Effect of Ischemia Duration and Protective Interventions on the Temporal Dynamics of Tissue Composition After Myocardial Infarction

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Rationale: The impact of cardioprotective strategies and ischemia duration on postischemia/reperfusion (I/R) myocardial tissue composition (edema, myocardium at risk, infarct size, salvage, intramyocardial hemorrhage, and microvascular obstruction) is not well understood.

Objective: To study the effect of ischemia duration and protective interventions on the temporal dynamics of myocardial tissue composition in a translational animal model of I/R by the use of state-of-the-art imaging technology.

Methods and Results: Four 5-pig groups underwent different I/R protocols: 40-minute I/R (prolonged ischemia, controls), 20-minute I/R (short-duration ischemia), prolonged ischemia preceded by preconditioning, or prolonged ischemia followed by postconditioning. Serial cardiac magnetic resonance (CMR)-based tissue characterization was done in all pigs at baseline and at 120 minutes, day 1, day 4, and day 7 after I/R. Reference myocardium at risk was assessed by multidetector computed tomography during the index coronary occlusion. After the final CMR, hearts were excised and processed for water content quantification and histology. Five additional healthy pigs were euthanized after baseline CMR as reference. Edema formation followed a bimodal pattern in all 40-minute I/R pigs, regardless of cardioprotective strategy and the degree of intramyocardial hemorrhage or microvascular obstruction. The hyperacute edematous wave was ameliorated only in pigs showing cardioprotection (ie, those undergoing short-duration ischemia or preconditioning). In all groups, CMR-measured edema was barely detectable at 24 hours postreperfusion. The deferred healing-related edematous wave was blunted or absent in pigs undergoing preconditioning or short-duration ischemia, respectively. CMR-measured infarct size declined progressively after reperfusion in all groups. CMR-measured myocardial salvage, and the extent of intramyocardial hemorrhage and microvascular obstruction varied dramatically according to CMR timing, ischemia duration, and cardioprotective strategy.

Conclusions: Cardioprotective therapies, duration of index ischemia, and the interplay between these greatly influence temporal dynamics and extent of tissue composition changes after I/R. Consequently, imaging techniques and protocols for assessing edema, myocardium at risk, infarct size, salvage, intramyocardial hemorrhage, and microvascular obstruction should be standardized accordingly. (Circ Res. 2017;121:439-450. DOI: 10.1161/CIRCRESAHA.117.310901.)

Key Words: edema ■ ischemic preconditioning ■ ischemic postconditioning ■ magnetic resonance imaging ■ reperfusion injury ■ translational medical research

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Novelty and Significance

What Is Known?

- Edema formation, inflammation, and microvascular injury contribute to irreversible loss of cardiac myocytes after myocardial I/R.
- The presence and extent of these processes and tissue healing are associated with outcome after MI.
- Evaluation of myocardial tissue composition changes by CMR imaging (ie, tissue characterization) serves as surrogate end points in experimental and clinical studies and clinical trials.

What New Information Does This Article Contribute?

- By using serial CMR exams in a translational large animal model of I/R, this study shows that the temporal dynamics and extent of post-MI tissue composition changes are greatly influenced by the application of cardioprotective interventions, the duration of the index ischemia, and the interplay between them.
- Rather than estimates of the ischemic area alone, the degree and spatial extent of myocardial edema seems to be a surrogate marker of ischemic insult severity.
- These findings highlight the need for protocol standardization when using post-MI imaging techniques to measure relevant end points in experimental and clinical studies.

Post-MI tissue composition is highly dynamic and could be characterized by CMR, which has been used to assess surrogate outcomes and efficacy end points in many experimental and clinical studies. However, there is paucity of studies tracking the temporal dynamics of these processes in a comprehensive manner. Our current work shows that the degree and extent of post-MI tissue composition changes (edema, necrosis, hemorrhage, and MVO) as assessed by CMR are greatly influenced by the time of image acquisition, duration of ischemia, cardioprotective strategies, and the interplay between them. Therefore, clinical and experimental imaging protocols for post-MI tissue characterization aiming at quantifying edema, myocardial area at risk, IS, myocardial salvage, IMH, and MVO should take into consideration these dynamics and be as standardized as possible.

Nonstandard Abbreviations and Acronyms

| Abbreviation | Definition |
|--------------|------------|
| CMR          | cardiac magnetic resonance |
| IMH          | intramyocardial hemorrhage |
| I/R          | ischemia/reperfusion |
| IR-TFE       | inversion recovery turbo field echo |
| IS           | infarct size |
| LAD          | left anterior descending |
| LGE          | late gadolinium enhancement |
| LV           | left ventricle |
| LVEF         | left ventricular ejection fraction |
| MaR          | myocardium at risk |
| MDCT         | multidetector computed tomography |
| MI           | myocardial infarction |
| MVO          | microvascular obstruction |
| ROI          | region of interest |
| STIR         | short-tau triple inversion-recovery |
| T2W          | T2-weighted |

Ischemia/reperfusion (I/R) results in a dramatic change of myocardial tissue composition, mainly in the first days after myocardial infarction (MI). Edema formation, inflammation, microvascular injury, and ultimately healing, among other phenomena, accompany irreversible loss of cardiac myocytes after MI. Cardiac magnetic resonance (CMR) can noninvasively identify most of these phenomena; however, accounting for the temporal dynamics and factors affecting post-MI tissue composition might be crucial for a precise assessment of related imaging end points. To date, this has not been studied in a comprehensive manner.

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For years, the intense edematous reaction appearing in the postischemic region early after MI was thought to persist in stable form for at least 1 week. On this basis, delineating the extent of post-MI edema by CMR rapidly became established as a measure of the original occluded coronary artery perfusion territory (myocardium at risk [MaR]). Thus, the amount of salvaged myocardium—a theoretically better surrogate of the effect of cardioprotective therapies—has been estimated from quantifications of the extent of delayed gadolinium enhanced myocardium (infarct size [IS]) and the extent of myocardial edema (assumed to delineate the MaR) in the same imaging session. Consequently, multiple clinical and experimental studies have used CMR-measured edema and IS as end points, within a single imaging session performed at a variety of post-MI time points. Recent evidence has challenged the view of a stable post-MI edema reaction, demonstrating dynamic changes in edema in a pig model of MI. In this model, an initial hyperacute edema reaction on reperfusion resolves within 24 hours and is followed by a deferred healing-related edema wave occurring a few days after reperfusion. Notably, intramyocardial hemorrhage (IMH) and microvascular obstruction (MVO) can counteract the high-intensity edema signal on T2 sequences by reducing tissue T2 relaxation time. The impact of IMH and MVO on the bimodal edematous reaction is, thus, a matter of intense debate. Moreover, the extent/degree of edema might be lower in patients undergoing infarct-limiting cardioprotective interventions. Another variable that might differently affect post-MI tissue composition is ischemia duration. However, CMR exams were performed at different time points in these studies, precluding a definitive interpretation.

In summary, temporal dynamics and extent of post-I/R key phenomena (edema, necrosis, MVO, and IMH) might be affected by duration of the index ischemic event, by the application of cardioprotective interventions and the interplay between them, but to date, this has not been studied in a comprehensive manner. To elucidate these, we designed an experimental study in which pigs underwent different myocardial I/R protocols. CMR was performed at baseline and at 120 minutes, 24 hours, 4 days, and 7 days after I/R for the serial noninvasive assessment of myocardial edema, extent of IMH and MVO, IS, and myocardial salvage. Histology and quantification of
myocardial water content by reference standard were performed at sacrifice immediately after the final CMR scan.

Methods

Study Design and MI Procedure
The study population comprised 25 Large White male pigs weighing 30 to 40 kg. The study was approved by the Institutional and Regional Animal Research Committees, and the study design is summarized in Online Figure I. Five pigs were euthanized with no intervention other than baseline CMR and served as reference healthy subjects. A group of 5 pigs underwent reperfused transmural acute MI by closed-chest I/R, consisting of 40-minute left anterior descending coronary artery occlusion followed by balloon deflation and reestablishment of blood flow, and were euthanized at 7 days after I/R serving as controls. Three additional groups of 5 animals each underwent modified I/R protocols incorporating different protective strategies followed by sacrifice on day 7: preconditioning, 40-minute I/R preceded by three 5-minute cycles of balloon inflation/deflation in the left anterior descending; postconditioning, 40-minute I/R followed by four 1-minute cycles of balloon inflation/deflation in the left anterior descending; and short-duration ischemia, 20-minute I/R. Arterial enhanced multidetector computed tomography (MDCT) was performed during the index coronary occlusion, between minute 10 and minute 20 of ischemia, to delineate the reference MaR (MDCT-MaR, hyperperfused region during coronary occlusion). Comprehensive CMR exams were performed at baseline and at 120 minutes, 24 hours, day 4, and day 7 after I/R. Animals were immediately euthanized after the final follow-up CMR scan, and myocardial tissue samples from ischemic areas were rapidly collected for histology and evaluation of water content. Full methods are provided in the Online Data Supplement.

Arterial Enhanced MDCT Protocol and Analysis
Arterial phase MDCT studies were performed on a 64-slice computed tomographic-scanner (Brilliance CT 64; Philips Healthcare, Cleveland, OH) after intravenous administration of iodinated contrast media. MDCT images were analyzed using dedicated software (MR Extended Work Space 2.6; Philips Healthcare, Best, The Netherlands) by 2 observers blinded to group allocation. Short axis orientation were obtained from volumetric computed tomographic images by multiplanar reconstruction using equivalent anatomic coordinates used for T2-weighted short-tau inversion-recovery (T2W-STIR) planning acquisition. MaR and remote areas were visually identified based on contrast enhancement differences, manually delineated, and expressed as a percentage of left ventricle (LV) area. Detailed information about imaging MDCT protocol and analysis is shown in the Online Data Supplement.

CMR Protocol and Analysis
Baseline CMR scans were performed immediately before MI, and scans were subsequently repeated at all postinfarction follow-up times until sacrifice. CMR examinations were conducted with a Philips 3-Tesla Achieva Tx whole body scanner (Philips Healthcare, Best, The Netherlands) equipped with a 32-element phased-array cardiac coil. The imaging protocol included a standard segmented cine steady-state free precession sequence to provide high quality anatomic references and assessment of LV mass, wall thickness, and LV ejection fraction (LVEF); a T2W-STIR sequence to assess the extent of edema and IMH; a T2-gradient-spin-echo mapping sequence to provide precise myocardial T2 relaxation time properties; and a T1-weighted inversion recovery turbo field echo sequence acquired 10 to 15 minutes after the administration of gadolinium contrast (late gadolinium enhancement [LGE]) to assess IS and MVO. To avoid any interference with T2 measures at immediate reperfusion, gadolinium contrast was not administered at baseline CMR scans. CMR images were analyzed using dedicated software (MR Extended Work Space 2.6; QMassMR 7.6; Medis, Leiden, The Netherlands) by 2 observers experienced in CMR analysis and blinded to group allocation. Detailed information about imaging protocol and parameters and CMR analysis can be found in the Online Data Supplement.

Myocardial Water Quantification and Histological Analysis
Paired myocardial samples were collected within minutes of sacrifice from the ischemic myocardium of all pigs. Myocardial water content was quantified by the desiccation technique and expressed as a percentage of wet weight. Histological sections of the at-risk myocardium were stained with hematoxylin and eosin, Masson trichrome, and antineutrophil antibody. Full methods are presented in the Online Data Supplement.

Statistical Analysis
Normal distribution of each data subset was checked using graphical methods and a Shapiro-Wilk test. Leven test was performed to check homogeneity of variances. For quantitative variables, data are expressed as mean±SD. A 1-way ANOVA was conducted for among-group comparison of myocardial water content and MDCT-MaR. To take account of repeated measures, generalized mixed models were applied for the study and comparison of the temporal evolution of T2, CMR-MaR, IS, IMH, MVO, and salvage myocardium within and between groups. Pairwise comparisons among groups and time points were performed and P value adjusted for multiple comparisons using the Hochberg method. Post hoc test of trends over time for imaging parameters by group was performed by using coefficients of orthogonal polynomials. A Kruskal–Wallis test was conducted to compare the number of inflammatory cells and collagen content among groups euthanized at day 7. Associations between measured T2 relaxation time and IMH, MVO, IS, and LVEF were evaluated by Pearson correlation coefficient. All statistical analyses were performed with Stata v12.0 (StataCorp, College Station, TX). Graphs were generated with Stata 12.0 or GraphPad-Prism v6.0 (GraphPad Software, Inc, La Jolla, CA).

Results

Impact of Cardioprotective Strategies and Ischemia Duration on T2 and Myocardial Water Content
T2 relaxation time determined at time points ≤7 days post-reperfusion are summarized in Table 1. As already reported, after 40-minute I/R, T2 relaxation times were significantly prolonged at reperfusion, almost completely normalized at

| Table 1. Time Profile of Myocardial T2 Relaxation in the Ischemic Porcine Myocardium During the First Week After Different I/R Protocols and Cardioprotective Strategies |
|---|---|---|---|---|---|
| **T2 Relaxation Time, ms** | **Baseline** | **R-120 min** | **R-24 h** | **R-Day 4** | **R-Day 7** |
| **40-min I/R (controls)** | 46.5 (3.0) | 74.8 (14.4) | 46.7 (4.9) | 66.4 (8.3) | 78.9 (11.7) |
| **40-min I/R+PostC** | 43.1 (4.1) | 73.4 (8.8) | 46.1 (2.5) | 72.5 (17.0) | 76.7 (16.8) |
| **PreC+40-min I/R** | 43.1 (1.0) | 62.5 (2.8)* | 49.1 (1.2) | 59.2 (10.8)* | 57.6 (10.3)* |
| **20-min I/R** | 46.6 (2.0) | 58.3 (1.5)* | 49.9 (1.9) | 50.1 (3.1)* | 48.7 (1.5)* |

Values are mean (SD). No significant differences were found between groups at baseline. A bimodal T2 time course was observed for all 40-min I/R pigs, with no significant differences detected between 40-min I/R and 40-min I/R+PostC. In contrast, PreC blunted both T2 peaks, whereas after 20-min I/R, the first peak was blunted and the second, absent. I/R indicates ischemia/reperfusion; PostC, postconditioning; and PreC, preconditioning.

*Statistically significant differences (P<0.05) compared with the same time point in the 40-min I/R (control) group. P value is adjusted for multiple comparisons among groups for each time point.
Preconditioning before 40-minute I/R resulted in a significantly shorter T2 relaxation time at reperfusion and an earlier and shorter deferred peak, occurring on day 4 rather than day 7. Conversely, postconditioning had no effect on edema dynamics, with animals showing the standard bimodal pattern with an initial peak at reperfusion and a deferred T2 prolongation peaking on day 7 (Figure 1A and 1B). Pigs subjected to short-duration ischemia (20-minute I/R) showed a unimodal pattern of edema, with T2 prolongation at reperfusion, albeit blunted compared with 40-minute I/R, and no deferred peak. True myocardial water content values in the

Figure 1. Time profile of T2 relaxation time and measurements of myocardial water content in the ischemic myocardium after different ischemia/reperfusion (I/R) protocols. A, Time profile of absolute T2 relaxation time (ms) in the ischemic myocardium of pigs undergoing different I/R protocols. Data are shown as mean±standard error of the mean. B, Representative serial cardiac magnetic resonance T2-mapping images from pigs that underwent 40-min I/R (control), 40-min I/R followed by postconditioning (PostC), 40-min I/R preceded by preconditioning (PreC), or 20-min I/R; examinations were made at baseline, 120 min, 24 h, 4 d, and 7 d after reperfusion. All T2 maps were scaled between 30 and 120 ms. C, Measurements of myocardial water content (%) in healthy pigs and in the ischemic myocardium of pigs subjected to different I/R protocols and euthanized at day 7 after I/R. Myocardial water content (mean±SD, %) in the healthy pig myocardium was 79.4±0.6%. Myocardial water content (mean±SD, %) in the ischemic myocardium of pigs undergoing 40-min I/R (controls), 40-min I/R+PostC, 40-min I/R preceded by PreC, and 20-min I/R (short-duration ischemia) at day 7 after I/R was 85.2±0.9%, 84.7±1.4%, 82.2±1.1%, and 79.8±0.2%, respectively. Between-group differences of water content in the ischemic myocardium remained statistically significant with the exception of 40-min I/R+PostC vs 40-min I/R and 20-min I/R vs healthy. Data represented are means±standard error of the mean.
ischemic myocardium were comparable with those estimated from T2 relaxation times on day 7 CMR (Figure 1C).

**Impact of Cardioprotective Strategies and Ischemia Duration on CMR-Measured MaR**

Mean MaR measured by MDCT in the different groups was as follows (% of LV): 40-minute I/R (controls), 28.3±4.3%; preconditioning plus 40-minute I/R, 31.7±6.9%; 40-minute I/R plus postconditioning, 31.3±4.0%; and short-duration ischemia (20-minute I/R), 24.6±6.7% (statistically nonsignificant differences as compared with controls). Measurements of MaR determined by T2W-CMR in these groups at different times after reperfusion are summarized in Table 2.

CMR-measured MaR and T2 relaxation time showed similar time course patterns. Pigs subjected to 40-minute I/R with or without postconditioning had significant swelling of the formerly ischemic myocardium at reperfusion (Online Table I; Online Movies I through IV). Consequently, at early reperfusion, CMR-measured MaR as delineated by T2W-STIR was significantly higher than MaR measured by the reference standard MDCT in these pigs (Figure 2A and 2B). Conversely, preconditioned and 20-minute I/R pigs developed edema at reperfusion but without myocardial swelling (Online Table I; Online Movies V through VIII), and, therefore, MaR measured by CMR was similar to the value recorded by MDCT at this time point in these groups. In all study groups, the initial edema wave dissipated by 24 hours postreperfusion, and CMR measurements systematically underestimated MaR at this time point. On days 4 and 7 postreperfusion, CMR-measured MaR in control and postconditioned pigs was similar to MaR measured by MDCT. Conversely, in preconditioned pigs, CMR- and MDCT-measured MaR values were similar on day 4, but on day 7, CMR underestimated MaR because of partial resolution of the deferred edema wave, which appeared earlier, peaking on day 4. In pigs undergoing short-duration ischemia (20-minute I/R), CMR underestimated MaR at all time points after the first reperfusion scan because of the absence of the deferred edema wave.

**Impact of Cardioprotective Strategies and Ischemia Duration on IS, Myocardial Salvage, IMH, and MVO**

CMR measurements of IS and myocardial salvage after reperfusion are summarized in Table 2. IS was maximal at reperfusion and progressively shrank during the first week after MI in all groups. Whereas postconditioning had no effect on IS, preconditioning significantly reduced IS at all time points (Figure 3A and 3B). In pigs undergoing short-duration ischemia, IS was negligible at all time points. In parallel with the temporal variations in CMR-measured MaR, myocardial salvage estimated by CMR dynamically changed over time and according to the I/R protocol applied (Table 2). The time profile of myocardium salvage as assessed by reference MDCT-measured MaR and CMR IS is presented in Online Table II.

CMR measurements of IMH and MVO after reperfusion are summarized in Online Table III. IMH was apparent at 24 hours, peaking on day 4 post-I/R (Figure 2C), whereas MVO was apparent early after reperfusion, peaking on day 1 post-I/R and progressively decreasing thereafter (Figure 3C). Presence and extent of IMH and MVO were unaffected by postconditioning, significantly reduced by preconditioning, and negligible after short-duration ischemia.

Table 2. Time Profile of MaR, IS and Myocardial Salvage as Assessed by CMR During the First Week After Reperfused Myocardial Infarction in Pigs Subjected to Different I/R Protocols and Cardioprotective Strategies

| Group               | CMR Measure                | Follow-Up          |
|---------------------|----------------------------|--------------------|
|                     | R-120 min | R-24 h | R-Day 4 | R-Day 7 |
| 40-min I/R (controls) | MaR, % of LV | 42.9 (5.7) | 2.2 (1.7) | 27.1 (3.4) | 30.1 (2.3) |
|                     | IS, % of LV | 39.2 (3.8) | 30.2 (3.1) | 28.2 (4.6) | 25.4 (4.0) |
|                     | Myocardial salvage, % | 8.3 (6.4) | −1310 (996) | −4.4 (14.2) | 15.7 (13.3) |
| 40-min I/R+PostC    | MaR, % of LV | 48.8 (6.2)* | 1.9 (2.0) | 33.9 (4.9)* | 32.5 (3.2) |
|                     | IS, % of LV | 46.2 (5.6)* | 33.1 (3.4) | 32.8 (4.5)* | 30.4 (3.6)* |
|                     | Myocardial salvage, % | 5.2 (4.8) | −1060 (513) | 3.3 (3.9) | 6.6 (3.0) |
| PreC+40-min I/R    | MaR, % of LV | 29.1 (1.4)* | 4.2 (1.4) | 24.6 (7.5) | 17.2 (5.7)* |
|                     | IS, % of LV | 21.0 (7.2)* | 18.5 (10.2)* | 7.5 (3.7)* | 6.1 (4.7)* |
|                     | Myocardial salvage, % | 28.2 (22.6)* | −389 (361)* | 68.0 (13.6)* | 64.0 (23.9)* |
| 20-min I/R          | MaR, % of LV | 27.7 (3.3)* | 3.6 (1.1) | 4.2 (2.2)* | 2.2 (1.9)* |
|                     | IS, % of LV | 6.1 (4.5)* | 3.0 (1.0)* | 2.2 (0.8)* | 1.5 (0.8)* |
|                     | Myocardial salvage, % | 78.8 (13.7)* | 17.3 (18.7)* | 25.2 (75.5) | 3.4 (79.0)* |

Values are mean (SD). A bimodal trend over time after reperfusion was observed for CMR-MaR and CMR-salvage ([CMR MaR−CMR IS]/CMR MaR, %) in all groups of pigs with the exception of the 20-min I/R group in which a negative linear trend or no clear trend was shown for CMR-MaR and salvage, respectively. In contrast, a strong negative linear trend over time was shown for IS in all groups. *P values for significant trends were all <0.01. CMR indicates cardiac magnetic resonance; I/R, ischemia/reperfusion; IS, infarct size; LV, left ventricle; MaR, myocardium at risk; PostC, postconditioning; PreC, preconditioning; and R, reperfusion.

*Statistically significant differences (\*P<0.05) as compared with the same time point in the 40-min I/R (control) group. P value is adjusted for multiple comparisons among groups for each time point and imaging parameter.
Impact of Cardioprotective Strategies and Ischemia Duration on Histological Features of MI

The histological analysis of pigs euthanized at day 7 postreperfusion are summarized in Online Table IV. Postconditioning had no discernable effect on lesion size, granulation tissue content, neutrophil infiltration, or collagen content (Figure 4). Preconditioning resulted in smaller lesion areas, a higher proportion of granulation tissue, and lower neutrophil infiltration.
and collagen content. Pigs undergoing short-duration ischemia showed no signs of tissue lesion and a low degree of neutrophil infiltration.

**Association Between T2 and IMH, MVO, IS, and LVEF**

Overall, T2 relaxation time in the ischemic region at 120 minutes post-I/R correlated positively with the degree of IMH and MVO: the longer the T2 at early reperfusion, the larger the extent of IMH and MVO. Conversely, the deferred T2 peak (highest value between day 4 and day 7) correlated positively with IS and inversely with LVEF: the greater the deferred T2 peak, the larger the infarct and the lower the LVEF on day 7 (Figure 5).

**Discussion**

In this study, we have comprehensively characterized the impact of cardioprotective strategies (preconditioning and post-conditioning) and ischemia duration on the temporal evolution and extent of myocardial tissue composition changes (edema, necrosis, IMH, and MVO) by CMR. In all instances of necrosis (positive LGE at day 7), a bimodal edematous response is seen, regardless of the presence and degree of IMH or MVO; however, when necrosis is absent (in animals undergoing short-duration ischemia with no day-7 LGE), the edematous reaction is unimodal, with a blunted reperfusion-related edema wave and no healing-related deferred wave. Preconditioning, which significantly reduces IS, modulates the intensity of both the initial and deferred edema waves. The deferred wave also peaks earlier in preconditioned than in nonpreconditioned infarctions. CMR-measured IS declined progressively after reperfusion in all groups, whereas the extent of IMH and MVO varied according to CMR timing and protocol applied. T2 relaxation time in the ischemic area early after reperfusion is correlated with the severity of IMH and MVO, whereas the deferred peak of T2 relaxation time is strongly associated with IS and LVEF.

Consequently, imaging protocols for post-MI tissue characterization aiming at quantifying edema, MaR, IS, myocardial salvage, IMH, and MVO should account for these dynamics and be as standardized as possible (Figure 6).

**Modulation of the Post-I/R Edema by Cardioprotective Strategies and Ischemia Duration**

The possibility of protecting the myocardium during an acute MI (cardioprotection) has interested the scientific community for several decades. Many studies use IS normalized to MaR as an acute end point, on the assumption that post-I/R edema is steady. However, recent studies suggest that the extent of edema might be influenced by cardioprotective conditioning interventions. However, these previous clinical studies were not designed to address the effect of conditioning interventions on edema formation, and subjects underwent a single CMR examination that was not at the same time point for all individuals.

Our results show that preconditioning (a potent cardioprotective strategy) and short ischemia duration have a major impact on the intensity and dynamics of post-MI edema (Figure 1). Preconditioning reduced the initial edema wave and also the deferred wave, which peaked early, on day 4, contrasting with the peak on day 7 in control animals. Shortening the duration of coronary occlusion to 20 minutes (20-minute I/R) also blunted initial edema wave, and in this case, the second edema wave was absent. These CMR results are consistent with histologically determined water content at day 7 after MI, which was reduced in pigs undergoing preconditioning and within the normal range in pigs undergoing short-duration ischemia (Figure 1C).
Impact of IMH and MVO on the Occurrence of the Bimodal Edematous Reaction

The extent of IMH and MVO varies as a function of time from the acute ischemic event. IMH is moreover closely associated with the development of MVO, conferring a worse prognosis when present. More importantly, in the present study, edema followed a bimodal T2 pattern in all pigs undergoing 40-minute I/R (with or without preconditioning or postconditioning) regardless of the degree of IMH or MVO (Figure 1), which was almost absent in preconditioned pigs (Figures 2C and 3C). Notably, all pigs, including those undergoing short-duration ischemia, showed a significant drop in T2 from the hyperacute phase to 24 hours postreperfusion (Figure 1). These findings suggest that the main driver of the drop in T2 is rapid resorption of the initial reperfusion-related edema, regardless of the degree of IMH and MVO, and reinforce the reality of bimodal post-MI edema. Nevertheless, IMH exerts some influence on T2 relaxation time, as we previously conceded, and might explain the slightly less noticeable drop in T2 at 24 hours in pigs with almost no IMH or MVO (pigs given preconditioning or short-duration ischemia; Figure 1).

Impact of Cardioprotective Strategies and Ischemia Duration on CMR-Measured Myocardial Area at Risk and Salvage

Parametric T2 mapping improves the detection and quantification of myocardial edema; however, this methodology is not universally available, contrasting with T2W-STIR, which is available from all major vendors. In this regard, CMR-measured MaR as delineated by the extent of edema on T2W-STIR imaging paralleled T2 relaxation time profile. The present study shows that the edema-sensitive T2W-STIR CMR sequence overestimates MaR (as compared with the reference standard, MDCT in this study) at early time points (120 minutes) after reperfusion in pigs subjected to 40-minute I/R with no protective strategies or undergoing postconditioning (which did not protect in this study). This overestimation is mainly driven by the massive swelling of the reperfused myocardium (Online Movies I through IV). By 24 hours, there is a systematic underestimation of MaR by CMR in all cases, mainly driven by the substantial resorption of edema and normalization of T2 relaxation time. This underestimation resulted in biologically implausible negative myocardial salvage data at 24 hours in most pigs (Table 2). This finding reinforces the idea that MaR (and consequently salvaged myocardium) may not reliably be quantified by CMR around this time point. Conversely, on days 4 and 7, CMR-measured MaR was similar to MaR measured by MDCT (Figure 2A). However, the dynamics of posts ischemia edema are altered by cardioprotective strategies and ischemia duration, with further implications for CMR-measured MaR and salvage.

Thus, the smaller extent of edema at reperfusion in pigs that underwent preconditioning or short-duration ischemia (Figure 2) was associated with less prolonged T2 relaxation times (Figure 1) and lower degree of myocardial swelling. Interestingly, these 2 groups of cardioprotected animals had

Figure 4. Impact of cardioprotection and ischemia duration on histological features of myocardial infarction. Representative histological images of porcine ischemic myocardium 7 d after 40-min ischemia/reperfusion (I/R; control), 40-min I/R followed by postconditioning (PostC), 40-min I/R preceded by preconditioning (PreC), and 20-min I/R. Images show staining with hematoxylin and eosin (H&E), antiPM1 antibody (PMN), and Masson trichrome. Stained sections were used to quantify lesion extent, proportion of necrosis and granulation tissue, neutrophil density, and percentage of collagen within granulation tissue. Note the scarce neutrophil infiltration in porcine myocardium subjected to PreC or 20-min I/R, accompanied by small patchy areas of granulation tissue (PreC) or its absence (20-min I/R). Scale bars, 100μM.
significantly smaller infarcts on day 7 (Figure 3). The cardioprotection (reduced IS) afforded by preconditioning or short-duration ischemia was demonstrated by CMR (LGE; Figure 3) and histology (Figure 4). In pigs undergoing short-duration ischemia, necrosis was barely detectable, and so was edema and CMR-measured MaR from day 1 on, supporting the notion that the deferred edema wave is related to post-MI healing.\textsuperscript{11} This would imply that interventions that protect the myocardium could affect edema dynamics, therefore, having an impact on CMR estimations of MaR and salvage, as suggested by indirect clinical evidence before.\textsuperscript{16,17} However, whether any intervention that reduces IS diminishes myocardial edema remains to be demonstrated. A paradigmatic example of such interplay may be seen in the group subjected to short duration of ischemia (Table 2). In this group, IS is small and stable on day 1 (3.0\% of LV), day 4 (2.2\% of LV), and day 7 (1.5\% of LV); however, a dramatic change is seen in CMR-measured salvage index (17.3\%, 25.2\%, and 3.4\% of MaR on days 1, 4, and 7, respectively). Of note, when using the MaR standard reference MDCT value to calculate salvage, salvage index remains stable (87.4\%, 90.0\%, and 94.4\% of MaR on days 1, 4, and 7, respectively; Online Table II). The latter measurements (based on MDCT-MaR) are consistent with the significant cardioprotective effect seen by early reperfusion (ie, short ischemia duration).

Decrease of CMR-Based IS During the First Week After MI

Previous studies show rapid resorption of LGE myocardium (a surrogate of IS) between ≈day 1 and day 7 after MI.\textsuperscript{4,22} Consistent with these observations, our data show a progressive decrease of CMR-based IS in all study groups (Figure 3). The massive swelling of the early postreperfused myocardium might explain the large IS detected in our study 120 minutes after reperfusion. Interestingly, animals that underwent preconditioning showed LGE-positive myocardial regions early after reperfusion that became LGE negative by day 4 or day 7 CMR (Figure 3). This phenomenon might indicate early and transient expansion of extracellular volume without irreversible myocardial injury.\textsuperscript{22} These data highlight the importance of performing CMR infarct imaging within a consistently defined and narrow time frame, preferably at the end of the first week, when using IS as an end point in clinical trials during

Figure 5. Association between T2, intramyocardial hemorrhage (IMH), microvascular obstruction (MVO), infarct size, and left ventricular ejection fraction (LVEF). Scatter plots showing positive association of T2 relaxation time in the ischemic myocardium during the hyperacute postreperfusion period (120 min) with (A) peak MVO (Pearson $r=0.68$) and (B) peak intramyocardial hemorrhage (Pearson $r=0.75$). C and D. The deferred T2 peak in the ischemic myocardium shows a strong positive association with day-7 infarct size (Pearson $r=0.87$) and a negative association with day-7 LVEF (Pearson $r=−0.85$). After adjustment for multiple testing, all $P$ values for the correlations shown remained significant ($P<0.05$). Graphs include 4 groups of 5 pigs each undergoing 40-min ischemia/reperfusion (I/R; controls), 40-min I/R followed by postconditioning, 40-min I/R preceded by preconditioning, or 20-min I/R. LV indicates left ventricle.
the acute post-MI period. Nevertheless, the fact that myocardial edema and LGE follow a disparate dynamic pattern after ischemia/reperfusion highlights the complexity of measuring myocardial salvage in real practice.

**T2 as a Maker of Post-I/R Myocardial Injury**

The ability to predict the fate of the myocardium early after MI would be of great clinical value. Myocardial edema contributes to impaired post-MI microvascular perfusion by increasing extravascular compression and might contribute to altered coronary physiology indices.1,2 In this study, we show that T2 relaxation time in ischemic myocardium in the hyperacute CMR examination (after 2 hours of reperfusion) correlates with IMH and MVO—events that develop days after MI (Figure 5A and 5B). Conversely, the deferred T2 peak correlated directly with IS and inversely with LVEF—parameters associated with post-MI healing (Figure 5C and 5D). Therefore, myocardial T2 relaxation time might be a quantitative surrogate of the ischemic insult or therapeutic effect, rather than a surrogate of MaR.

However, these findings might apply only to the reperfused MI because the bimodal edema response is less pronounced in nonreperfused MI.11 These findings warrant further experimental and clinical studies specifically addressing the prognostic significance of T2 at different time points and the perfusion status of the myocardium.

**Limitations**

Caution is needed when extrapolating experimental results to the clinic. However, the pig is one of the most clinically translatable large animal models for the study of reperfused MI.23,24 The nature of our experimental study, that is, longitudinal follow-up, precluded individual histological gold standard measurements of area at risk (ie, Evans Blue by coronary reocclusion), IS (ie, triphenyl tetrazolium chloride), or MVO (thioflavin-S) at each time point. In addition, the heart was harvested at sacrifice to perform reference standard myocardial water content measurements and histological quantification of different tissue components, which otherwise might be altered by such procedures. It is fair to acknowledge that although previous studies have used histological standards to validate the use of CMR to measure IS and MVO22,25 and MDCT to measure MaR,19,26 there is probably no perfect noninvasive method for such purposes. In this regard, acutely detected LGE does not necessarily equate to irreversible injury and may contribute to severely distort estimates of salvaged myocardium when comparing against a prereperfusion reference standard to assess MaR.22 Other reasons that might contribute to inaccurate estimations include damage extent beyond the boundaries of the actual MaR (defined as the hypoperfused region during coronary occlusion);19 slightly shrinking of MaR in MDCT performed during coronary occlusion because of lack of perfusion in animal models with poor collateral circulation; and the presence of residual edema in salvaged myocardium.

![Figure 6. Effect of timing of evaluation, ischemia duration, and protective interventions on cardiac magnetic resonance (CMR)-measured individual outcomes after myocardial infarction (MI).](https://circres.ahajournals.org/)

According to the data presented, establishing the appropriate timing to measure IMH and MVO could occur somewhere around the first 24 to 48 h post-MI; whereas the optimal timing to measure edema, myocardium at risk (MaR), IS, and salvage could occur somewhere between day 4 and day 7 after MI. Nevertheless, caution should be exercised when addressing post-MI tissue characterization by any means given that several components and factors might additionally impact the dynamics and complex changes which undergo the heart after MI. y axis numbers and graphs represent real data relative to the highest point on the chart for the given imaging outcome (MaR, IS, MVO, and IMH). A value of 100 is the highest value for the (Continued)
myocardium which might contribute to overestimate of IS early after reperfusion. Hybrid imaging techniques have been proposed to overcome some of these issues; however, further validation is needed.

Our study examined only anterior MI. The reasons for this choice include the avoidance of possible magnetic-field nonhomogeneity related to the inferolateral wall, and adherence to recommendations for patient selection in clinical trials of cardioprotective interventions. The observed changes in tissue composition are likely to occur regardless of MI location; however, caution should be exercised when extrapolating other results, especially those regarding the association of T2 with ventricular outcome.

Our results do not necessarily invalidate the reported protective effects of the majority of preclinical postconditioning studies published. However, our findings are in line with some recent experimental and clinical data. In this regard, ischemic postconditioning did not reduce the incidence of clinical events in ST-segment–elevation MI (STEMI) patients in the DANAMI-3-iPOST trial (The Third Danish Study of Optimal Acute Treatment of Patients With ST Elevation Myocardial Infarction–Ischemic Postconditioning). Moreover, this intervention did not reduce IS, myocardial salvage index, extent of MVO, or improve LVEF in the subset of patients who underwent CMR. Interspecies differences, small sample size, the lack of comorbidities and comedications in animal experiments, and the diversity of postconditioning protocols applied might explain the ambivalent preclinical and clinical data. Nevertheless, our goal was not to test the cardioprotective effect of postconditioning, but rather to use different protective strategies to characterize post-MI tissue over a range of injury severities. Indeed, the use of strategies that did not achieve cardioprotection reinforced our earlier results obtained by the standard I/R procedure.

The regions of interest used to quantify T2 relaxation time in this study included the full wall thickness. The regions of interest might, therefore, include different myocardial states (such as hemorrhage or MVO). We took this approach to match our previous experimental work and because the differentiation of such small areas might be challenging. This might have contributed to the differences in absolute T2 relaxation time between our study and others taking a different approach to region of interest selection.

Conclusions

In a translational large animal model of myocardial I/R and by using CMR, we have shown that the temporal dynamics and extent of post-MI tissue composition changes are greatly influenced by application of cardioprotective interventions and the duration of the index ischemia. Post-MI LGE areas do not necessarily equate to necrotic myocardium because the extent of hyperenhanced ischemic tissue evolves rapidly during the first week after reperfusion. The magnitude of IS and MVO varies according to ischemia duration and cardioprotection. The greatest divergences are seen in the degree and spatial extent of myocardial edema during the first week after MI, which seem to be a quantitative surrogate marker of ischemic damage or therapeutic effect.

These data highlight the need for protocol standardization when using post-MI imaging techniques to measure edema, MaR, IS, myocardial salvage, IMH, and MVO in experimental and clinical studies.

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Disclosures

J. Sánchez-González is a Philips Healthcare employee. The other authors report no conflicts.

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Effect of Ischemia Duration and Protective Interventions on the Temporal Dynamics of Tissue Composition After Myocardial Infarction

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SUPPLEMENTAL MATERIAL

DETAILED METHODS

Study design
Experiments were performed on castrated male Large-White pigs weighing 30 to 40 kg. A total of 25 pigs completed the full protocol and comprised the study population. The study was approved by the Institutional and Regional Animal Research Committee. The study design is summarized in Online Figure I. Five pigs were sacrificed with no intervention other than baseline CMR, and served as reference healthy subjects. A group of 5 pigs underwent reperfused transmural acute myocardial infarction by closed-chest I/R, consisting of 40-minute left anterior descending (LAD) coronary artery occlusion followed by balloon deflation and reestablishment of blood flow, and were sacrificed at 7 days after I/R serving as controls. Three additional groups of 5 animals each underwent modified I/R protocols incorporating different protective strategies followed by sacrifice on day 7: preconditioning, 40min-I/R preceded by three 5-minute cycles of balloon inflation/deflation in the LAD; postconditioning, 40min-I/R followed by four 1-minute cycles of balloon inflation/deflation in the LAD; and short-duration ischemia, 20min-I/R. In all pigs, arterial enhanced multidetector computed tomography (MDCT) was performed during the index coronary occlusion, between minute 10 and minute 20 of ischemia, to delineate the reference MaR (hypoperfused region during coronary occlusion). Comprehensive CMR exams were performed at baseline, and at 120min, 24h, day4, and day7 after I/R. Animals were immediately sacrificed after the final follow-up CMR scan, and myocardial tissue samples from ischemic areas were rapidly collected for histology and evaluation of water content.

Myocardial infarction procedure
The MI protocol is detailed elsewhere. Anesthesia was induced by intramuscular injection of ketamine (20 mg/kg), xylazine (2 mg/kg), and midazolam (0.5 mg/kg), and maintained by continuous intravenous infusion of ketamine (2 mg/kg/h), xylazine (0.2 mg/kg/h), and midazolam (0.2 mg/kg/h). Animals were intubated and mechanically ventilated with oxygen (fraction of inspired O2: 28%). Central venous and arterial lines were inserted and a single bolus of unfractioned heparin (300 IU/kg) was administered at the onset of instrumentation. The left anterior descending coronary artery, immediately distal to the origin of the first diagonal branch, was occluded for 40 or 20 minutes with an angioplasty balloon introduced via the percutaneous femoral route using the Seldinger technique. Balloon location and maintenance of inflation were monitored angiographically. After balloon deflation, a coronary angiogram was recorded to confirm patency of the coronary artery. A continuous infusion of amiodarone (300 mg/h) was maintained during the procedure in all pigs to prevent malignant ventricular arrhythmias. In cases of ventricular fibrillation, a biphasic defibrillator was used to deliver non-synchronized shocks.

Conditioning strategies
In the postconditioned group, at the end of the 40 minutes coronary occlusion, coronary flow was reestablished for 1 min by deflating the angioplasty balloon. After 1 min of reflow, the balloon was re-inflated to occlude the coronary artery (low pressure, 4-6 atmospheres) at the same level for 1 minute. This procedure was repeated four times and then chronic coronary reperfusion was allowed. In the pre-conditioning group, the angioplasty balloon was placed immediately distal to the first diagonal branch, and inflated 3 times for 5 min with low pressure (4 to 6 atmospheres) inflations, each separated by 5 min of reflow. At the end of the pre-conditioning protocol, the balloon was inflated for 40 minutes and then deflated to allow chronic reperfusion. Balloon location and maintenance of inflation were monitored angiographically in all cases for both procedures.

Arterial enhanced MDCT protocol
Arterial enhanced multidetector computed tomography (MDCT) was performed during the index coronary occlusion in all pigs, between minute 10 and minute 20 of ischemia, to delineate the reference MaR (hypoperfused region during coronary occlusion). All MDCT studies were performed
on a 64-slice CT-scanner (Brilliance CT 64, Philips Healthcare, Cleveland, Ohio). The pigs were positioned supine, and all scans were performed in the crano-caudal direction during free-breathing. Arterial phase MDCT was performed after intravenous administration of 60 ml iomeprol 400 mgI/ml (Iomeron 400, Bracco Imaging, Milano, Italy) at a flow rate of 3 ml/s followed by a 20-ml saline chaser bolus at the same flow rate. The scan delay was determined using a bolus tracking technique. Data acquisition started 15 seconds after a threshold of 180 Hounsfield Units was reached in a region of interest placed in the descending aorta. MDCT examinations were acquired using retrospective cardiac triggered at the 75% of the cardiac cycle with 64 x 0.625 mm collimation and a pitch of 0.2, 120 kV tube voltage, 800 mA tube current and tube rotation time of 400 ms. Image reconstruction was performed with a 512x512 matrix size over a 273x273mm² FOV and 0.45mm slice thickness by using high resolution filter (Xres Sharp).

**Arterial enhanced MDCT analysis**

MDCT images were analyzed using dedicated software (MR Extended Work Space 2.6, Philips Healthcare, Best, The Netherlands). Short axes orientation were obtained from volumetric CT images by multi-planar reconstruction using equivalent anatomical coordinates used for T2W-STIR planning acquisition. In order to have equivalent LV sections, MDCT studies had to be reconstructed in slices equivalent in thickness and level to the CMR ones. Thus, T2W-STIR and multi-planar reconstructed (MPR) short axis CT images were co-registered in 13 to 15 short-axis LV slices by one observer. To ensure CT as independent reference for MaR, endocardial and epicardial borders from MPR CT short-axis images were manually traced by a different observer blinded to the co-registration information; and MaR and remote areas were visually identified based on contrast enhancement differences, manually delineated, and expressed as a percentage of LV area.

**CMR protocol**

Baseline CMR scans were performed immediately before myocardial infarction and scans were subsequently repeated at all post-infarction follow-up times until sacrifice. CMR examinations were conducted with a Philips 3-Tesla Achieva Tx whole body scanner (Philips Healthcare, Best, the Netherlands) equipped with a 32-element phased-array cardiac coil. The imaging protocol included a standard segmented cine steady-state free-precession (SSFP) sequence to provide high quality anatomical references, and assessment of LV mass, wall thickness and left ventricular ejection fraction (LVEF); a T2-weighted short-tau triple inversion-recovery (T2W-STIR) sequence to assess the extent of edema and intramyocardial hemorrhage; a T2-gradient-spin-echo mapping (T2-GraSE map) sequence to provide precise myocardial T2 relaxation time properties; and a T1-weighted inversion recovery turbo field echo (T1-IR-TFE) sequence acquired 10 to 15 minutes after the administration of gadolinium contrast to assess infarct size and microvascular obstruction. To avoid interference with T2 measures at immediate reperfusion, gadolinium contrast was not administered at baseline CMR scans. All sequences were acquired in free-breathing mode. The imaging parameters for the SSFP sequence were FOV 280 x 280 mm, slice thickness 6 mm with no gap, TR 2.8 ms, TE 1.4 ms, flip angle 45°, cardiac phases 30, voxel size 1.8 x 1.8 mm, and 3 NEX. The imaging parameters for the T2W-STIR sequence were FOV 280 x 280, slice thickness 6 mm, TR 2 heartbeats, TE 80 ms, voxel size 1.4 x 1.9 mm, delay 210 ms, end-diastolic acquisition, echo-train length 16, and 2 NEX. The imaging parameters for the T2-GraSE mapping were FOV 300x300 with an acquisition voxel size of 1.8x2.0 mm² and slice thickness 8 mm, TR 2 heartbeats, eight echo times ranging from 6.7 to 53.6ms, and EPI factor 3. Late gadolinium enhancement imaging was performed 10 to 15 min after intravenous administration of 0.20 mmol of gadopentetate dimeglumine contrast agent per kg of body weight using a T1 inversion-recovery spoiled turbo field echo (T1-IR-TFE) sequence with the following parameters: FOV 280 x 280 mm, voxel size 1.6 x 1.6 mm, end-diastolic acquisition, thickness 6 mm with no gap, TR 5.6 ms, TE 2.8 ms, inversion delay time optimized to null normal myocardium, and 2 NEX. SSFP, T2W-STIR, and T1-IR-TFE sequences were performed to acquire 13 to 15 contiguous short-axis slices covering the heart from the base to the apex, whereas T2-maps were acquired in a mid-apical ventricular short axis slice corresponding to the same anatomical level in all acquisitions in order to track T2 relaxation time changes over time.
CMR analysis

CMR images were analyzed using dedicated software (MR Extended Work Space 2.6, Philips Healthcare, The Netherlands; and QMassMR 7.6, Medis, Leiden, The Netherlands) by two observers experienced in CMR analysis and blinded to protocol allocation. Left ventricular (LV) mass, myocardial T2 relaxation time, and extent of edema, necrosis, intramyocardial hemorrhage (IMH) and microvascular obstruction (MVO) were determined.

LV cardiac borders were automatically traced with manual adjustment in each cine image to obtain LV end-diastolic volume (LVEDV), end-systolic volume (LVESV), and LVEF. In the tracing convention used, the papillary muscles were included as part of the LV cavity volume. LVEF was computed as LVEF (%) = (LVEDV-LVESV)/LVEDV. LV epicardial borders were also traced on the end-diastolic images to measure end-diastolic wall thickness at the myocardium at risk and remote areas, with LV mass computed as the end-diastolic myocardial volume (ie, the difference between the epicardial and endocardial volumes) multiplied by myocardial density (1.05 g/mL). Values of LV mass normalized to body surface area were calculated with the modified Brody’s formula.6

T2-maps were automatically generated on the acquisition scanner by fitting the signal intensity of all echo times to a monoexponential decay curve at each pixel with a maximum likelihood expectation maximization algorithm. T2 relaxation maps were quantitatively analyzed by placing a wide transmural region of interest (ROI) at the ischemic area of the corresponding slice in all studies. Hypointense areas suggestive of microvascular obstruction or hemorrhage were included in the ROI for T2 quantification purposes.1, 3, 5

The extent of edema, expressed as a percentage of LV mass, was defined after manually tracing the endocardial and epicardial contours of T2W-STIR short-axis images. Abnormal areas were initially identified using the full-width at half-maximum (FWHM) method.7, 8 Given that the solely use of FWHM may be prompt to inaccurate patchy estimations,9, 10 extensive manual correction and visual border delineation were performed. Areas corresponding to slow-flow artifacts were carefully excluded from edematous area. Hypointense areas within the edematous zone, corresponding to intramyocardial hemorrhage (IMH), were included within the edematous region.11, 12 Additionally, the size of the area of IMH was calculated by manual delineation of the hypointense areas on T2W-images,11 and expressed as a percentage of LV mass. Manual delineation of clear hypointense areas was permitted in the absence of discernible hyperintense myocardium.

Myocardial necrosis (infarct size, IS), expressed as a percentage of LV mass, was defined according the extent of late gadolinium enhancement (LGE) after manually tracing the endocardial and epicardial contours on T1-IR-TFE short axis images. Abnormal areas were defined using the FWHM, with manual correction if needed. Hypointense black areas within the necrotic zone, corresponding to microvascular obstruction (MVO), were included within the necrotic area.11, 12 Additionally, the size of the area MVO was calculated by manual delineation of the hypointense areas on LGE images,11 and expressed as a percentage of LV mass.

Quantification of myocardial water content

Paired myocardial samples were collected within minutes of euthanasia from the ischemic myocardium of all pigs. Tissue samples were immediately blotted to remove surface moisture and introduced into laboratory crystal containers previously weighed on a high-precision scale. The containers were weighed before and after drying for 48 hours at 100°C in a desiccating oven. Tissue water content was calculated as follows: water content (%) = [(wet weight−dry weight)/wet weight] ×100. An empty container was weighed before and after desiccation as an additional calibration control.

Histological and immunnohistochemical analysis

Myocardial samples were collected within minutes of euthanasia from the ischemic (anteroseptal) mid-apical ventricular wall and processed as previously described.4 Briefly, tissue samples were fixed in 10% neutral buffered formalin for 48 hours and processed by dehydrating the tissue in increasing concentrations of ethanol. Samples were then cleared in xylene, embedded in paraffin wax and cut into 4 micron sections.
For histopathological analysis sections were stained with Hematoxylin and Eosin (H&E) and Masson’s Trichrome. Necrotic tissue was defined by the presence of typical signs of coagulative necrosis including marginal contraction bands, fading and eventually loss of nuclei and striation in cardiomyocytes. Granulation tissue was defined by the presence of loose collagen and abundant capillaries with a nearly complete removal of necrotic myocytes by infiltrated macrophages. Lesion area was defined as the sum of necrotic and granulation areas.

For immunohistochemical analysis sections were deparaffinized and antigen unmasking was performed using heat induced epitope retrieval with citrate buffer at pH6. Before incubation with primary antibodies, endogenous peroxidase was blocked by incubation with H₂O₂ for 5 minutes and endogenous antigens were blocked with fetal bovine serum for 20 minutes. Primary antibody used was mouse monoclonal anti-PM1 (BMA biomedicals; T-3503) to detect neutrophils. As secondary antibodies we used a HRP-conjugated goat anti-mouse (Dako; P0447) for PM1. Bound antibody was revealed by staining with diaminobenzidine and nuclei were counterstained with hematoxylin. All immunohistochemical procedures were performed using an automated autostainer (Autostainer Plus®, Dako).

For analysis, the images were digitalized with a scanner (Nanozoomer-RS C110730®, Hamamatsu) and examined with image analysis software (Tissuemorph®, Visiopharm) by an experienced veterinary pathologist blinded to experimental procedure performed. For the Trichrome staining analysis, we used the AEC-DAB 35% protocol instead of the H&E 12% protocol used before, to increase sensitivity to changes in collagen content.
SUPPLEMENTAL FIGURES AND FIGURE LEGENDS

Online Figure I. Study design.
The study population comprised 25 pigs weighing 30-40 kg. Five pigs were sacrificed with no intervention other than baseline cardiac magnetic resonance (CMR), and served as healthy reference subjects. Four 5-pig groups underwent different closed-chest I/R protocols: 40-min-I/R (prolonged ischemia, controls), prolonged ischemia followed by postconditioning, prolonged ischemia preceded by preconditioning, or 20-min-I/R (short-duration ischemia). Arterial enhanced multidetector computed tomography was performed during coronary occlusion in all pigs as a reference standard for measuring the myocardial area at risk. Serial CMR-based tissue characterization was done in all pigs at baseline, and at 120-min, day1, day4, and day7 after I/R. Animals were sacrificed immediately after the final follow-up CMR scan, and myocardial tissue samples from ischemic areas were rapidly collected for histology and determination of water content.

CMR: cardiac magnetic resonance; I/R: ischemia/reperfusion; MDCT: multidetector computed tomography; PostC: postconditioning; PreC: preconditioning.
**SUPPLEMENTAL TABLES**

**Online Table I.** Temporal evolution of left ventricular mass, thickness, and positive- and negative-late gadolinium enhanced mass as assessed by cardiac magnetic resonance in pigs subjected to different I/R protocols and followed up to day 7 after I/R.

| Group               | CMR measure | Baseline | R-120min | R-24hours | R-Day4 | R-Day7 |
|---------------------|-------------|----------|----------|-----------|--------|--------|
| 40min-I/R (Control) | LV mass, g/m² | 64.0 (7.4) | 92.6 (6.0) | 71.1 (4.7) | 73.6 (5.2) | 74.3 (4.7) |
|                     | Thickness ratio | 1.06 (0.18) | 2.08 (0.36) | 1.23 (0.13) | 1.17 (0.15) | 1.08 (0.06) |
|                     | LGE (+) mass, g | - | 25.9 (4.7) | 17.2 (2.9) | 16.8 (2.4) | 14.6 (2.3) |
|                     | LGE (-) mass, g | - | 39.9 (3.4) | 39.6 (5.1) | 43.0 (4.5) | 43.5 (6.1) |
| 40min-I/R + PostC   | LV mass, g/m² | 53.3 (4.4) | 77.7 (11.7) | 54.1 (4.7) | 59.7 (4.8) | 60.5 (6.1) |
|                     | Thickness ratio | 0.97 (0.20) | 2.62 (0.62) | 1.31 (0.13) | 1.23 (0.13) | 1.10 (0.23) |
|                     | LGE (+) mass, g | - | 24.1 (6.6) | 13.3 (1.7)* | 16.1 (2.3) | 14.8 (1.7) |
|                     | LGE (-) mass, g | - | 27.5 (2.5) | 27.1 (2.7) | 33.1 (3.8) | 34.0 (2.1) |
| PreC + 40min-I/R    | LV mass, g/m² | 54.0 (9.5) | 58.5 (8.6) | 56.7 (6.2) | 66.1 (9.9) | 60.2 (6.2) |
|                     | Thickness ratio | 1.13 (0.15) | 1.34 (0.34) | 1.22 (0.20) | 1.21 (0.07) | 1.24 (0.18) |
|                     | LGE (+) mass, g | - | 8.8 (2.1)* | 7.2 (3.6)* | 3.1 (1.3)* | 2.5 (1.8)* |
|                     | LGE (-) mass, g | - | 34.8 (9.0) | 34.0 (8.7) | 39.9 (7.0) | 41.0 (9.0) |
| 20min-I/R           | LV mass, g/m² | 56.30 (1.5) | 57.60 (3.2) | 57.10 (0.5) | 59.60 (2.5) | 60.80 (1.9) |
|                     | Thickness ratio | 1.05 (0.10) | 1.04 (0.14) | 1.06 (0.08) | 1.11 (0.04) | 1.03 (0.12) |
|                     | LGE (+) mass, g | - | 2.4 (1.8)* | 1.2 (0.5)* | 0.9 (0.3)* | 0.6 (0.3)* |
|                     | LGE (-) mass, g | - | 36.6 (2.6) | 39.9 (2.4) | 39.4 (3.9) | 40.5 (4.1) |

Values are mean (standard deviation). Table includes data from 4 groups of 5 pigs each undergoing 40min-I/R (controls), 40min-I/R plus postconditioning, 40min-I/R preceded by preconditioning, or 20min-I/R (short-duration ischemia). Total LV mass (g/m²) and end-diastolic wall thickness ratio (MaR/remote) was assessed in standard segmented cine SSFP sequence; while LV mass showing positive or negative late gadolinium enhanced was assessed in T1-IR-TFE sequence.

No significant differences were found between time-points within same group in regards to total LV mass or wall thickness ratio, except for the R-120 min CMR in 40min-I/R and 40min-I/R plus postconditioning, in which LV mass and thickness ratio was significantly higher than at the other time-points and groups due to the intense swelling of the ischemic myocardium at early reperfusion (p<0.05). Note that pigs subjected to 20min-I/R or preconditioning plus 40min-I/R did not show a significant increase of LV mass or thickness ratio at early reperfusion. P-value is adjusted for multiple comparisons.

Assessment of absolute LGE positive mass confirmed a cardioprotective effect in preconditioning and short-duration ischemia groups as compared to controls. A strong negative linear trend over time was shown for absolute LGE positive mass in all groups (p<0.01). *statistically significant differences (p<0.05) in regards to LGE positive areas as compared with the same time point in the 40min-I/R (control) group.

CMR: cardiac magnetic resonance; I/R: ischemia/reperfusion; PostC: post-conditioning; PreC: preconditioning; LV: left ventricle; g: grams; g/m²: grams normalized for body surface area; LGE (+): late gadolinium enhanced positive; LGE (-): late gadolinium enhanced negative.
Online Table II. Time profile of myocardium salvage as assessed by multidetector computed tomography and cardiac magnetic resonance during the first week after reperfused myocardial infarction in pigs subjected to different I/R protocols and cardioprotective strategies.

| Group               | Follow up   | R-120min | R-24hours | R-Day4  | R-Day7  |
|---------------------|-------------|----------|-----------|---------|---------|
| 40min-I/R (Controls)|             | -40.0 (21.2) | -7.8 (14.2) | -0.9 (20.8) | 9.1 (19.9) |
| 40min-I/R + PostC   |             | -50.7 (36.0) | -6.6 (13.7) | -5.1 (9.8)  | 2.3 (11.8)  |
| PreC + 40min-I/R    |             | 34.1 (18.3)* | 43.1 (28.6)* | 76.3 (10.5)* | 80.8 (14.2)* |
| 20min-I/R           |             | 77.0 (12.4)* | 87.4 (2.6)*  | 90.0 (4.3)* | 94.4 (3.2)*  |

Values are mean (standard deviation).
The extent of myocardium at risk as assessed by MDCT reference standard during the index coronary occlusion (MDCT-MaR) was 28.3±4.3%, 31.3±4.0%, 31.7±6.9%, and 24.6±6.7% of LV; for pigs subjected to 40min-I/R (controls), 40min-I/R plus postconditioning, preconditioning plus 40min-I/R, and 20min-I/R, respectively (statistically non-significant differences as compared to controls).
Myocardial salvage as assessed by MDCT/CMR [(MDCT MaR – CMR infarct size) / MDCT MaR, %] is presented in the table. Note that MDCT was performed only once (during the index ischemic event), while CMR was performed at all follow-up time-points.
A strong negative linear trend over time was shown for myocardial salvage as assessed with MDCT as reference standard for MaR in all groups (p<0.01).
*statistically significant differences (p<0.05) as compared with the same time point in the 40min-I/R (control) group. P-value is adjusted for multiple comparisons among groups for each time-point and imaging parameter.
CMR: cardiac magnetic resonance; MaR: myocardium at risk; I/R: ischemia/reperfusion; R: reperfusion; PostC: postconditioning; PreC: preconditioning.
**Online Table III.** Time course of intramyocardial hemorrhage and microvascular obstruction assessed by cardiac magnetic resonance up to day 7 after I/R in pigs subjected to different I/R protocols.

| Group          | CMR measure | Follow up          |        |        |        |        |
|----------------|-------------|--------------------|--------|--------|--------|--------|
|                |             | R-120min          | R-24hours | R-Day4 | R-Day7 |
| 40min-I/R (Control) | IMH, % of LV | 0.2 (0.5)         | 4.3 (1.8) | 4.4 (1.9) | 1.4 (1.4) |
|                | MVO, % of LV | 5.1 (4.5)         | 6.5 (2.1) | 2.3 (2.8) | 1.8 (2.9) |
| 40min-I/R + PostC | IMH, % of LV | 0.7 (0.9)         | 4.6 (3.1) | 6.2 (4.0) | 3.2 (2.3) |
|                | MVO, % of LV | 4.5 (1.7)         | 5.1 (2.8) | 2.7 (2.1) | 1.9 (2.8) |
| PreC + 40min-I/R | IMH, % of LV | 0.2 (0.4)         | 0.9 (1.2)* | 0.9 (0.9)* | 0.2 (0.3) |
|                | MVO, % of LV | 0.5 (0.8)*         | 0.8 (1.1)* | 0.3 (0.6) | 0.1 (0.2) |
| 20min-I/R      | IMH, % of LV | 0.2 (0.3)         | 0.7 (0.5)* | 0.7 (0.2)* | 0.4 (0.4) |
|                | MVO, % of LV | 0*                  | 0*                 | 0                 | 0                 |

Values are mean (standard deviation). Table includes data from 4 groups of 5 pigs each undergoing 40min-I/R (controls), 40min-I/R plus postconditioning, 40min-I/R preceded by preconditioning, or 20min-I/R (short-duration ischemia). Note that the degree of intramyocardial hemorrhage and MVO was significantly lower at crucial time-points evaluated in pigs subjected to 20min-I/R or preconditioning plus 40min-I/R than in pigs subjected to 40min-I/R or 40min-I/R plus post-conditioning. *statistically significant differences (p<0.05) as compared with the same time point in the 40min-I/R (control) group. P-value is adjusted for multiple comparisons among groups for each time-point and imaging parameter.

CMR: cardiac magnetic resonance; I/R: ischemia/reperfusion; PostC: post-conditioning; PreC: preconditioning; IMH: intramyocardial hemorrhage; MVO: microvascular obstruction; LV: left ventricle.
**Online Table IV.** Histological features in the ischemic myocardium of pigs sacrificed at day 7 after I/R.

|               | Healthy | 40min-I/R (Control) | 40min-I/R + PostC | PreC + 40min-I/R | 20min-I/R |
|---------------|---------|---------------------|-------------------|------------------|-----------|
| **Lesion, %** | 0.0 (0.0) | 45.0 (39.7) | 69.3 (25.7) | 10.1 (8.0)* | 0.0 (0.0)* |
| **Necrosis†, %** | 0.0 (0.0) | 8.9 (12.7) | 9.0 (10.8) | 0.3 (0.6) | 0.0 (0.0) |
| **Granulation tissue†, %** | 0.0 (0.0) | 91.1 (12.7) | 91.0 (10.8) | 99.7 (0.6) | 0.0 (0.0)* |
| **Neutrophils, number/mm²** | 6.3 (1.5) | 39.4 (43.4) | 63.2 (45.5) | 5.5 (3.3) | 11.2 (2.3) |
| **Collagen, %** | 0.0 (0.0) | 30.4 (6.1) | 31.2 (13.1) | 18.8 (8.0)* | 0.0 (0.0)* |

Neutrophils are reported as the mean (standard deviation) number of cells per square millimeter in the histological section. Collagen content is reported as the mean (standard deviation) percentage within the granulation tissue area. Given that a more sensitive protocol was used to quantify collagen content, the value obtained for pigs subjected to 40min-I/R (control) differs from that previously published for the same group of pigs.3

*statistically significant differences (p<0.05) as compared to the 40min-I/R (control) group. Healthy animals were excluded from comparisons. P-value is adjusted for multiple comparisons among groups for each one of the histological features.

†Necrosis and granulation tissue as defined within the lesion area (see detailed methods).

I/R: ischemia/reperfusion; PostC: post-conditioning; PreC: pre-conditioning.
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LEGENDS FOR VIDEO FILES

**Online Videos I and II.** Short-axis view of a standard segmented cine steady-state free-precession (SSFP) from a pig subjected to 40min-I/R (control) at baseline (Video I) and within the first two hours after reperfusion (Video II).

**Online Videos III and IV.** Short-axis view of a standard segmented cine steady-state free-precession (SSFP) from a pig subjected to 40min-I/R plus post-conditioning at baseline (Video III) and within the first two hours after reperfusion (Video IV).

**Online Videos V and VI.** Short-axis view of a standard segmented cine steady-state free-precession (SSFP) from a pig subjected to preconditioning plus 40min-I/R at baseline (Video V) and within the first two hours after reperfusion (Video VI).

**Online Videos VII and VIII.** Short-axis view of a standard segmented cine steady-state free-precession (SSFP) from a pig subjected to 20min-I/R at baseline (Video VII) and within the first two hours after reperfusion (Video VIII).