Dynamic Response to Aspirin in Patients with ACS, Clinical and Prognostic Implications Spectre, Galia ${ }^{1}$; Mosseri, Morris ${ }^{2}$; Loncar, Sasa ${ }^{3}$; Varon, David ${ }^{1}$; Alcalai, Ronny ${ }^{3}$ ${ }^{1}$ Hadassah Hebrew University Medical Center, Coagulation Unit, Hematology, Jerusalem, Israel; ${ }^{2}$ Meir Medical Center, Cardiology Division, Kfar-Sava, Israel; ${ }^{3}$ Hadassah Hebrew University Medical Center, Heart Institute, Jerusalem, Israel

Background: Increased platelet reactivity and reduced response to anti-platelet drugs may result in recurrent ischemic events after acute coronary syndrome (ACS).
Aim: To evaluate the laboratory response to aspirin in patients with ACS before and after percutaneous coronary intervention (PCI) and assess its effect on major adverse clinical events. Methods and Results: Sixty three consecutive patients with ACS were tested for the response to aspirin by light transmittance aggregometry (LTA) and the IMPACT-R test (both with arachidonic acid- AA) before and 2-4 days after PCI and clopidogrel loading. Patients were followed for clinical events up to 15 month from PCI. Response to aspirin improved significantly after PCI and clopidogrel treatment: mean AA-induced LTA decreased from $34.9 \pm 3.35 \%$ before PCI to $15.2 \pm 2.2 \%$, and surface coverage increased from $2.2 \pm 0.27 \%$ to $6.2 \pm 0.6 \%$ ( $\mathrm{p}<0.0001$ for both methods). The improved response to aspirin after PCI correlated with the response to clopidogrel (LTA and IMPACT-R, $\mathrm{p}<0.01$ ). Patients with good laboratory response to aspirin before but not after PCI had significantly lower major cardiovascular event rate during 15 months follow up, in multivariate analysis.
Conclusion: The laboratory response to aspirin is highly dynamic in patients with ACS. The improved response to aspirin following PCI may result from stabilization of coronary artery disease and/or clopidogrel treatment. The laboratory response to aspirin before PCI and clopidogrel loading is a sensitive marker for platelet reactivity that correlates with clinical outcome in patients with ACS.

