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Late gadolinium enhancement in cardiac sarcoidosis predicts ICD implantation and appropriate discharge

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Introduction

Cardiac sarcoidosis is associated with sudden arrhythmic death, conduction abnormalities and heart failure. Occult disease is frequently found at autopsy although its significance remains unclear. Late gadolinium enhancement (LGE) on cardiac magnetic resonance (CMR) imaging permits detection of granulomatous cardiac infiltration and myocardial fibrosis.

Purpose

We sought to establish whether CMR diagnosis of cardiac sarcoidosis correlates with clinical outcome in this complex disease.

Methods

96 consecutive patients with sarcoidosis were referred to our centre for CMR between 2002 and 2007 (46 M, 50 F; mean age 57 years). Outcomes in two cohorts distinguished by evidence of LGE were retrospectively assessed (mean follow-up 3.5 years). Left ventricular (LV) volumes, ejection fraction (EF) and mass were evaluated using semi-automated analysis software and indexed to body surface area. Continuous variables were reported as mean ± standard deviation; groups (based on the presence or absence of LGE) were compared using a t-test or Fisher's exact test.

Results

LGE was observed in 31 out of 96 (32%) patients. No significant difference was seen in all-cause death or hospitalisation for a cardiovascular event between the LGE+ and LGE- cohort. The LGE+ cohort had a significantly greater rate of ICD implantation (7 vs 1, p < 0.0005) and appropriate ICD discharge (4 vs 0, p < 0.005). Notably, only four patients had right ventricular LGE+; all required defibrillator (ICD) implantation and sustained appropriate ICD discharge. There was no difference in other secondary outcomes such as pre-ICD implantation sustained ventricular tachycardia, pacemaker implantation rate or escalation of heart failure therapy. Amongst LGE+ individuals, there was a significant reduction in biventricular EF (LVEF $50 \pm 13.0\%$ vs. $64 \pm 10\%$, p < 0.05; RVEF 49 ± 13 vs. $58 \pm$ 9%, p < 0.05). RVEDV was also significantly greater (173 \pm 14 vs.137 \pm 5 mls, p < 0.0065), although not LVEDV $(105 \pm 5.9 \text{ vs. } 132 \pm 7.9 \text{ mls, p} < 0.0066)$. The distribution of LGE did not appear to correlate with outcome. The two groups did not differ in the incidence of biopsy-proven sarcoidosis (48/96 cases), pre-referral cardiac symptoms or duration of follow-up. Within this cohort, only five patients had STIR imaging clearly suggestive of active cardiac sarcoidosis.

Conclusion

In this cohort, presence of cardiac LGE in sarcoidosis is associated with increased ICD implantation and appro-

priate discharge but not worse mortality or hospitalization outcome, despite an associated biventricular reduction in ejection fraction. RV LGE may predict a requirement for ICD implantation. Further prospective research is needed to improve understanding of this complex disease.

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