
















Improving cardiovascular magnetic resonance access in low- and middle-income countries for cardiomyopathy assessment: rapid cardiovascular magnetic resonance

Katia Devorha Menacho ^{1,2}, Sara Ramirez ³, Aylene Perez⁴, Laura Dragonetti ⁵, Diego Perez de Arenaza⁶, Diana Katekaru ⁷, Violeta Illatopa⁸, Sara Munive⁸, Bertha Rodriguez⁹, Ana Shimabukuro¹⁰, Kelly Cupe¹⁰, Rajiv Bansal¹¹, Vivek Bhargava¹², Ivonne Rodriguez¹³, Andreas Seraphim ^{1,2}, Kris Knott ^{1,2}, Amna Abdel-Gadir¹, Salomon Guerrero¹⁴, Marco Lazo ¹⁵, David Uscamaita⁹, Marco Rivero ³, Neil Amaya⁹, Sanjiv Sharma¹⁶, Amelia Peix ⁴, Thomas Treibel^{1,2}, Charlotte Manisty ^{1,2}, Sam Mohiddin^{1,2}, Harold Litt ^{17,18}, Yuchi Han^{17,18}, Juliano Fernandes ¹⁹, Ron Jacob²⁰, Mark Westwood ², Ntobeko Ntusi ²¹, Anna Herrey ^{1,2}, John Malcolm Walker ^{1,22}, and James Moon^{1,2*}

¹Institute of Cardiovascular Science, University College London, London, UK; ²St Bartholomew's Hospital, Barts Heart Centre, London EC1A 7BE, UK; ³Peruvian Air Force Hospital, Lima, Peru; ⁴Cardiology and Cardiovascular Surgery National Institute, La Havana, Cuba; ⁵High Technology Medical Institute IMAT, Buenos Aires, Argentina; ⁶Italian Hospital of Buenos Aires, Buenos Aires, Argentina; ⁷Military National Hospital, Cardiac Imaging Department, Lima, Peru; ⁸National Cardiovascular Institute—INCOR, Lima, Peru; ⁹Edgardo Rebagliati Hospital, MRI and CT Department, Lima, Peru; ¹⁰Guillermo Almenara Irigoyen Hospital, National Hospital, Lima, Peru; ¹¹Santokba Durlabhji Memorial Hospital Cum Medical Research Institute, Jaipur, India; ¹²Okay Diagnostic Centre, Jaipur, India; ¹³Sermedial Hospital, Arequipa, Peru; ¹⁴Almanzor Aguinaga National Hospital, Chiclayo, Peru; ¹⁵Ramiro Priale National Hospital, Huancayo, Peru; ¹⁶All India Institute of Medical Sciences, New Delhi, India; ¹⁷Department of Medicine (Cardiovascular Division), Perelman School of Medicine, University of Pennsylvania, Philadelphia, PA, USA; ¹⁸Department of Radiology, Perelman School of Medicine, University of Pennsylvania, Philadelphia, PA, USA; ¹⁹Jose Michel Kalaf Research Institute, Campinas, Brazil; ²⁰Lancaster General Health Hospital, Lancaster, USA; ²¹Department of Medicine, University of Cape Town, Cape Town, South Africa; and ²²The Hatter Cardiovascular Institute, University College London Hospital, London, UK

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Aims

To evaluate the impact of a simplified, rapid cardiovascular magnetic resonance (CMR) protocol embedded in care and supported by a partner education programme on the management of cardiomyopathy (CMP) in low- and middle-income countries (LMICs).

Methods and results

Rapid CMR focused particularly on CMP was implemented in 11 centres, 7 cities, 5 countries, and 3 continents linked to training courses for local professionals. Patients were followed up for 24 months to assess impact. The rate of subsequent adoption was tracked. Five CMR conferences were delivered (920 attendees—potential referrers, radiographers, reporting cardiologists, or radiologists) and five new centres starting CMR. Six hundred and one patients were scanned. Cardiovascular magnetic resonance indications were 24% non-contrast T2* scans [myocardial iron overload (MIO)] and 72% suspected/known cardiomyopathies (including ischaemic and viability). Ninety-eight per cent of studies were of diagnostic quality. The average scan time was 22 ± 6 min (contrast) and 12 ± 4 min

* Corresponding author. Tel: +44 203 8870566, Email: j.moon@ucl.ac.uk

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(non-contrast), a potential cost/throughput reduction of between 30 and 60%. Cardiovascular magnetic resonance findings impacted management in 62%, including a new diagnosis in 22% and MIO detected in 30% of non-contrast scans. Nine centres continued using rapid CMR 2 years later (typically 1–2 days per week, 30 min slots).

Conclusions

Rapid CMR of diagnostic quality can be delivered using available technology in LMICs. When embedded in care and a training programme, costs are lower, care is improved, and services can be sustained over time.

Key question

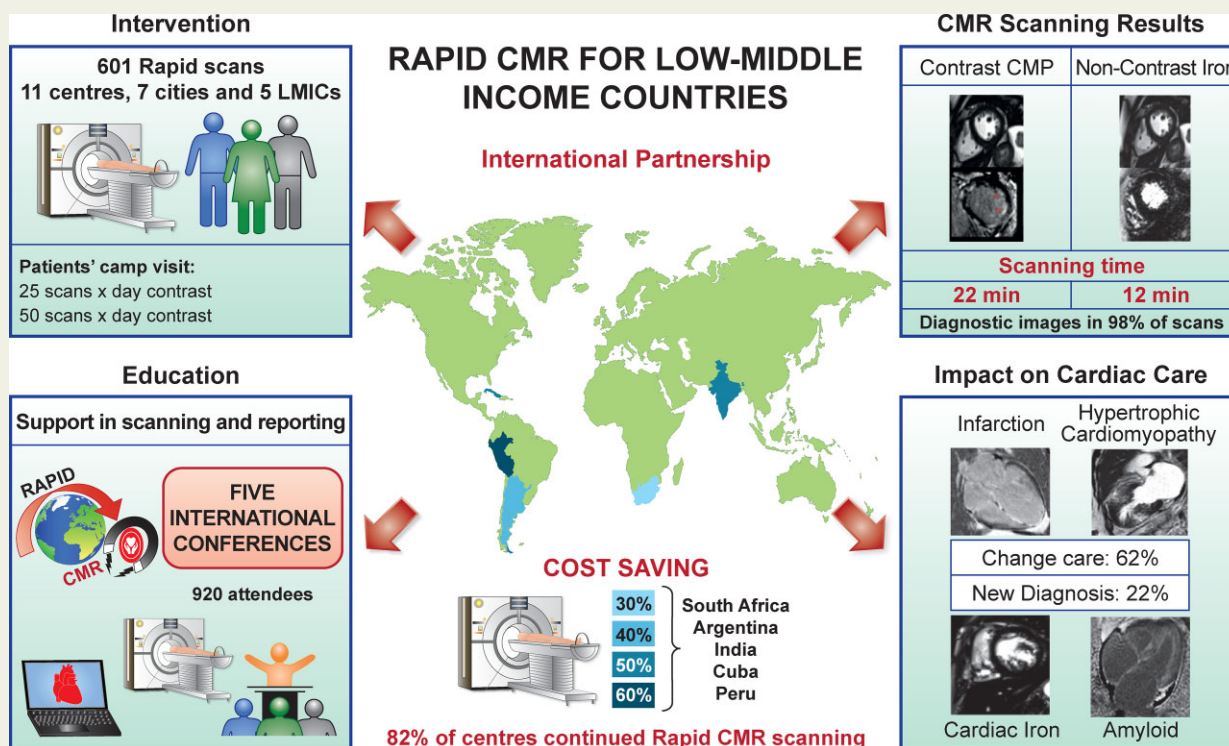
- Can cardiovascular magnetic resonance (CMR) be delivered with a simplified and rapid core protocol in low- and middle-income countries (LMICs) with available technology?
- Can rapid CMR improve patient's care if embedded with a training programme for local professionals in LMICs and incorporated into clinical care?

Key finding

- Five CMR conferences (920 attendees), 11 scanning centres, 7 cities, 5 low- and middle-income countries (LMICs).
- Scanning time: 22 min (contrast) and 12 min (non-contrast).
- Cost reduction: 30–60%.
- CMR findings impacted management in 62% of patients.

Take-home message

- Rapid CMR can be provided quickly and cheaply with diagnostic quality in LMICs.
- When embedded in an education programme, this improves patient care and can be sustainable.



Structured Graphical Abstract Rapid cardiovascular magnetic resonance (CMR) is an international partnership project that aimed to evaluate the impact of a simplified and less expensive CMR protocol embedded with an educational programme to assess a wide spectrum of cardiomyopathies in 11 centres from national/state capital in five low- and middle-income countries (LMICs) with available technology. Over a 5-year period, we delivered an education programme to 920 referrers. Six hundred and one patients were scanned, reducing the time of scanning to <22 min, potentially saving costs and impacting on patients' care. Our project highlights the potential and beneficial role of rapid CMR in LMICs. CMR, cardiovascular magnetic resonance; LMICs, low- and middle-income countries; CMP, cardiomyopathy.

Keywords

Rapid cardiac MRI • Education • Abbreviated protocols • Low–middle-income countries • Cardiomyopathy • Impact on management

Introduction

Optimizing cardiovascular care relies on the availability of diagnostic testing. Test appropriateness depends on factors, including disease prevalence, pre-test probability, therapeutic options, test performance, availability, and cost.¹ Limited testing may be a barrier to optimal care in low- and middle-income countries (LMICs), where cardiovascular disease is a leading cause of mortality and morbidity.² Echocardiography is the first-line diagnostic test for cardiac patients,³ but in isolation may not be sufficient. Cardiovascular magnetic resonance (CMR) complements echocardiography^{4–6} and is recommended in multiple international guidelines, including the majority from the European Society of Cardiology (ESC) (14 guidelines, 39 Class I, and 22 Class II recommendations).⁷ It is the gold standard for the evaluation of ventricular function,⁸ scar imaging using late gadolinium enhancement (LGE)⁹ and iron quantification (CMR-T2*¹⁰). Where CMR-T2* has been adopted to guide chelation therapy for transfusion-dependent thalassaemic patients, a >70% reduction in cardiovascular mortality has been reported.¹¹ Cardiovascular magnetic resonance also plays a critical role in quantifying changes in myocardial composition based on T1, T2 relaxation times, and extracellular volume,¹² and it detects ischaemic heart disease, having shown to be superior to other imaging techniques.¹³

Cardiovascular magnetic resonance has been economically limited to high-income countries (HICs).^{14,15} Access is very limited in LMICs despite magnetic resonance imaging (MRI) availability,¹⁶ though CMR would likely have utility in the major LMIC tertiary healthcare centres, for example, where device therapy, angioplasty, or cardiac surgery are available but need optimal targeting to optimize resource utilization. There are several barriers to adoption, including negative perception (that CMR scans are expensive, time-intensive, and complex), a lack of local expertise (poor access to training for service deliverers and referrers), insufficient investment (overall health sector, private/public sector distribution, poor inter-sector co-ordination),¹⁷ a lack of radiology and cardiology co-ordination, and competition for MRI access.

We aimed to make CMR available in LMICs and to assess its utility. Over 5 years in two pilot studies (TIC-TOC¹⁸ and INCA-Peru¹⁹), we developed and demonstrated rapid CMR in LMICs, initially for cardiac iron quantification in thalassaemic patients (Bangkok, Thailand) with an 8 min protocol¹⁸ and then contrast CMR for cardiomyopathy (CMP) evaluation (two centres, Lima, Peru) with an 18 min protocol.¹⁹ These demonstrated the technical feasibility of rapid CMR and highlighted the importance of system-wide change, embedding imaging in cardiac care with an educational programme and a supportive mentoring environment for emerging services. To explore this further, we developed the rapid CMR in LMICs project (www.rapidcmr.com) and performed abbreviated CMR scans in 11 centres, 7 cities, 5 countries, and 3 continents. We used large centres in national capital cities but also involved smaller centres in state capitals. We aimed to embed rapid CMR within clinical care pathways in order to improve sustainability and utilization by supplementing scanning with an educational programme and mentoring local staff who tracked patient outcomes.

Methods

This is a multicentre, prospective observational cohort study (*Figure 1*). The research complied with the Declaration of Helsinki and received ethical approval in the UK—University College London (REC 11255/001 and 11255/002) and by the relevant local ethics committees in participating centres. An international partnership was established between UK-UCL and Barts Heart Centre, London, UK, and each participating centre, with the support of the local cardiology and radiology societies (that supported advertising the rapid CMR in their channels to local professionals), local scientific councils, and British embassies in Argentina, Peru, and Cuba (who co-financed the UK team visit), and the Society for Cardiovascular Magnetic Resonance (SCMR) (who sponsored one to two SCMR educators/trainers and provided no-fee SCMR certification for participants). A team of five doctors, experts in the field (UK, SCMR expert delegates) travelled for each visit to the participating countries to deliver an educational programme to local doctors and train/partner with local personnel to help perform assessments and support rapid CMR scanning. When possible, we additionally included experts from nearby countries with native language capabilities and encouraged the development of local networks for sustainability and mentorship.

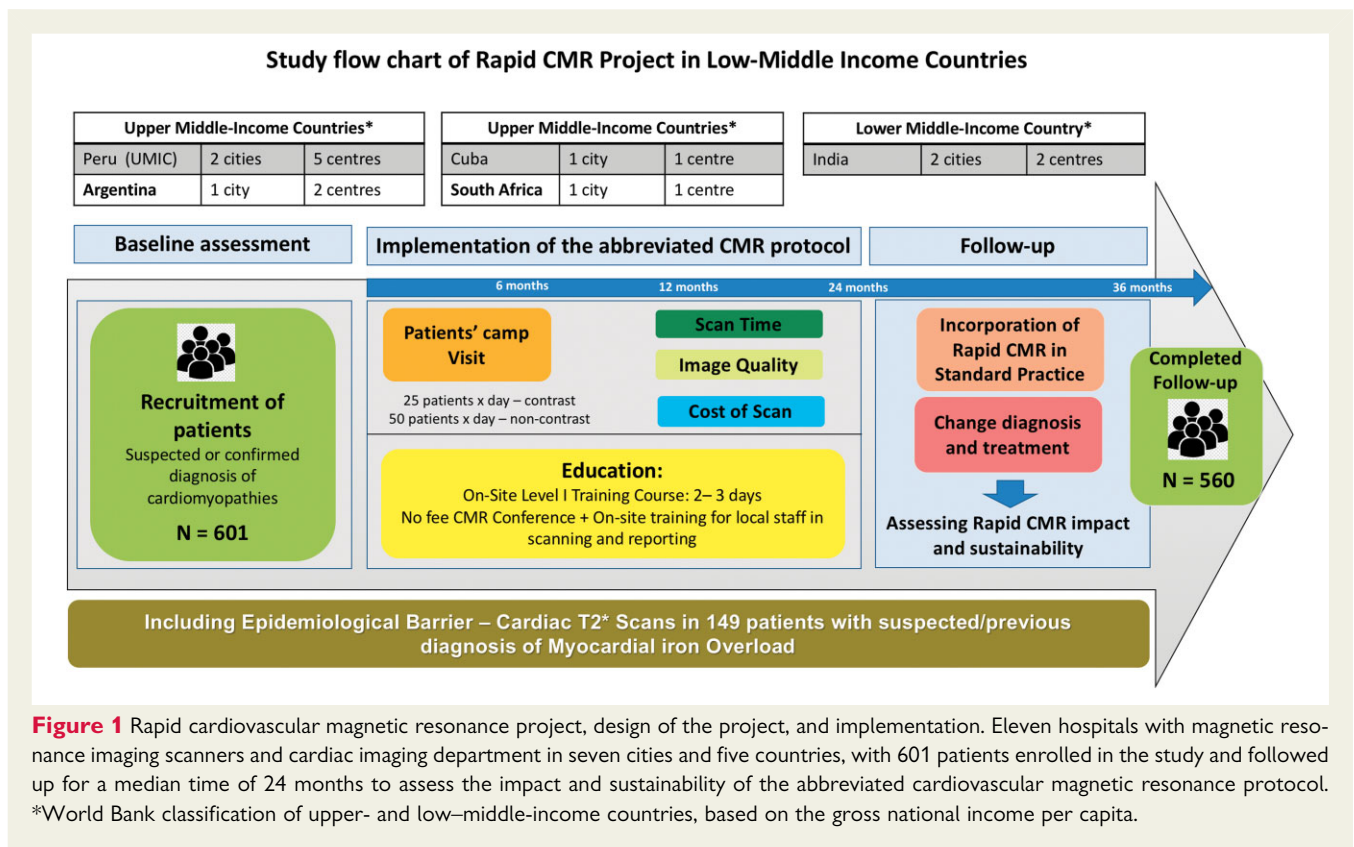
Study implementation

Patients were recruited by the local research teams. All participants provided written consent. The rapid CMR contrast demonstrator study of 98 patients scanned in two centres in Peru were previously published¹⁹ who remain included but with their 36-month follow-up and in multiple cases (iron overload, cardiotoxicity) interval scanning. Following this pilot project, 10 additional participating centres contacted our research group as they wanted to initiate/improve their efficiency in CMR scans. There was one dedicated cardiac CMR scanner (Cuba, 3 days per week), five scanners with one-day-a-week CMR (Peru—one centre, South Africa—one Centre, Argentina—two centres, India—one centre), and five scanners performing no CMR before initiation of the project (Peru—four centres and India—one centre). Scanner time was secured mainly during unused magnet time (weekends, national holidays) and the international delegation visit, educational programme, and ‘patients’ camp’ visits were scheduled accordingly. Scans were at no cost to participants and the costs were borne by the participating centre (scanner availability, staff support) and the research funding (providing contrast agent and ECG). The patients’ inclusion criteria are cited in [Supplementary material online, Appendix—Table S1](#).

Education and training programme

Education was a key component of our project, and it was delivered at two levels:

- (1) *CMR International Conferences*: These were 2–3 days on CMR scanning and reporting per country for local cardiologists, radiologists, and radiographers. All were free of charge. Society for Cardiovascular Magnetic Resonance provided free Level 1 SCMR certification²⁰ for participants and a free 1-year SCMR membership. CMR teaching included lectures by local/national/international speakers, direct CMR scanning observation, and a Level 1 reporting course.
- (2) *Training for those delivering CMR*: Six CMR centres were already established pre the rapid CMR intervention. Five were novice. For these, one to two international visiting doctors/radiographers delivered training in scanning and reporting to the local team days before the main intervention. For both, there was support during



scanning days. Follow-up continuous support was given to local doctors remotely (technical/interpretation of difficult cases). We assessed the impact by measuring sustainability—the number of scans (and duration) delivered after the intervention period.

Abbreviated cardiovascular magnetic resonance protocol and implementation

The rapid CMR protocol was published previously (INCA-Peru study).¹⁹ Here, we modified for specific patients (e.g. patient unable to perform breath holds adequately, patients with arrhythmia), and additional sequences were added for specific cardiac pathologies following SCMR sequence recommendations dependent on the scan indication⁸ (Figure 2). We followed a standardized workflow to maximize efficiency surrounding CMR (see [Supplementary material online, Appendix—Table S2](#)). One to two weeks before the main intervention, the rapid CMR protocol was shared remotely and installed in the local MRI scanner (with the support of the local radiographers/MRI engineering service). Preliminary scanning was done in healthy volunteers in each participating centre and mentored via live webcasting.

Standard cardiomyopathy cardiovascular magnetic resonance protocol

- (1) *Localizers*: A pilot three-plane localizer, a transverse bright blood stack for anatomic evaluation (optional ungated); pilots two-chamber and short-axis (SAX) stack.
- (2) *Longitudinal axis*: Steady-state free precession (SSFP) cine (25 phases) in four-, two-, and three-chamber views.
- (3) *Hand contrast injection* (dose 0.1 mmol/kg) gadoterate meglumine/gadobutrol.²¹

- (4) *SAX cine stack and aortic valve cine*.
- (5) *LGE*: repeating cine views as needed, followed by an inversion time scout if needed, then a segmented k-space LGE acquisition in multiple planes with a phase-sensitive inversion recovery sequence and magnitude reconstructions.
- (6) *Additional sequences* (if available in the scanners): for specific clinical indications including:⁸
- (7) *For all cardiomyopathies*: a single T1 mapping mid-SAX slice and four-chamber pre- and post-contrast (10 min after contrast administration).
- (8) *For arrhythmogenic right ventricular CMP (ARVC)*:
 - (a) Transaxial or oblique transaxial bSSFP cine images (slice thickness 5–6 mm) covering the right ventricle, including the right ventricular outflow tract.
 - (b) Selected black blood images (four-chamber and three SAX slices: base, mid, and apex) + repeat same geometry with fat suppression.
- (9) *For myocarditis*: Selected T1 mapping, T2 mapping, or T2w STIR sequences at four-chamber and three SAX slices: base, mid, and apex.
- (10) *For pericardial disease*:
 - (a) T₁ or T₂-weighted fast/turbo spin echo (FSE) images in one representative long-axis and one representative SAX slice to measure pericardial thickness.
 - (b) Real-time cine imaging during dynamic breathing manoeuvres for evaluation of ventricular interdependence (mid-ventricular SAX).
- (11) *For myocardial and liver iron overload (non-contrast scan)*: a T2* multi-echo gradient-echo transversal single slice trying to cover the maximum liver parenchyma and one single mid-cardiac SAX slice.

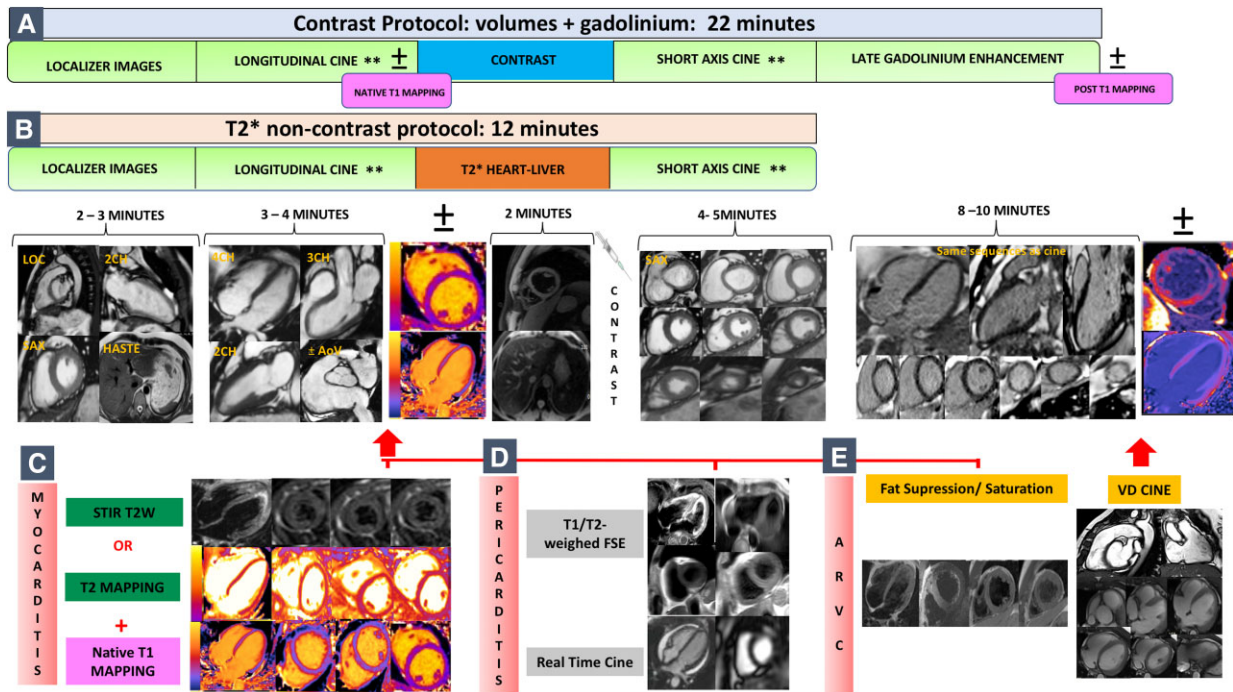


Figure 2 Abbreviated cardiovascular magnetic resonance protocol: (A) 22 min contrast cardiovascular magnetic resonance protocol for the assessment of cardiomyopathies, with the option to add T1 mapping sequence if available in the scanner; (B) 12 min T2* non-contrast protocol to assess myocardial iron overload. Extra sequences added for specific pathologies: (C) short tau inversion recovery T2w or T2 mapping + T1 mapping for myocarditis (27 min scan); (D) T1 or T2w FSE + real-time cine for pericarditis (27 min scans); and (E) fat suppression and saturation + right ventricle cine acquisition for arrhythmogenic right ventricular cardiomyopathy (30 min scan). **If poor breath-holders, modification of the protocol using parallel imaging for cine images acquisition and in patients with arrhythmia, prospective triggering, and real-time cine imaging sequences acquired. CMR, cardiovascular magnetic resonance; MIO, myocardial iron overload; STIR, short tau inversion recovery; FSE, fast/turbo spin echo; RV, right ventricle; ARVC, arrhythmogenic right ventricular cardiomyopathy.

In patients with arrhythmia and poor breath-holder patients, we modified the protocol and used parallel imaging, prospective triggering, and real-time cine imaging. All these sequences were available in the participating MRI centres.

Image reporting

Images were analysed using cvi42 (Circle Cardiovascular Imaging Inc., Version 5.11.4–1559, Calgary, Canada) with a provided typically 3-month ongoing licence to participating centres (two centres had pre-existing Cvi42 licences). Scans were reported the same day by the local doctor, supported, and reviewed by the international visiting experts (final report signed-off by a doctor with a Level 2 or 3 CMR EACVI—ESC certification or equivalent).²⁰ After the onsite intervention, the participating centres used other software (e.g. Osirix, ARGUS, and free online post-processing software for cardiac and iron T2* analysis—<http://www.isodense.com/mcdcm/mviewer.html>). Reporting was used as a training exercise during the educational conference, where indications for the scan and the anonymized images were shown and discussed with the attendees after the patients' consent. Reports were translated into the local language where needed and incorporated into local medical records.

Image quality

Image quality was assessed by a CMR expert (Level 3 EACVI-ESC certified).²⁰ The criteria used were not the presence or absence of

artefacts but whether images acquired could answer the clinical question posed: *poor quality* (unable to answer the clinical question), *moderate quality* (artefact but still interpretable), and *good quality* (optimal quality, definitive assessment).

Study outcome measures

Patients were followed up for a median time of 24 months by their local physician. Cardiovascular magnetic resonance-induced management changes were reported if CMR resulted in: (i) new diagnosis (of a pathology not suspected before), (ii) medication change, (iii) interventional treatment, and (iv) hospital admission or discharge.

Cost evaluation

Local CMR reference costs were assessed individually, as healthcare infrastructure, systems, and costs differed among countries. In general, we considered as reference the average cost from public and private hospitals offering CMR at the time of scanning. Costs included: hospital CMR charges, contrast, intravenous cannula/tubing, ECG electrodes, physician (reporting), nurse, and radiographer fees.

Statistical analysis

Data were analysed using SPSS (version 24-0, Statistical Package for the Social Sciences, International Business Machines, Inc., Armonk, NY, USA). Continuous data were expressed as mean with standard deviation, and categorical data are presented as absolute numbers and percentages. Normal distribution was formally tested using the

Shapiro–Wilk test. Continuous data were compared using two-sided Student's *t*-tests, and categorical variables were compared using χ^2 -test or Fisher's exact test as appropriate. Statistical significance was defined as a two-sided *P*-value of <0.05.

Results

Baseline characteristics

Overall, 601 patients were referred from 14 hospitals, seven cities in five countries as follows: Peru, Lima, and Arequipa, seven centres, *n* = 261; Argentina, Buenos Aires, three centres, *n* = 69; Cuba, La Havana, one centre, *n* = 114; South Africa, Cape Town, one centre, *n* = 34 and India, Delhi, and Jaipur, two centres, *n* = 123. A total of 449 (74%) were referred by cardiologists, 131 (22%) by haematologists [myocardial iron overload (MIO) indication], and 20 (3%) by other specialists. The mean age of the patients was 46 ± 19 years with suspected iron overload patients being younger (28 ± 13 vs. 53 ± 16 years, *P* < 0.001, 95% of MIO scans were of patients with beta-thalassaemia); 56% of participants were male (Table 1).

Despite the intention to concentrate on the assessment of cardiomyopathies, referrals included other cardiac conditions deemed urgent, usually involving economically disadvantaged patients who would normally not have access to this resource.

Reasons for CMR referrals were: cardiomyopathies in 433 (72%), assessment of viability in 98 (16%), myocarditis/pericarditis in 35 (6%), valvular heart disease in 9 (1%), cardiac tumours in 6 (1%), aortic disease in 5 (1%), and other indications (e.g. congenital) in 15 (2%). Within the group of cardiomyopathies, the indications were: dilated CMP in 259 (60%), hypertrophic CMP (HCM) in 126 (29%), ARVC in 15 (3%), restrictive CMP in 15 (3%), and unclassified in 18 (4%). Eighteen different pathologies were assessed (Figure 3).

Cardiovascular magnetic resonance results

Patients were scanned in 11 hospitals with MRI units, using 14 different scanners (six models from three manufacturers at two field strengths, with scanner age at study time of 12 ± 3.4 years) (see Supplementary material online, Appendix—Table S3).

One hundred and fifty-four (26%) patients had a non-contrast scan, 149 (96%) for MIO, four other patients had chronic kidney disease, and hence contrast was not administered, and one patient declined contrast. For contrast CMR scanning, 410 (94%) had undergone transthoracic echocardiography prior to the CMR scan, 10% computed tomography coronary angiography, 12% single photon emission computed tomography, and 8% had a previous CMR. Within the group of patients referred for a non-contrast T2* CMR scan to assess MIO, 31% had previously undergone CMR to assess cardiac iron. Among the 601 scans, 98% were of diagnostic quality.

Good image quality was reported in 506 scans (84%), moderate in 85 (14%), and poor in 10 (2%). Poor image quality was due to atrial fibrillation with rapid ventricular response (*n* = 5), very poor breath-holding (*n* = 3), and frequent ventricular ectopic beats (*n* = 2). Moderate image quality was noted with arrhythmia (*n* = 58) and poor breath holds (*n* = 27).

The delivered throughput was 50 scans per scanner day for iron T2* CMR with typically ~10 h day and 25 scans per scanner day for contrast CMP. Scan time (average \pm standard deviation) was 12 ± 4 min for T2* for MIO and 22 ± 6 min for the basic CMP CMR protocol (function + contrast). In 32 (5%) of the CMP scans, pre- and post-T1 mapping sequences were included with an average scan time of 25 ± 5 min. For other specific clinical indications, CMR scanning time took longer as additional sequences were added: myocarditis/pericarditis protocol (STIR and T2 FSE or

Table 1 Baseline characteristics of the patients enrolled in the rapid cardiovascular magnetic resonance project

Clinical data	Total	Non-contrast CMR scans	Contrast CMR scans	P-value
Demographic				
Total	601 (100%)	154 (26%)	446 (74%)	–
Age, years (mean \pm SD)	46 ± 19	28 ± 13	53 ± 16	<0.001
Female sex	267 (44%)	82 (53%)	185 (41%)	0.01
BMI, kg/m ² (mean \pm SD)	25.2 ± 5.4	21.4 ± 3.7	26.1 ± 5.2	0.02
BSA, m ² (mean \pm SD)	1.7 ± 0.3	1.5 ± 0.2	1.8 ± 0.2	0.9
Imaging modalities used before CMR				
Echocardiography	471 (78%)	51 (33%)	420 (94%)	<0.001
CTCA	46 (8%)	0	46 (10%)	<0.001
SPECT/bone scan	55 (9%)	0	55 (12%)	<0.001
CMR	85 (14%)	48 (31%)	37 (8%)	<0.001
Referral				
Cardiologist	449	21	428	<0.001
Haematologist	131	131	0	<0.001
Other	20	0	20	<0.001

CMR, cardiovascular magnetic resonance; BMI, body mass index; BSA, body surface area; SD, standard deviation; CTCA, computed tomography coronary angiography; SPECT, single photon emission computed tomography.

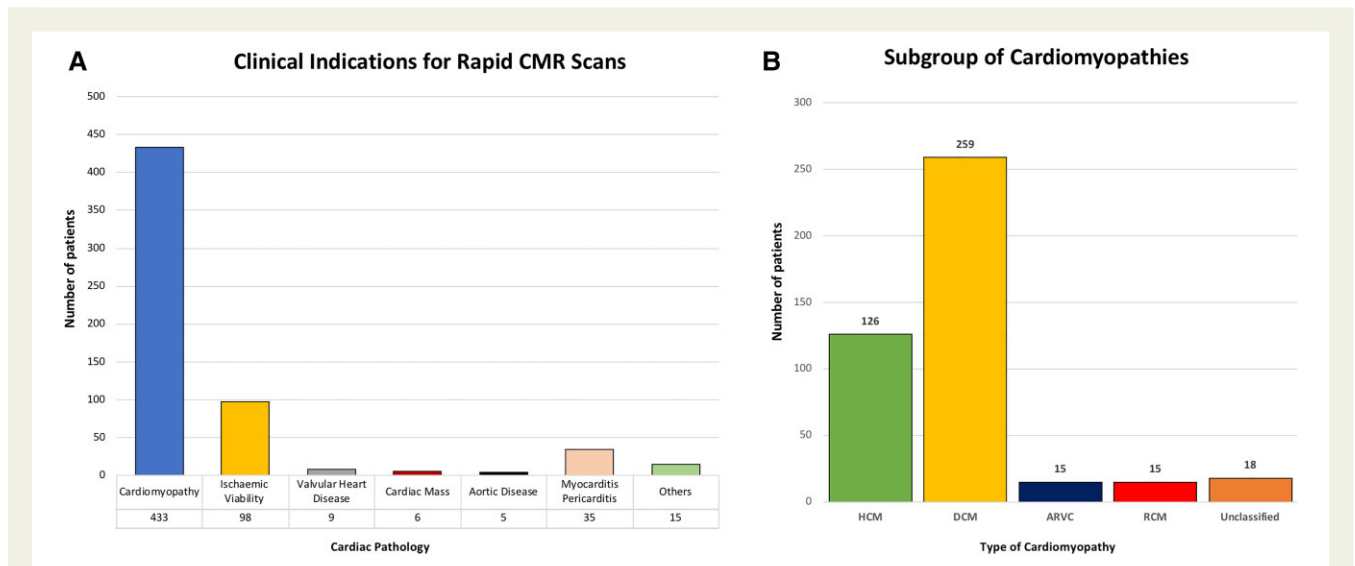


Figure 3 Clinical indications for the use of the abbreviated cardiovascular magnetic resonance protocol in (A) all cardiac pathologies and (B) within the group of cardiomyopathy sub-types. ARVC, arrhythmogenic right ventricular cardiomyopathy; DCM, dilated cardiomyopathy; HCM, hypertrophic cardiomyopathy; RCM, restrictive cardiomyopathy.

T1/T2 mapping if available) 27 ± 4 min and ARVC protocol (with RV cine + fat suppression sequences) 30 ± 10 min. Within the T2* CMR scans, 45 (30%) of patients had cardiac iron (cardiac T2* <20 ms); 36 (80%) previously undocumented. For contrast CMR scans, 288 (64%) patients had positive LGE (for some diseases, such as HCM, LGE was often extensive on qualitative assessment). Thirty-six (8%) were considered normal or non-conclusive studies (Table 2).

Cardiovascular magnetic resonance impact on diagnosis and management

A total of 560 (93%) patients completed follow-up (median 24 months post-CMR). Forty-one participants were not contactable.

The abbreviated CMR protocol changed clinical management in 62%, either by revealing a new diagnosis (22%) or by leading to a change in therapy (40%) (Figure 4, for new diagnosis—contrast rapid CMR scans and Supplementary material online, Appendix S4 for three clinical case examples). The T2* CMR scans (cardiac and liver assessment) largely led to a change/start of chelation therapy in 55% of patients, as the diagnosis of transfusion-dependent thalassaemia was already established. After rapid CMR, 12% with cardiomyopathies required further invasive tests to arrive at their final diagnosis (e.g. angiography, cardiac biopsy) or to therapeutic interventions (e.g. implantable-cardioverter defibrillator) (Table 3). Although non-blinded, rapid CMR did not miss any findings previously made by echocardiography.

Table 2 Time of scanning for the rapid cardiovascular magnetic resonance protocol

	Type of rapid CMR protocol for the detection of cardiomyopathies				
	Iron CMR (cine + T2*) n = 149	CMP (cine + LGE) n = 350	CMP mapping (cine + LGE + T1 mapping) n = 32	Myocarditis/pericarditis (STIR/T1 FSE or mapping if available) n = 35	ARVC (RV cine fat Sat/Sup.) n = 15
Age, years (mean ± SD)	26 ± 11	54 ± 16	47 ± 16	48 ± 19	30 ± 10
Time, min (mean ± SD)	12 ± 4	22 ± 6	25 ± 5	27 ± 4	30 ± 10
Image quality					
Good	129 (87%)	294 (84%)	28 (87%)	29 (82%)	12 (82%)
Moderate	17 (11%)	49 (14%)	4 (13%)	6 (18%)	3 (18%)
Poor	3 (2%)	7 (2%)	0 (0%)	0	0
Arrhythmia	2 (1%)	48 (14%)	4 (13%)	0 (0%)	4 (27%)

Contrast CMR protocol for cardiomyopathies and non-contrast CMR scan to assess myocardial iron overload. Modification of the protocol and time of scanning for specific cardiac pathologies. CMR, cardiovascular magnetic resonance; ARVC, arrhythmogenic right ventricular cardiomyopathy; CMP, cardiomyopathy; LGE, late gadolinium enhancement; STIR, short tau inversion recovery; FSE, fast/turbo spin echo; RV, right ventricle; SD, standard deviation; Sat/Sup, saturation and suppression.

Diagnosis Pre – Contrast Rapid CMR

Total: 601

20 patients no-cardiomyopathy protocol
149 patients for Cardiac Iron non-contrast CMR

Diagnosis Post – Contrast Rapid CMR

Total Follow-up: 560

18 patients non-cardiomyopathy protocol
129 patients for Cardiac Iron non-contrast CMR

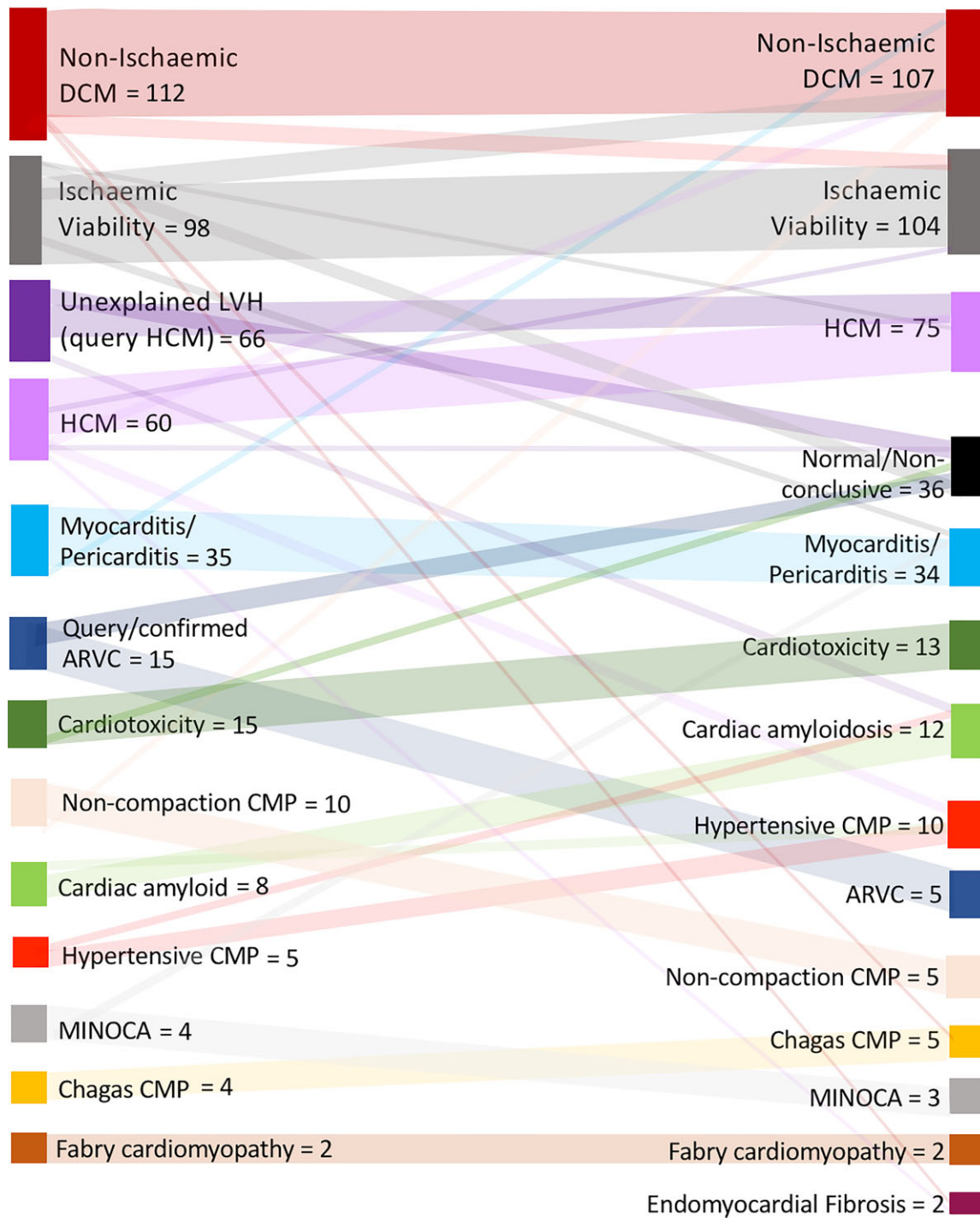


Figure 4 Clinical indications for the use of the abbreviated contrast rapid cardiovascular magnetic resonance protocol to assess ischaemic/non-ischaemic cardiomyopathies and myocarditis/pericarditis. After a 24-month follow-up period, 89 new diagnoses were made. CMR, cardiovascular magnetic resonance; CMP, cardiomyopathy; HCM, hypertrophic cardiomyopathy; LVH, left ventricular hypertrophy; MINOCA, myocardial infarction with non-obstructive coronary arteries; ARVC, arrhythmogenic right ventricular cardiomyopathy; DCM, dilated cardiomyopathy.

Table 3 Impact of the rapid cardiovascular magnetic resonance protocol on patient care

CMR findings	Cohort	Cardiac T2* iron scans	Contrast CMR scans	P-value
Abnormal baseline CMR results				
No.	601 (100%)	149 (25%)	452 (75%)	–
Cardiac iron overload (T2* <20 ms)	45 (7%)	45 (30%)	N/A	–
Presence of LGE	288 (48%)	N/A	288 (64%)	–
Inconclusive CMR	10 (2%)	0 (0%)	10 (2%)	<0.001
Follow-up—impact on patient care				
No.	560 (100%)	129 (23%)	431 (77%)	
New diagnosis	125 (22%)	36 (28%)	89 (21%)	0.19
Change/addition of new medication	134 (24%)	36 (28%)	98 (23%)	<0.001
Intervention or surgery	46 (8%)	0 (0%)	46 (8%)	<0.001
Coronary angiography or biopsy	21 (4%)	0 (0%)	21 (4%)	<0.001
Hospital discharge and admission	25 (4%)	0 (0%)	25 (6%)	<0.001
Other imaging modalities requested after CMR to support diagnosis and therapy				
No.	560 (100%)	129 (23%)	431 (77%)	–
Echocardiography	45 (8%)	3 (2%)	42 (9%)	<0.001
CTCA	17 (3%)	0 (%)	17 (3%)	<0.001
SPECT	12 (2%)	0 (%)	12 (2%)	<0.001

CMR findings during patients' camp scanning and assessment of the impact of the abbreviated CMR protocol on patients' care and further additional cardiac imaging requested to support patients' care. CMR, cardiovascular magnetic resonance; LGE, late gadolinium enhancement; CTCA, computed tomography coronary angiography; SPECT, single photon emission computed tomography.

Cost evaluation

Estimated cost savings for rapid compared with conventional contrast CMR were between 30 and 60% (equivalent to one-third to two-thirds cheaper) with a range of 25–30% in Cape Town, 30% in Argentina, 40% in Cuba, and 50–60% in Peru. For non-contrast T2* CMR, the saving ranged between 40% in India and 50% in Peru (Table 4).

Cardiovascular magnetic resonance education programme

Five international conferences were delivered in Peru (November 2016 and January 2019), Argentina (June 2018), Cuba (November 2019), and South Africa (June 2018). Each conference had at least 180 attendees, with a total of 920 professionals. All completing participants received no-fee Level 1 SCMR certification and free SCMR membership. See www.rapidcmr.com for prior course details.

A total of 18 professionals were trained in CMR scanning and reporting (1–2 professionals per centre—7 doctors and 11 radiographers in total). Thirteen out of 18 professionals are still using their skills, with three doctors achieving a formal CMR training (equivalent to Level 2 CMR certification), but five professionals (three radiographers, one cardiologist, and one radiologist) did not continue CMR as the MRI units in two hospitals in Peru sustained permanent scanner breakdowns.

Subsequent adoption of rapid cardiovascular magnetic resonance

Nine out of 11 participating centres continued using rapid CMR (two hospitals in Peru ceased due to scanner breakdowns), with

services scanning 1–2 days per week. From the entire list of CMR indications, the participating centres continue using the rapid CMR protocol to assess cardiomyopathies in 2–8 patients per day (Table 4). We followed up sites and found that the main services were maintained and even grew (Table 4) with a variety of different local funding solutions as the healthcare systems are complex and different for each country: (i) public (e.g. Peru, South Africa with CMR provided directly by public hospitals to government/non-government employees), (ii) Army/Navy (e.g. Peru), (iii) governmental (e.g. Cuba), (iv) private with third party payers (e.g. Argentina, Peru, with private hospital charging CMR scans to public hospitals), (v) Charity Organization Societies (e.g. India, with T2* CMR scans costs co-financed by Thalassaemics India Patient Society—<https://www.thalassaemicsindia.org/>).

Two new centres in two non-capital cities in Peru (Chiclayo and Huancayo) and one centre in La Havana, Cuba, initiated a CMR service (scanning 1/2 day per week) led by a CMR specialist physician with remote support from the international team of experts.

Discussion

We report the ability to perform rapid CMR in multiple LMICs in three continents across different clinical environments utilizing a variety of different scanners. These scans provided diagnostic image quality in the majority of cases at greatly reduced cost with an important impact on the care of the patient (Structured Graphical Abstract). This was achieved without having the most current scanner and software in many of the centres. To achieve this, four key strategies were employed: (i) a collaborative partnership between

Table 4 Cardiovascular magnetic resonance practice in the participating centres before and after the implementation of the rapid cardiovascular magnetic resonance protocol

Country	Argentina		Cuba	Peru		South Africa		India (non-contrast T2* CMR)					
	Buenos aires (n = 69)	Centre 1	Centre 2	Centre 3	Centre 4	Centre 5 ^a	Centre 6	Centre 7 ^a	Centre 8	Cape town (n = 34)	Centre 9	Delhi (n = 53)	Jaipur (n = 70)
Before rapid CMR scan													
Number of days per week	1	2	3	0	1	0	0	0	0	2	1	1	0
Number of hours × day	5	5	5	0	5	0	0	0	0	5	2	2	0
CMR scans per day	4	5	3–4	0	4	0	0	0	0	5	2	2	0
Cost	\$400–\$500		\$600	\$400–\$600						\$350–\$500	\$70–\$100		
After rapid CMR scan													
Number of days per week	1	2	3	1	1	1	1	1	1	2	1	1	1
Number of hours × day	5	5	5	4	5	4	4	4	4	5	2	2	2
CMR scans per day	5–7	6–7	7–9	6	6–8	6	6	6	6	6–7	4–6	4–6	4–6 ^b
Rapid CMR scans (3.0 min × contrast scan) (2.0 min × T2* scan)	5–6	3–4	6–8	4	4–6	4	4	4	4	5	4–6	4–6	4–6 ^b
Cost rapid CMR scan	\$280–\$380		\$360	\$160–\$300						\$200–\$400	\$42–\$60		
% saving	30%		40%	50%–60%						25–30%	40%		

Argentina—Buenos Aires: Centre 1: Instituto IMAT. Centre 2: Hospital Italiano de Buenos Aires.

Cuba—La Havana: Centre 3: Instituto de Cardiología y Cirugía Cardiovascular.

Peru—Lima: Centre 4: Hospital ESSALUD Edgardo Rebagliati. Centre 5: Hospital Central Militar. Centre 6: Hospital Central de la Fuerza Aerea del Peru. Centre 7: Hospital ESSALUD Guillermo Almenara.

Peru—Arequipa: Centre 8: Centro Salud Cerema.

South Africa—Cape Town: Centre 9: Groote Schuur Hospital in Cape Town.

India—Jaipur: Centre 10: Mahajan Imaging Centre.

India—Jaipur: Centre 11: OK Diagnostic Imaging Centre.

^aFaulty scanners, stopped CMR service (the number representing post-study to pre-breakdown).

^bTen scans done after rapid CMR scan visit. Starting that number of CMR scans this year.

the investigators and local teams at multiple levels (a local health-care champion, hospital team, political authorities, scientific societies, and embassies, (ii) CMR training and education at two levels: first, referring physician and second, to those providing CMR services (cardiologists, radiologists, and technologists—acquisition and reporting) in the local language or English with translation, (iii) a focused, rapid, and less expensive CMR scan with sequences adapted to local scanners and software, (iv) the integration of results into the care of the patient was with follow-up to assess the impact. The rate of subsequent adoption in the participating centres was tracked.

This is the first study that tackles the education barrier of CMR in LMICs, where most professionals need to travel overseas and self-fund for expensive courses to receive certified training.²⁰ We provided training at the local participating sites ensuring that the appropriate technology is available. It is important to emphasize that while we provided training for practitioners (doctors and radiographers to deliver CMR), we made a large effort to train potential *referrers* by organizing the five Level 1 CMR international conferences, aiming that they can be enthusiastic about CMR, to learn more cardiology, and as a gateway for the (few) who want to take CMR further. Here, we had a ratio of basic advanced training of ~50:1. The Level 1 training here is in no way wasted if they do not directly deliver CMR services but is perhaps the foundation of CMR incorporation into clinical practice.

Ninety-four per cent of patients had undergone previous echocardiography but needed CMR to complete their evaluation as there was equipoise regarding the diagnosis or appropriate therapy, which was not answered by previous imaging testing. Advanced tissue characterization with LGE allows for important risk stratification²² and estimation of prognosis.²³ The extension of LGE seen in some cardiomyopathies such as HCM patients was far more than is typically seen in an HIC with an established CMR service. We believe this is a marker of a lack of access to advanced imaging. In LMICs, patients with more advanced stages of HCM were typically seen, suggesting that cases that might have been picked up by CMR during the early stages of the disease may have been missed. The referral indications for rapid CMR were mainly to assess cardiomyopathies and differed compared with HICs where heart failure, cardiomyopathies, function and viability, and stress perfusion are commonly indicated.²⁴

The core structure of the abbreviated CMR protocol previously published in the INCA-Peru study¹⁹ to assess specific cardiomyopathies was modified for this study. Adding these sequences went out of the scope of our ‘keep it simple’ approach and this happened due to different challenges: individual patients outside our planned remit (congenital, masses, valvular), where to turn them down would give them no CMR option; also referrers and local champions some of who had trained overseas who demanded guideline appropriate scans,⁸ and in some cases of course ourselves. Even though we were delivering a research project, it proved difficult at times to not provide the same level of care we would in HIC parent institutions. It is interesting that the added sequences in many cases provided little incremental value (as discussed below for T1 mapping). Despite this, we showed that CMR can be acquired in 22 min on average for a core CMP protocol. Even when adding other sequences for specific pathologies, we

were able to scan patients in <30 min. Overall, this corresponds to a reduction in scanning time of around 30–45% compared with reported CMR exam durations.⁸ The scanning time improved progressively once local radiographers gained experience (e.g. our second visit in Peru, the average scan time was 19 vs. 24 min for the first visit in Cuba).

Arrhythmias were detected in 10%, requiring protocol modification, but despite this, 98% of the scans were of diagnostic quality, similar to reports of conventional CMR scans from international surveys.¹⁴

In three centres (two in Peru and one in South Africa), T1 mapping was available, and we incorporated one mid-SAX and four-chamber view native and post-contrast T1 mapping—adding 3 min to the core protocol. T1 mapping provides incremental value in specific pathologies such as Fabry disease²⁵ and cardiac amyloidosis²⁶ without the need for contrast. Here, the (untargeted) yield was low, with none of these pathologies detected in 32 patients, although cardiac amyloidosis was found in some patients in the rapid core contrast CMR protocol.

The clinical utility of rapid CMR in affecting patient care was 62%, defined by a new or changed diagnosis, change in therapy, or subsequent procedure, similar to reports from international registries.^{14,15} Six per cent of our cohort was lost to follow-up. Many of those lost to follow-up lived in remote rural areas; we believe the impact of CMR in LMICs could be potentially higher than in HICs if these disadvantaged groups could be better tracked, although barriers to ongoing clinical management, including access to costly medications or devices, may limit this. Only 13% of our patients had additional cardiac imaging investigations triggered, mainly as part of the confirmation of a new diagnosis revealed by CMR, e.g. bone scintigraphy test to confirm cardiac amyloidosis.

Cost implications can only be broadly estimated due to different healthcare systems in the five countries. Our best estimate is that rapid CMR could reduce costs of contrast scans between one-third and two-thirds (equivalent to 30–60% of cost reduction). This does not include any cascading savings of better targeting of treatment. Further study would be needed to estimate this.

Further barriers remain: the lack of engineering and MRI maintenance service remains a barrier for service continuity (2 of the 11 centres ceased MRI because of this).

Although we focused on LMICs, there is potential utility in HICs. In the current COVID-19 pandemic,²⁷ as an example, CMR needs to be done in a way that minimizes exposure to the staff without compromising the diagnostic yield of the study, as recently evaluate.²⁸

Study limitations

This was not a randomized study. No assessment was made of prior services when present. The study was of the willing and able—included centres all had had a local, supported champion. The study was in either capital cities or state capitals within LMICs: results are not expected to generalize beyond this. Scanners were those that were available. Advanced CMR techniques (e.g. perfusion) were out of scope,¹³ even though recent data suggest these can also be done rapidly.²⁹ Latest techniques were not used (e.g. compressed sensing).³⁰ The team performed immediate CMR viewing, post-processing, and reporting in a way

that would not necessarily be applicable to clinical services. No assessment on physician reimbursement, workloads, or whole care pathway costs was made. Our study does not fully address the barriers to a sustainable service, although 'mission creep' and a gradual increase in scanning times once the initial implementation is over was observed.

In conclusion, CMR core clinical information can be provided more quickly, easily, and less expensive. Our rapid CMR protocol can be delivered globally to major centres in LMICs (national/state capitals) and HICs alike. When incorporated into a clinical service and linked to an education/training programme with a supporting international network and partnership, it can deliver high diagnostic quality and improve patient care. The initial data suggest that these changes can be sustainable.

Authors' contributions

K.D.M., J.M., and J.M.W. contributed to the idea, design of the study, and drafted the manuscript. All authors contributed to data collection, analysis, interpretations, and re-drafting of this manuscript. K.D.M., A.H., and J.M. drafted the study protocol and analysis plan. K.D.M., J.M.W., and J.M. drafted the manuscript and checked data for accuracy of information of the study.

Supplementary material

Supplementary material is available at *European Heart Journal* online.

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Data availability

Data used for the current study are all anonymized and are available to other researchers upon reasonable request and approval of the collaborative groups.

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