The role of cardiac magnetic resonance (CMR) in the diagnosis of cardiomyopathy: A systematic review

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Abstract

Background
Myocardial pathologies are significant causes of morbidity and mortality in patients worldwide. Ischemic and non-ischemic cardiomyopathies have become a worldwide epidemic of the 21st century with an increasing impact on health care systems. The 2012 European Society of Cardiology and 2013 American College of Cardiology Foundation/American Heart Association guidelines provide current therapy guidance to reduce mortality and morbidity.

Methods
This was a systematic review involving cardiac magnetic resonance (CMR) studies for the diagnosis of cardiomyopathy from January 2013 to April 2017. Out of 62 reviewed studies, only 12 were included in our study.

Results
The average sensitivity and specificity of CMR in the diagnosis of cardiomyopathy was 86.75% (95% confidence interval [CI], 70.30% to 92.58%) and 81.75% (95% CI, 73.0% to 87.6%), respectively, and the positive predictive and negative predictive values were 80.17% and 86.75%, respectively.

Conclusion
Despite some limitations, our study shows that CMR has high sensitivity, specificity, and positive predictive value in diagnosing different types of cardiomyopathy. CMR may be used to differentiate types of cardiomyopathy, accurately quantify the chamber dimensions, volumes, and cardiac function, which make it useful for prognosis as well.

Keywords: Cardiac magnetic resonance, dilated cardiomyopathy, hypertrophic cardiomyopathy, ischemic cardiomyopathy, late gadolinium enhancement

Introduction
Myocardial pathologies are significant causes of morbidity and mortality in patients worldwide. Ischemic and non-ischemic cardiomyopathies have become a worldwide epidemic of the 21st century with an increasing impact on health care systems. The 2012 European Society of Cardiology and 2013 American College of Cardiology Foundation/American Heart Association guidelines have served a purpose as another imaging tool.

In ischemic cardiomyopathy, the sub-endocardium is always enhanced on DE-MRI, while a patchy mid-myocardial enhancement is observed in dilated cardiomyopathy (DCM). Furthermore, patients with restrictive cardiomyopathy showed delayed myocardial improvement over the entire sub-endocardial circumference. Heart failure represents the final stage in the continuum of cardiovascular diseases. Cardiac remodelling is a key component of heart failure that progresses from adaptive to maladaptive as the disorder worsens. Increased myocardial wall stress during diastole contributes to the development and progression of adverse cardiac remodeling. Cardiac magnetic resonance is superior to other cardiac imaging modalities such as echocardiography, computed tomography angiography and coronary angiography in determining the type of cardiomyopathy, and cardiac function.

Methods
Patient characteristics
The age of participants in the included 12 cardiomyopathy studies ranged from 18 to 87 years, with dilated cardiomyopathy being a more seen pathology in the studies followed by ischemic cardiomyopathy. The patient characteristics are summarized in Table 1.

Identification of studies and journals
We identified published studies using CMR in the diagnosis of different types of cardiomyopathy in original and review articles by systematic searches of PubMed, MedLine, Cochrane database and Embase, and by manual searches of listed references in the papers, we found. We limited our search to studies published from January 2013 to April 2017 because, during this period, we noted an increase in the application of
Role of CMR in diagnosis of cardiomyopathy

The keywords used were: “dilated cardiomyopathy”, “ischemic cardiomyopathy”, “hypertrophic cardiomyopathy”, “myocarditis”, “cardiac amyloidosis”, “cardiac sarcoidosis”, and “cardiac MR”, or “LGE CMR” (late gadolinium enhancement cardiovascular magnetic resonance), “Cine CMR”, and “sensitivity” or “specificity”. We identified 62 studies through this search strategy. We further screened the reference list of the retrieved studies for any additional publications. There were no restrictions on studies based on their sample size.

Eligibility criteria

We considered all eligible studies that evaluated the role of CMR in the diagnosis of cardiomyopathy. We also included studies with sufficient information to allow the calculation of sensitivity and specificity. We excluded meeting reports, abstracts, and reviews whose final stories were unavailable.

Data extraction

For each eligible study, we extracted the following information: author names, journal, year of publication, number of enrolled patients, the age of study patients, study design, and CMR protocol.

Statistical analysis

Statistical analysis was done using MedCalc for Windows version 64 bits; we calculated the sensitivity, specificity, positive predictive value, negative predictive value, positive likely-hood ratio, and negative likelihood ratio.

Results

Eligible studies

The search yielded 35 relevant studies. Of these, due to limited data, 23 studies were excluded, and 12 studies were available for systematic review. Figure 1 summarizes the flow diagram of how eligible studies were obtained. These studies were selected because they related to our analysis, they were recent, and they had likely extractable data as shown in figure 1.

Figure 1: The flow diagram of study selection process for systematic review

| Table 1: Characteristics of patients |
|------------------------------------|
| Category                            |
| Total number of patients included   |
| Number of studies reporting on sample size |
| Range of sample size reported       |
| Age of participants                 |
| Number of studies reporting on age  |
| Age range, yrs                      |
| Type of cardiomyopathy             |
| Ischemic cardiomyopathy             |
| Dilated cardiomyopathy              |
| Dual pathologies (NICM and DCM)     |
| Hypertrophic cardiomyopathy         |
| Tachycardia-induced cardiomyopathy  |
| Myocarditis                         |
| Cardiac amyloidosis                 |
| Cardiac sarcoidosis                 |
| Left ventricle non-compaction       |
| Other tests                         |
| Histopathology                      |
| Echocardiography                    |
| Coronary angiography                |

| Table 2: individual studies with sample sizes and types of cardiomyopathy |
|---------------------------------------------------------------|
| Study              | Sample size | CMR                  | Findings                                      |
|--------------------|-------------|----------------------|-----------------------------------------------|
| Goebel et al.      | 150         | T1 mapping CMR       | Does not differentiate healthy and diffusely  |
|                    | 2016        |                      | diseased myocardium                           |
| Makoto et al.      | 44          | LGE CMR              | LE in CS predominantly basal, mid septum and  |
|                    | 2016        |                      | through the LV, while in DCM, LE was         |
| Mikami et al.      | 118         | LGE CMR              | Septal fibrosis                               |
| Okada et al.       | 102         | LGE CMR              | TIC had a significant lower RVEF, and a       |
|                    | 2016        |                      | larger RVEDV and RVEDS                        |
| Kwong et al.       | 81          | Cine SSFP and LGE    | CA mean proportion of atrial enhancement     |
|                    | 2015        |                      | was significantly greater compared to SH and  |
| Maurizio et al.    | 77          | LGE CMR              | Hypertrophied septum indicating regional      |
|                    | 2015        |                      | fibrosis in HCM                              |
| Schwab et al.      | 43          | CMR                  | Wall motion abnormalities                     |
| Nguyen et al.      | 23          | CMR/LGE CMR          | Diffuse myocardial fibrosis                   |
| Dungu et al.       | 97          | LGE CMR              | Distinguished ATTR from AL cardiac amyloidosis|
| Perreira et al.    | 50          | T1 mapping CMR       | T1 mapping is a criterion for detection of    |
|                    | 2013        |                      | acute myocarditis with a higher sensitivity   |
| Choi et al.        | 114         | Cine CMR/LGE CMR     | Classification of distribution of trabeculation, |
|                    | 2016        |                      | 43.9% global type, 56.2% apical type          |
| Gulsin et al.      | 100         | Cine CMR/LGE CMR     | Global LV hypokinesis                         |

Abbreviations: NICM= nonischemic cardiomyopathy, DCM= dilated cardiomyopathy
Table 3: Showing individual studies with sensitivity, specificity and confidence intervals

| Study            | Sensitivity | 95% CI  | Specificity | 95% CI | PPV   (95% CI) | NPV   (95% CI) |
|------------------|-------------|---------|-------------|--------|----------------|----------------|
| Goebel et al.2016| 0.85        | 0.75-0.91| 0.75        | 0.66-0.83| 0.15            |                 |
| Makoto et al.2016| 0.68        | 0.57-0.78| 0.63        | 0.53-0.72| 0.32            |                 |
| Mikami et al.2016| 0.70        | 0.60-0.79| 0.71        | 0.61-0.80| 0.30            |                 |
| Okada et al.2016 | 0.88        | 0.79-0.95| 0.75        | 0.66-0.82| 0.12            |                 |
| Kwong et al.2015 | 0.93        | 0.85-0.97| 0.80        | 0.71-0.87| 0.07            |                 |
| Maurizio et al.2015| 0.99       | 0.93-1.00| 0.83        | 0.75-0.89| 0.01            |                 |
| Schwab et al.2015| 0.94        | 0.86-0.95| 1.0         | 0.92-1.00| 0.14            |                 |
| Nguyen et al.2015| 0.95        | 0.87-0.99| 0.88        | 0.80-0.93| 0.04            |                 |
| Ferreira et al.2013| 0.91       | 0.83-0.96| 0.90        | 0.82-0.95| 0.09            |                 |
| Choi et al.2016  | 0.85        | 0.76-0.92| 0.76        | 0.64-0.83| 0.15            |                 |
| Gulsin et al.2017| 0.96        | 0.90-0.99| 1.00        | 0.96-1.00| 0.04            |                 |

Table 4: Showing individual studies with positive predictive value and negative predictive value

| Study            | PPV  | NPV  |
|------------------|------|------|
| Goebel et al.2016| 0.71 | 0.87 |
| Makoto et al.2016| 0.57 | 0.73 |
| Mikami et al.2016| 0.72 | 0.69 |
| Okada et al.2016 | 0.69 | 0.91 |
| Kwong et al.2015 | 0.76 | 0.94 |
| Maurizio et al.2015| 0.80 | 0.99 |
| Schwab et al.2015| 1.00 | 0.85 |
| Nguyen et al.2015| 0.81 | 0.85 |
| Dungu et al.2015 | 0.87 | 0.96 |
| Ferreira et al.2013| 0.97 | 0.90 |
| Choi et al.2016  | 0.72 | 0.88 |
| Gulsin et al.2017| 1.00 | 0.96 |

Study description and patient characteristics

The 12 studies had a total of 999 patients. The sample size of the studies ranged from 23 to 150. The age range of the study subjects, 18 to 87 years, was reported in 10 reviews. Dilated cardiomyopathy was more prevalent in the studies, followed by ischemic cardiomyopathy and the least pervasive was cardiac sarcoidosis. The average sensitivity and specificity of CMR in the diagnosis of cardiomyopathy was 86.75% (95% confidence interval [CI], 70.30% to 92.58%) and 81.75% (95% CI, 73.0% to 87.6%), respectively. The positive predictive and negative predictive values were 80.17% and 86.75%. Tables 1, 2, 3, and 4 summarize the results.

Discussion

Cardiomyopathy has been diagnosed and assessed by echocardiogram or cardiac computed tomography for many years. With technological advancements and further research, several studies have addressed the role of cardiac magnetic resonance (CMR) as a functional modality in the diagnosis and quantification of cardiac function in different types of cardiomyopathy. CMR can measure and quantify chamber sizes and left ventricle (LV) systolic function accurately. Therefore, Cardiac Magnetic Resonance has potential as a tool to assess patient prognosis.

In our systematic review of 12 studies, we found moderately high sensitivity and specificity values for CMR, which implies that CMR is a standard valuable imaging modality for diagnosing different types of cardiomyopathy. According to this study, we appreciate the ability of CMR to diagnose and differentiate types of cardiomyopathies, through its high spatial resolution and tomographic image capabilities. Late gadolinium magnetic resonance was able to distinguish cardiac sarcoidosis and dilated cardiomyopathy. Furthermore, Late Gadolinium Enhancement cardiac magnetic resonance was also used to diagnose non-ischemic Dilated cardiomyopathy by revealing septal fibrosis and other studies identified a mid-wall septal strain pattern of Late gadolinium enhancement to be the most reliable predictor of future events. Moreover, CMR was also used to assess the diagnostic value of early right ventricular dysfunction to predict tachycardia-induced cardiomyopathy, in which the studies revealed that CMR imaging assessing right ventricular function might be valuable compared with echocardiography.

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Ischemic cardiomyopathy can easily be missed in routine screening of suspected coronary heart disease patients particularly in microvascular coronary artery disease whereby coronary angiogram may be the standard investigation expected to be done. But with the current new technology of Positron emission magnetic resonance imaging (PETMRI), it will be easy to know which areas are poorly perfused and confirmed by measuring coronary flow reserve. Among other non-invasive imaging modalities, CMR is emerging as a highly sensitive and specific test for myocardial ischemia and infarction. Resting perfusion in CMR is used to evaluate microvascular obstruction, which is shown to predict adverse left ventricular remodeling. Thus, as previous studies have indicated, the significance of magnetic resonance in diagnosing different types of cardiomyopathies especially in differentiating ischemic versus nonischemic cardiomyopathy, we concur with our results of MRI having high sensitivity and specificity.
**Clinical implications**

CMR is superior in evaluating cardiac function, LV dimensions, and capable of differentiating types of cardiomyopathies with a specificity of 81.75% and sensitivity of 86.75%. In the clinical setting, the ability to diagnose the form of cardiomyopathy helps in choosing the specific treatment. In a study of Choi et al., they used cardiac magnetic resonance imaging to establish refined diagnostic criteria for left ventricle non-compaction. As a quantitative approach, we have shown that a trabeculated left ventricle volume of >35% of the LV myocardial volume is diagnostic for left ventricle non-compaction with high specificity. Our study has also shown the diagnostic accuracy of late gadolinium-enhanced cardiac magnetic resonance for establishing the etiology of heart failure. Late gadolinium enhancement–cardiac magnetic resonance (LGE-CMR) was able to differentiate between ischemic cardiomyopathy and non-ischemic cardiomyopathy. Furthermore, the addition of adenosine stress perfusion-cardiac magnetic resonance (SP-CMR) to cine and LGE-CMR provided minimal incremental diagnostic yield for determining the etiology of heart failure in patients with severe left ventricle systolic dysfunction.

**Limitations**

The inclusion of studies with small sample sizes may influence the statistical power of the individual research and lead to imprecise and inconclusive results. Other limitations include bias through selection, publication, and verification of the studies.

**Conclusion**

Despite some limitations, our study shows that cardiac magnetic resonance (CMR) has high sensitivity, specificity, and positive predictive value in diagnosing different types of cardiomyopathies. CMR may be used to differentiate types of cardiomyopathy, accurately quantify the chamber dimensions, volumes, and cardiac function which make it useful for prognosis as well.

**Disclosure**

The authors declare no conflict of interest.

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