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Guidelines acc aha dapt 2016 acc/aha

2015 ACC/AHA/SCAI focused update on primary percutaneous coronary intervention for patients with ST-elevation myocardial Infarction: An update of the 2011 ACCF/AHA guideline for the management of ST-elevation myocardial infarction: A report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines and the Society for Cardiovascular Angiography and Interventions. 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Mar 29, 2016 | Debabrata Mukherjee, MD, FACC Authors: Levine GN, Bates ER, Bittl JA, et al. Citation: 2016 ACC/AHA Guideline Focused Update on Duration of Dual Antiplatelet Therapy in Patients With Coronary Artery Disease: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. An Update of the 2011 ACC/AHA/SCAI PCI Guideline, 2011 ACC/AHA CABG Guideline, 2012 ACCF/AHA/ACP/AATS/PCNA/SCAI/STS SIHD Guideline, 2013 ACC/AHA STEMI Guideline, 2014 ACC/AHA NSTE-ACS Guideline, 2016; Mar 29: [Epub ahead of print]. The following are key points to remember about the updated guideline on duration of dual antiplatelet therapy (DAPT) in patients with coronary artery disease (CAD). The scope of this focused update is limited to addressing recommendations on duration of antiplatelet therapy, with the addition of a P2Y12 inhibitor to aspirin monotherapy, and prolongation of DAPT, necessitate a fundamental tradeoff between decreasing ischemic risk and increasing bleeding risk. Decisions regarding treatment with and duration of study data, and patient preference. Recommendations in the document apply specifically to duration of P2Y12 inhibitor therapy in patients with CAD treated with DAPT. Aspirin therapy should almost always be continued indefinitely in patients treated with DAPT, are associated with DAPT. Aspirin therapy should almost always be continued indefinitely in patients treated with DAPT. aspirin. The recommended daily dose of aspirin in patients treated with DAPT is 81 mg (range 75-100 mg). In patients with SIHD treated with DAPT after drug-eluting stent (DES) implantation, P2Y12 inhibitor therapy with clopidogrel should be given for at least 6 months (Class I). In patients with SIHD treated with DAPT after bare-metal stent (BMS) implantation, P2Y12 inhibitor therapy (clopidogrel) should be given for a minimum of 1 month (Class I). In patients with DAPT after BMS or DES implantation who have tolerated DAPT without a bleeding complication and who are not at high bleeding risk (e.g., prior bleeding on DAPT, coagulopathy, oral anticoagulant use), continuation of DAPT with clopidogrel for longer than 1 month in patients treated with DES may be reasonable (Class IIb). In patients with acute coronary syndrome (ACS) (non-ST elevation [NSTE]-ACS or ST elevation myocardial infarction [STEMI]) treated with DAPT after BMS or DES implantation, P2Y12 inhibitor therapy (clopidogrel, prasugrel, or ticagrelor) should be given for at least 12 months (Class I). In patients with ACS (NSTE-ACS or STEMI) treated with coronary stent implantation who have tolerated DAPT without a bleeding complication and who are not at high bleeding risk (e.g., prior bleeding on DAPT, coagulopathy, oral anticoagulant use), continuation of DAPT (clopidogrel, prasugrel, or ticagrelor) for longer than 12 months may be reasonable (Class IIb). A new risk score (the "DAPT score"), derived from the Dual Antiplatelet Therapy study, may be useful for decisions about whether to continue (prolong or extend) DAPT in patients treated with coronary stent implantation. In patients with ACS (NSTE-ACS or STEMI) treated with DAPT after coronary stent implantation and in patients with NSTE-ACS treated with medical therapy alone (without revascularization), it is reasonable to use ticagrelor in preference to clopidogrel for maintenance P2Y12 inhibitor therapy (Class IIa). Among those who are not at high risk for bleeding complications and who do not have a history of stroke or transient ischemic attack, it is reasonable to choose prasugrel over clopidogrel for maintenance P2Y12 inhibitor therapy (Class IIa). In patients with ACS (NSTE-ACS or STEMI) being treated with DAPT who undergo coronary artery bypass grafting (CABG), P2Y12 inhibitor therapy should be resumed after CABG to complete 12 months of DAPT therapy after ACS (Class I). In patients with STEMI treated with DAPT in conjunction with fibrinolytic therapy, P2Y12 inhibitor therapy (clopidogrel) should be continued for a minimum of 14 days and ideally at least 12 months (Class I). Elective noncardiac surgery should be delayed 30 days after BMS implantation and optimally 6 months after DES implantation. In patients treated with DAPT after coronary stent implantation who must undergo surgical procedures that mandate the discontinuation of P2Y12 inhibitor therapy, it is recommended that aspirin be continued if possible and the P2Y12 platelet receptor inhibitor be restarted as soon as possible after surgery (Class I). Clinical Topics: Acute Coronary Syndromes, Anticoagulation Management, Cardiovascular Angiography and Intervention, Atherosclerotic Disease (CAD/PAD), Anticoagulation Management and ACS, Interventions and ACS, Interventions and Coronary Artery Disease Keywords: Acute Coronary Syndrome, Coronary Artery Disease, Adenosine, Anticoagulants, Aspirin, Blood Platelets, Coronary Artery Disease, Adenosine, Anticoagulants, Aspirin, Blood Platelets, Coronary Artery Disease, Stroke, Stroke, Interventions and Coronary Artery Disease, Adenosine, Anticoagulants, Aspirin, Blood Platelets, Coronary Artery Disease, Adenosine, Anticoagulants, Aspirin, Surgical Procedures, Operative, Thrombolytic Therapy < Back to Listings