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## **TCT@ACC-i2: The Interventional Learning Pathway**

## IMPACT OF DAILY GLUCOSE FLUCTUATION ON CORONARY PLAQUE VULNERABILITY IN PATIENTS PRETREATED WITH LIPID LOWERING THERAPY

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**Background:** There has been growing evidence that the glucose fluctuation is an important contributing factor to the development of coronary artery disease (CAD) as residual risk beyond dyslipidemia.

**Objective:** This study sought to investigate the impact of daily glucose fluctuation on coronary plaque character in CAD patients pretreated with lipid lowering therapy.

**Methods:** In this prospective study, 60 consecutive CAD patients referred to percutaneous coronary intervention whose low density lipoprotein (LDL) cholesterol level < 120 mg/dl with statin or < 100 mg/dl without statin were enrolled. In addition to culprit lesions, non-culprit angiographically mild-to-moderate narrowing lesions were evaluated with virtual histology intravascular ultrasound (VH-IVUS), and the volume percentage of necrotic core within the plaque (%NC) and the presence of thin-cap fibroatheroma (TCFA) were evaluated. Daily glucose fluctuation was analyzed by continuous glucose monitoring system (CGMS), and the standard deviation of the 24-hour blood glucose level (glycemic SD) was calculated. The impact of glycemic SD as well as other coronary risk factors on %NC and the presence of TCFA were assessed.

**Results:** Among the study subjects, LDL cholesterol was  $87.0 \pm 18.2 \text{ mg/dl}$  and Hemoglobin A1c was  $6.4 \pm 1.0 \text{ \%}$ . Oral glucose tolerance test revealed the proportion of diabetes and impaired glucose tolerance were 56.7% and 28.3%, respectively, and only 15% showed normal glucose tolerance. A total of 143 plaques were evaluated in 60 patients: 88 plaques in 34 diabetic patients, and plaques of culprit lesion were 32.2%. %NC by VH-IVUS was well correlated with the glycemic SD by CGMS (r=0.511, P<0.001). A multiple linear regression analysis showed that the glycemic SD had the most strong effect on the %NC (standardized coefficient  $\beta$ =0.479, P <0.001). Moreover, after multiple logistic regression analysis, the glycemic SD remained the only independent risk factor of the presence of TCFA (Odds Ratio 1.083; 95% Cl 1.025-1.145, P=0.005).

**Conclusion:** The daily glucose fluctuation may have an impact on coronary plaque vulnerability in CAD patients pretreated with lipid lowering therapy.