Cardiovascular Magnetic Resonance (CMR) for the Evaluation of Myocardial Infarction in Patients with Non-obstructive Coronary Artery Disease (MINOCA)

F. De Michele¹ ² · F. S. Guerra¹ ² · V. Forte³ · A. Carrieri¹ · D. R. R. Chieppa⁴ · G. Guglielmi¹ ² ⁵

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Abstract

Purpose of Review Myocardial infarction in the absence of coronary artery disease is caused by a variety of clinical conditions, so it is important to detect the specific causes in order to perform a better prognostic stratification of these patients. The aim of this review is to summarize the most important findings that established the role of CMR (cardiovascular magnetic resonance) to detect the MINOCA (myocardial infarction with non-obstructive arteries) patients and the importance to differentiate them from myocardial infarction patients.

Recent Findings The role of CMR is crucial to diagnose the principal diseases involved in MINOCA, as demonstrated. The several MR sequences used in all the MINOCA patients showed different results for all the different causes of MINOCA and, surely, high-resolution MR with gadolinium enhancement has been considered the best method to differentiate the transmural lesions.

Summary Another fundamental aspect to be considered is the experience of CMR radiologists, which represent the most important element for the right diagnosis of MINOCA. Surely, in the future, CMR will be the most important technique of choice for MINOCA patients, playing a key role in their management.

Keywords MINOCA · Microvascular disease · CMR · Coronary artery disease · Myocardial infarction · Non-obstructive · Takotsubo · Myocarditis

Introduction

Myocardial infarction in the absence of obstructive coronary artery disease is found in 5% to 6% of all patients with acute infarction, who are referred for coronary angiography [1]. The term MINOCA indicates patients with evidence of atherosclerosis that is not hemodynamically compromising [2]. The management of patients with MINOCA is strictly dependent on the cause, so an evaluation should be mandatory in all these patients. The demographic and clinical characteristics of MINOCA patients differ from other patients with AMI (acute myocardial infarction). MINOCA patients are usually younger than patients with AMI—CAD (acute myocardial infarction–coronary artery disease). Women are disproportionately represented among individuals with MINOCA [3–5]; they make up close to 50% of the MINOCA population but only 25% of the population with AMI—CAD [5]. MINOCA patients have a lower prevalence of dyslipidemia than their counterparts with AMI—CAD [3–6]. Finally, other CAD risk factors, such as hypertension, diabetes mellitus, tobacco abuse, and family history of myocardial infarction, are less frequent in MINOCA patients [3–6].

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G. Guglielmi giuseppe.guglielmi@unifg.it

¹ Department of Clinical and Experimental Medicine, Foggia University School of Medicine, Viale L. Pinto 1, 71121 Foggia, Italy
² Radiology Unit, “Dimiccoli” Hospital, Barletta, Italy
³ Radiology Unit, “San Nicola Pellegrino” Hospital, Trani, Italy
⁴ Cardiology Unit, “Bonomo” Hospital, Andria, Italy
⁵ Radiology Unit, Hospital “Casa Sollievo Della Sofferenza”, San Giovanni Rotondo, Foggia, Italy
MINOCA Classification

The European Society of Cardiology proposed the following MINOCA criteria [7]:

- AMI criteria as defined by the “Third Universal Definition of Myocardial Infarction” [8]
- Non-obstructive coronary arteries as per angiographic guidelines with no lesions >50% in a major epicardial vessel (plaque disruption) [9].
- No other clinical specific cause that can be an alternative cause for the acute presentation [1].

Fundamental to the definition of MINOCA is the diagnosis of AMI with an elevated cardiac biomarker, typically a cardiac troponin > 99th percentile of the upper reference level on serial assessment [1]. To limit the diagnostic mistakes due to indiscriminate use of troponin, the “Fourth Universal Definition of Myocardial Infarction” redefined the concept of myocardial injury [10]. Similar to myocardial infarction, the myocardial injury is an elevated troponin beyond the 99th percentile of the upper reference level. The diagnostic challenge is to define these entities, because patients with myocardial injury can show symptoms that mimic myocardial infarction at the initial presentation. In the evaluation of patients with a suspected AMI, it is mandatory to exclude clinical causes for the elevated troponin (sepsis, pulmonary embolism), obstructive disease (complete occlusion of a small coronary artery), and non-ischemic mechanisms of myocyte injury that can mimic myocardial infarction (myocarditis). The elevation of troponin level and the elevation of ST segment, as expression of myocardial injury, are also found in Myocarditis and Takotsubo syndrome that are involved in MINOCA causes, despite the myocardial injury is not referable to a coronary obstruction [1]. Once these causes and pathologies have been considered, a diagnosis of MINOCA can be made (Table 1).

Diagnosis

Many studies have demonstrated the utility of cardiovascular magnetic resonance (CMR) in providing a reclassification of an AMI diagnosis. So CMR is considered a crucial multiparametric diagnostic tool by the European Society of Cardiology working group on MI (myocardial infarction) with non-obstructive coronary arteries [11•]. Based on a meta-analysis of studies using CMR as a diagnostic tool in MINOCA, myocarditis is the leading diagnosis, and when CMR is performed early, it can secure a diagnosis in up to 87% of cases [1–11•]. Other causes of MINOCA, as diagnosed by CMR, commonly include Takotsubo cardiomyopathy, acute MI, hypertrophic or dilated cardiomyopathy, and non-cardiac causes (such as pulmonary embolism). Although CMR can provide an evaluation of the myocardium in MINOCA, additional diagnostic testing of the coronary arteries, such as intravascular ultrasonography (IVUS) and optical coherence tomography (OCT), may elucidate the mechanism of MINOCA, including atherosclerotic plaque rupture, erosion, and spontaneous coronary dissection. Furthermore, provocative testing may support coronary vasospasm as a potential cause of MINOCA [1].

Radiological Features

CMR Study Protocol

In MINOCA patients, the CMR study should be performed within 7 days from symptom onset. A CMR study protocol to evaluate MINOCA patients include the evaluation of cardiac structure and function of LV (left ventricle) with cine imaging, presence of myocardial edema with T2-weighted (T2w-STIR) image, and presence of myocardial injury with late gadolinium enhancement (LGE) imaging. Moreover, the use of new tissue characterization techniques, T1 and extracellular volume (ECV), and T2 mapping are recommended, due to their diagnostic accuracy in the detection of myocardial damage [12, 13]. Recently, Pierre Francoise Lintigre et al. had showed the diagnostic yield of CMR with high-resolution late gadolinium enhancement, in order to investigate the intramural myocardial lesions in MINOCA patients [14••]. This retrospective study involved 229 patients categorized as MINOCA. The patients were evaluated after lab tests, electrocardiogram, echocardiogram, and conventional CMR, and were performed cine sequences, T2-weighted, first pass perfusion, conventional LGE at 10 min, and high-resolution LGE (HR LGE) free breathing. Was used CMR 1.5 T and the images were interpreted by two readers with 5 and 15 years of experience in a random order, months or years after the acquisitions, and they were blinded to the initial CMR report. Left ventricular volume and ejection fraction were quantified using Argus software (Siemens Medical System). Ventricular dilatation and systolic dysfunction were defined on the basis of previously reported normal values [15]. Cine images were used to look for the abnormality of wall motion or pericardial effusion. T2 weighted were analyzed to look for myocardial edema. LGE imaging was analyzed to look for myocardial perfusion defects and the HR LGE imaging volume could be reviewed in multiplanar reformation of any orientation, depending on LGE location. In each patient, the distribution of LGE was described as subendocardial,
subepicardial, and midwall. The criterion to diagnose the myocardial infarction on CMR was the presence of subendocardial or transmural LGE [16]. Myocarditis was instead diagnosed with the presence of definite midwall or subepicardial LGE in the absence of subendocardial LGE [12]. Takotsubo syndrome was diagnosed when wall motion abnormality was found evolving in the apical or basal in the absence of myocardial LGE [17]. In these patients were performed a follow-up at 3 months using CMR or TTE. A total of 229 patients were recruited for MINOCA. Electrocardiography showed ST elevation for myocardial infarction in 85 patients, and TTE were negative in 124 patients, coronary angiography showed normal arteries in 125 patients, and non-obstructive CAD in 104. All patients underwent CMR, and a subpopulation of 172 patients had HR LGE as added protocols when diagnosis remained inconclusive. These patients (86) had a definite diagnosis: myocardial infarction in 39, acute myocarditis in 32, Takotsubo cardiomyopathy in 13, and other diagnosis in 2. In this study, HR LGE could reveal the infarction in 14% and rule out the diagnosis of infarction in 12%. The interpretation of myocardial injury with HR LGE led to a modification of diagnosis in 45 of 172 patients, with a modification of prognosis, and a different decision making on therapy [14*]. HR LGE had showed in this study how it could be more incisive for the outcome of MINOCA patients when conventional CMR is negative.

Acute Myocarditis

Inflammation of the myocardium may occur as a result of exposure to antigens such as viruses, bacteria, protozoa, drugs, toxins, or as an autoimmune condition. CMR is indicated for patients with new presentation or persisting of symptoms suggestive of myocarditis (dyspnea, orthopnea, palpitations, chest pain, effort intolerance), if viral etiology is suspected with no evidence of coronary stenosis. According to the European Society of Cardiology Working group and to the last guidelines, myocarditis diagnosis is performed by CMR as a Class I procedure for the identification of myocarditis [18]. The CMR protocol used for these patients consisted in evaluation of: Presence and pattern of myocardial edema with T2w-STIR images: Patchy areas of high signal intensity; Subepicardial or septal high signal intensity; Transmural high signal intensity (not specific for myocardial inflammation); Global high signal intensity evaluated through the T2-ratio technique; Evaluation of myocardial early gadolinium enhancement ratio that explore the regional vasodilation as a feature of tissue inflammation. The presence and pattern of necrosis and fibrosis with LGE imaging: Patchy areas of enhancement; Subepicardial enhancement; and transmural enhancement. These criteria are evaluated in parallel to the cardiac structure and function with cine imaging due to the evaluation of the presence of LV dysfunction with regional or global systolic dysfunction or pericardial effusion which were considered supportive criteria. According to LLC (Lake Louise criteria) CMR findings of T2 STIR sequences for myocardial edema and early gadolinium sequences for hyperemia are predictive of myocarditis, if at least 2 of 3 of the previous criteria are present. The mapping techniques can derive T1 and T2 relaxation time and permit ECV (extracellular volume) calculation. These techniques provide better diagnosis than traditional sequences in the
diagnosis of acute myocarditis, in particular the reported AUC (area under the curve) for T1 mapping was 0.95, for T2 mapping was 0.88, and for ECV (extracellular volume) 0.81 [18] (Fig. 1). To assess the role of CMR and to detect the non-obstructive causes of Minoca, Heidecker Bettina et al., in a study of 2019, had investigated the myocarditis patients in a population of 556 MINOCA patients with high levels of Troponin and angina-like symptoms, without significant coronary artery obstruction [19]. The study was a retrospective chart review between 2011 and 2017. CAD was excluded using a coronary angiography or a coronary CT in all the patients with a rise in Troponin level and angina-like symptoms. All the patients with an obstruction < 50% had a CMR to exclude other causes of MINOCA. CMR 1.5 T was used and all the images were performed under the supervision of the head of cardiac imaging, who had several years of experience. Were performed, with this purpose, cine-images, T2-weighted imaging and LGE (Late Gadolinium Enhancement) images evaluated in 10 min. Myocarditis was considered when there was an epicardial or midwall distribution of LGE, associated with a patchy pattern and non-coronary distribution [20, 21]. 556 patients were admitted, and CMR was able to diagnose 76 (13.7%) myocarditis, 34 (6.1%) dilated cardiomyopathy, 41 ischemia during stress, 15 hypertrophic, 13 pericarditis, 8 amyloidosis, 7 Takotsubo, 5 sarcoidosis, 3 Fabry’s disease, others 22, and 332 normal heart. For these reasons, a great number of myocarditis was investigated, and this study had showed the central role of CMR to detect another important cause of MINOCA, often unremarkable by systematic work up of the patients with angina-like symptoms, rise of troponin, and no significant coronary artery disease [22].

**Takotsubo Syndrome**

CMR is able to identify LV (left ventricle) wall motion abnormalities, and provides the evaluation of right ventricle (RV) motility [17, 23]. The most common presentation of TS (Takotsubo) is LV apical akinesia with preserved function of the remaining segments, which causes the typical “apical ballooning” appearance. Mid-ventricular

Fig. 1 30-year-old patient with myocarditis symptoms. The diagnosis was made with these significant images according to Lake Louise Criteria: a T2 STIR 3-chamber view with subepicardial hyperintensity of inferior and inferolateral walls and edema. b Delayed enhancement in short-axis view with subepicardial hyperintensity of inferior and inferolateral walls and edema of LV. c Delayed enhancement in 3-chamber view with subepicardial hyperintensity of inferolateral wall with edema.
type, basal or inverted type, focal variants, and isolated or concomitant RV (right ventricle) akinesia have also been described but with lower frequency [24]. Wall motion abnormalities are reversible and complete recovery of systolic function has been demonstrated at 3 months of follow-up [25]. CMR represents a widely established method to non-invasively assess myocardial tissue, within a scanning time of 30 min [26]. CMR myocardial edema can be detected by means of T2w-STIR images or assessed either on T1- or T2-mapping sequences, with increased values in both ballooning and non-ballooning segments [27]. Usually, no LGE (late gadolinium enhancement) is seen in Takotsubo patients. In conclusion, wall motion abnormalities and absence of LGE represent the findings that permit CMR for providing a diagnosis of Takotsubo in patients with high serum troponin levels, and for differentiating this entity from myocardial infarction and myocarditis [27]. The epidemiological data of MINOCA showed that women were disproportionately represented among MINOCA patients, in fact Harmony R. Reynolds et al., in a recent prospective study of November 2020, had involved 301 women at 16 sites, with a clinical diagnosis et al., in a recent prospective study of November 2020, had involved 301 women at 16 sites, with a clinical diagnosis of myocardial infarction [28]. When coronary angiography showed a < 50% stenosis in all major artery, an OCT was performed, followed by CMR. CMR was performed one week after the symptoms, and the images were interpreted by different laboratories blinded to the results of other laboratories. OCT was performed using an OCT ILUMIEN OPTIS, and CMR was 1.5 T or 3.0 T and T2-weighted, fat-suppressed, fast-spin echo, LGE at 10 or 15 min, and T1 mapping were performed [29]. Although patients involved were 301, only 147 had an OCT evaluation and 67 had a culprit lesion (rupture plaque, intra-plaque cavity, and thrombus without rupture plaque). Three patients had coronary spasm. CMR was interpretative in 116 of the 147 participants with an interpretable OCT and showed regional injury in 24 women, myocarditis in 17, Takotsubo cardiomyopathy in 4, non-ischemic cardiomyopathy in 3, and normal heart in 30 patients [30]. Furthermore, this study had demonstrated that among 116 women who had OCT and CMR evaluation, 98 had an abnormality on one or both methodic, and surely the prevalence of abnormalities indicating a cause of MINOCA presentation after the combination of OCT and CMR was 84.5%, significantly higher than OCT alone or CMR alone.

Ischemic Causes of MINOCA

CMR imaging is a diagnostic tool to be employed in MINOCA patients with suspected acute myocardial infarction. Myocardial edema is evaluated on T2w-STIR images and LGE that permit the evaluation of myocardial damage and gives information about the mechanisms. Transmural LGE indicates an ischemic cause of injury, but it does not specify the particular mechanism of ischemia, while a non-ischemic appearance of LGE (mesocardial or subepicardial localization) is indicative of other myocardial disorders such as myocarditis and other cardiomyopathies [30] (Figs. 2, 3). To this purpose, Amerdeep Gosh Dastidar et al., in a study of 2018, investigated the prognostic impact of CMR in these patients [31] in a retrospectively study of 388 MINOCA. The protocols provided the use of 1.5 T CMR and included cine imaging, T2 weighted for myocardial edema, and early and late gadolinium enhancement imaging [32]. Patients were grouped in 4 categories: myocardial Infarction, myocarditis, cardiomyopathies, and normal CMR. CMR was performed with a median of 37 days from clinical presentations, and the images were analyzed by radiologists with 15 years of experiences, all with an ESC CMR level 3 certification. The cases of myocarditis were diagnosed by LLC: T2 STIR sequences for myocardial edema and early gadolinium sequences for hyperemia [19]. Myocardial infarction was diagnosed by territorial subendocardial and transmural LGE. Cardiomyopathies, in particular Takotsubo, were diagnosed by T2 STIR for myocardial edema and regional wall motion in mid-cavity or apical distribution with no myocardial LGE [24]. Normal heart was defined based on no regional wall motion abnormality, no myocardial edema, and no myocardial LGE. In this study, CMR was able to identify the causes of troponin rise in 74% of patients, which were so distributed: 25% myocarditis, 25% patients with myocardial infarction, 25% cardiomyopathy, and 26% normal heart. The cardiomyopathy group, represented by Takotsubo syndrome, had showed a worst prognosis over a median of follow-up of 3.5 years with a 15% of death. In particular, in these patients, CMR could better identify the cases by delineating the myocardial edema, RV involvement, presence of thrombus caused by the stunning of apical wall, and LV outflow tract obstruction. The study had also showed the importance of CMR and ST elevation on ECG, as best predictors of mortality. All these data had led a better stratification of all the MINOCA patients involved in order to assess the right treatment and change the clinical history of the patients.

Other Causes of MINOCA

Hypertrophic Cardiomyopathy

HCM (Hypertrophic cardiomyopathy) patients present with increased ventricular myocardial thickness and could have hyperintense myocardial areas at T2w-STIR images, altered first pass perfusion and presence of LGE. T1- and
T2-mapping MR imaging techniques are able to identify myocardial injuries associated with HCM without gadolinium-based contrast agents\(^{33–37}\). Therefore, T1 mapping without contrast may identify myocardial scarring as well as interstitial fibrosis in HCM. The T2 alterations are associated with edema and may represent acute ischemic damage so hypertrophic cardiomyopathy could be the expression of MINOCA. Furthermore, the presence of LGE seems to be a marker of increased risk of non-sustained ventricular tachycardia episodes as a prognostic
factor. Cine SSFP (steady-state free precession sequences) MR imaging is used for morphological assessment of HCM and evaluation of cardiac function. These sequences are able to define all phenotypes of HCM because of its no limited view. Using retrospective gating, cine SSFP is the most important method to quantify the myocardial thickness and mass, which are related to the prognosis of HCM [33–37]. Late gadolinium enhancement is the most valuable MR imaging sequence for HCM, because it identifies the myocardial fibrosis that contributes to risk stratification. Other cardiomyopathies with similar symptoms and LVH enhancement is also useful for differentiating HCM from other cardiomyopathies with similar symptoms and LVH [35, 36].

Conclusions

These studies highlight the potential benefit of using CMR to evaluate MINOCA patients for a better treatment based on the detected cause. CMR can confirm MI (myocardial infarction) for accurately categorizing patients according to the contemporary definition of MINOCA. Additional tests of the coronary arterial system could provide further insights into the cause of MINOCA in patients who have an MI (Figs. 2 and 3), and although invasive, remain the first step to diagnose MINOCA. CMR is non-invasive and can differentiate MI from myocarditis (Fig. 1) and various cardiomyopathies, led to personalized treatment pathways and led to a better prognostic stratification of these patients. A multi-parametric CMR protocol may include functional imaging and advanced tissue characterization techniques, such as T1 mapping, LGE imaging, and HR LGE, that if performed early within 1 week of the acute presentation could change statistically significant the outcome of MINOCA patients, changing clinical assessment, therapy, and prognosis. The advantages of using CMR to guide management in MINOCA patients include a true diagnosis of MI, to distinguish from non-ischemic causes, and evaluate the extent and functional impact of the acute myocardial injury. Different and combined methodical as OCT and CMR had showed the importance of this approach to detect the obstructive causes of MINOCA. CMR has demonstrated clinical utility in diagnosis and prognosis of MINOCA and this has led to insert CMR in the guidelines, as a first method to investigate MINOCA in all inconclusive diagnoses.

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