OPEN ACCESS

ORIGINAL RESEARCH ARTICLE

Daily remote ischaemic conditioning following acute myocardial infarction: a randomised controlled trial

Andrew Peter Vanezis,¹ Jayanth Ranjit Arnold,¹ Glenn Rodrigo,¹ Florence Y Lai,¹ Radek Debiec,¹ Sheraz Nazir,¹ Jamal Nasir Khan,¹ Leong L Ng,¹ Kamal Chitkara,² John G Coghlan,³ Simon Lee Hetherington,⁴ Gerry P McCann,¹ Nilesh J Samani¹

► Additional material is published online only. To view please visit the journal online (<http://dx.doi.org/10.1136/heartjnl-2018-313091>).

¹Department of Cardiovascular Sciences, University of Leicester and the NIHR Leicester Biomedical Research Centre, University Hospitals of Leicester NHS Trust, Glenfield Hospital, Leicester, UK

²Royal Derby Hospital, Derby, UK

³Royal Free Hospital, London, UK

⁴Kettering General Hospital, Kettering, UK

Correspondence to

Dr Andrew Peter Vanezis, Department of Cardiovascular Sciences, University of Leicester, BHF Cardiovascular Research Centre Glenfield Hospital, Leicester LE3 9QP, UK; andrew.vanezis@nhs.net

Received 3 February 2018

Revised 21 April 2018

Accepted 23 April 2018

Published Online First

10 May 2018

ABSTRACT

Background Remote ischaemic conditioning (rIC) is a cardioprotective tool which has shown promise in preclinical and clinical trials in the context of acute ischaemia. Repeated rIC post myocardial infarction may provide additional benefits which have not previously been tested clinically.

Methods The trial assessed the role of daily rIC in enhancing left ventricular ejection fraction (LVEF) recovery in patients with impaired LVEF (<45%) after ST segment elevation myocardial infarction (STEMI) treated with primary percutaneous coronary intervention (P-PCI). Patients were recruited from four UK hospitals and randomised to receive either 4 weeks of daily rIC or sham conditioning using the autoRIC Device (CellAegis) starting on day 3 post P-PCI. The primary endpoint was the improvement in LVEF over 4 months assessed by cardiac MRI (CMR). Seventy-three patients (38 cases, 35 controls) completed the study.

Results The treatment and control groups were well matched at baseline including for mean LVEF (42.8% vs 44.3% respectively, $p=0.952$). There was no difference in the improvement in LVEF over 4 months between the treatment and control groups ($4.8\pm 7.8\%$ vs $4.6\pm 5.9\%$ respectively, $p=0.924$). No differences were seen in the secondary outcome measures including changes in infarct size and left ventricular end-diastolic and systolic volumes, major adverse cardiac and cerebral event, mean Kansas City Cardiomyopathy Questionnaire score and change in N-terminal pro-brain natriuretic peptide levels.

Conclusions Daily rIC starting on day 3 and continued for 4 weeks following successful P-PCI for STEMI did not improve LVEF as assessed by CMR after 4 months when compared with a matched control group.

Trial registration number NCT0166461.

The mechanisms through which rIC confers cardioprotection have not been fully elucidated but numerous animal and human models have highlighted the importance of humoral signalling acting in conjunction with neural and inflammation pathways.³ When used after the onset of ST segment elevation myocardial infarction (STEMI) and either prior to revascularisation or at the onset of reperfusion with primary percutaneous coronary intervention (P-PCI), rIC has been shown in early clinical studies to reduce the final infarct size as assessed by biomarker release, reduce the extent of left ventricular (LV) dysfunction as well as reduce major adverse cardiac and cerebral events (MACCE)^{4–8} although it should be noted that not all studies have been positive.⁹

Recently, experimental animal studies have demonstrated that in addition to its acute effect, repeated bouts of daily rIC applied after a myocardial infarction (MI) may also be beneficial. In a rat model of acute MI, Wei *et al*¹⁰ induced MI by left anterior descending coronary artery (LAD) ligation and then applied rIC for 28 days, either daily or intermittently (every 3 days). They did not see any difference in infarct size but observed a dose-dependent improvement in LV remodelling as well as survival to 84 days compared with a control group given rIC only at the time of MI.¹⁰ This suggests that chronic rIC applied after an MI may confer distinct and separate benefits, especially on LV remodelling, to those from acute application at the time of MI.

The trial was designed as the first clinical study to test the hypothesis that repeated daily rIC for 4 weeks following successful revascularisation of a STEMI with P-PCI can reduce maladaptive cardiac remodelling as assessed by cardiac MRI (CMR).

METHODS

Trial design and governance

The trial was a multicentre, double-blind, randomised clinical study performed at four P-PCI centres in the UK (Leicester Glenfield Hospital, Kettering General Hospital, Royal Derby Hospital and The Royal Free Hospital, London). The study was approved by the National Health Service Research Ethics Committee (12/EM/0304), registered with ClinicalTrials.gov (NCT0166461) and was conducted in accordance with the principles of Good Clinical Practice and according to the Declaration of Helsinki, under the oversight of University of Leicester (UNOLE 0306). The trial

INTRODUCTION

Remote ischaemic conditioning (rIC) is a non-invasive cardioprotective therapy in which serial bouts of short lived periods of ischaemia/reperfusion (I/R) in a distant organ or tissue bed confers protection to the heart during ischaemia.¹ It can be performed by the inflation of a standard sphygmomanometer cuff placed on the upper arm and inflated above systolic pressure to temporarily halt blood flow, typically utilising four cycles of 5 min of cuff inflation interspersed by 5 min of complete cuff deflation.² More recently, this process has been simplified by the development of an automated rIC device (the autoRIC Device).

© Author(s) (or their employer(s)) 2018. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

To cite: Vanezis AP, Arnold JR, Rodrigo G, *et al.* *Heart* 2018;**104**:1955–1962.

Figure 1 Participant flow diagram—CONSORT (Consolidated Standards of Reporting Trials) diagram of recruitment to the trial. CMR, cardiac MRI; LVEF, left ventricular ejection fraction.

was coordinated by the National Institute for Health Research Leicester Cardiovascular Biomedical Research Unit.

Participant recruitment

The Consolidated Standards of Reporting Trials diagram for the study is shown in [figure 1](#). The study was conducted between November 2012 and October 2016. The aim was to recruit 72 subjects who completed the study protocol (see the Statistical Analysis section). A total of 1855 patients were screened for inclusion, of which 99 were randomised. Twenty-six participants withdrew or were withdrawn after randomisation and 73 patients (38 patients in the treatment group and 35 patients in the control group) completed the study.

Participants were recruited from patients presenting with a first STEMI who had undergone a successful P-PCI and were not deemed to require any further intervention. Potential participants over the age of 18 were approached on the day or the day after the procedure if they met all the inclusion criteria and did not meet any exclusion criteria ([box 1](#)). Transthoracic echocardiography was performed after obtaining consent and left ventricular ejection fraction (LVEF) was calculated

according to the modified Simpson Biplane method. This echocardiographic-derived LVEF assessment was used purely for assessing eligibility to the trial and was not used in the final analysis.

Randomisation

Participants were block randomised according to age (<45 and ≥45), gender and infarct location (anterior vs all others) with a 1:1 allocation to the rIC or the sham group. Randomisation was conducted by using a secure randomisation web service (Sealed Envelope, London, UK). The participants, cardiologists involved in their care and study investigators analysing all outcome data were blinded to the assigned treatment group.

Implementation of treatment

All participants were given and trained in the use of a device that automatically delivers rIC or a sham treatment—the autoRIC Device, a CE marked licenced product developed by CellAegis Devices (Toronto, Canada). The device allowed

self-implementation of therapy by participants in their own homes by the simple press of a button. In the active treatment (rIC) group, the device delivered 4×5 min cycles of inflation at 200 mm Hg interspersed with 5 min of deflation between each cycle with a total treatment time of 35 min. In the control group, the sham device went through the same timed cycles but with inflation only to 10 mm Hg. During the inflation period, both devices made identical vibrating noises. With ethics approval, participants were not told of the level of inflation required to deliver active treatment. Specifically, although the inflation pressure in the sham treated group was much lower, since they had not experienced a device inflated to a higher pressure, and their device appeared to deliver a 'treatment', they were unaware that they were in the control group. Starting on the third day following their P-PCI, participants implemented the treatment for 28 consecutive days. Participants were instructed to use the autoRIC Device at the same time of day and on the same arm. No stipulation was made as to a certain time of day, so long as the intervals between treatments were approximately 24 hours. Participants were asked to keep a diary of use of the device and were removed from the trial if they returned an incomplete diary sheet, missed more than three treatment sessions or missed more than 2 consecutive days of treatments. For those participants still in hospital at day 3, use of the device was started predischarge.

Box 1 Trial eligibility criteria—full inclusion and exclusion criteria

Inclusion criteria

- ▶ LVEF <45% on baseline echocardiogram.
- ▶ First STEMI.
- ▶ Successful revascularisation by P-PCI.
- ▶ Able to attend research centre for baseline and 4-month follow-up appointment.
- ▶ Competent to consent.

Exclusion criteria

- ▶ <18 years of age.
- ▶ ICD or CRT/D in situ.
- ▶ History of heart failure.
- ▶ Haemoglobin <11.5 g/dL, creatinine >200 µmol/L and/or GFR <30 mL/min/m².
- ▶ Known malignancy or other comorbid condition which in the opinion of the investigator was likely to have significant negative influence on life expectancy.
- ▶ Significant complications or illness following MI including cardiogenic shock, ischaemic mitral regurgitation, ventricular wall rupture, severe pericarditis or recurrent malignant arrhythmias.
- ▶ CMR contraindicated or unobtainable due to claustrophobia or significantly raised BMI.
- ▶ Further planned coronary interventions including surgery.
- ▶ Enrolment in another clinical trial.

BMI, body mass index; CMR, cardiac MRI; CRT/D, cardiac resynchronisation therapy/defibrillator; GFR, glomerular filtration rate; HF-PEF, heart failure with preserved ejection fraction; HF-REF, heart failure with reduced ejection fraction; ICD, implantable cardiac defibrillator; LVEF, left ventricular ejection fraction; MI, myocardial infarction; P-PCI, primary percutaneous coronary intervention; STEMI, ST segment elevation myocardial infarction.

CMR outcomes

The primary endpoint was change in LVEF from participant's baseline to 4-month follow-up scan. Secondary CMR endpoints included: changes in final LV infarct size, left ventricular end-diastolic volume index (LVEDVI), left ventricular end-systolic volume index (LVESVI) and left ventricular stroke volume index, all indexed to body surface area (BSA) between baseline and 4 months.

Participants recruited at Glenfield Hospital, Kettering General Hospital and Royal Derby Hospital had CMR performed at Glenfield Hospital and locally for those recruited at the Royal Free Hospital. All scans were supervised by cardiologists with an interest in CMR, and consisted of localisers, long axis cine images, contrast administration, a complete short axis stack for calculation of LV volumes, mass and ejection fraction (EF) and finally late gadolinium enhancement for quantification of infarct size as previously described.¹¹ CMR analysis was undertaken at the University of Leicester core laboratory using CVI42 (Circle Cardiovascular Imaging, Calgary, Canada) by an experienced CMR observer (JRA) blinded to randomisation or temporal order of scans. Endocardial and epicardial borders were manually contoured on contiguous short axis LV slices, excluding papillary muscles and trabeculae at end-diastole and at end-systole. The following LV parameters were determined: end-diastolic volume, end-systolic volume, stroke volume, EF and myocardial mass, using well-established protocols.¹² Illustrative CMR late gadolinium images at baseline and follow-up from a patient are shown in the online supplementary figure.

Clinical outcomes and follow-up

At 4 months, MACCE comprising recurrent MI, hospitalisation for heart failure, hospitalisation not for heart failure, stroke or transient ischaemic attack, unstable angina and ischaemia driven revascularisation were documented and the Kansas City Cardiomyopathy Questionnaire (KCCQ) was completed by surviving participants to assess quality of life measures in relation to heart failure. All participants had blood samples taken before the start of rIC or sham therapy on day 3 post-P-PCI and then again on the same day as their 4-month follow-up CMR. Samples were collected in EDTA tubes, processed for plasma and stored at -80°C. A non-competitive N-terminal pro-brain natriuretic peptide (NT-proBNP) assay was used to analyse all collected plasma samples in a single batch as previously described.¹³

Statistical analysis

Analysis was carried out on an intention-to-treat basis. The primary outcome was the change in LVEF from baseline CMR to 4-month follow-up CMR. To inform sample size, reported changes in LVEF after MI from recent trial and registry data in the P-PCI era^{14–17} were combined and an estimated increase of 7.7%±7.5% in LVEF reflecting the natural recovery after MI was calculated. On the basis that rIC would increase LVEF by 5% above this natural recovery (ie, a 12.7% increase in the active treatment group) a sample of 36 in each group was needed for an alpha level 0.05% and 80% power using a two-sided test. To allow for a drop-out rate of 20%, we aimed to recruit 45 patients in each group or 90 patients in total. During the trial, we found a higher-than-expected drop-out and eventually recruited 50 patients in each group. The study was not powered to detect any differences in clinical outcomes between the two groups.

Continuous data are expressed as a mean (SD) or a median (IQR) and compared with Student's t-test or Mann Whitney test as appropriate. Binary data are expressed as number (percentage).

of patients, and comparisons were performed with χ^2 or Fisher exact test. For CMR variables and NT-proBNP, we calculated changes from baseline for each patient and the comparison of changes between two treatment groups was conducted with Mann-Whitney or Student's t-test as appropriate. We confirmed that the data were normally distributed before undertaking analysis. A linear regression model was also fitted to compare the changes of LVEF with adjustment for important baseline variables and known predictors. We included age, gender, diabetes status, smoking status, infarct size, LVEF and presence/absence of microvascular obstruction at baseline in the initial model, and selected the final model using a backward approach. All statistical analyses were conducted using SAS V.9.4.

RESULTS

Participants

Participants in the treatment and control groups were well matched in terms of baseline characteristics and demographics (table 1). There was a non-statistically higher proportion of smokers in the treatment group and diabetics in the control group. In both groups, the majority of infarcts were caused by LAD occlusion. All participants received drug-eluting stents; thrombus evacuation was infrequently performed. Mean post-MI echocardiography estimated LVEF before recruitment was 40% for both groups. Discharge medication was very similar between the two groups with high use of optimal medical therapy. There was no significant difference between length of hospital stay

Table 1 Demographics and clinical characteristics at baseline*

	Control (n=35)	Treatment (n=38)	P values
Demographic			
Age (years)	58.9 (11.9)	58.7 (9.9)	0.920
Sex (% male)	29/35 (82.9)	31/38 (81.6)	0.999
Ethnicity (% Caucasian)	30/35 (85.7)	34/38 (89.5)	0.729
Body mass index (BMI)	27.6 (4.0)	27.7 (4.5)	0.876
Body surface area (BSA) (m ²)	2.0 (0.2)	1.9 (0.2)	0.102
Risk factors			
Current smoker	7/35 (20.0%)	15/37 (40.5%)	0.076
Hypertension	12/35 (34.3%)	11/38 (28.9%)	0.801
Hypercholesterolaemia	6/35 (17.1%)	9/38 (23.7%)	0.570
Diabetes mellitus	10/35 (28.6%)	4/38 (10.5%)	0.074
Previous angina	1/35 (2.9%)	1/38 (2.6%)	1.000
Clinical characteristics			
Ischaemic time (min)	158.2±50.1	167.2±39.1	0.761
Anterior STEMI	28/35 (80.0%)	27/38 (71.1%)	0.425
Culprit vessel—LAD	28/35 (80.0%)	25/38 (65.8%)	0.199
Cx	2/35 (5.7%)	4/38 (10.5%)	0.095
RCA	5/35 (14.2%)	8/38 (21.1%)	0.133
LMS	0/35 (0%)	1/38 (2.6%)	0.710
Multivessel disease	5/33 (15.2%)	7/37 (18.9%)	0.758
Mean TIMI flow pre-PCI	0.12±0.06	0.22±0.1	0.811
Mean TIMI flow post-PCI	2.78±0.08	2.57±0.13	0.851
Intervention			
DES	35/35 (100.0%)	38/38 (100.0%)	0.999
Thrombus evacuation	5/35 (14.3%)	8/38 (21.1%)	0.547
Systolic blood pressure	122.7 (20.9)	120.2 (19.3)	0.592
Total cholesterol	5.0 (0.9)	5.1 (1.6)	0.793
Haemoglobin (g/dL)	14.6 (1.5)	14.1 (1.6)	0.199
Renal function (GFR) mL/min	78 (60–90)	81.5 (60–90)	0.496
Ejection fraction by echo (%)	40 (38–43)	40 (35–42)	0.452
Length of hospital stay (days)	2.9 (2.4–3.7)	3.0 (2.4–3.5)	0.699
Medication on discharge			
Beta blocker	34/35 (97.1%)	34/38 (89.5%)	0.360
Statin	34/35 (97.1%)	37/38 (97.4%)	0.999
ACEi/ARB	35/35 (100.0%)	37/38 (97.4%)	0.999
MRA	9/35 (25.7%)	11/38 (28.9%)	0.798
Diuretic	1/35 (2.9%)	0/38 (0.0%)	0.480
Aspirin	34/35 (97.1%)	38/38 (100.0%)	0.480
P2Y ₁₂ inhibitor	34/35 (97.1%)	37/38 (97.4%)	0.999

*Results are shown as mean (SD) or median (Q1–Q3) for continuous variables and as number of patients (percentage) for categorical variables.

ACEi, ACE inhibitor; ARB, angiotensin receptor blocker; Cx, circumflex artery; DES, drug-eluting stent; GFR, glomerular filtration rate; LAD, left anterior descending coronary artery; LMS, left main stem artery; MRA, mineralocorticoid receptor antagonist; PCI, percutaneous coronary intervention; RCA, right coronary artery; STEMI, ST elevation myocardial infarction; TIMI, thrombolysis in myocardial infarction.

Figure 2 Cardiac MRI (CMR) assessment of left ventricular ejection fraction (LVEF) at baseline and 4 months—difference in LVEF as assessed by CMR from baseline scan to 4-month scan in the control (blue) versus treatment (orange) groups.

with a mean of 2.9 days in the control group and 3.0 days in the treatment group.

Tolerability of the autoRIC Device

Of the 26 participants who withdrew, only one (in the treatment arm) did so because they could not tolerate the device and withdrew after 2 days. No serious adverse effects were reported. There were three reports of mild discomfort when using the device, but in all cases the participants stated it was bearable and less noticeable with time. There was one report of tingling in the fingers or hands. There were no incidents of prolonged pain or of permanent marks, bruises or spots after application of the device.

CMR outcomes

Baseline CMR was performed at a median of 4 days (IQR: 4–5 days) post-P-PCI in the treatment group and 4 days (IQR: 3–6 days) in the control group ($p=0.826$). Follow-up CMR was performed at a median of 123 days (IQR: 122–125 days) in the treatment group and 122 days (IQR: 120–126 days) in the control group ($p=0.299$) after P-PCI. Of the total 73 participants who completed the study, all had LVEF assessments but two participant's follow-up CMRs had insufficient data to allow for infarct size estimation.

Mean baseline LVEF was not significantly different between treatment group (42.8%, IQR: 38.5–48.6) and control group (44.3%, IQR: 38.3–46.8) ($p=0.952$). There was no statistically significant difference in the primary endpoint of CMR assessed LVEF recovery from baseline to 4 months between the treatment and control groups (figure 2 and table 2). The mean increase in LVEF over 4 months was $4.8\% \pm 7.8\%$ in the treatment group and $4.6\% \pm 5.9\%$ in the control group ($p=0.924$). In multivariable regression analysis, infarct size ($p=0.0003$) and baseline LVEF ($p<0.001$) predicted change in LVEF at 4 months, and the effect of rIC treatment remained insignificant ($p=0.862$) after adjusting for these two covariates.

With regard to secondary CMR outcomes, there were no differences at baseline in infarct size, LVESVI or prevalence of microvascular obstruction (MVO) between the treatment and control groups (table 2). Mean baseline LVEDVI was significantly higher in the treatment group (99.8 mL/m^2 (84.9–114.4)) than control group (86.3 mL/m^2 (76.9–95.9)) ($p=0.012$). However, this may mainly reflect the effect of indexing (because of the small difference in BSA between the groups, table 1) as LVEDV was not significantly different (189.5 (158.8–218.4) vs 182.0 (135.6–194.8), $p=0.128$). There were no significant differences between the two groups in the changes in infarct size, LVEDVI or LVESVI at 4 months (table 2). Specifically, in a multivariable analysis similar to that carried out for the change in LVEF, the change in LVEDVI between the treatment and control groups at 4 months remained not significant ($p=0.972$).

Clinical outcomes, quality of life and biomarkers

There was no statistically significant difference in MACCE at 4 months between the two groups, although overall numbers were small (table 3). Furthermore, there were no differences in the KCCQ outcome scores at 4 months or the change in NT-proBNP levels from baseline to 4 months (table 3).

DISCUSSION

This trial has shown that in patients who have undergone successful P-PCI for STEMI, daily rIC for 4 weeks did not result in a greater improvement in LVEF at 4 months as assessed by CMR compared with a control group that underwent sham conditioning.

rIC extends the pre-existing paradigm of local ischaemic conditioning, which has been demonstrated in numerous studies to attenuate I/R injury via a number of mechanisms including modulation of key intracellular pathways such as the reperfusion injury salvage kinase and survivor activating factor enhancement pathways, which act predominantly to inhibit opening of the mitochondrial permeability transition pore.¹⁸ A number of trials have shown that one-off rIC at around the time of P-PCI for STEMI can increase the myocardial salvage index and reduce

Table 2 CMR outcomes

	Baseline			4 months			Change		
	Control group	Treatment group	P values	Control group	Treatment group	P values	Control group	Treatment group	P values
LVEF (%)	44.3 (38.3–46.8)	42.8 (38.5–48.6)	0.952	48.9 (42.0–53.1)	49.4 (41.7–55.6)	0.830	4.6 (5.9)	4.8 (7.8)	0.924
LVEDVI (mL/m^2)	86.3 (76.9–95.9)	99.8 (84.9–114.4)	0.012	92.4 (78.2–104.2)	97.6 (84.1–116.6)	0.090	5.1 (13.1)	2.4 (13.6)	0.390
LVESVI (mL/m^2)	50.4 (40.9–57.5)	53.3 (45.9–70.5)	0.055	46.6 (38.9–55.7)	52.6 (37.4–70.0)	0.334	−1.0 (10.7)	−2.9 (12.3)	0.489
Infarct size (% of LVM)	21.9 (16.9–35.4)	21.4 (13.0–32.3)	0.526	17.6 (13.7–22.6)	15.2 (8.5–23.3)	0.345	−7.9 (9.6)	−6.7 (8.4)	0.588
Presence of MVO	31 (88.6%)	27 (71.1%)	0.085	NA	NA	NA	NA	NA	NA

Results are shown as mean (SD) or median (Q1–Q3).

CMR, cardiac MRI; LVEDVI, left ventricular end-diastolic volume index; LVEF, left ventricular ejection fraction; LVESVI, left ventricular end-systolic volume index; LVM, left ventricular mass; MVO, microvascular obstruction; NA, not applicable.

Table 3 Clinical, quality of life and biomarker outcomes

	Control group	Treatment group	P values
MACCE	8 (22.9%)	8 (21.1%)	0.852
Non-heart failure hospitalisation	6	6	
Stroke/TIA	0	1	
NSTEMI/UA	2	1	
Heart failure hospitalisations/death	0	0	
KCCQ score	92 (83–98) (n=34)	88 (72–97) (n=37)	0.237
NT-proBNP (baseline)	336.2 (40.9–855.2) (n=31)	188.0 (74.0–535.0) (n=36)	0.965
NT-proBNP (4 months)	256.5 (96.1–488.4) (n=30)	225.6 (46.6–1000.5) (n=36)	0.699
NT-proBNP (change)	18.9 (–313.0 to 202.1) (n=31)	2.7 (–374.9 to 316.5) (n=36)	0.954

Results are shown as mean (SD) or median (Q1–Q3). There were missing data for one participant in each arm in the collection of the KCCQ scores and missing data in six participants (four in the control arm and two in treatment arm) for baseline NT-proBNP data and in seven participants (five in the control arm and two in treatment arm) for 4-month NT-proBNP data.

KCCQ, Kansas City Cardiomyopathy Questionnaire; MACCE, major adverse cardiac and cerebral event; NSTEMI, non-ST segment elevation myocardial infarction; NT-proBNP, N-terminal pro-brain natriuretic peptide; TIA, transient ischaemic attack; UA, unstable angina.

biomarker release and that this translates to better clinical outcomes in terms of MACCE and LVEF.^{4–8}

The rationale for this trial was provided by preclinical data,¹⁰ showing that repeatedly administered rIC over 4 weeks produced additional beneficial changes in LV function after an MI over and above that observed with a single application at the time of MI. These functional changes were accompanied by positive modulation of key remodelling processes such as a reduction in oxidative stress, attenuation of the expression of genes associated with fibrosis and hypertrophy, and blunting of the inflammatory response with reduced levels of neutrophil and macrophage infiltration in the myocardium.¹⁰ Previously, the same group had demonstrated that repetitive rIC significantly altered the behaviour of neutrophils after MI with reduced levels of adhesion at days 1 and 10 as well as a reduction in phagocytosis at day 10, apoptosis at days 1 and 10 and an overall change in the profile of cytokine release.¹⁹ Yamaguchi *et al*²⁰ reinforced the potential of repeated rIC post-MI and implicated exosomes as mediators for signalling in rIC, possibly by their action of transferring antifibrotic microRNAs such as miR29a as well as insulin-like growth factor-1, which is known to be protective in the context of remodelling. We have shown in a rat myoblast cell culture model that that humoral modulation induced by rIC benefits a number of the maladaptive processes that contribute to post-MI remodelling including cardiomyocyte hypertrophy, differentiation of fibroblasts to pathological myofibroblasts and the upregulation of prohypertrophic and profibrotic genes.²¹ In human studies, Jones *et al*²² have shown that daily rIC for 1 week improved endothelial function and microcirculation both locally and remotely possibly via increased STAT-3 activation triggering an increase in endothelial progenitor cells.

The use of CMR rather than echocardiography allowed for more accurate quantification of LVEF and volumes as well as allowing for the measurement of infarct size and MVO which is not possible with standard echocardiography. With regard to participant selection, we specifically looked at the effects of rIC on naïve myocardium, excluding patients with a history of MI or heart failure to examine any effect of the intervention on acute remodelling. As far as we are aware, the trial is one of the first studies where rIC was delivered chronically using an automated device. In addition to consistency, the autoRIC Device also allowed for better patient blinding, as patients in the control arm were also issued with the device and instructed in its use in an identical manner to the patients in the treatment arm. Their device was also programmed to cycle but not inflate and

none of the participants were aware of the assigned treatment. Finally, the trial illustrates the challenge of undertaking studies in patients with post-STEMI LV dysfunction in the modern era of P-PCI. Over 1800 subjects are needed to be screened to identify 99 patients with an EF <45% with a substantial proportion excluded because of preserved LV function (figure 1).

This trial was designed as a proof of concept phase II study and was powered to detect a substantial increase in LVEF (5%) over the natural recovery that occurs post-STEMI. The extent of the latter was estimated from a collation of recent trial and registry studies.^{14–17} Despite targeting individuals with impaired LV function, the observed improvement in LVEF in the control group ($4.6\% \pm 5.9\%$) was less than expected ($7.7\% \pm 7.5\%$), and therefore it is possible that the level to which the study was powered or any more modest improvement due to rIC could have been missed. A post hoc power calculation based on the observed increase in LVEF in the control group suggested that to detect a 3% greater increase in the treatment arm would require a total sample size of 560 patients. It should be noted that while we did not find an effect of rIC on changes in LVEF between baseline and 4 months, in the multivariable analysis we did see a significant impact of two well-established determinants of LV remodelling, namely infarct size and baseline LVEF,²³ indicating that the study was adequately powered to detect such effects. However, ultimately the goal of any therapy is to impact on clinical outcomes. There were very few such events in this trial (table 3) and it should be emphasised that the trial was not powered to detect any differences in clinical outcomes.

The trial did not implement rIC until day 3 post-MI. This is different from the experimental protocol of Wei *et al*¹⁰ where the benefits of chronic rIC were observed on top of rIC applied at the time of the MI. Our rationale for delaying the implementation of rIC was to determine any potential effects of rIC on ventricular remodelling, independently of any effects on the initial infarct. This may also have impacted on the null outcome as some studies have suggested that acute remodelling begins very early postinfarct.^{24–25} Therefore, by instigating treatment on day 3, a number of the upstream mechanisms that can lead to maladaptive remodelling and heart failure may have already been upregulated or downregulated and the potential for rIC to modulate them may have been attenuated. Pryds *et al*²⁶ recently demonstrated that the use of a once daily rIC regimen for 4 weeks similar to that applied in this trial did not improve LVEF in patients with chronic ischaemic heart failure. This supports the theory that to reap the maximum benefits from repeated

rIC post MI in terms of remodelling, treatment should be instigated early in the acute phase. The currently recruiting Chronic Remote Ischemic Conditioning to Modify Post-MI Remodelling trial (CRIC-RCT) (<http://clinicaltrials.gov/NCT01817114>) may answer this question as it uses a similar protocol to this trial but commences rIC at the time P-PCI.

Another potential reason for the lack of clinical translation of the beneficial effects on LV function of chronic rIC seen in the post-MI rodent model¹⁰ is the use of adjunctive therapies (including beta blockers, ACE inhibitors/angiotensin receptor blockers and mineralocorticoid receptor antagonists) in the post-MI setting in patients. These therapies have been shown to attenuate maladaptive remodelling.^{27–29} In the animal model, the effect of rIC was assessed naïve of these treatments, whereas in this trial any benefit needed to be shown on top of these established therapies. Similarly, the effect of chronic rIC in the animal model was studied in the absence of any comorbidities. For example, diabetes mellitus has been shown to attenuate the effect of rIC³⁰ and patients with this condition were not excluded from the study.

To enable participant blinding, patients in the control group were also given and instructed in the use of a device. Although every effort was made to make the sham devices look and behave like the treatment devices and participants were not made aware of what constituted active treatment (see the Methods section), because the sham device only inflated to a maximum pressure of 10 mm Hg it is possible that some of the control patients may have deduced that they were in the control arm. This was unavoidable. However, it should be noted that the assessment of

the primary outcome (change in LVEF after 4 months with MRI) was done entirely blinded to treatment assignment and indeed the scans were assessed in a random order.

Finally, daily oversight of correct application of daily rIC was limited due to the nature of the participants' implementing their own treatment at home. While the autoRIC Device is simple and easy to use even for very frail participants, the generation of device used during this trial had no interrogative capability to assess compliance; a subsequent device that can monitor compliance has been developed (CRICtrac, CellAegis Devices). Although it was standard practice to telephone participants regularly to monitor device use, we also relied on participant diaries, similar to ones used in drug trials, to encourage and evaluate concordance.

CONCLUSIONS

In this phase II, double-blinded, multicentre randomised controlled trial, daily rIC for 4 weeks starting on day 3 in patients with significantly impaired LV function following successful P-PCI for STEMI did not substantially improve LVEF at 4 months. Although this trial does not itself provide support for further investigation of repeated application of rIC in this setting, given its limited power it does not exclude the possibility of chronically applied rIC causing clinically meaningful improvement in LVEF post-STEMI. This trial provides valuable information on the scale of the trial that would be necessary to demonstrate such an effect in the context of contemporary treatment of STEMI and highlights other issues that may affect the outcome of such a study.

Acknowledgements Glenfield Hospital: Jayshree Gracey, Sue Parish, Ellie Clarke, Emma Beeston, Susan Sterland, Sadat Edroos. Royal Derby Hospital: Praveen Sadarmin, Marie Appleby, Eithne Heneghan. Royal Free Hospital: Angelique Smit, Karl Salazar, Tushar Kotecha. Kettering General Hospital: Dinu Johns, Charmaine Beirnes, Lissy Jose. CMR scans were performed in the NIHR Leicester Clinical Research Facility. The authors acknowledge the financial support from the NIHR Leicester Cardiovascular Biomedical Research Unit and the Masonic Charitable Foundation. The authors also thank CellAegis Devices for providing the autoRIC Devices for the study.

Contributors APV conceived and designed the trial and drafted the manuscript. NJS provided overall supervision and was the chief investigator. All other authors were involved in either recruitment of participants or analysis of CMR or secondary outcome investigation results. All authors were involved in the final draft of the manuscript. NJS acts as the guarantor for the overall content of the work.

Funding This research received financial support from the NIHR Leicester Cardiovascular Biomedical Research Unit and the Masonic Charitable Foundation.

Competing interests None declared.

Patient consent Not required.

Ethics approval NHS Research Authority-NRES Committee East Midlands-Northampton.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement The original data to all the results outlined in the manuscript can be provided in raw format on request in an anonymised form.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>.

REFERENCES

- 1 Heusch G, Bøtker HE, Przyklenk K, *et al*. Remote ischemic conditioning. *J Am Coll Cardiol* 2015;65:177–95.
- 2 Kharbanda RK, Mortensen UM, White PA, *et al*. Transient limb ischemia induces remote ischemic preconditioning in vivo. *Circulation* 2002;106:2881–3.

Key messages

What is already known on this subject?

- Remote ischaemic conditioning (rIC) has been shown to reduce infarct size when used at the time of primary percutaneous coronary intervention (P-PCI) in ST segment elevation myocardial infarction (STEMI).
- Preclinical studies have suggested the protective benefits of rIC may extend to the remodelling phase following myocardial infarction and thus may positively impact ventricular remodelling.

What might this study add?

- This first-in-man phase II clinical trial compared daily rIC for 4 weeks post-STEMI treated with P-PCI with sham treatment to assess the effect of rIC on ventricular remodelling in patients with impaired left ventricular function (left ventricular ejection fraction (LVEF) <45%).
- rIC was started 3 days after STEMI.
- rIC did not significantly impact the primary endpoint of improvement of LVEF at 4 months postinfarct.

How might this impact on clinical practice?

- Our study did not find evidence of a substantial impact on ventricular remodelling of chronic rIC administered after STEMI.
- However, because of the size of the study a modest but clinically relevant effect was not ruled out.
- Furthermore, rIC was started 3 days postinfarct and earlier initiation might have been more beneficial.
- Future studies also need to establish any impact on clinical outcomes.

- 3 Kleinbongard P, Skyschally A, Heusch G. Cardioprotection by remote ischemic conditioning and its signal transduction. *Pflugers Arch* 2017;469:159–81.
- 4 Sloth AD, Schmidt MR, Munk K, *et al*. Improved long-term clinical outcomes in patients with ST-elevation myocardial infarction undergoing remote ischaemic conditioning as an adjunct to primary percutaneous coronary intervention. *Eur Heart J* 2014;35:168–75.
- 5 Munk K, Andersen NH, Schmidt MR, *et al*. Remote Ischemic Conditioning in Patients With Myocardial Infarction Treated With Primary Angioplasty: Impact on Left Ventricular Function Assessed by Comprehensive Echocardiography and Gated Single-Photon Emission CT. *Circ Cardiovasc Imaging* 2010;3:656–62.
- 6 Elbadawi A, Ha LD, Abuzaid AS, *et al*. Meta-Analysis of Randomized Trials on Remote Ischemic Conditioning During Primary Percutaneous Coronary Intervention in Patients With ST-Segment Elevation Myocardial Infarction. *Am J Cardiol* 2017;119:832–8.
- 7 Schmidt MR, Rasmussen ME, Bøtker HE. Remote Ischemic Conditioning for Patients With STEMI. *J Cardiovasc Pharmacol Ther* 2017;22:302–9.
- 8 Bøtker HE, Kharbanda R, Schmidt MR, *et al*. Remote ischaemic conditioning before hospital admission, as a complement to angioplasty, and effect on myocardial salvage in patients with acute myocardial infarction: a randomised trial. *Lancet* 2010;375:727–34.
- 9 Verouhis D, Sörensson P, Gourine A, *et al*. Effect of remote ischemic conditioning on infarct size in patients with anterior ST-elevation myocardial infarction. *Am Heart J* 2016;181:66–73.
- 10 Wei M, Xin P, Li S, *et al*. Repeated remote ischemic postconditioning protects against adverse left ventricular remodeling and improves survival in a rat model of myocardial infarction. *Circ Res* 2011;108:1220–5.
- 11 Nazir SA, McCann GP, Greenwood JP, *et al*. Strategies to attenuate micro-vascular obstruction during P-PCI: the randomized reperfusion facilitated by local adjunctive therapy in ST-elevation myocardial infarction trial. *Eur Heart J* 2016;37:1910–9.
- 12 Hudsmith LE, Petersen SE, Tyler DJ, *et al*. Determination of cardiac volumes and mass with FLASH and SSFP cine sequences at 1.5 vs. 3 Tesla: a validation study. *J Magn Reson Imaging* 2006;24:312–8.
- 13 Omland T, Persson A, Ng L, *et al*. N-terminal pro-B-type natriuretic peptide and long-term mortality in acute coronary syndromes. *Circulation* 2002;106:2913–8.
- 14 Ottervanger JP, van 't Hof AW, Reiffers S, *et al*. Long-term recovery of left ventricular function after primary angioplasty for acute myocardial infarction. *Eur Heart J* 2001;22:785–90.
- 15 Baks T, van Geuns RJ, Biagini E, *et al*. Recovery of left ventricular function after primary angioplasty for acute myocardial infarction. *Eur Heart J* 2005;26:1070–7.
- 16 Solomon SD, Glynn RJ, Greaves S, *et al*. Recovery of ventricular function after myocardial infarction in the reperfusion era: the healing and early afterload reducing therapy study. *Ann Intern Med* 2001;134:451–8.
- 17 Klug G, Metzler B. Assessing myocardial recovery following ST-segment elevation myocardial infarction: short- and long-term perspectives using cardiovascular magnetic resonance. *Expert Rev Cardiovasc Ther* 2013;11:203–19.
- 18 Bell RM, Bøtker HE, Carr RD, *et al*. 9th Hatter Biannual Meeting: position document on ischaemia/reperfusion injury, conditioning and the ten commandments of cardioprotection. *Basic Res Cardiol* 2016;111:41.
- 19 Shimizu M, Saxena P, Konstantinov IE, *et al*. Remote ischemic preconditioning decreases adhesion and selectively modifies functional responses of human neutrophils. *J Surg Res* 2010;158:155–61.
- 20 Yamaguchi T, Izumi Y, Nakamura Y, *et al*. Repeated remote ischemic conditioning attenuates left ventricular remodeling via exosome-mediated intercellular communication on chronic heart failure after myocardial infarction. *Int J Cardiol* 2015;178:239–46.
- 21 Vanezis AP, Edroos SA, Samani NJ, *et al*. The inhibitory effect of remote ischaemic conditioning on a cellular model of cardiac hypertrophy. *Heart* 2012;98(Suppl 5):A4.4–A5.
- 22 Jones H, Hopkins N, Bailey TG, *et al*. Seven-day remote ischemic preconditioning improves local and systemic endothelial function and microcirculation in healthy humans. *Am J Hypertens* 2014;27:918–25.
- 23 Hombach V, Grebe O, Merkle N, *et al*. Sequelae of acute myocardial infarction regarding cardiac structure and function and their prognostic significance as assessed by magnetic resonance imaging. *Eur Heart J* 2005;26:549–57.
- 24 Konstam MA, Kramer DG, Patel AR, *et al*. Left ventricular remodeling in heart failure: current concepts in clinical significance and assessment. *JACC Cardiovasc Imaging* 2011;4:98–108.
- 25 Brenner S, Ertl G. Remodelling and adverse remodelling in CAD. *Herz* 2012;37:590–7.
- 26 Pryds K, Nielsen RR, Jorsal A, *et al*. Effect of long-term remote ischemic conditioning in patients with chronic ischemic heart failure. *Basic Res Cardiol* 2017;112:67.
- 27 Pfeffer MA, Lamas GA, Vaughan DE, *et al*. Effect of captopril on progressive ventricular dilatation after anterior myocardial infarction. *N Engl J Med* 1988;319:80–6.
- 28 Dargie HJ. Effect of carvedilol on outcome after myocardial infarction in patients with left-ventricular dysfunction: the CAPRICORN randomised trial. *Lancet* 2001;357:1385–90.
- 29 Tsutamoto T, Wada A, Maeda K, *et al*. Effect of spironolactone on plasma brain natriuretic peptide and left ventricular remodeling in patients with congestive heart failure. *J Am Coll Cardiol* 2001;37:1228–33.
- 30 Miki T, Itoh T, Sunaga D, *et al*. Effects of diabetes on myocardial infarct size and cardioprotection by preconditioning and postconditioning. *Cardiovasc Diabetol* 2012;11:67.