

Identifying post-myocardial infarction patients at risk by imaging techniques

Nico Bruining*

Thoraxcenter, Erasmus MC, Rotterdam, The Netherlands

Most myocardial infarction patients will undergo emergency percutaneous intervention (PCI) today. However, most of these patients will have diffuse cardiovascular disease and will often show more disease than a single culprit lesion. It is important to identify those patients at risk after a myocardial infarction (MI). The current guidelines suggest that the resting left ventricular (LV) function must be assessed as part of the risk stratification by both the ESC (ESC ST-elevation myocardial infarction (STEMI) in 2012)¹ as well by the ACCF/AHA (2013)². The guidelines suggest that patients with an LV ejection fraction (LVEF) <30-40% and New York Heart Association (NYHA) functional class I or II should receive an implantable cardioverter-defibrillator (ICD) treatment.

However, there are two major concerns to this classification and those are: 1) Are the current imaging methods accurate enough to measure this threshold in LVEF? And 2) the great majority of patients with a sudden cardiac death (SCD) have an LVEF >30%.³ So the major question is how we can identify the patient at risk and whether we have any other possibilities to identify them by imaging⁴?

There are currently many additional imaging methods available which are aimed at identifying vulnerable coronary lesions⁵,

such as: intravascular ultrasound (IVUS), optical coherence tomography (OCT) and near infrared spectroscopy (NIRS), to name a few. Some of these imaging methods can be used for in-depth analysis of plaque components as by example IVUS-Virtual Histology. Most of these intracoronary imaging techniques are used to identify the so-called thin-cap fibro-atheroma's (TCFA's)^{6,7}. Also functional measurements of coronary blood flow, e.g. fractional flow reserve (FFR)⁸ or even virtual FFR by multi-slice computed tomography (MSCT)⁹.

However, not a single imaging method could identify these vulnerable plaques at itself, the results up until now are somewhat disappointing. We expect that combination of the results of the individual methods by multi-modality imaging, might improve this¹⁰. The ultimate multi-modality assessment of the LV and the heart might be "electro-mechanical imaging"¹¹.

Identifying the vulnerable patient at risk after MI is a difficult task³. Although the imaging guidelines today recommend to measure the LVEF at rest to identify patients at risk and who might benefit from additional treatment, there is still a large scientific debate if this is appropriate enough. More recent imaging methods are necessary and perhaps multi-modality imaging could provide better insight into the very important topic of identifying patients at risk.

Received: 15th Feb 2014

*Address for correspondence: Erasmus MC, Thoraxcenter, Department of Cardiology, Room BA-571, Rotterdam, The Netherlands.

Phone: +31107033934

Email: n.bruining@erasmusmc.nl

KEYWORDS: myocardial infarction, risk stratification, imaging, left ventricular function.

CITATION: *Cardiol Croat.* 2014;9(3-4):92.

Literature

1. The Task Force on the management of ST-segment elevation acute myocardial infarction of the European Society of Cardiology (ESC); Steg PG, James SK, Atar D, et al. ESC guidelines for the management of acute myocardial infarction in patients presenting with ST-segment elevation. *Eur Heart J.* 2012;33:2569-619.
2. O'Gara PT, Kushner FG, Ascheim DD, et al; American College of Cardiology Foundation/American Heart Association Task Force on Practice G. 2013 accf/aha guideline for the management of st-elevation myocardial infarction: A report of the American College of Cardiology Foundation/American Heart Association Task Force on Practice Guidelines. *Circulation.* 2013;127:e362-425.
3. Dagues N, Hindricks G. Risk stratification after myocardial infarction: is left ventricular ejection fraction enough to prevent sudden cardiac death? *Eur Heart J.* 2013;34:1964-71.
4. Bruining N, Knaapen M, de Winter S, et al. A histological "fly-through" of a diseased coronary artery. *Circ Cardiovasc Imaging.* 2009;2:e8-9.
5. Libby P. Current concepts of the pathogenesis of the acute coronary syndromes. *Circulation.* 2001;104:365-72.
6. Virmani R, Kolodgie FD, Burke AP, Farb A, Schwartz SM. Lessons from sudden coronary death: A comprehensive morphological classification scheme for atherosclerotic lesions. *Arterioscler Thromb Vasc Biol.* 2000;20:1262-75.
7. Choi BJ, Prasad A, Gulati R, et al. Coronary endothelial dysfunction in patients with early coronary artery disease is associated with the increase in intravascular lipid core plaque. *Eur Heart J.* 2013;34:2047-54.
8. Tonino PA, Fearon WF, De Bruyne B, et al. Angiographic versus functional severity of coronary artery stenoses in the fame study fractional flow reserve versus angiography in multivessel evaluation. *J Am Coll Cardiol.* 2010;55:2816-21.
9. Koo BK, Erglis A, Doh JH, et al. Diagnosis of ischemia-causing coronary stenoses by noninvasive fractional flow reserve computed from coronary computed tomographic angiograms. Results from the prospective multicenter discover-flow (diagnosis of ischemia-causing stenoses obtained via noninvasive fractional flow reserve) study. *J Am Coll Cardiol.* 2011;58:1989-97.
10. Bruining N, de Winter S, Serruys PW. Intravascular ultrasound registration/integration with coronary angiography. *Cardiol Clin.* 2009;27:531-40.
11. Galeotti L, van Dam PM, Loring Z, Chan D, Strauss DG. Evaluating strict and conventional left bundle branch block criteria using electrocardiographic simulations. *Europace.* 2013;15:1816-21.