

WALKING POSTER PRESENTATION

Open Access

Clinical impact of cardiovascular magnetic resonance in evaluation for possible arrhythmogenic right ventricular dysplasia/cardiomyopathy

Andrew D Choi^{1,2*}, Sujata M Shanbhag¹, Peter Kellman¹, Christine Mancini¹, Marcus Y Chen¹, Andrew E Arai¹, Patricia W Bandettini¹

From 18th Annual SCMR Scientific Sessions
Nice, France. 4-7 February 2015

Background

This study examined the impact of CMR on clinical management in patients with undergoing evaluation for arrhythmogenic right ventricular dysplasia/cardiomyopathy (ARVD/C).

Methods

Patients referred for assessment of ARVD/C were evaluated. Using 2010 ARVD/C Task Force criteria, clinical history, family history, ECG and other test results were evaluated with and without CMR findings to determine definite, borderline or possible ARVD/C. CMR included

assessment of right ventricular (RV) size, function, and regional wall motion (RWM). For alternative diagnoses, tissue characterization and late gadolinium enhancement were routinely performed. Qp:Qs was performed when intracardiac shunt was suspected by the supervising physician.

Results

311 consecutive patients (mean age 45 ± 14 years, 53% male) were included. Prior to CMR, patients were classified as definite (n=1), borderline (n=1) or possible ARVD (n=18, Table). After CMR, 6(2%) were diagnosed with

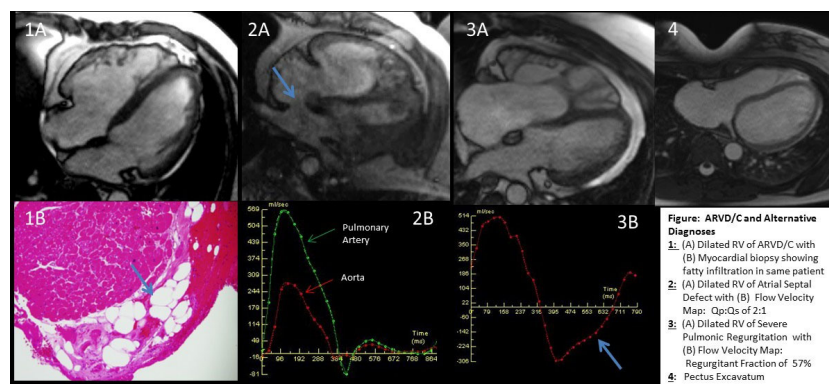


Figure 1

¹Advanced Cardiovascular Imaging Laboratory, National Heart, Lung and Blood Institute, National Institutes of Health, Bethesda, MD, USA
Full list of author information is available at the end of the article

Table 1 Clinical Impact of CMR on Diagnosis of ARVD/C vs. Alternative Diagnoses

n=311	2010 Guidelines without CMR Findings, n (%)	2010 Guidelines with CMR Findings, n (%)
Definite Criteria for ARVD/C	1 (0.3)	6 (2)
Borderline Criteria for ARVD/C	1 (0.3)	5 (2)
Possible Criteria for ARVD/C	18 (5.8)	9 (3)
Patients with 1 or no Minor Criteria, not meeting 2010 Guidelines Definition of "Definite", "Borderline" or "Possible"	291 (93.6)	51 (16) Alternate Diagnosis* 76 (24) RV Enlargement Alone** 164 (53) Normal RV***

* Alternative Diagnosis Resulting in Change in Management included 6 (1.9%) Intracardiac shunts, other Cardiomyopathy or RV Overload State 36 (11.5%), or Other Diagnosis 9 (2.8%)

**RV Enlargement Alone with normal RV Function and Regional Wall Motion

*** Normal RV Function, Size and Regional Wall Motion by CMR

definite ARVD/C and underwent defibrillator implantation, 5(2%) were classified as borderline ARVD/C, and 9 (3%) remained possible ARVD/C. 51(16%) had alternative diagnoses (Figure/Table), resulting in a management change: 6(1.9%) patients had intracardiac shunt, 36(11.5%) had another cardiomyopathy or RV overload state, and 9(2.8%) had other diagnoses. 76(24%) had RV enlargement alone with normal RV function and absent RWM by CMR while 164(53%) without other major criteria had normal RV function, size, and RWM.

Conclusions

CMR impacted clinical management by contributing to the diagnosis of definite or borderline ARVD/C in 4% of patients and by excluding the presence of significant RV dysfunction, enlargement, and RWM in over half of patients. CMR identified important alternative diagnoses in 16%.

Funding

No conflicts of interest to disclose.

Authors' details

¹Advanced Cardiovascular Imaging Laboratory, National Heart, Lung and Blood Institute, National Institutes of Health, Bethesda, MD, USA. ²Division of Cardiology, Medstar Washington Hospital Center, Washington, DC, USA.

Published: 3 February 2015

doi:10.1186/1532-429X-17-S1-Q53

Cite this article as: Choi et al.: Clinical impact of cardiovascular magnetic resonance in evaluation for possible arrhythmogenic right ventricular dysplasia/cardiomyopathy. *Journal of Cardiovascular Magnetic Resonance* 2015 **17**(Suppl 1):Q53.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

