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**IN VIVO VIRTUAL HISTOLOGY INTRAVASCULAR  
ULTRASOUND COMPARISON OF NEOINTIMAL  
HYPERPLASIA WITHIN DRUG-ELUTING-VERSUS BARE  
METAL STENTS IN PATIENTS WITH STEMI**

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**Objectives** The process of in-stent neointimal hyperplasia (NIH) between drug-eluting stents (DES) and bare metal stents (BMS) might be different. We compared in vivo composition of in-stent NIH between DES and BMS using virtual histology-intravascular ultrasound (VH-IVUS) in Patients with ST segment elevation myocardial infarction (STEMI).

**Methods** From May 2009 to Dec 2011, 63 patients were prospectively included in the protocol. Volumetric VH-IVUS was used to compare in-stent NIH between 45 DES and 33 BMS in 63 patients who underwent PCI because of STEMI. The inner and outer VH-IVUS contours were drawn in a way to avoid the stent strut artefacts. Cross-sectional analysis was done at every VH-IVUS frame within the stent, thereby allowing volumetric measurement of stent, lumen, and NIH and its components.

**Results** Baseline characteristics and IVUS measurements were similar between DES and BMS groups. The duration of follow-up was similar between DES (median 9 months (IQR, 3–15)) vs BMS (median 10 months (IQR, 4–15)), ( $p=0.32$ ). %necrotic core (NC) volume was significantly higher in DES than BMS: 19.5 (16.3, 25.6) vs 12.1 (8.2, 18.5) ( $p=0.006$ ), %NC volume significantly increased with time in BMS ( $p=0.007$ ), but not in DES ( $p=0.24$ ) so that at any given time point, %NC in DES was greater than in BMS. After adjustment for baseline differences, only DES ( $p=0.003$ ) and stent age ( $p=0.043$ ) were independent predictors of %NC volume. VH-IVUS in-stent thin-cap fibroatheromas were detected only in the DES group: 34.8% vs 1.5%,  $p=0.013$ .

**Conclusions** in vivo composition of in-stent NIH between DES and BMS was different, suggesting that the process of in-stent NIH in DES and BMS is diverse in patients with STEMI