

# POLYMER-FREE YUKON<sup>DES</sup>, 2-YEAR FOLLOW-UP

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## **Study Title:**

Catch-up in anti-restenotic efficacy in the drug-eluting stent era.  
Results of systematic angiographic follow-up at 6-8 months and  
2 years after coronary stenting

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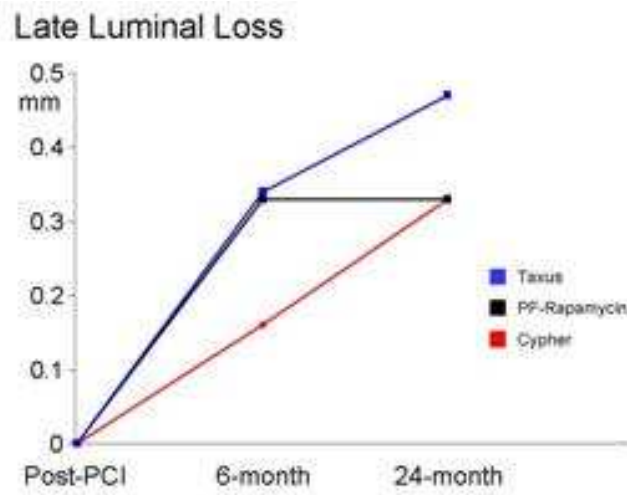
## **Abstract:**

**Background:** Drug-eluting stent (DES) therapy is associated with low levels of late luminal loss (LLL) at 6-8 months. Whether these findings endure over a more prolonged period is unknown. The objective of this study was to assess changes in the anti-restenotic efficacy of DES by restudying subjects at 2 time points post coronary stenting, namely 6-8 months and 2 years. Furthermore, we sought to assess relative differences in time courses of luminal loss between 3 different stent platforms in clinical use at our institution.

**Methods:** Systematic 6-8-month and 2-year angiographic follow-up post index stenting procedure was planned for patients undergoing coronary stenting with Cypher, polymer-free rapamycin Yukon DES (PF rapa) or Taxus stents. Eighty percent had valid 6-8 month data. Of these, patients not requiring re-intervention were scheduled for further follow-up at 2 years. The primary endpoint was interval in-stent LLL between 6-8 months and 2 years.

**Results:** A total of 2341 patients were enrolled: 1036 patients received Cypher, 565 Yukon DES (PF rapa) and 740 Taxus stents. Overall mean LLL at 6-8-month angiographic follow-up was  $0.37\pm 0.56$  mm [ $0.25\pm 0.50$  mm in the Cypher group,  $0.46\pm 0.57$  mm in the PF rapa stent group and  $0.46\pm 0.59$  mm in those receiving a Taxus stent ( $p < 0.001$ )]. At 2 year re-angiography, data was available for 1580 patients (82.0% of eligible patients). Overall mean interval LLL was  $0.12\pm 0.49$  mm. Interval LLL in the Cypher, Yukon DES (PF rapa) and Taxus groups was  **$0.17\pm 0.50$  mm**,  **$0.01\pm 0.42$  mm** and  **$0.13\pm 0.50$  mm** respectively ( $p < 0.001$ ).

**Conclusion:** Ongoing erosion of luminal calibre beyond 6-8-months post index procedure may be observed following DES therapy. There appears to exist a device-specificity in this late reduction in anti-restenotic efficacy in favour of a platform devoid of permanent polymer.



**Late luminal loss of the polymer-based Cypher and Taxus stents in comparison with the polymer-free Translumina Yukon DES stent, 2%Rapamycin-coating (PF-Rapamycin).**