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Villa Doria D'Angri - Via F. Petrarca 80, Napoli



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UOC Cardiologia-Utic-Emodinamica



A.O.R.N A. Cardarelli

STATE-OF-THE-ART REVIEW

2024 Update

International Consensus Statement on Platelet Function and Genetic Testing in Percutaneous Coronary Intervention





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TABLE 2	Clinical	Phenotypes	Identified	by	Genetic	Testing
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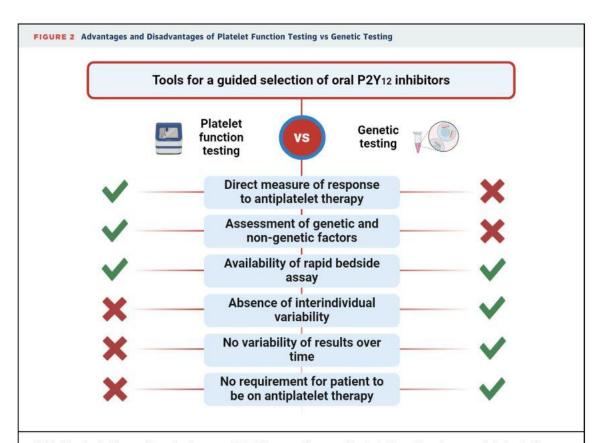
Metabolizer Phenotypes	Prevalence (%)	Response to Clopidogre		
Ultrarapid *17/*17	1-5	Normal or increased		
Rapid *1/*17	10-30	Normal or increased		
Normal *1/*1	25-60	Normal		
Intermediate *1/*2, *1/*3, *2/*17, *3/*17	20-45	Reduced		
Poor	2-15	Significantly reduced		

The prevalence of loss-of-function alleles is highest in Oceanian biogeographical group may present up to metabolizer.

*2/*2, *3/*3, *2/*3

FIGURE 1 Therapeutic Window and Platelet Reactivity Levels Commonly Associated With Specific P2Y12 Inhibitor Therapy in Patients **Undergoing Percutaneous Coronary Interventions** Risk Bleeding risk Ischemic risk **Platelet** reactivity **Low Platelet Reactivity Optimal Platelet Reactivity High Platelet Reactivity ↓** Lower Upper Commonly associated with Commonly associated with Commonly associated with Prasugrel/ Clopidogrel Clopidogrel responders **Ticagrelor** poor-responder

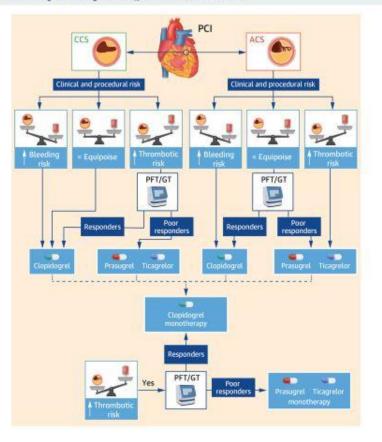




Platelet function testing provides a direct measure of platelet aggregation accounting for both genetic and non-genetic factors but is characterized by a large interindividual variability, often requiring multiple testing over time and needing to be performed while the patient is on antiplatelet treatment. Genetic testing has the advantage that the genetic makeup of an individual remains unvaried and does not require a patient to be on antiplatelet treatment. However, genetic testing contributes only in part to antiplatelet drug response, with other variables, including clinical factors, playing a contributing role.



CENTRAL ILLUSTRATION Consensus Recommendations on the Use of Platelet Function and Genetic Testing for Guiding Oral P2Y12 Inhibitor Treatment in PCI



Angiolillo DJ, et al. JACC Cardiovasc Interv. 2024;17(22):2639-2663.

Among chronic coronary syndrome (CCS) patients undergoing percutaneous coronary intervention (PCI), patients should undergo the guideline-recommended dual antiplatelet therapy (DAPT) with clopidogrel. However, among patients at high clinical and/or procedural technic risk and without increased bleeding risk, or in patients with a history of recurrent bithemic events, platelet function testing (PFT) or genetic testing (GTI) should be used to escalate P2Y12 inhibition with prasugrel or ticagretor among clopidogrel poor responders. Among acute coronary syndrome (ACS) patients undergoing PCI, patients at high bleeding and low ischemic risk should undergo DAPT with clopidogrel, while those with high ischemic and low bleeding risk should undergo guideline-recommended DAPT with prasugrel or ticagrelor. PFT or GT should be considered in the subgroup of patients who are at both increased risk of bleeding and ischemic events, aiming at descalating P2Y12 inhibition (ie, switching from prasugrel or ticagrelor to clopidogrel among clopidogrel responders). Among patients treated with P2Y12 inhibitor monotherapy, PFT or GT should be considered in patients treated with clopidogrel monotherapy, particularly if at high clinical and/or procedural ischemic risk or with a history of recurrent ischemic events.



Management of Coronary Vulnerable Plaque With Medical Therapy or Local Preventive Percutaneous Coronary Intervention





Hoyun Kim, MD, Jung-Min Ahn, MD, Do-Yoon Kang, MD, Jinho Lee, MD, Yeonwoo Choi, MD, Seung-Jung Park, MD, Duk-Woo Park, MD





Preventive percutaneous coronary intervention versus optimal medical therapy alone for the treatment of vulnerable atherosclerotic coronary plaques (PREVENT): a multicentre, open-label, randomised controlled trial

Seung-Jung Park*, Jung-Min Ahn*, Do-Yoon Kang, Sung-Cheol Yun, Young-Keun Ahn, Won-Jang Kim, Chang-Wook Nam, Jin-Ok Jeong, In-Ho Chae, Hiroki Shiomi, Hsien-Li Kao, Joo-Yong Hahn, Sung-Ho Her, Bong-Ki Lee, Tae Hoon Ahn, Ki-Yuk Chang, Jei Keon Chae, David Smyth, Gary S Mintz, Gregg W Stone, Duk-Woo Park, for the PREVENT Investigators†



> Eur Heart J Cardiovasc Imaging. 2024 Nov 8:jeae289. doi: 10.1093/ehjci/jeae289. Online ahead of print.

High-risk features in non-culprit lesions and clinical outcome after NSTEMI versus STEMI

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Figure 1 Study flowchart

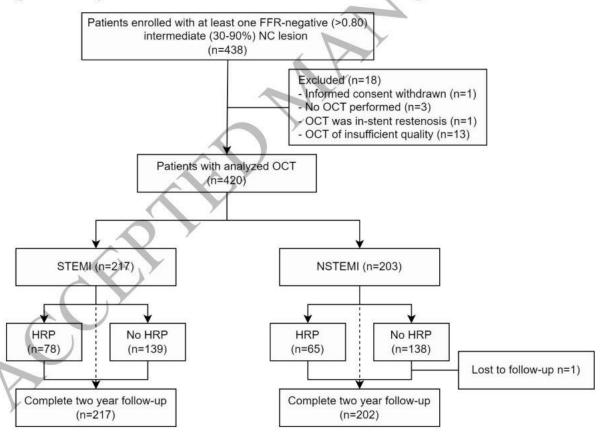
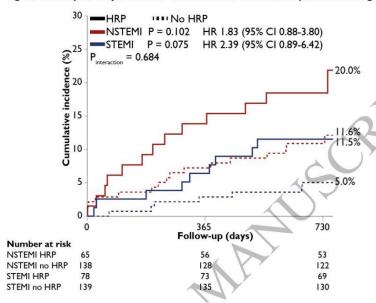
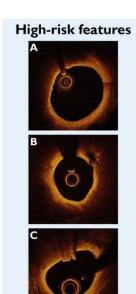
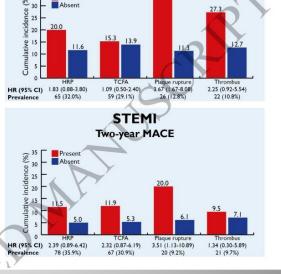




Figure 3 Two-year major adverse cardiovascular events and presence of high-risk plaques







NSTEMI Two-year MACE

Present



Icosapent ethyl following acute coronary syndrome: the REDUCE-IT trial

Neila Sayah (1) 1*, Deepak L. Bhatt (1) 2, Michael Miller (1) 3, Eliot A. Brinton (1) 4, Terry A. Jacobson (1) 5, Steven B. Ketchum (1) 6, Lixia Jiao (1) 6, Armando Lira Pineda (1) 6, Ralph T. Doyle Jr. (1) 6, Jean Claude Tardif (1) 7, Christie M. Ballantyne (1) 8, and Ph. Gabriel Steg (1) 1,9; on behalf of the REDUCE-IT Investigators †

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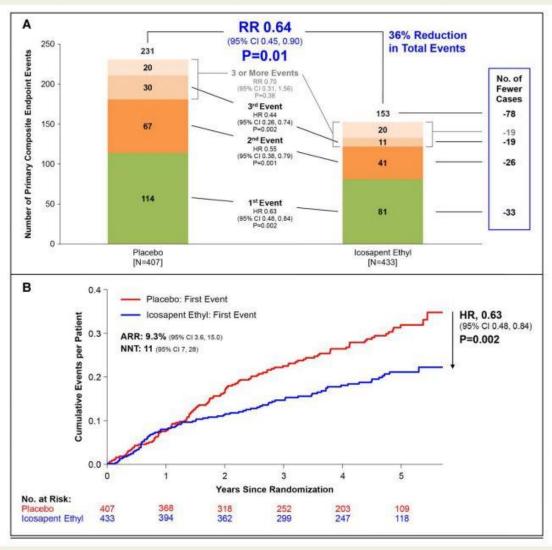


Figure 1 (A) First and subsequent events for the primary composite outcome in patients with recent ACS. (B) Cumulative incidence curves of the primary composite outcome in patients with recent ACS



Coronary bypass surgery for multivessel disease after percutaneous coronary intervention in acute coronary syndromes: why, for whom, how early?

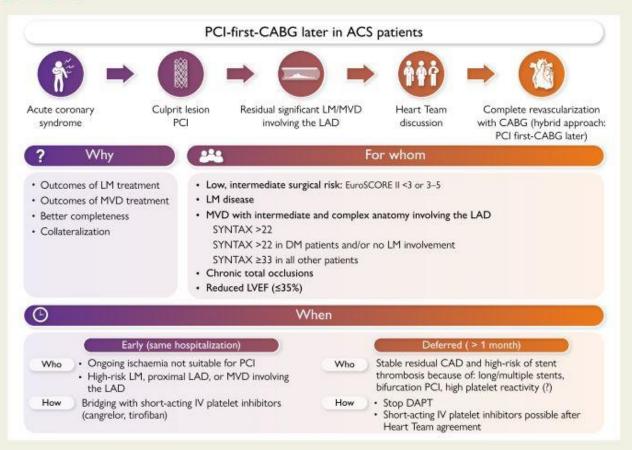
Laura Besola 1, Andrea Colli 1 1, and Raffaele De Caterina 1 2*

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Graphical Abstract



'PCI first-CABG later' in acute coronary syndrome patients. Patients with an acute coronary syndrome should receive immediate treatment of the culprit lesion with percutaneous coronary intervention. In cases of residual significant left main disease or multivessel disease involving the left



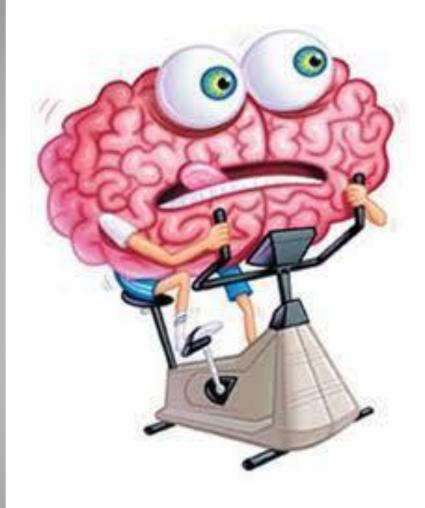
Moderate-intensity aerobic exercise inhibits cell pyroptosis to improve myocardial ischemia-reperfusion injury

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Yu Wang, Yushan Li, Chaofan Chen, Hailong Zhang, Weili Liu, Chao Wu, Haonan Chen, Ran Li, Jinghan Wang, Yingchao Shi, Shengfang Wang & Chuanyu Gao





Mens sana in corpore sano







