

Oral Communication Abstract

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Associations between urinary 11-dehydrothromboxane B2 and laboratory parameters in obese and non-obese aspirin-treated patients with cardiorenal syndrome

Kseniya Lukyanets, Ivan Pchelin

Department of Faculty Therapy, Saint Petersburg State University, Russia

Objectives: Measurement of urinary levels of 11-dehydrothromboxane B2 is one of the methods for assessing high platelet reactivity in aspirin-treated patients. There are many potential causes for increased platelet reactivity including overweight. The aim of the present study was to evaluate the associations between urinary 11-dehydrothromboxane B2 and laboratory parameters in obese and non-obese aspirin-treated patients with cardiorenal syndrome.

Methods: 82 patients with stable coronary artery disease and CKD (stages 2-4) were included in the study. Urinary levels of 11-dehydrothromboxane B2 (a stable metabolite of thromboxane A2) were assessed using ELISA kit by Enzo Life Sciences (Switzerland). Spearman's correlation coefficient (r_s) was used for statistical analysis.

Results: The median of urinary concentrations of 11-dehydrothromboxane B2 in non-obese patients was 488.36 [360.84 – 672.14] pg/mg creatinine. The median of urinary levels of 11-dehydrothromboxane B2 in obese patients amounted to 488.94 [364.63 – 602.50] pg/mg creatinine. The analysis showed a positive correlation between the urinary level of 11-dehydrothromboxane and fasting plasma glucose level in the subgroup of patients with obesity ($r_s=0.397$, $p=0.009$). There were significant correlations between urinary 11-dehydrothromboxane B2 and left ventricular mass index ($r_s=0.714$, $p=0.047$), end-systolic size of the left ventricle ($r_s=0.648$, $p=0.019$) and erythrocyte sedimentation rate ($r_s=0.347$, $p=0.019$) in subgroups of non-obese patients. In subgroups of patients with obesity, we identified statistically significant correlations between urinary 11-dehydrothromboxane B2 and platelet count ($r_s=-0.714$, $p=0.047$) and platelet indices (platelet crit ($r_s=0.498$, $p=0.008$) and mean platelet volume ($r_s=0.833$, $p=0.010$)).

Conclusions: Our study revealed the relationships between the urinary concentration of 11-dehydrothromboxane B2 and laboratory parameters in cardiorenal syndrome. Hence, the obtained results reflect the varying response to aspirin in subgroups of non-obese and obese patients. Further studies should contribute to the development of the personalization of antiplatelet therapy