The Effects of 80 mg Loading Dose Atorvastatin Pretreatment on Myocardial Protection During Percutaneous Coronary Intervention

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Objectives This study was designed to evaluate the peri-procedural myocardial injury and safety profile of atorvastatin in patients over 70 years old with non-ST segment elevation acute coronary syndromes (NSTEACS).

Methods A total of 157 patients over 70 years old with NSTE-acute coronary syndromes were randomly divided into the pretreatment with loading dose atorvastatin group or control atorvastatin group. The serum levels of Creatine kinase-MB (CK-MB) and cardiac troponin I (cTnI) were measured at the baseline and at 8, 12 and 24 h after the procedure. Clinical outcome was measured by taking the main end points of major adverse cardiac events including cardiac death, nonfatal acute myocardial infarction, or revascularisation during 30 days follow-up.

Results The proportion of patients with the serum levels of CK-MB and cTnI above normal up-limit value and that with PCI-related myocardial infarction at 24 h was significantly lower in high dose atorvastatin group than in the control dose group ((37.9% vs 50.5%, p=0.002, 5.4% vs 22.6%, p=0.041; 12.6% vs 33.4%, p=0.035, 4.3% vs 11.8%, p=0.034). At multivariable analysis, pretreatment with atorvastatin conferred an 67% risk reduction of PCI-related myocardial infarction. (p<0.05 No deaths and revascularisation were recorded at 30-day follow-up in both groups.

Conclusions Short-term pretreatment with a high dose of atorvastatin significantly reduces PCI-related myocardial infarction in elderly patients with NSTEACS and enjoys similar safety profile.