

Letters

Prognostic Role of CMR Imaging After Myocardial Infarction



We read with great interest the recent review by El Aidi and colleagues (1) highlighting the prognostic role of cardiac magnetic resonance imaging (CMR) in patients with myocardial infarction (MI) and patients with suspected or known coronary artery disease (CAD). However, several limitations of that review, especially regarding the prognostic value of CMR in the setting of acute MI, should be also brought to the reader's attention.

The authors conclude that left ventricular ejection fraction (LVEF) is the only CMR predictor of hard clinical events. This finding is inconsistent with previous CMR evidence (2-5) and is mainly a result of the arbitrary definition of an independent prognostic CMR variable chosen by the authors (CMR variable studied in <1,000 patients was classified as not enough evidence to draw definitive conclusions). Almost all CMR studies that were performed demonstrated a superior prognostic value of infarct size (IS) and/or microvascular obstruction (MO) over and above LVEF (2,3). As LVEF is influenced both by residual stunning of the viable myocardium and the necrotic, nonviable myocardium, IS can be a more specific marker for determining the extent of irreversible myocardial damage. Consequently, the primary goal of any infarction therapy is to reduce the amount of infarcted tissue.

The authors do not account for several important factors, issues, and confounders. First, they included a heterogeneous population of patients with ST-segment elevation myocardial infarction (STEMI) and also non-STEMI. However, the prognostic significance of CMR markers of myocardial damage has been only convincingly demonstrated in previous studies for patients with STEMI. Second, the authors included infarction patients reperfused by both primary percutaneous coronary intervention and thrombolysis. Third, it has been clearly demonstrated that late MO has a stronger impact on future cardiovascular events as compared with early MO and

should, therefore, be the only included CMR measure of microvascular injury (4). Finally, the authors do not mention the potential prognostic role of myocardial salvage at all, which has been also shown to be independently associated with hard clinical endpoints (5).

In our opinion, there is emerging and strong evidence that CMR markers of myocardial damage (especially IS, myocardial salvage, and MO) add incremental prognostic information to clinical, electrocardiographic, biomarker, angiographic and mere functional outcome markers, including LVEF (2-5). However, we agree with the authors that further, preferably multicenter, investigations are welcome and necessary to further underscore the prognostic significance of myocardial damage and reperfusion injury assessed by CMR.

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