

## عنوان مقاله:

Lectin-Like OLR<sub>1</sub> 3'UTR Rs1050286 Gene Polymorphism and Plasma Oxidized-LDL in Coronary Artery Disease and Their Relation to Cardiovascular Risk and Outcomes

## محل انتشار:

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## خلاصه مقاله:

**Background:** Oxidized low-density lipoprotein (ox-LDL) has an important role in the genesis of coronary atherosclerosis. Lectin-like ox-LDL receptor 1 (OLR<sub>1</sub>) contributes to the uptake and internalization of ox-LDL. Genetic polymorphisms have been associated with coronary artery disease (CAD). Here we explore the association of plasma levels of ox-LDL and 3' UTR OLR<sub>1</sub> (rs1050286) SNP with CAD risk and inhospital adverse outcomes. **Methods:** A case-control study enrolled 192 patients with ST-segment elevation myocardial infarction (STEMI), 100 patients with unstable angina, and 100 healthy controls. Baseline, clinical characteristics, and risk scores of the patients were determined. Plasma ox-LDL and other biochemical variables were measured. All subjects are genotyped for OLR<sub>1</sub> (rs1050286) by RT-PCR with TaqMan SNP genotyping assay. **Results:** Plasma ox-LDL was higher with enhanced sensitivity and specificity in identifying patients with STEMI and was found as a significant independent risk factor for CAD in those two groups. Levels of ox-LDL were increased with increasing poor prognostic factors in STEMI patients that are associated with an increased incidence of some adverse events and in-hospital mortality. Elevated STEMI risk was associated with T allele of OLR<sub>1</sub> (rs1050286) (odds ratio of ۴.۹, ۹۵% CI: ۲.۶-۹.۴, p < ۰.۰۰۱). STEMI patients who have T allele exhibited higher risk scores, coronary multivessel narrowing, and elevated incidence of in-hospital major adverse clinical events. **Conclusions:** These results suggest that plasma ox-LDL, as well as T allele of OLR<sub>1</sub> (rs1050286), is associated with the increased risk for developing STEMI and the associated adverse clinical outcomes

