ENDEAVOR IV

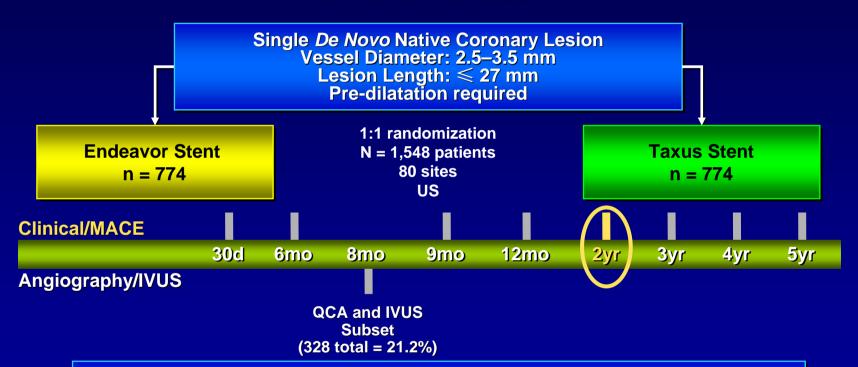
Two-Year Follow-up from a Prospective Randomized Trial Comparing a Zotarolimus-Eluting Stent and a Paclitaxel-Eluting Stent in Patients with Coronary Artery Disease

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ENDEAVOR IV – 2yr FU

Clinical Trial Design PI: Martin B. Leon



Primary Endpoint: TVF at 9 months

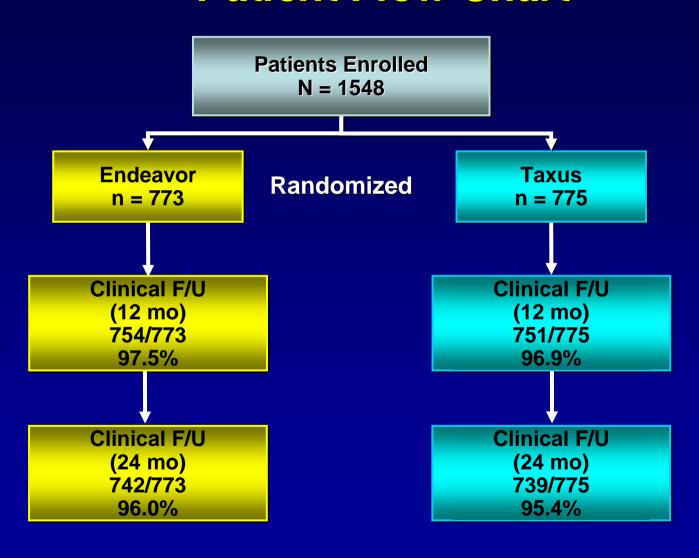
Secondary Endpoints: In-segment % DS at 8 months; TLR and TVR at 9 months
Drug Therapy: ASA and Clopidogrel/Ticlid >6 months
Zotarolimus Dose: 10 µg per mm stent length

ENDEAVOR IV – thru 1yr FU Summary Findings

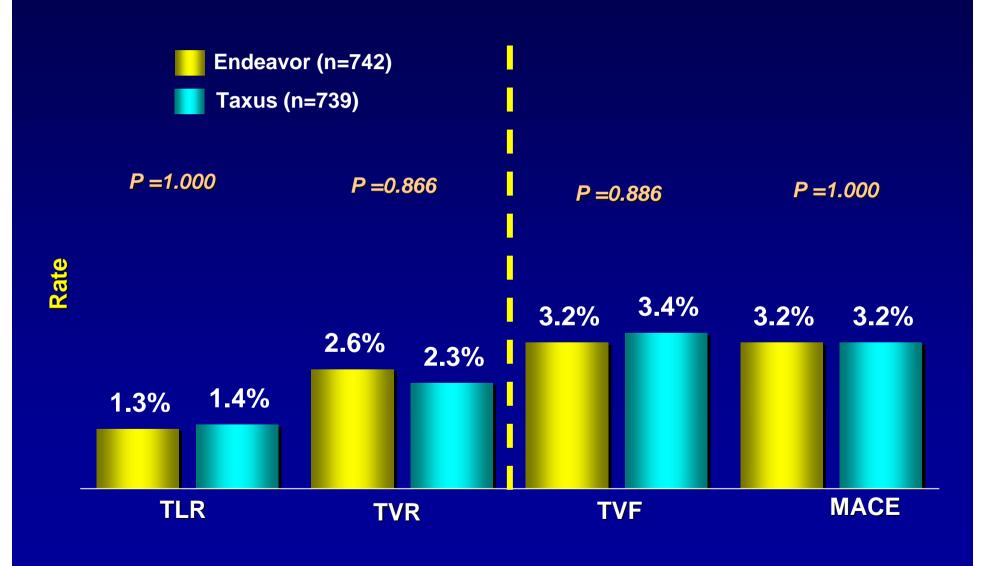
The Endeavor DES, compared with the Taxus DES, demonstrated...

- Angiographic results
 - Higher angio late loss at 8 mos FU
- Clinical outcomes
 - Reduced peri-procedural non-Q MIs, and a similar overall safety profile (death, Q-MI, and stent thrombosis) thru 12 mos FU
 - Similar TVF (1^{ry} endpoint) thru 12 mos FU
 - Similar TVR/TLR in subsets of interest (including diabetics) thru 12 mos FU, especially in the patient cohort with only clinical FU

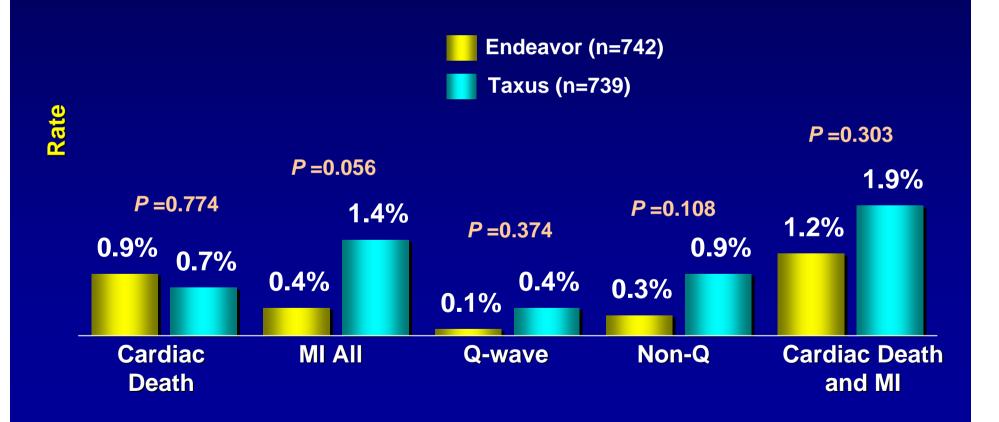
ENDEAVOR IV – 2yr FU Patient Flow Chart



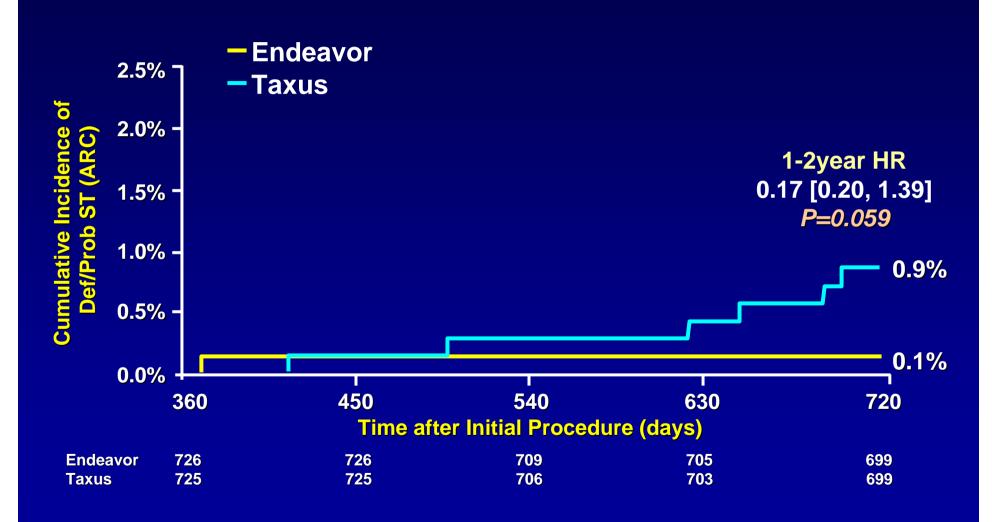
ENDEAVOR IV – 2yr FU Efficacy Endpoints 12-24 months



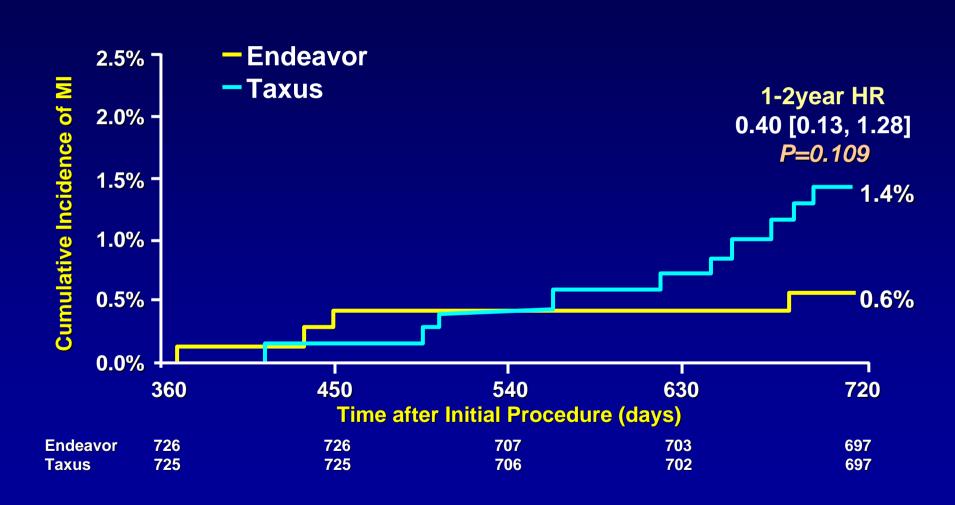
ENDEAVOR IV – 2yr FU Safety Endpoints 12-24 months



ENDEAVOR IV – 2yr FU ARC Def/Prob ST 12-24 mos (VLST)



ENDEAVOR IV – 2yr FU Myocardial Infarction 12-24 mos



ENDEAVOR IV – 2yr FU Conclusions

Between 12 and 24 mos FU, the Endeavor DES, compared with the Taxus DES, demonstrated...

Efficacy

 Continued similar TLR and TVR, even in high restenosis risk subgroups (e.g. diabetics) and especially in the patients with only clinical FU

Safety

- Reduced ARC def/prob very late stent thrombosis (P=0.059; 1 vs. 6 events), even in the setting of frequent DAPT use
- Reduced all MIs (P=0.023), linked to the decrease in very late stent thrombosis events
- Safety findings consistent with the larger pooled analysis of Endeavor studies