



Favorable Long Term Outcome Following Drug Eluting Stent Implantation in Non Bifurcation Lesions Involving Unprotected Left Main Coronary Artery: a Multicenter European and Asian Registry

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Disclosures



The authors have nothing to disclose regarding this presentation



Backgrounds I



· The presence of a lumen narrowing at the ostium and/or the body of an unprotected left main coronary artery (LMCA), which does not require bifurcation treatment, is a class I indication to surgical revascularization despite the fact that does not appear technically very demanding and can treated with a single stent implantation



Backgrounds II



 Recently, a multicenter registry have reported favorable outcome following DES implantation in in the ostium and/or body of unprotected LMCA stenosis



Aim of the Study



 The aim of the study was to assess the safety and long-term efficacy of drug-eluting stent (DES) implantation in the ostium and/or body of unprotected LMCA stenosis



Methods



 All consecutive patients (pts) who had a stenosis in the ostium and/or the mid-shaft of an unprotected LMCA (not requiring the treatment of the bifurcation) electively treated, from April 2002 to May 2005, with PCI and SES (Cypher, Cordis, Johnson and Johnson Company, Warren, NJ) or PES (Taxus, Boston Scientific, Natick, MA) in 7 Centres were analysed.



Baseline Clinical Characteristics I



	(n = 208)		
Age, years	64.0 ±11.9		
Female gender, n (%)	74(35.6)		
Hypertension, n (%)	124 (59.6)		
Hyperchol, n (%)	108 (51.9)		
Smoking, n (%)	70 (33.6)		
Diabetes Mellitus , n (%)	43 (20.7)		



Baseline Clinical Characteristics II

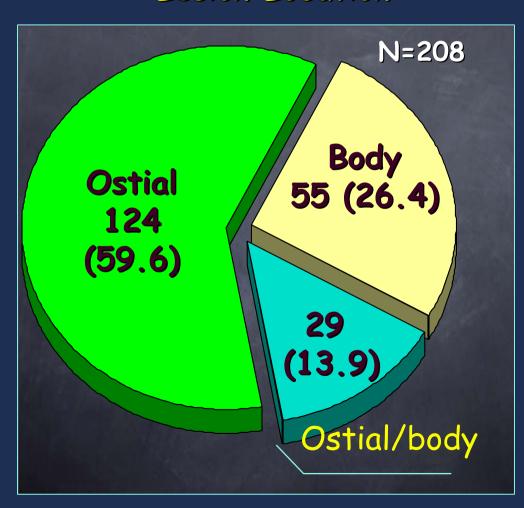


	(n = 208)		
Unstable Angina, n (%)	84 (40.4)		
LVEF, %	56.9 ±11.7		
Parsonnet	5.5 (2.5-13.8)		
Euroscore	4 (2-7)		
Euroscore >6 and/or Parsonnet>13 , n (%)			
RCA disease, n (%)	70 (33.6)		
RCA concomitant treatment	22		

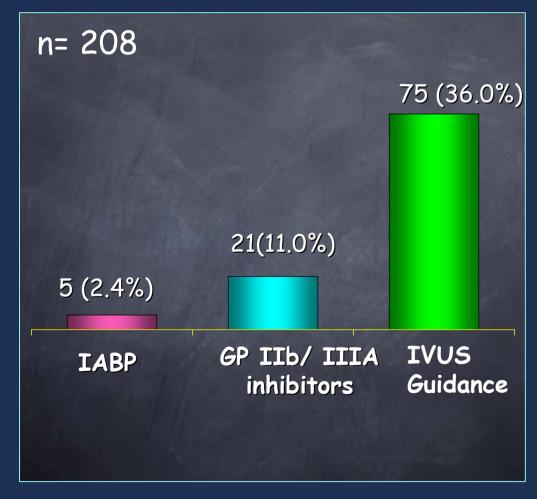




Lesion Location



Procedural Characteristics

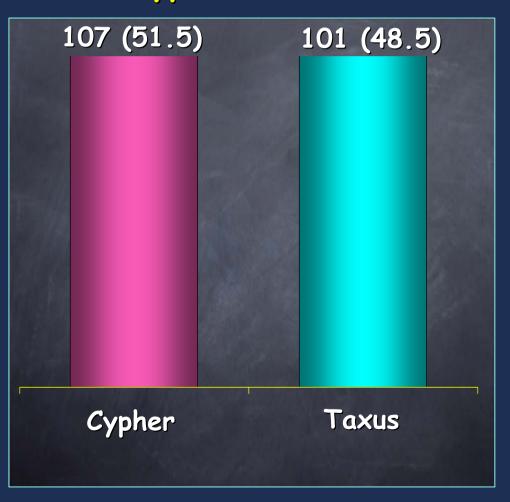






n= 208

Type of Stent



Predilatation/Direct Stenting









	(n = 208)		
Stent Length	14.5±7.5		
Max pressure, atm	17.0± 4.3		
Number vessel treated	1.33±0.9		
Number lesions treated	1.77±1.5		





QCA Measurements

	(n = 208)		
Reference Vessel Diam ,mm	3.6±0.5		
Pre MLD, mm	1.5±0.5		
Lesion Length, mm	7.9±4.5		
Final reference diameter, mm	3.7±0.5		
Final MLD, mm	3.3±0.7		



In-Hospital Outcome



	n = 208		
Q wave MI, n (%)	0		
Non Q wave MI*	6 (2.9)		
TLR or TVR, n (%)**	2 (1.0)		
Death, n (%)	O		
Acute Thrombosis, n (%)	0		
MACE, n (%)	7 (3.3)		

^{*}defined as 3 times CK-MB ULN; **1 because of the presence of more than 30% residual stenosis after stenting and 1 for procedural complication





6-Month Angiographic Follow-Up

Angiographic Fup was performed in 146 (70.2%) of the pts

	n=146		
Restenosis, %	4 (1.9)		
Late Loss, mm	-0.01		





873±297 days Clinical Follow-Up

	n = 208
Death, n (%)	10 (4.8)
Cardiac Death	6 (2.9)
TLR, n (%)	4 (1.9)
TVR, n (%)*	18 (8.6)
MI, n (%)	2 (0.9)
MACE, n (%)	24 (11.5)

*16 re-PCIs and 2 CABG



Stent Thrombosis



ARC Definitions

	n = 208		
Definite Stent Thrombosis	0		
Probable Stent Thrombosis	0		
Possible Stent thrombosis	5 (2.4%)		



Five Patients Died of Unexplained Cause



	P† 1	Pt2	Pt3	Pt4	Pt5
Age, ys	76	41	75	84	51
LVEF, %	30	25	40	35	40
Parsonnet	34	19	18	26	1
Euroscore	16	6	7	10	2
Time of death, ds	208	606	1149	1360	480





Conclusions

- In this multicenter registry evaluating PCI with DES implantation in non-bifurcation unprotected LMCA stenosis this procedure confirmed to be safe and effective.
- · At 6 month angiographic follow-up the restenosis rate is 1.9%.
- The benefit is maintained at long-term clinical follow-up with a low MACE rate.