NYHA functional class is an indicator of clinical status and outcomes (28). Hence, NYHA class worsening is linked to increasing fatigue, dyspnoea and other symptoms of failing heart, and may be regulated in positive manner by pharmacological and electrical therapy in responders (26).

A clinically relevant result emerging from our data analysis is that TM is predictive of heart failure hospitalisation rate. CRT-D recipients are hospitalised for dyspnoea worsening, for an increasing weight and for all symptoms related to heart failure disease worsening. TM may represent a useful monitoring system to follow heart failure CRT-D recipients, who may receive an appropriate, safe to use for physicians and patients (8), continuous follow-up monitoring and consequently the care that they need.

The main limitation of this study is the small size of our population of patients treated by CRT-D, monitored or not by TM, also attributable to loss of patients during follow-up, and to the low adherence of patients to the study protocol, as mentioned in the results section. Unfortunately, we do not have sufficient data on sympathetic nervous activity, which plays a crucial role in the pathophysiology of heart failure (29–34) and can be modulated by CRT (35). Besides, our conclusions remain linked to the relatively short follow-up duration and should not be extrapolated to long-term clinical outcomes.

Acknowledgement

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References

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