Clinical utility of cardiac innervation imaging in patients with heart failure

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WHAT IS THE PROBLEM?

Sudden cardiac death is an important problem in clinical cardiology. In the USA; the annual incidence of sudden cardiac death is estimated at 110.8 individuals per 100,000 population per year.¹ In the year 2015, it was estimated that 356,461 patients died suddenly of underlying cardiac disease.¹ Moreover, in patients with successful resuscitation, the recurrence rate of sudden death is high.

Particularly, patients with previous myocardial infarction and reduced left ventricular (LV) function are at elevated risk of sudden death.² With the introduction of the implantable cardioverter defibrillator (ICD), a therapy has become available that has been demonstrated to prevent sudden cardiac death in patients with ischemic and non-ischemic cardiomyopathy.

In the Multicenter Automatic Defibrillator Implantation Trial-I (MADIT-I) study, 196 patients with a history of myocardial infarction (≥ 3 weeks), reduced LV function ($\text{LVEF} \leq 35\%$) and documented episode of ventricular tachyarrhythmias, were randomized to either prophylactic ICD implantation ($n = 95$) or standard medical therapy ($n = 101$).² Over a 27-month follow-up period, 15 patients (15.8%; 11 cardiac death) died in the defibrillator group as compared to 39 patients (38.2%; 27 cardiac death) in the medical group (hazard ratio for overall mortality, 0.46; 95% confidence interval, 0.26 to 0.82; $P = 0.009$). Importantly, the use of amiodarone or beta-blockers did not influence these observations.

In a subsequent study (MADIT-II), 1232 patients with previous infarction (≥ 1 month) and $\text{LVEF} \leq 30\%$ were randomized in a 3:2 ratio to receive an ICD or medical therapy.³ During a 20-month follow-up period,

mortality was 19.8% in the patients receiving medical therapy versus 14.2% in patients receiving an ICD (hazard ratio for the risk of all-cause mortality, 0.69 (95% confidence interval, 0.51 to 0.93; $P = 0.016$). Importantly, age, gender, LVEF, New York Heart Association class, and QRS duration did not influence the results.

Both these studies paved the way for the European Society of Cardiology (ESC) to indicate in their 2015 guidelines that an ICD is indicated in patients with symptomatic heart failure (NYHA class II-III) and reduced LVEF ($\leq 35\%$)⁴ (class IA in ischemic heart disease, and class IB in non-ischemic cardiomyopathy). Similar recommendations have been issued by the ACC/AHA guidelines.⁵

It has also become evident from the MADIT-II trial that a limited number of patients received appropriate ICD therapy during an average of 21 months of follow-up.⁶ More specifically, of the 720 patients receiving an ICD in the MADIT-II trial, 169 (23.5%) patients received antiarrhythmic therapy (anti-tachycardia pacing in 139 patients, and defibrillator shocks in 30) and 551 (76.5%) patients did not receive ICD therapy.

More recently, Kober et al.⁷ demonstrated that in patients with non-ischemic cardiomyopathy, prophylactic ICD implantation did not improve long-term outcome as compared to usual clinical care. Specifically, 1116 patients with non-ischemic cardiomyopathy ($\text{LVEF} \leq 35\%$) were randomized to ICD implantation ($n = 556$) or usual care ($n = 560$); all-cause death was 21.6% in the ICD group versus 23.4% in the control group (hazard ratio, 0.87; 95% confidence interval [CI], 0.68 to 1.12; $P = 0.28$). Still, sudden cardiac death was more frequent in the usual care group (8.2%) versus the ICD group (4.3%) (hazard ratio, 0.50; 95% CI, 0.31 to 0.82; $P = 0.005$).

Moreover, ICD implantation is associated with procedural complications, device/lead infections, and inappropriate shocks.⁸ Thus, there is significant benefit from ICDs to prevent sudden cardiac death, but the criterium of $\text{LVEF} \leq 35\%$ may be suboptimal for

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selecting the patients who may benefit most from ICD implantation, and a more personalized risk stratification may be of potential use.⁹

Various ECG-based criteria have been proposed, including microvolt T-wave alternans, signal-averaged ECG, QRS fragmentation, and various measures of autonomic (dys-)function such as heart rate variability.¹⁰ In addition, different imaging techniques have attempted to better identify the substrate underlying sudden cardiac death, specifically advanced echocardiographic and CMR techniques, as well as nuclear imaging with PET and SPECT. The different imaging techniques reflect different aspects of the LV substrate underlying the lethal ventricular arrhythmias; the anatomical substrate is probably the presence of central homogenous scar tissue (secondary to myocardial infarction), with a peripheral, heterogeneous border zone with fibrosis, viable but jeopardized myocardium and normal tissue. Contrast-enhanced CMR is focused on anatomical imaging (specifically scar and fibrosis assessment), whereas echocardiography with strain reflects functional imaging (LV deformation assessment) and nuclear imaging with PET and SPECT addresses biological imaging (specifically innervation and denervation assessment).⁹ Over the recent years, the use of cardiac innervation imaging with ¹²³I-metaiodobenzylguanidine (MIBG) and planar imaging or SPECT has been evaluated for risk stratification of heart failure patients at risk of sudden cardiac death. In the current issue of the Journal, two articles are published regarding the pros and cons of MIBG imaging, mostly focusing on risk stratification in patients with heart failure and elevated risk of sudden cardiac death.^{11,12}

MIBG FOR RISK STRATIFICATION: THE EVIDENCE OF NEURO-CARDIAC IMAGING

The first article, by Travin, addresses the pros of MIBG imaging.¹¹ He introduces the concept of the heart-to-mediastinum (H/M) ratio to quantify the tracer uptake in the heart as compared to the mediastinum on planar images, reflecting cardiac innervation and denervation. Many smaller studies have used this parameter to demonstrate the use of MIBG for risk stratification of heart failure patients. Agostini and colleagues¹³ performed a large analysis including 290 heart failure patients who previously underwent MIBG imaging (between 1993 and 2002). During a 2-year follow-up, 67 patients (26%) experienced a major cardiac event (death, cardiac transplantation, potentially lethal arrhythmias). These patients had a lower H/M ratio (1.51 ± 0.30) as compared with the patients without events (1.97 ± 0.54 , $P < 0.001$).

Dr Travin then referred to the ADMIRE-HF (AdreView Myocardial Imaging for Risk Evaluation in Heart

Failure) trial and its sub-studies.¹⁴ The ADMIRE-HF study included 961 heart failure patients (NYHA II-III) and LVEF $\leq 35\%$ who underwent MIBG imaging, and the H/M ratio was assessed from the 4-hour delayed image; a value of 1.60 was used as the normal value. Patients were followed up for 2 years and the end-points included time to first occurrence of NYHA functional class progression, potentially life-threatening arrhythmic event, or cardiac death. During a median follow-up, 237 patients (25%) experienced events; the 2-year event rate was 15% for H/M ≥ 1.60 and 37% for H/M < 1.60 , and the H/M ratio was predictive of all end-points. On multivariate analysis, H/M ratio, LVEF, B-type natriuretic peptide, and NYHA class were predictive of outcome. Addition of the H/M ratio enabled further stratification in patients with high B-type natriuretic peptide (140 ng/l) and low LVEF ($\leq 30\%$).

More recently, a pooled analysis from Japan was reported, including 6 cohorts with 1322 heart failure patients, with a follow-up of 78 months, and primary outcome being all-cause mortality.¹⁵ The mortality was 5.6% at 1 year, and 19.7% at 5-year follow-up. On the multivariate analysis, age, NYHA class, LVEF and the late MIBG H/M ratio were predictive of outcome.

All these data have built a strong case supporting the use of MIBG for risk stratification in heart failure patients, and have shown the incremental value of the H/M ratio over the routine risk markers. One important issue is the predictive value. This remains to be further explored in future studies. As with any technique, a high negative predictive value is important: “to rule out.” This could imply not providing a certain therapy or less close monitoring. In the ADMIRE-HF study, the all-cause mortality was 1% (2 patients of 201) in the patients with an H/M ratio ≥ 1.6 . Moreover, in the high-risk groups (according to BNP > 140 ng/l and LVEF $\leq 30\%$) with a H/M ratio ≥ 1.6 there were no cardiac deaths. These data are supportive of the conception that MIBG imaging could serve as gate-keeper for ICD implantation.

WHAT ARE THE UNCERTAINTIES

Drs Liga and Scholte discuss the limitations of MIBG imaging hampering the widespread acceptance of this technique in clinical practice.¹² The first issue discussed by these authors concerns the need for a standardized protocol. The main parameter used in the clinical setting is the H/M ratio, which reflects cardiac innervation and denervation. At present, an early (15 min after tracer injection) and late MIBG scan (4 hour after tracer injection) are obtained, and the H/M ratio can be derived from both scans. It is currently not clear which scan provides the optimal prognostic

information. From a practical point of view, it would be ideal to use the scan obtained within the first hour after tracer injection. Finally, the cut-off value of 1.6 needs further, prospective evaluation.

Other parameters from planar MIBG imaging have also been used, such as the washout rate, and these are less well validated as compared to the H/M ratio.

Moreover, with planar imaging, the global sympathetic innervation of the heart can be evaluated, but regional information may be more important. SPECT provides information on regional variations in sympathetic innervation, but regional physiological variations also occur, with less tracer uptake in the inferior wall in normal individuals. Also, in patients with severe heart failure, tracer uptake may be low resulting in poor SPECT image quality, and in general, data acquisition with SPECT is more time-consuming than planar imaging.

WHAT EVIDENCE IS MISSING AND NEEDED?

Prospective, randomized controlled trials will be needed to establish the role of MIBG in risk stratification of heart failure patients. From a clinical point of view, the ADMIRE-HF study provided strong prognostic data in a prospective cohort of heart failure patients, but an important question remains whether MIBG could direct therapy in these patients and specifically guide ICD implantation. As outlined above, the current selection criterium of LVEF $\leq 35\%$ does not optimally include and exclude patients who need an ICD for primary prevention in ischemic and non-ischemic cardiomyopathy. As Dr Travin points out, this trial is planned, including >2,000 heart failure patients (LVEF between 30% and 35%, and NYHA class II-III) who will receive MIBG-guided ICD implantation or guideline-directed ICD implantation; this trial will compare the outcomes (primary outcome all-cause mortality) over 2.75 to 3 years, but will only be completed in 2019.¹⁶ However, Dr Travin also points out, this trial will take time to be completed, but there is currently significant evidence that MIBG contributes to risk stratification in heart failure patients, probably more evidence than has been obtained with other imaging techniques that are used in clinical practice.

Disclosure

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