Myocardial Infarct Size by CMR in Clinical Cardioprotection Studies
Insights From Randomized Controlled Trials

Heerajnarain Bulluck, PhD,a,b,c Matthew Hammond-Haley, MSc,a Shane Weinmann, MBBS,a Roberto Martinez-Macias, MSc,a Derek J. Hausenloy, PhD a,b,c,d

ABSTRACT

OBJECTIVES The aim of this study was to review randomized controlled trials (RCTs) using cardiac magnetic resonance (CMR) to assess myocardial infarct (MI) size in reperfused patients with ST-segment elevation myocardial infarction (STEMI).

BACKGROUND There is limited guidance on the use of CMR in clinical cardioprotection RCTs in patients with STEMI treated by primary percutaneous coronary intervention.

METHODS All RCTs in which CMR was used to quantify MI size in patients with STEMI treated with primary percutaneous coronary intervention were identified and reviewed.

RESULTS Sixty-two RCTs (10,570 patients, January 2006 to November 2016) were included. One-third did not report CMR vendor or scanner strength, the contrast agent and dose used, and the MI size quantification technique. Gadopentetate dimeglumine was most commonly used, followed by gadoterate meglumine and gadobutrol at 0.20 mmol/kg each, with late gadolinium enhancement acquired at 10 min; in most RCTs, MI size was quantified manually, followed by the 5 standard deviation threshold; dropout rates were 9% for acute CMR only and 16% for paired acute and follow-up scans. Weighted mean acute and chronic MI sizes (≤12 h, initial TIMI [Thrombolysis in Myocardial Infarction] flow grade 0 to 3) from the control arms were 21 ± 14% and 15 ± 11% of the left ventricle, respectively, and could be used for future sample-size calculations. Pre-selecting patients most likely to benefit from the cardioprotective therapy (≤6 h, initial TIMI flow grade 0 or 1) reduced sample size by one-third. Other suggested recommendations for standardizing CMR in future RCTs included gadobutrol at 0.15 mmol/kg with late gadolinium enhancement at 15 min, manual or 6-SD threshold for MI quantification, performing acute CMR at 3 to 5 days and follow-up CMR at 6 months, and adequate reporting of the acquisition and analysis of CMR.

CONCLUSIONS There is significant heterogeneity in RCT design using CMR in patients with STEMI. The authors provide recommendations for standardizing the assessment of MI size using CMR in future clinical cardioprotection RCTs. (J Am Coll Cardiol Img 2017;10:230–40) © 2017 The Authors. Published by Elsevier on behalf of the American College of Cardiology Foundation. This is an open access article under the CC BY license (http://creativecommons.org/licenses/by/4.0/).
reperfusion injury to reduce MI size in patients with STEMI treated with PPCI (3).

Late gadolinium enhancement (LGE) by CMR is considered the gold standard for MI size quantification (4). MI size (5), microvascular obstruction (MVO) (6), and myocardial salvage (7) assessed by CMR performed in the first few days post-PPCI have all been shown to be strongly prognostic. As a result, CMR is increasingly being used for surrogate endpoints in RCTs (3). A recent meta-analysis of 2,622 patients from 10 RCTs found that MI size measured by CMR or single-photon computed tomography within 1 month post-PPCI showed that for every 5% increase in MI size, there was a 20% increase in the relative hazard ratio for 1-year hospitalization for heart failure and all-cause mortality (5).

Despite CMR endpoints being quite robust and their ability to keep sample size small (3), there is limited guidance on its use. In this study, we reviewed all published RCTs in this field so far, and we provide recommendations for standardizing the use of CMR in future clinical cardioprotection RCTs.

METHODS

We performed a comprehensive systematic search in the MEDLINE and Embase databases via Ovid up to November 23, 2016. PubMed and the Web of Science, editorsials, and the reference lists of included RCTs were also screened. Further details of the search terms and the study screening and selection as per the Preferred Reporting Items for Systematic Reviews and Meta-Analyses are provided in the Online Appendix. The flow diagram is shown in Figure 1.

The inclusion criteria were as follows: 1) RCTs investigating cardioprotective strategies aimed at reducing MI size; 2) including patients presenting within 12 h of symptom onset; 3) MI size measured by CMR; and 4) RCTs with full-text reports in English. The exclusion criteria were as follows: 1) RCTs selecting patients on the basis of admission ejection fraction; 2) RCTs performing post hoc analysis on other included RCTs; 3) observational studies; and 4) RCTs only reporting left ventricular (LV) ejection fraction by CMR.

STUDY SELECTION AND DATA EXTRACTION. Four authors (H.B., M.H.-H., S.W., R.M.M.) independently identified suitable RCTs from the screened reports and extracted all data. Data on study characteristics, patient eligibility criteria, CMR scanner and contrast agent used, MI size quantification technique used, and mean MI size and SD in the control arm were collected. RCTs reporting both acute MI size and follow-up MI size (≥1 month) were recorded. The primary outcome of interest was MI size, expressed as a percentage of LV volume or mass (%LV).

STATISTICAL ANALYSIS. Categorical data are reported as frequencies and percentages. For sample size calculation, to ensure normally distributed MI size in the control arms, only RCTs reporting mean MI size as %LV were included, and an unpaired Student t test was used. Only RCTs including STEMI in all coronary territories were included, and they were grouped according to duration of symptoms (<6 or <12 h) and TIMI (Thrombolysis In Myocardial Infarction) flow grade 0 or 1 or 1 to 3 in the culprit arteries pre-PPCI if at least 3 RCTs were present in the subgroups. The mean and SD from each control arm were weighted against its respective sample size to obtain representative pooled mean acute and follow-up MI sizes using RevMan version 5.2 (Nordic Cochrane Center, Copenhagen, Denmark). Sample-size calculation was performed for 90% power, a 2-sided alpha value of 0.05, and expected effect sizes of 20%, 25%, and 30% using Stata/IC version 12.1 (StataCorp LP, College Station, Texas). The expected sample sizes were provided for each group after accounting for potential dropouts.

RESULTS

Our initial search identified 399 reports and 62 RCTs, involving 10,570 patients, that met the inclusion criteria. Details of the 62 included RCTs are provided in Online Table 1.

CHARACTERISTICS OF THE RCTs. The number of RCTs using CMR has steadily increased since 2009 (Figure 2). There was an average of 8 RCTs reported per year between 2010 and 2016. The majority of RCTs came from Europe (41 of 62 [66%], 6,880 patients), and 35 of 62 (56%) were single-center RCTs.

MI size as %LV was reported in 44 RCTs (71%). Five RCTs (8%) reported MI size as percentage of the area at risk (AAR), 5 (8%) in grams, and 7 RCTs (12%) reported myocardial salvage index (MSI).

The majority of RCTs included: 1) patients presenting within 12 h of symptom onset (32 of 62 [52%]); 2) STEMI in all epicardial territories (49 of 62 [79%]); and 3) patients with TIMI flow pre-PPCI (33 of 62 [53%]), as shown in Figure 3.

Forty-six of the RCTs (74%) used 1.5-T scanners, and a minority (2 RCTs [3%]) used 3-T scanners. Four RCTs (7%) used a combination of 1.5- and 3-T scanners. Ten RCTs (16%) did not specify the field...
strength of the CMR scanners used, and 8 of 10 (80%) of those were multicenter RCTs.

Thirty-six of the RCTs (58%) used single CMR vendors, and 7 RCTs (11%) used multivendor scanners at different recruitment sites. Nineteen RCTs (31%) did not specify the vendors used, the majority of which (15 RCTs [75%]) were multicenter RCTs.

Paired acute and follow-up CMR was performed in 26 RCTs (42%), acute CMR only in 24 RCTs (39%), and follow-up CMR only in 12 RCTs (19%).

**GADOLINIUM-BASED CONTRAST AGENT AND DOSE.** Gd-dopentetate dimeglumine (Gd-DTPA) (Magnevist, Bayer Healthcare, Berlin, Germany), gadoterate meglumine (Gd-DOTA) (Dotarem, Guerbet, Roissy Cedex, France), and gadobutrol (Gadovist, Bayer Healthcare) were used in 18 (29%), 12 (19%), and 9 (14%) of the 62 RCTs, respectively. Nineteen of 62 RCTs (31%) did not specify the gadolinium-based contrast agent (GBCA) used, of which 15 of 19 (79%) were multicenter RCTs. Online Table 2 summarizes the doses of the GBCAs and the reported timing for LGE acquisition post-contrast. The most common doses for Gd-DTPA (Magnevist), gadobutrol (Gadovist), and Gd-DOTA (Dotarem) were similar at 0.20 mmol/kg, with LGE acquisition starting at 10 min onward.

**TIMING OF CMR ACQUISITION.** The distribution of the timings to acquire acute and follow-up CMR is shown in Figure 4. Of 50 RCTs with acute CMR acquisition, only 1 did not specify the exact timing of the scan. There was a wide range of timings to acquire the acute CMR scan, with the majority (82%) acquired within the first 7 days post-PPCI and the most common timing being 3 to 5 days post-STEMI (15 of 50 RCTs [30%]). There was also a wide range of timings (1 to 9
months) to acquire follow-up MI size, with the most common timing being 6 months (31 of 37 RCTs [30%]).

**MI Size Analysis.** Seventeen of 25 of the multicenter RCTs (68%) specified using a CMR core laboratory. The majority of the RCTs (29 of 62 [47%]) used specialist software (Qmass, Medis Medical Imaging Solutions, Leiden, the Netherlands; CVI42, Circle Cardiovascular Imaging Inc., Calgary, Canada; Segment, Medviso, Lund, Sweden; and CAAS, Pie Medical Imaging B.V., Maastricht, the Netherlands) for MI size quantification, 7 RCTs (11%) used the scanners’ own software, and 5 of 62 (8%) used shareware (ImageJ, National Institutes of Health, Bethesda, Maryland; OsiriX, Geneva, Switzerland). Twenty-one of 62 RCTs (34%) did not specify the analysis tool used, of which 9 of 21 (43%) were multicenter RCTs.

Manual contouring was the most commonly used method (22 of 62 RCTs [36%]), followed by 5-SD (8 of 62 RCTs [13%]) and full width half maximum (FWHM) (6 of 62 RCTs [10%]). Sixteen of 62 RCTs (27%) did not specify the method used for MI quantification, of which 9 of 16 (56%) were multicenter RCTs. Figure 5 shows the distribution of the quantification techniques for MI size used in these RCTs.

**Distribution of Culprit Vessel and TIMI Flow Pre-PPCI.** Among the RCTs including all culprit vessels (n = 6,020), 45% of the patients had STEMI in the left anterior descending coronary artery territory. Among the RCTs including all TIMI flow grades (n = 3,229), 73% of patients had TIMI flow grades of 0 or 1 pre-PPCI.

**Effect Size for Reduction in MI Size in RCTs With Positive Results.** Seventeen RCTs were positive for a reduction in MI size. Of the RCTs reporting MI size as %LV, the median reduction in acute MI size was 34% (mode 35%; range 14% to 38%). As for chronic MI size, both the median and mode reduction in MI size were 18% (range 14% to 46%).

**MI Size in the Control Arm.** RCTs reporting mean ± SD, including STEMI presenting within 6 or 12 h, in all coronary territories, and with TIMI flow grades of 0 or 1 or 0 to 3 pre-PPCI were selected to obtain representative weighted mean MI size, expressed as %LV in the control arm for both acute (20 RCTs) and follow-up (18 RCTs) CMR (summarized in Online Tables 3 and 4). The expected sample sizes on the basis of 20%, 25%, and 30% reductions in MI size are summarized in Table 1. The expected sample sizes are provided on the basis of 90% power and a 2-sided alpha of 0.05.

The estimated sample size was smallest for RCTs including patients with STEMI within 6 h of symptom onset and with pre-PPCI TIMI flow grades of 0 or 1 and was largest for RCTs recruiting patients within 12 h of symptom onset and with any TIMI flow grades pre-PPCI, and this was driven by a wider SD in the latter group. For the same effect size, RCTs planning to use chronic MI would require on average 30% more patients than if acute MI were chosen as the endpoint.

**Dropout Rates.** In RCTs using CMR for the primary endpoint, the average dropout rate was 9% for RCTs using acute CMR only, 13% for RCTs acquiring follow-up CMR only, and 16% for those with paired acute and follow-up scans.

**Discussion.** On the basis of the 62 RCTs reviewed, substantial heterogeneity exists in trial design with respect to patient selection, timing of acute and follow-up scans, GBCA dose, timing of LGE acquisition, and method for MI quantification. Among those RCTs not reporting scanner strength and vendor, GBCA used, and software used for MI size quantification, the majority were multicenter RCTs.

The other major findings were as follows: 1) acute CMR was most commonly performed at 3 to 5 days and follow-up CMR at 6 months; 2) Gd-DOTA, gadobutrol, and Gd-DTPA were most commonly used at a similar dose of 0.20 mmol/kg, with LGE acquired at 10 min; 3) MI size was quantified manually in most RCTs,
followed by the 5-SD threshold; 4) STEMI in the left anterior descending coronary artery territory accounted for one-half and those with TIMI flow grades of 0 or 1 accounted for three-quarters of all STEMIs entering these RCTs and could be taken into consideration for sample-size calculation in specific circumstances; 5) for positive RCTs, the most commonly seen effect sizes were 34% for acute MI and 18% for chronic MI (<1 month) size reduction; 6) using the control arms from RCTs reporting mean MI size, theoretical sample sizes for future clinical cardioprotection studies were provided; and 7) the dropout rate was highest for RCTs performing paired acute and follow-up scans, followed by those performing follow-up CMR only, and smallest in the RCTs performing acute CMR only.

**ROBUSTNESS OF CMR- DERIVED SURROGATE ENDPOINTS.** CMR has recently emerged as a robust tool not only for MI size quantification (5) but also to provide additional information on the AAR (to derive MSI)(8), LV ejection fraction(9), and MVO(6,10) from a single scan. MI size, myocardial salvage, and LV ejection fraction by CMR are highly reproducible (reducing sample size) (8,9,11,12) and are strongly linked to prognosis (5,7,13–15). Furthermore, with recent advances in mapping techniques, multiparametric information can be obtained from native (for the AAR) (16) and post-contrast T1 maps (to derive extracellular volume fraction maps to interrogate the remote
myocardium) (17,18), T2 maps (AAR) (16) and T2* maps (intramyocardial hemorrhage and residual myocardial iron) (19,20). These additional mapping parameters provide more robust measurement of the edema-based AAR (when compared with angiographic scores) (21), pathophysiological insights in post-STEMI LV remodeling (17–19,22), and prognostic information (20). Unlike single-photon emission computed tomography, CMR has superior spatial resolution (14), does not involve radiation, and requires only a single examination, when the patient is relatively stable. Furthermore, MVO and MI size by CMR have been shown to be more prognostic compared with MI size by single-photon emission computed tomography (14). The superior spatial resolution of CMR also allows detection of small MIs that could be missed by relying on wall motion abnormalities on echocardiography alone (23), interrogation of the peri-infarct zone (24), and hypointense core of the MVO (25), which are all prognostic (24–26). Therefore, it is not surprising that CMR endpoints have gained popularity for use in several RCTs (3).

OPTIMAL TIMING OF ACUTE CMR POST-STEMI. Preclinical studies have shown that performing CMR too early post-reperfusion (day 1) leads to an overestimation of MI size because of a combination of edema and partial volume effect (27). In the clinical setting, acute MI size has also been shown to be dynamic and to decrease significantly in size between days 1 and 7 (28,29) but is stable between days 3 and 4 (8). Recently, Carrick et al. (30) showed that acute MI size was stable between days 1 and 3 and subsequently reduced in size by day 10. Furthermore, late MVO has also been shown to be stable between days 1 and 3 and to reduce in size by day 10 (20), and the persistence of late MVO at 1 week following STEMI was more prognostic (31). The detection of intramyocardial hemorrhage has been shown to peak at day 3, and reduced in size and incidence by day 10, and intramyocardial hemorrhage was more prognostic than MVO (20).

There is no established method to assess the AAR (a pre-requisite to calculate MSI) in the clinical setting by CMR. T2 mapping has recently emerged as more robust than T2-weighted imaging for assessing edema-based AAR (32). Some controversies exist as to whether edema within the AAR follows a bimodal pattern (33) and whether T2-weighted imaging delineates the AAR at all (34) in the preclinical setting. It was previously believed that edema was stable in the first week of a MI (28) using T2-weighted imaging. However, Carrick et al. (30) recently showed that the extent of myocardial edema followed a unimodal pattern and peaked at day 3 in patients.

On the basis of the clinical research published so far, if the CMR scan is performed at <3 days, MI size would be overestimated and AAR would be underestimated. If the CMR scan is performed at >5 days, the edema-based AAR may be underestimated. Therefore, acquiring the acute CMR scan at 3 to 5 days following STEMI, as performed in most cardioprotection RCTs in this review, may be the optimal time to undertake the acute CMR scan, as illustrated in Figure 6.

OPTIMAL TIMING OF FOLLOW-UP CMR. Chronic MI size has been shown to be stable when performed between 1 month and up to 1 year (31). Paired acute and follow-up scans also provide information on post-STEMI LV remodeling, which occurs by 2 months, although the process may continue for up to 1 year (31). However, waiting too long between the acute and follow-up CMR scans may increase the likelihood of patients’ dropping out of the studies. Therefore, in RCTs aiming to assess both chronic MI size and LV remodeling, performing the follow-up scan at 6 months, as done in most of the RCTs reviewed in this study, would be optimal.

The most common myocardial infarct (MI) size quantification method was manual delineation, followed by 5-SD and full width half maximum (FWHM). However, 27% of randomized controlled trials did not specify the method used.

**FIGURE 5** Quantification Techniques for MI Size

![Quantification Techniques for MI Size](image-url)
OPTIMAL GBCA DOSE AND TIMING OF LGE ACQUISITION. Gadobutrol has been shown to delineate the infarcted myocardium better from the blood pool (better contrast-to-noise ratio) compared with Gd-DTPA (35) and Gd-DOTA (36) in chronic MI. Furthermore, gadobutrol has been shown to differentiate the infarcted myocardium from the LV blood pool as early as 9 min. The recommended relaxivity-adjusted standard doses are 0.22 mmol/kg for Gd-DOTA (36), 0.15 mmol/kg for gadobutrol (35,36), and 0.20 mmol/kg for Gd-DTPA (35).

Acquiring LGE too early (<8 min) post-GBCA administration has been shown to result in overestimation of MI size (37), and acquiring LGE at 25 min for acute MI size was a better predictor of LV recovery (38). In most RCTs included in this study, LGE imaging was performed at 10 min. Performing comprehensive CMR in patients with STEMI can be time-consuming, and to minimize patient discomfort and prevent dropout of patients, every attempt is made to keep scan time to a minimum. Therefore, acquiring LGE images 15 min post-GBCA administration in future RCTs would be a good compromise.

METHOD FOR QUANTIFYING MI SIZE. There is currently no established gold-standard semiautomated technique for MI size quantification. Manual contouring is considered the reference standard (4,11), but it can be time-consuming and may be subjective. The 5-SD approach is currently recommended, as it may improve reproducibility (4).

Although FWHM has emerged as being the most reproducible (39,40), it has been shown to underestimate acute and chronic MI size (40). Some studies showed that 5-SD was promising (39-41), but others

TABLE 1 Sample-Size Estimation for Future RCTs Investigating Therapy for Reducing MI Size by CMR

| Patient Inclusion Criteria | Pooled MI Size by CMR in the Control Group From Previous RCTs (as %LV) | Sample Size per Group After Accounting for Potential Dropouts (9% for Acute CMR and 13% for Follow-Up CMR) for 90% Power With a 2-Sided Alpha of 0.05 |
|----------------------------|-----------------------------------------------------------------------|--------------------------------------------------------------------------------------------------|
|                            | MI Size | Sample Size | MI Size | Sample Size | MI Size | Sample Size |
| TIMI Flow                  |         |             |         |             |         |             |
| Symptom Onset (h)          |         |             |         |             |         |             |
| Acute scan only (including STEMI in all coronary territories) | | | | | | |
| 0 or 1                     | ≥6       | 22 (20-24)  | 12      | 3 RCTs (136 patients): Chan (2012) García-Dorado (2014) Waltenberg (2014) | 17.6  | 171  | 16.5  | 110  | 15.4  | 76   |
| O or 1                     | ≤12      | 24 (22-26)  | 13      | 3 RCTs (189 patients): Freixa (2011) Mewton (2013) Siddiqi (2014) White (2015) | 19.2  | 169  | 18.0  | 108  | 16.8  | 75   |
| O-3                        | ≤12      | 21 (19-22)  | 14      | 5 RCTs (305 patients): Yoon (2013) Ko (2014) Hoole (2015) McCann (2015) Liu (2016) | 16.8  | 265  | 15.8  | 164  | 14.7  | 85   |
| Chronic scan only (including STEMI in all coronary territories) | | | | | | |
| 0 or 1                     | ≥6       | 15 (14-17)  | 9       | 4 RCTs (152 patients): Tarantini (2012) Atar (2015) Waltenberg (2014) Roos (2016) | 12.0  | 215  | 11.3  | 215  | 10.5  | 96   |
| O or 1                     | ≤12      | 15 (13-16)  | 9       | 4 RCTs (186 patients): Lonborg (2010) Freixa (2011) Lonborg (2012) Siddiqi (2014) | 12.0  | 215  | 11.3  | 216  | 10.5  | 96   |
| O-3                        | ≤12      | 15 (14-17)  | 11      | 4 RCTs (284 patients): Song (2009) Ranchord (2012) Roolvink (2016) Ko (2014) | 12.0  | 320  | 11.3  | 205  | 10.5  | 142  |

The references for the RCTs included in each row are provided in Online Table 5. CI = confidence interval; CMR = cardiac magnetic resonance; MI = myocardial infarction; %LV = percentage of left ventricular mass or volume; PPCI = primary percutaneous coronary intervention; RCT = randomized controlled trial; TIMI = Thrombolysis in Myocardial Infarction.
showed that it overestimated MI size (42,43). The n-SD technique requires the remote myocardium to be appropriately nulled and free of artifacts. A manual region of interest is required in the remote myocardium, and this can be a source of variability. The Otsu technique does not require a region of interest as a reference and has been shown to accurately delineate MI size (40). But 2 subsequent studies showed that Otsu overestimated MI size (42,43).

The most common method used to quantify MI size in the RCTs was manual contouring, followed by 5-SD and FWHM. We recently showed that FWHM underestimates chronic MI size in those with MVO on the acute scan because of very high extracellular volume in the area previously occupied by MVO and should be avoided for RCTs assessing chronic MI size (42).

The 6-SD method appears the most promising and has been shown to have the highest accuracy to predict segment wall recovery in patients with chronic myocardial infarction (44) and is similar to manual quantification in patients with both acute and chronic myocardial infarction (39,42). The 6-SD approach also performed well when using 2 different LGE sequences (42).

IMPACT OF PATIENT SELECTION AND TIMING OF CMR ON SAMPLE SIZE. In Table 1 we provide guidance for expected sample sizes, depending on the desired effect size, after accounting for expected dropout rates when planning future RCTs. For example, if an RCT includes patients with all TIMI flow grades pre-PPCI and presenting within 12 h of symptom onset, for an expected 25% reduction in acute MI size, 164 patients would be needed in each arm. If the endpoint were to be changed to chronic MI size, then for the same effect size, the sample size for each arm would increase to 205. However, we have observed that the realistic effect size for chronic MI size on average is lower (18%), and this would increase the sample size further.

If only patients with pre-PPCI TIMI flow grades of 0 or 1 and within 6 h of symptom onset were included, the sample size for each arm would be reduced by about one-third to 110 patients. However, only three-quarters of patients with STEMI present with TIMI flow grades of 0 or 1, and therefore at least one-third more patients would need to be screened to achieve the desired sample size.

RECOMMENDATIONS FOR FUTURE CARDIOPROTECTION RCTs IN PATIENTS WITH STEMI. Careful patient selection to include those patients most likely benefit from a novel cardioprotective therapy for reducing MI size (ischemic time <6 h and pre-PPCI TIMI flow grade 0 or 1) can reduce sample size by one-third, but more patients would need to be screened.

Acute MI size should be preferred to chronic MI size as a surrogate endpoint, because acute MI size is already prognostic (5), and this would reduce sample size and result in fewer dropouts.

Acute CMR should ideally be performed on day 3, 4, or 5. When a follow-up scan is planned, 6 months would be the optimal timing for data on both chronic MI size and LV remodeling.

The preferred GBCA should be gadobutrol (superior contrast-to-noise ratio) (35) at 0.15 mmol/kg (recommended relaxivity-adjusted standard dose) (35,36), with LGE performed at 15 min.

Manual quantification of MI size by experienced operators at a core laboratory level is considered the gold standard (4). When this is not practical, the 6-SD threshold would be an alternative option. However, any semiautomated technique is likely to be influenced by the LGE image quality, and therefore, each
center may need to validate the performance of these semiautomated techniques at the respective center or core laboratory.

In the absence of a gold-standard method for the AAR, MI size should be reported as %LV, which has also been shown to be prognostic (5). Although MSI is reported to be a better surrogate, in adequately powered RCTs, the AAR should be adequately balanced in both arms, and therefore expressing MI size as %LV would be acceptable.

Finally, all RCTs, in particular multicenter RCTs, should provide adequate details on the execution and quantification of MI size in RCTs to allow fair interpretation and comparison of study results and provide more reliable MI size in the control arms for future sample-size calculation.

**STUDY LIMITATIONS.** We included only RCTs in this review, but some of the recommendations could also be applied to observational studies to standardize the conduct of CMR, making comparison among studies easier.

We concentrated only on MI size. Using MSI as a surrogate endpoint is considered more sensitive to assess the effectiveness of cardioprotective therapies (45) and has recently been shown to reduce sample size (46). However, given the lack of consensus (33,34) and ongoing validation of the AAR methodologies, we did not evaluate the use of MSI in the RCTs published so far.

MVO has been shown to be more prognostic than MI size (15), but we did not assess the definition of MVO in the included RCTs. However, if the performance and quantification of LGE for MI size were standardized, this would also standardize MVO (late) quantification.

The recommended dose for gadobutrol is based on the recommended relaxivity-adjusted standard dose, and ideally a head-to-head comparison of the different doses should be performed. However, this is challenging in the acute MI setting, given the dynamic nature of MI size within the first week. The most commonly seen effect sizes for acute and chronic MI size are related to the intervention they were subjected to and are meant to serve as a guide for realistic effect sizes in the acute and chronic MI setting.

**CONCLUSIONS**

There is significant heterogeneity in the design of RCTs using CMR to quantify MI size in clinical cardioprotection studies in reperfused patients with STEMI. Here, we have provided insights from RCTs published so far and have offered recommendations for standardizing the assessment of MI size by CMR to optimize the design of future studies assessing the efficacy of cardioprotective strategies.

**ADDRESS FOR CORRESPONDENCE:** Prof. Derek J. Hausenloy, Cardiovascular and Metabolic Diseases Program, Duke-National University of Singapore Graduate Medical School Singapore, 8 College Road, Singapore 169857, Singapore. E-mail: derek.hausenloy@duke-nus.edu.sg.

**REFERENCES**

1. Jimenez-Candil J, Diaz-Castro O, Barrabes JA, et al. Update on ischemic heart disease and critical care cardiology. Rev Esp Cardiol 2013;66:198-204.
2. Torabi A, Cleland JG, Khan NK, et al. The timing of development and subsequent clinical course of heart failure after a myocardial infarction. Eur Heart J 2008;29:859-70.
3. Desch S, Eitel I, de Waha S, et al. Cardiac magnetic resonance imaging parameters as surrogate endpoints in clinical trials of acute myocardial infarction. Trials 2011;12:204.
4. Schulz-Menger J, Bluemke DA, Bremerich J, et al. Standardized image interpretation and post processing in cardiovascular magnetic resonance: Society for Cardiovascular Magnetic Resonance (SCMR) Board of Trustees Task Force on standardized Post Processing. J Cardiovasc Magn Reson 2013;15:35.
16. Carberry J, Carrick D, Haig C, et al. Remote zone extracellular volume and left ventricular remodeling in survivors of ST-elevation myocardial infarction. Hypertension 2016;68:385-91.

17. Bulluck H, Rosmini S, Abdel-Gadir A, et al. Residual myocardial iron following intramyocardial hemorrhage during the convalescent phase of reperfused ST-segment-elevation myocardial infarction and adverse left ventricular remodeling. Circ Cardiovasc Imaging 2016;9:e004940.

18. Carrick D, Haig C, Ahmed N, et al. Myocardial hemorrhage after acute reperfused ST-segment-elevation myocardial infarction: relation to microvascular obstruction and prognostic significance. Circ Cardiovasc Imaging 2016;9:e004148.

19. Wright J, Adlaenssens T, Dymarkowski S, Desmet W, Bogert J. Quantification of myocardial area at risk with T2-weighted CMR: comparison with contrast-enhanced CMR and coronary angiography. J Am Coll Cardiol Img 2009;2:825-31.

20. Carrick D, Haig C, Rauhalammi S, et al. Pathophysiology of LV remodeling in survivors of STEMI: inflammation, remote myocardium, and prognosis. J Am Coll Cardiol Img 2015;8:779-89.

21. Jaarsma C, Schalla S, Cheriex EC, et al. Incremental value of cardiovascular magnetic resonance over echocardiography in the detection of acute and chronic myocardial infarction. J Cardiovasc Magn Reson 2013;15:5.

22. Van AT, Shayne AJ, Brown KA, et al. Characterization of post-infarct zone by contrast-enhanced cardiac magnetic resonance imaging is a powerful predictor of post-myocardial infarction mortality. Circulation 2006;114:3227-34.

23. Carrick D, Haig C, Rauhalammi S, et al. Prognostic significance of infarct core pathology revealed by quantitative non-contrast in comparison with contrast cardiac magnetic resonance imaging in reperfused ST-elevation myocardial infarction survivors. Eur Heart J 2016;37:1044-59.

24. Kwong RY, Sattar H, Wu H, et al. Incidence and prognostic implication of unrecognized myocardial scar characterized by cardiac magnetic resonance in diabetic patients without clinical evidence of myocardial infarction. Circulation 2008;118:1011-20.

25. Jablonskiwski R, Enblom H, Kanski M, et al. Contrast-enhanced CMR overestimates early myocardial infarct size: mechanistic insights using ECV measurements on day 1 and day 7. J Am Coll Cardiol Img 2015;8:1379-89.

26. Dell’Armellina E, Karia N, Lindsay AC, et al. Dynamic changes of edema and late gadolinium enhancement in post-infarct cardiac functional recovery after an acute myocardial infarction. J Cardiovasc Magn Reson 2016;18:26-36.

27. Ibrahim T, Hackl T, Nekolla SG, et al. Acute myocardial infarction: serial cardiac MR imaging shows a decrease in delayed enhancement of the myocardium during the 1st week after reperfusion. Radiology 2010;254:88-97.

28. Cormier R, Dansby S, Adler B, et al. Temporal evolution of myocardial hemorrhage and edema in patients after acute ST-segment elevation myocardial infarction: pathophysiological insights and clinical implications. J Am Heart Assoc 2016;5:e002834.

29. Orn S, Manhenke C, Greve OJ, et al. Microvascular obstruction is a major determinant of infarct healing and subsequent left ventricular remodelling following primary percutaneous coronary intervention. Eur Heart J 2009;30:1978-85.

30. Verhaert D, Thavendiranathan P, Giri S, et al. Direct T2 quantification of myocardial edema in acute ischemic injury. J Am Coll Cardiol Img 2011;4:269-78.

31. Bellenger NG, Groothues F, Smith GC, Pennell DJ. Quantification of right and left ventricular function by cardiovascular magnetic resonance. Herz 2000;25:392-9.

32. van Kranenburg M, Magni M, Thiele H, et al. Prognostic value of microvascular obstruction and infarct size, as measured by CMR in STEMI patients. J Am Coll Cardiol Img 2014;7:930-9.

33. Thiele H, Kappl MJ, Conradi S, Niebauer J, Hambrecht R, Schuler G. Reproducibility of chronic and acute infarct size measurement by delayed enhancement-magnetic resonance imaging. J Am Coll Cardiol Img 2014;7:643-5.

34. Lonborg J, Veijstrup N, Mathiasen AB, Thomsen C, Jensen JS, Engstroem T. Myocardial area at risk and salvage measured by T2-weighted cardiovascular magnetic resonance: reproducibility and comparison of two T2-weighted protocols. J Cardiovasc Magn Reson 2011;13:50.

35. White HD, Norris MA, Brandt PW, Whitlock RML, Wild CJ. Left ventricular end-systolic volume as the major determinant of survival after recovery from myocardial infarction. Circulation 1987;76:44-51.

36. Hadamitzky M, Langhans B, Hausleiter J, et al. Prognostic value of late gadolinium enhancement in cardiovascular magnetic resonance imaging after acute ST-elevation myocardial infarction in comparison with single-photon emission tomography using Tc99m-sestamibi. Eur Heart J Cardiovasc Imaging 2014;15:216-25.

37. Eitel I, de WAHA S, Wohrle J, et al. Comprehensive prognosis assessment by CMR imaging after ST-segment elevation myocardial infarction. J Am Coll Cardiol Img 2014;7:1217-26.

38. Bulluck H, White SK, Rosmini S, et al. T1 mapping and T2 mapping at 3T for quantifying the area-at-risk in reperfused STEMI patients. J Cardiovasc Magn Reson 2015;17:73.

39. Bulluck H, Rosmini S, Abdel-Gadir A, et al. Automated extracellular volume fraction mapping provides insights into the pathophysiology of left ventricular remodeling post-reperfusion ST-elevation myocardial infarction. J Am Heart Assoc 2016;5:e003555.
automated techniques for quantifying acute and chronic myocardial infarction by cardiovascular magnetic resonance. Open Heart 2016;3:e000535.

43. Khan JN, Nazir SA, Horsfield MA, et al. Comparison of semi-automated methods to quantify infarct size and area at risk by cardiovascular magnetic resonance imaging at 1.5T and 3.0T field strengths. BMC Res Notes 2015;8:52.

44. Beek AM, Bondarenko O, Afsharzada F, van Rossum AC. Quantification of late gadolinium enhanced CMR in viability assessment in chronic ischemic heart disease: a comparison to functional outcome. J Cardiovasc Magn Reson 2009;11:6.

45. Botker HE, Kaltoft AK, Pedersen SF, Kim WY. Measuring myocardial salvage. Cardiovasc Res 2012;94:266-75.

46. Engblom H, Heiberg E, Erlinge D, et al. Sample size in clinical cardioprotection trials using myocardial salvage index, infarct size, or biochemical markers as endpoint. J Am Heart Assoc 2016;5:e002708.