The evolving role of cardiac magnetic resonance imaging in the assessment of cardiovascular disease

Michael B Stokes, Nitesh Nerlekar, Stuart Moir, Karen S Teo

Background

Imaging of the heart is important in the diagnosis and follow-up of a broad range of cardiac pathology. The authors discuss the growing role of cardiac magnetic resonance imaging (CMR) in cardiology practice and its relevance to primary healthcare.

Objective

In this article we discuss the advantages of CMR over other imaging modalities, and give a brief description of the common CMR techniques and cardiac pathologies where CMR is especially useful.

Discussion

CMR provides specific advantages over other cardiac imaging modalities when evaluating pathology in congenital heart disease, cardiac masses, cardiomyopathies, and in some aspects of ischaemic and valvular heart diseases. The strength of CMR in these pathologies includes its precise anatomical delineation of structures, characterisation of myocardial tissue, and accurate, reproducible measurements of blood volume and flow. CMR is used in inpatient and outpatient settings, and is available primarily in major hospitals.

Cardiac imaging

A number of imaging modalities are used by clinicians in the assessment and care of patients with cardiovascular disease. Echocardiography is the firstline cardiac imaging investigation for many cardiac conditions. It is widely available, provides structural and functional information, and reliably reports the required clinical information in the majority of cases. However, the limitations of echocardiography include difficulty obtaining ultrasound images in specific patients and differentiating specific pathologies (eg cardiac infiltration versus hypertrophy). Delineation of the endocardial border can also be challenging with echocardiography when measuring ventricular volumes used to calculate ejection fraction.¹ Cardiac computed tomography (CT) and nuclear cardiac perfusion studies are modalities used predominantly in the investigation of ischaemic heart disease.

Cardiac magnetic resonance imaging (CMR) has grown as an imaging modality

Box 1. Advantages and disadvantages of CMR compared with imaging modalities

Advantages

- · Improved image quality
- · No ionising radiation
- Accurate ejection fraction measurement
- Investigation for suspected myocarditis that cannot be confirmed on echocardiography
- Assessment of newly diagnosed cardiomyopathy with diagnostic and prognostic advantages over echocardiography
- Assessment of myocardial viability and perfusion in work-up for coronary artery bypass surgery
- Assessment of congenital heart disease

Disadvantages

- Cost and accessibility
- Contraindicated in patients with metallic implants (eg aneurysm clips, neurostimulators, implanted pacemakers/defibrillators), metal retained fragments in eyes or gunshot injuries
- May be unsuitable for patients with claustrophobia
- A specific cut-off weight for patients varies among individual MRI scanners. Large abdominal and shoulder girths may limit the physical ability of patients to fit in an MRI scanner
- Gadolinium is contraindicated in significant renal disease (eGFR <30 mL/minute)

CMR, cardiac magnetic resonance imaging; eGFR, estimated glomerular filtration rate; MRI, magnetic resonance imaging

to provide additive and complementary information to echocardiography. It provides high-guality, cross-sectional images that enable accurate anatomical delineation. It also enables highly reproducible measurement of ventricular volumes and mvocardial mass, and quantifies flow across heart valves.1,2 CMR can assess myocardial structure and assist in the diagnosis of cardiac masses. Sophisticated electrocardiogram (ECG) gating and respiratory motion suppression methods are used to ensure the acquisition of high-quality images. CMR can overcome patient-specific limitations, such as obesity, presence of respiratory disease, or presence of soft tissue, lungs and/or scar tissue following cardiac surgery, which can limit the ability to obtain high-quality echocardiographic images.3,4

Advantages and disadvantages of CMR

Box 1 summarises the advantages and disadvantages of CMR.

There is now an increasing number of cardiac devices that are considered MRI-conditional. Patients in whom these devices have been implanted can have CMR performed with appropriate safety precautions and device checks.⁵

Techniques in CMR

A number of MRI sequences are used in CMR studies. These are tailored in each individual study for a particular pathology or the assessment of a specific aspect of cardiac function:

- still-frame images that delineate cardiac structures and great vessels in the assessment of congenital heart disease
- high-quality moving images that measure cardiac volumes and myocardial mass, and visualise regional wall motion abnormalities, which may indicate underlying ischaemic heart disease (Figure 1)
- velocity flow imaging that directly measures blood flow in assessment of valvular abnormalities, such as calculation of regurgitant volumes in aortic regurgitation⁶

- others
 - quantification of myocardial iron content: used in patients with beta-thalassemia major, where iron chelation therapy is introduced or intensified to prevent the development of heart failure⁷
 - myocardial fibrosis estimates in cardiomyopathy⁸
 - myocardial oedema detection: indicative of myocarditis or recent myocardial injury.⁹

Gadolinium use in CMR

Gadolinium is an intravenous contrast agent used in CMR. It is a useful adjunct in CMR to aid the diagnosis of specific cardiac pathologies. Delayed or late myocardial enhancement imaging (where images are acquired 10–20 minutes after gadolinium administration), allows assessment for specific patterns of uptake (Figure 2).¹⁰ Gadolinium is contraindicated in patients with significant renal disease (estimated glomerular filtration rate <30 mL/minute) as it increases the risk of developing the rare, progressive skin condition called nephrogenic systemic fibrosis.¹¹

Pathologies where CMR can be considered

Aortic and valvular disease

CMR visualises the aortic course and precisely measures aortic dimensions. It is advantageous in patients who require repeated interval imaging, avoiding the repeated exposure to ionising radiation that occurs with CT. Aortic coarctation, bicuspid aortic valve disease and aortopathies (eg Marfan syndrome) may result in progressive aortic enlargement and the severity may influence the timing of surgical intervention. Flow velocity– encoded mapping enables more accurate quantification of the severity of valvular lesions.⁶

Pericardial constriction or pericarditis

Assessment of the pericardium on CMR by measuring pericardial thickness, detection of pericardial inflammation and/or masses,



Figure 1. Utility of CMR in ischaemic heart disease and measurement of ventricular volumes on short axis CMR images

A and B. Short axis imaging allowing LV and RV end diastolic, and end systolic volume measurement. C and D. A positive adenosine stress perfusion study with no rest perfusion defect at rest (image C), while the stress image (image D) shows a subendocardial perfusion defect in the anterior and anteroseptal walls (arrowed). This corresponded to a significant left anterior descending (LAD) artery stenosis at coronary angiography. E and F. Detection of LV apical thrombus and transmural late gadolinium enhancement indicative of full thickness apical infarction

CMR, cardiac magnetic resonance imaging; LV, left ventricle; RV, right ventricle

and assessment of pericardial effusion can provide important additive information to echocardiography. Interventricular interdependence (indicative of constrictive physiology) can be detected by visualising movement of the interventricular septum during respiration.¹²

Cardiomyopathy

CMR is important in the diagnostic workup and management of newly diagnosed cardiomyopathy because it can accurately measure ejection fraction, calculate myocardial mass, and detect scar tissue, fibrosis and oedema.¹⁰

In hypertrophic cardiomyopathy (HCM), CMR-measured septal wall thickness is a prognostic indicator of sudden death. The burden of myocardial fibrosis on CMR is also an emerging marker of risk in HCM. Information obtained through a CMR study in a patient with HCM may directly impact upon the decision for a patient to undergo placement of an implantable cardiac defibrillator (Figure 3).¹³

Variable patterns of late gadolinium uptake can indicate myocardial infiltration (eg cardiac amyloidosis, sarcoidosis), myocardial scar and viability (in ischaemic heart disease), and myocardial fibrosis (in non-ischaemic cardiomyopathy).¹⁰

ARVC is a rare but important cause of sudden cardiac death in athletes and young people.¹⁴ It is characterised by ventricular arrhythmias and fibrous or fibro-fatty replacement of the myocardium that typically involves the right ventricle (RV). There are complex diagnostic criteria for the diagnosis of ARVC, which includes cardiac imaging. CMR has significant benefits over echocardiography in the assessment of the right heart in this condition and is therefore the imaging modality of choice, and in the screening of relatives of those diagnosed with ARVC.¹⁵

Ischaemic heart disease

CMR is useful in ischaemic heart disease (IHD) in the following settings:

 distinguishing ischaemic versus non-ischaemic aetiology of a newly diagnosed cardiomyopathy¹⁶

- detection of complications following myocardial infarction, such as detection of left ventricular (LV) thrombus (when suspected in large left anterior descending artery territory myocardial infarction)
- assessment of myocardial viability prior to potential revascularisation¹⁶
- detection of ischaemia: CMR imaging is combined with dobutamine or a vasodilator such as adenosine, which allows the detection of LV wall segments that become ischaemic (with reduced wall motion) with stress or define a myocardium 'at risk.'¹⁷¹⁸

Myocarditis

In suspected myocarditis, CMR is extremely useful. It allows detection of inflammatory hyperaemia, oedema and myocyte necrosis. Clinically, this may have an impact on the decision to perform endomyocardial biopsy or the indication for corticosteroid therapy.⁹ CMR may identify myocarditis in more than 30% of patients with chest pain, elevated troponin levels and normal coronary arteries. This has important long-term therapeutic implications.¹⁹

Congenital heart disease and pulmonary hypertension

The ability of CMR to calculate accurate right ventricular volumes and provide optimal imaging planes for the assessment of RV size has a growing role in assessment, prognostication and for monitoring treatment efficacy in pulmonary arterial hypertension.²⁰ Congenital heart disease encompasses a broad range of cardiac pathologies, and CMR can provide accurate data regarding ventricular size



Figure 2. Patterns of late gadolinium enhancement

A. LGE image showing subendocardial enhancement of the anterior and anteroseptal walls (marked with *) corresponding to a LAD territory myocardial infarction. B. Mid-wall LGE in dilated cardiomyopathy. C. Sub-epicardial LGE in myocarditis or sarcoidosis

LAD, left anterior descending; LGE, late gadolinium enhancement





Figure 3. Hypertrophic cardiomyopathy on CMR

A. Significant septal hypertrophy

B. Corresponding patchy mid-wall myocardial fibrosis (indicated by LGE) post-gadolinium administration in a patient with hypertrophic cardiomyopathy

Ao, aorta; CMR, cardiac magnetic resonance imaging; LA, left atrium; LGE, late gadolinium enhancement; LV, left ventricle

and function, the follow-up of previous surgery, and the assessment and quantification of shunts. CMR overcomes many of the limitations of a transthoracic echocardiogram (TTE) in this setting. Three-dimensional magnetic resonance angiography (3D-MRA) provides excellent anatomical assessment of the great vessels, vascular abnormalities and vascular shunts.

Accessibility of CMR

CMR is generally accessed through a cardiologist and it is normal practice for echocardiography to be performed first. In each major Australian city, there are a small number of CMR services that are usually attached to the cardiology department of major public hospitals. A Medicare rebate for CMR can be accessed for congenital heart disease, aortic disease and for the assessment of cardiac masses in Australia.²¹ When a patient does not meet the Medicare requirements, the individual CMR service may fund the scan or patients may pay an out-of-pocket fee, which may vary between departments. Some private health insurance companies may provide a rebate for inpatient CMR testing.

Future directions

New CMR techniques are constantly undergoing research and validation that includes further development of sequences that characterise myocardial tissue content. Techniques that allow for single acquisition-based measurements of ventricular volumes, and evaluation of valvular and aortic disease, may assist in reducing the scan times required for CMR studies.^{22,23}

Conclusion

CMR is primarily used as a complementary tool to other imaging modalities, such as echocardiography, in the investigation and diagnosis of specific cardiac diseases. Current availability of CMR is limited by the expense of the study, a limited number of indications with a Medicare rebate, and the time and resources required for an individual CMR study. However, the number of CMR studies being performed in Australia will inevitably increase with improved MRI techniques, increased clinical utility and wider availability.

Authors

Michael B Stokes MBBS FRACP, Cardiac Imaging Fellow, Department of Cardiology, Royal Adelaide Hospital, North Terrace, Adelaide, SA. mbstokes83@ gmail.com

Nitesh Nerlekar MBBS FRACP, Cardiac Imaging Fellow and PhD Student, Monash Heart, Monash Health, Clayton, Vic

Stuart Moir MBBS PhD FRACP, Consultant Cardiologist and Lead of Cardiac MRI, Monash Heart, Monash Health, Clayton, Vic

Karen S Teo MBBS PhD FRACP, Consultant Cardiologist, Department of Cardiology, Royal Adelaide Hospital, North Terrace, Adelaide, SA Competing interests: None.

Provenance and peer review: Not commissioned, externally peer reviewed.

References

- Bellenger NG, Burgess MI, Ray SG, at al. Comparison of left ventricular ejection fraction and volumes in heart failure by echocardiography, radionuclide ventriculography and cardiac magnetic resonance. Are they interchangeable? Eur Heart J 2000;21:1387–96.
- Teo KS, Carbone A, Piantadosi C, et al. Cardiac MRI assessment of left and right ventricular parameters in healthy Australian normal volunteers. Heart Lung Circ 2008;17(4):313–17.
- Saeed M, Van TA, Krug R, Hetts SW, Wilson MW. Cardiac MR Imaging: Current status and future directions. Cardiovasc Diagn Ther 2015;5(4):290–310.
- Hamilton-Craig CR, Slaughter RE, Maki JH. Cardiovascular magnetic resonance from basics to clinical applications. Appl Radiol 2010;39(11):42–53.
- Van Der Graaf AWM, Phagirath P, Gotte MJW. MRI and cardiac implantable electronic devices: Current status and required safety conditions. Netherlands Heart J 2014;22:269–76.
- Cawley PJ, Maki JH, Otto C. Valvular heart disease: Changing concepts in disease management. Cardiac magnetic resonance imaging for valvular heart disease. Circulation 2009;119:468–78.
- Pennell DJ. T2* Magnetic resonance: Iron and gold. JACC: Cardiovasc Imaging 2008;1(5):579–81.
- Jellis CL, Kwon DH. Myocardial T1 mapping: Modalities and clinical applications. Cardiovasc Diagn Ther 2014;4(2):126–37.
- 9. Friedrich MG, Marcotte F. Cardiac magnetic reasonance assessment of myocarditis. Circ Cardiovasc Imaging 2013;6:833–39.
- Parsai C, O'Hanlon R, Prasad SK, Mohiaddin RH. Diagnostic and prognostic value of cardiovascular magnetic resonance in non-ischaemic cardiomyopathies. J Cardiovasc Magn Reson 2012;14:54.
- The Royal Australian and New Zealand College of Radiologists. Guidelines for use of gadoliniumcontaining MRI contrast agents in patients with renal impairment. Sydney: RANZCR, 2013.

- Verhaert, D, Gabriel RS, Johnston D, Lytle BW, Desai MY, Klein AL. The role of multimodality imaging in the management of pericardial disease. Circ Cardiovasc Imaging 2010;3(3):333–43.
- Maron MS, Maron BM. Clinical impact of contemporary cardiovascular magnetic resonance imaging in hypertrophic cardiomyopathy. Circulation 2015;132:292–98.
- Sen-Chowdhry S, Lowe MD, Sporton SC, McKenna W. Arrhythmogenic right ventricular cardiomyopathy: clinical presentation, diagnosis, and management. Am J Med 2004;117(9):685–95.
- Riele ASJM, Tandri H, Bluemke DA. Arrhythmogenic right ventricular cardiomyopathy: Cardiovascular magnetic resonance update. J Cardiovasc Magn Reason 2014;16:50.
- McCrohon JA, Moon JCC, Prasad SK, et al. Differentiation of heart failure related to dilated cardiomyopathy and coronary artery disease using gadolinium-enhanced cardiovascular magnetic resonance. Circulation 2003;108: 54–59.
- Kim RJ, Wu E, Rafael A, et al. The use of contrast-enhanced magnetic resonance imaging to identify reversible myocardial dysfunction. N Engl J Med 2000;343:1445–53.
- Nagel E, Lehmkuhl HB, Bocksch W, et al. Noninvasive diagnosis of ischemia-induced wall motion abnormalities with the use of high-dose dobutamine stress MRI: Comparison with dobutamine stress echocardiography. Circulation 1999;99(6):763.
- Assomull RG, Lyne JC, Keenan N. The role of cardiovascular magnetic resonance in patients presenting with chest pain, raised troponin, and unobstructed coronary arteries. Eur Heart J 2007;28:1242–49.
- Bradlow WM, Gibbs JSR, Mohiaddin RH. Cardiovascular magnetic reasonance in pulmonary hypertension. J Cardiovasc Magn Reson 2012;14:6.
- Department of Health. Medicare Benefits Schedule book category 5. Canberra: DoH, 2013. Available at www.mbsonline.gov.au/internet/ mbsonline/publishing.nsf/Content/85A357DC1C D8BA24CA257D6600087FAD/\$File/201407-Cat5. pdf [Accessed 30 June 2016].
- Claus P, Omar AM, Pedrizzetti G, Sendgupta PP, Nagel E. Tissue tracking technology for assessing cardiac mechanics: Principles, normal values, and clinical applications. JACC Cardiovasc Imaging 2015;8(12):1444–60.
- Dyverfeldt P, Bissell M, Barker AJ, Bolger AF, Carlhall CJ, Ebbers T. 4D flow – Cardiovascular magnetic resonance consensus statement. J Cardiovas Magn Reson 2015;17:72.

correspondence afp@racgp.org.au