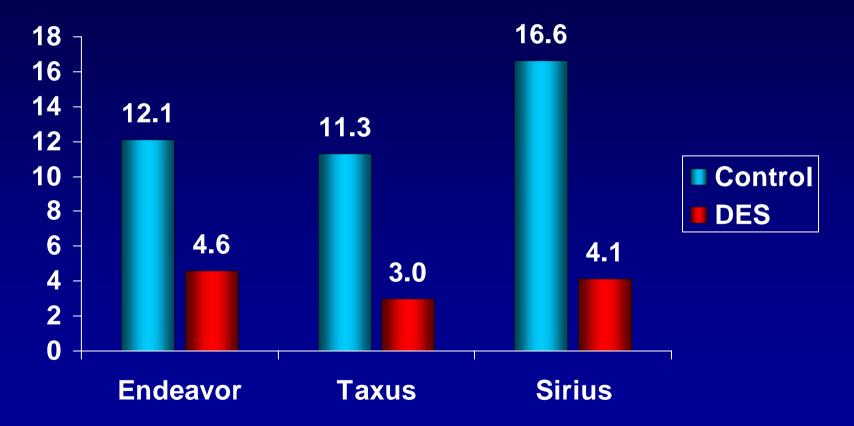
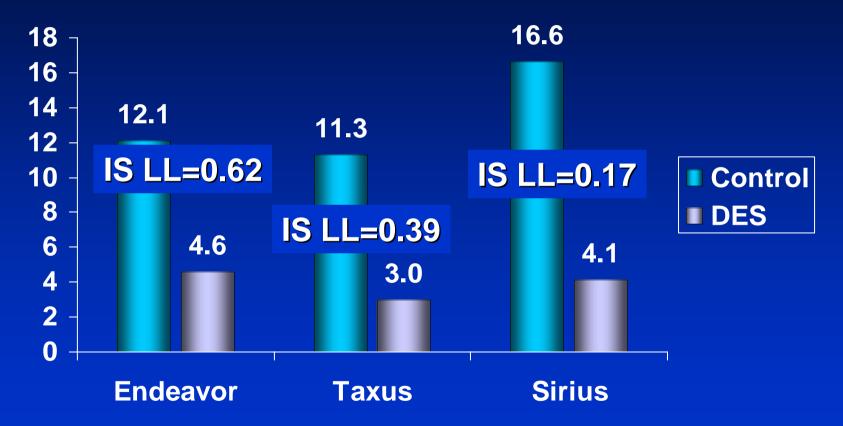
Drug-eluting Stents, Late Loss, and Restenosis DES to Prevents MIs

Richard Kuntz Brigham and Women's Hospital Harvard Medical School

Pivotal Trial Comparisons TLR to 9 Months

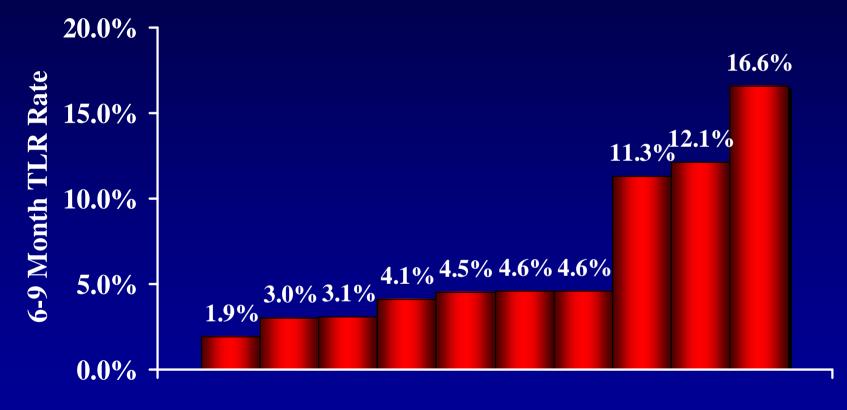


Pivotal DES Trial Comparisons *TLR to 9 Months*



Risk and Restenosis

Some Contemporary Clinical Restenosis Rates

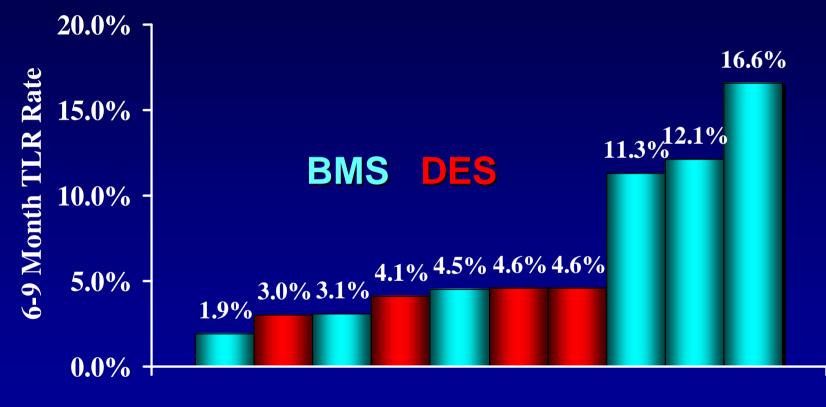


Recent BMS and DES Trials

Mauri L, Kuntz R submitted for publication

Risk and Restenosis

Some Contemporary Clinical Restenosis Rates

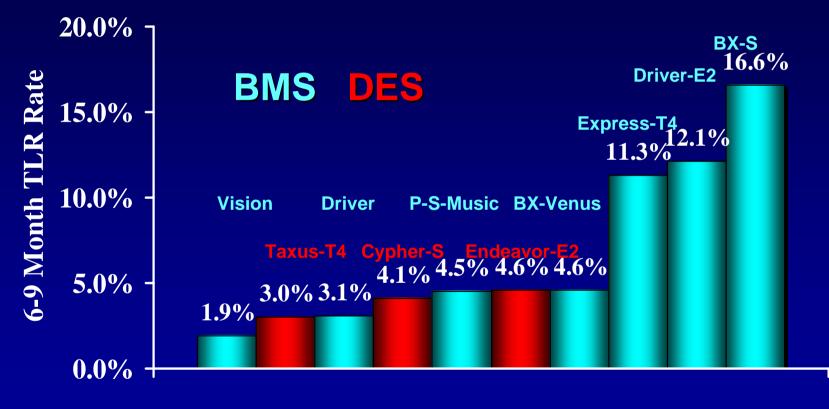


Recent BMS and DES Trials

Mauri L, Kuntz R submitted for publication

Risk and Restenosis

Some Contemporary Clinical Restenosis Rates



Recent BMS and DES Trials

Mauri L, Kuntz R submitted for publication

Restenosis Endponts

Target Lesion Revascularization

- Best endpoint in a randomized Trial
- Needs large sample size for stable Estimation
- High level of influence by case-mix confounders renders it almost meaningless in comparison across trials.
- Late Loss (In-stent version only)
 - Stable and efficient estimate for any stent-type
 - Less influenced by case-mix confounders, and provides a "signature" value for any particular stent.

Restenosis Endponts The Noise Factor

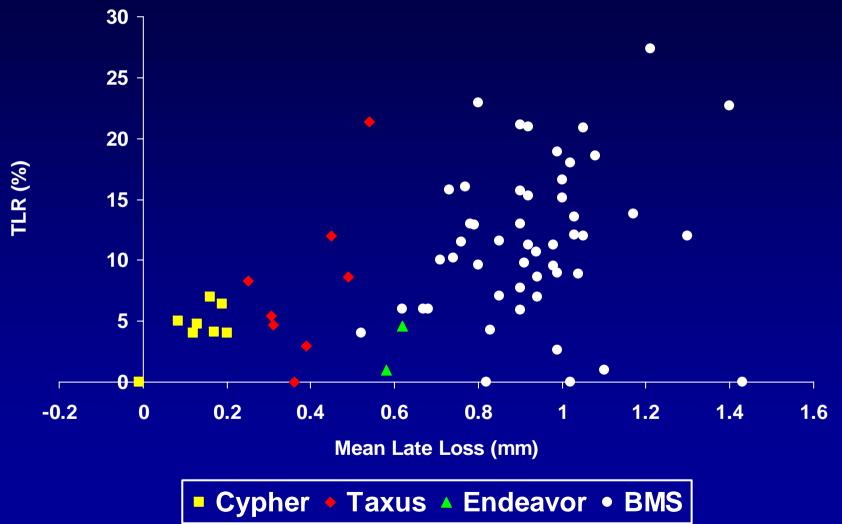
Target Lesion Revascularization

- Affected by
 - Lesion length
 - Diabetes prevelance
 - Reference vessel size
 - Threshold for revascularization (50-70% renarrowing)
- Estimates are wide ranging for BMS and DES

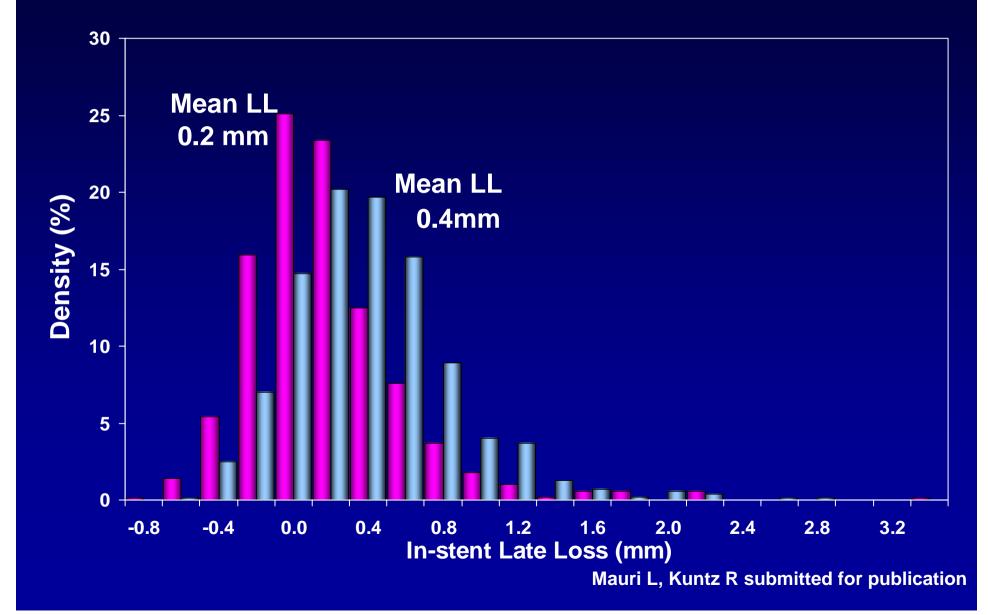
In-Stent Late Loss

- Affected by
 - Diabetes
 - Lesion length
- Relatively more stable across trials

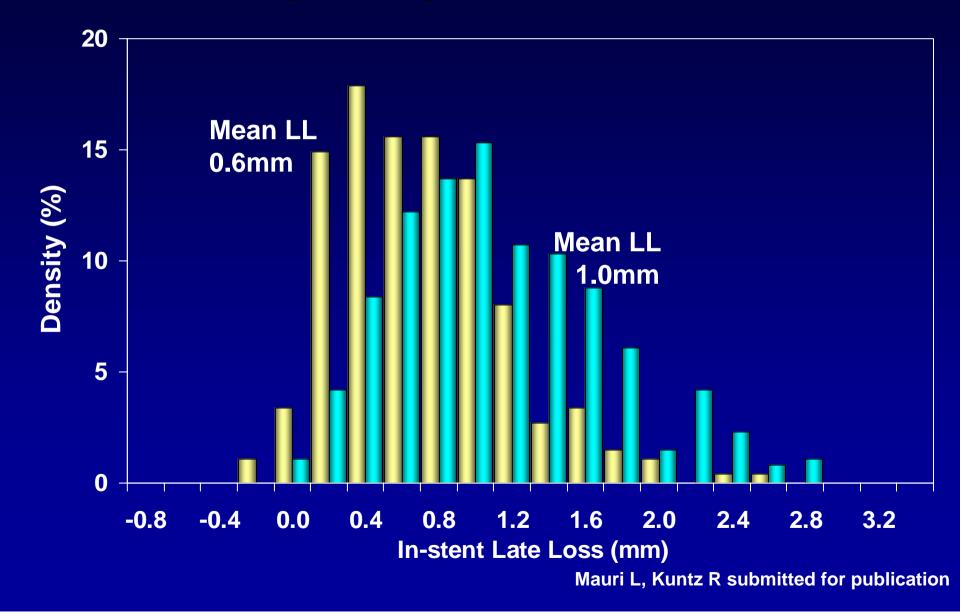
In-Stent Late Loss and TLR Current DES and BMS Results

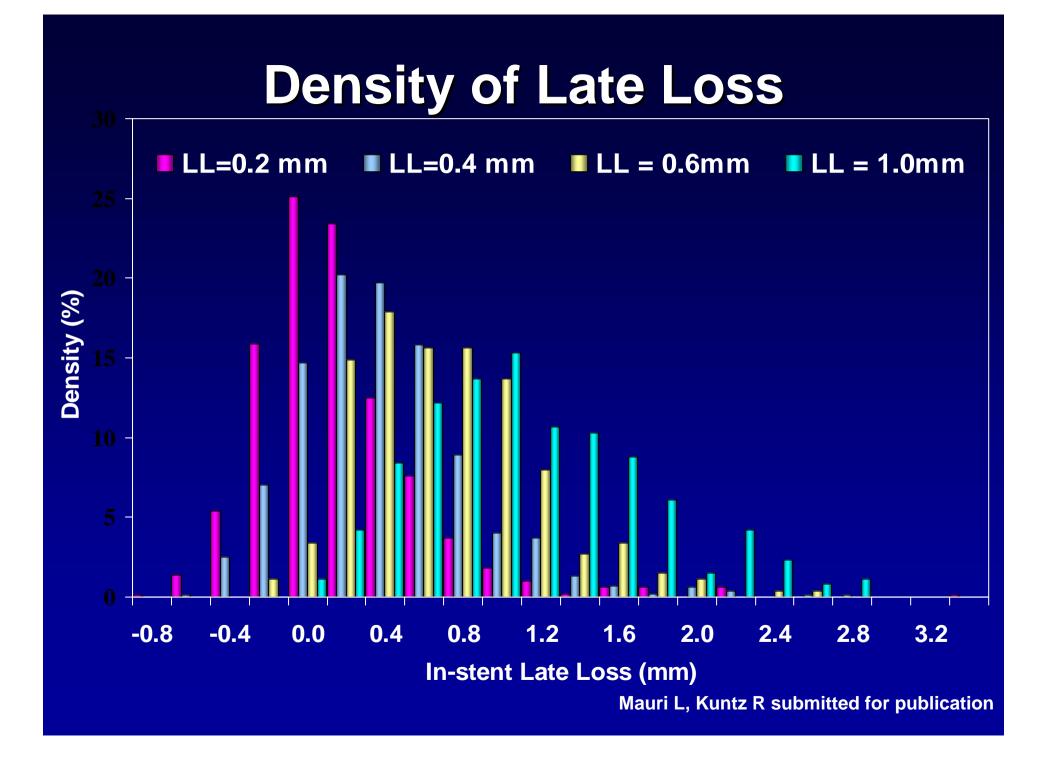


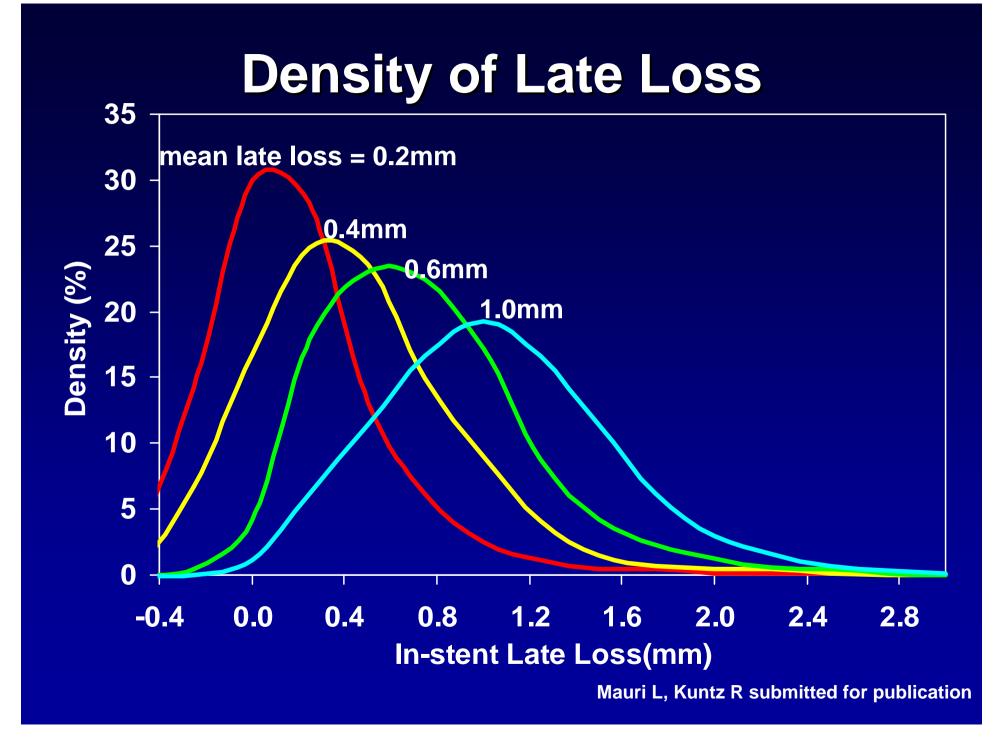
Frequency of Late Loss



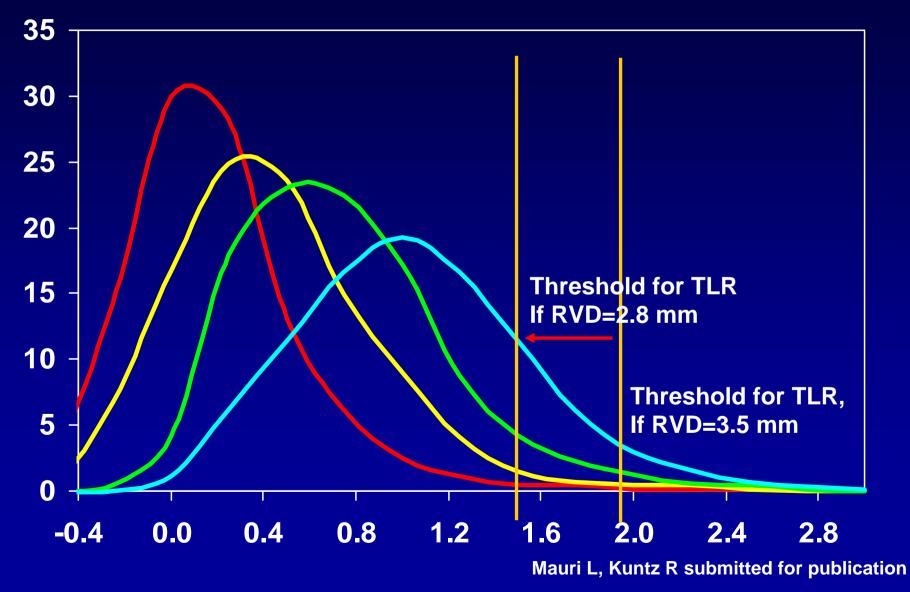
Frequency of Late Loss



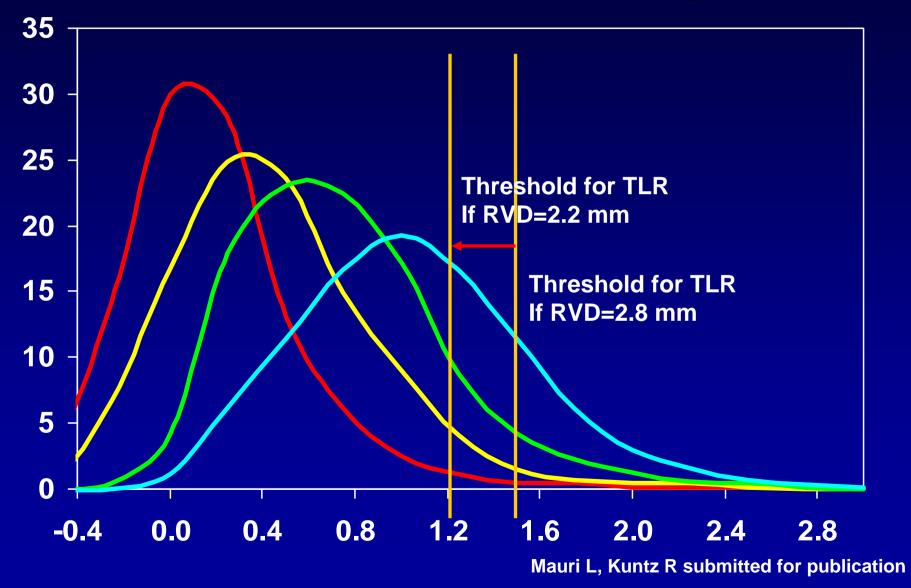




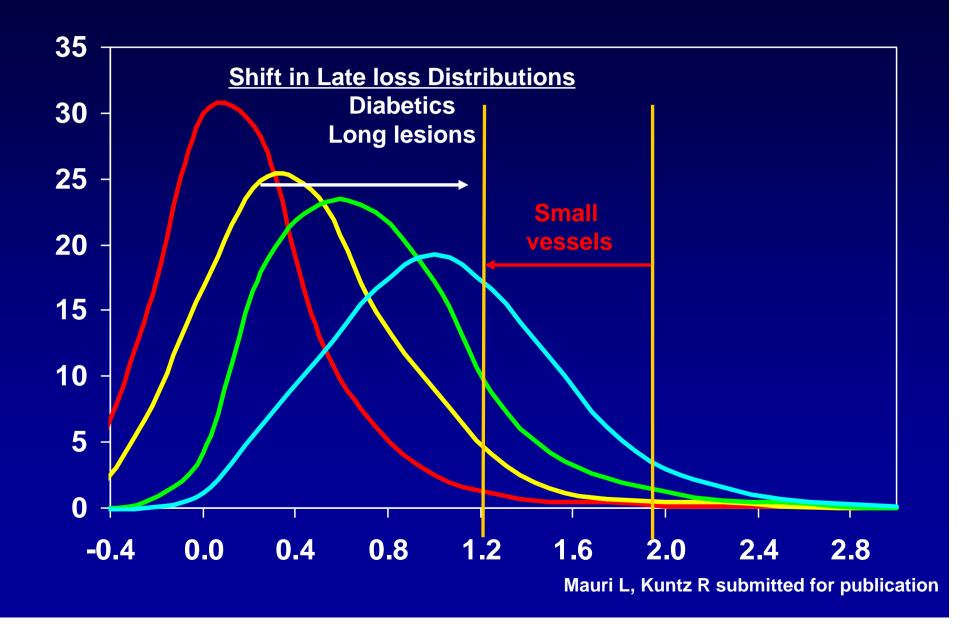
Late Loss and TLR Effect of mean reference vessel diameter



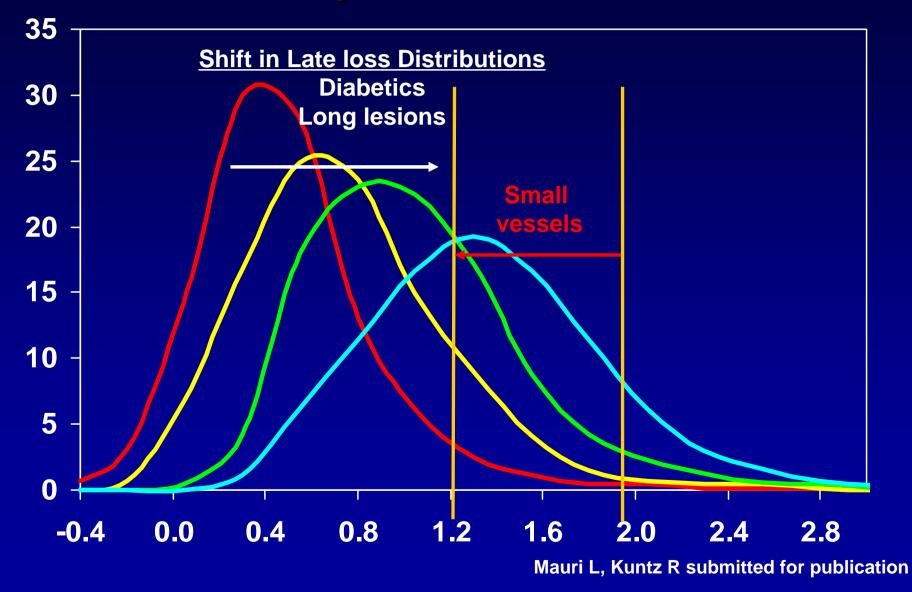
Late Loss and TLR Effect of small vessel stenting



Density of Late Loss



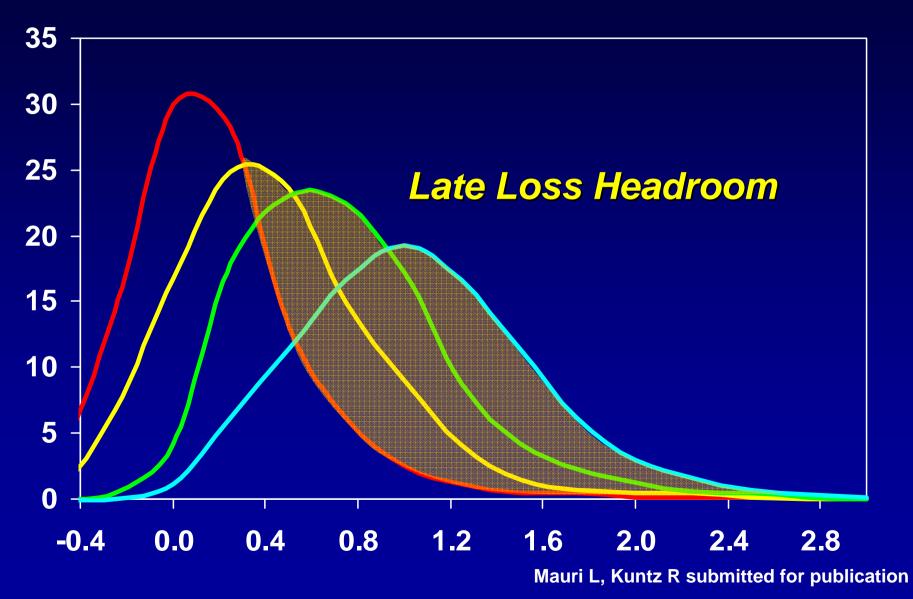
Late Loss and TLR Effect of High Risk Characteristics



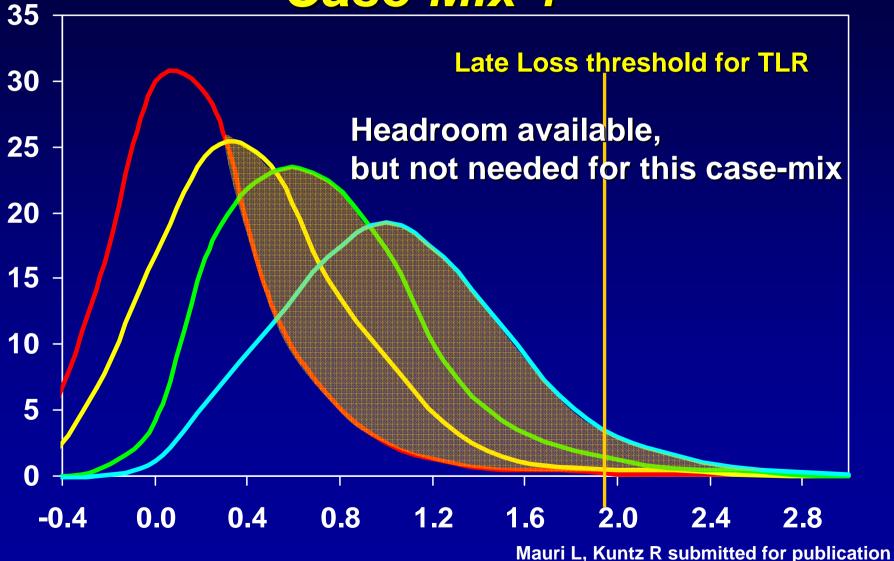
Late Loss Headroom

- Late Loss headroom is the space of extra late loss available for high risk restenosis case-mix cohorts
 - Headroom highest for low in-stent late loss stent systems

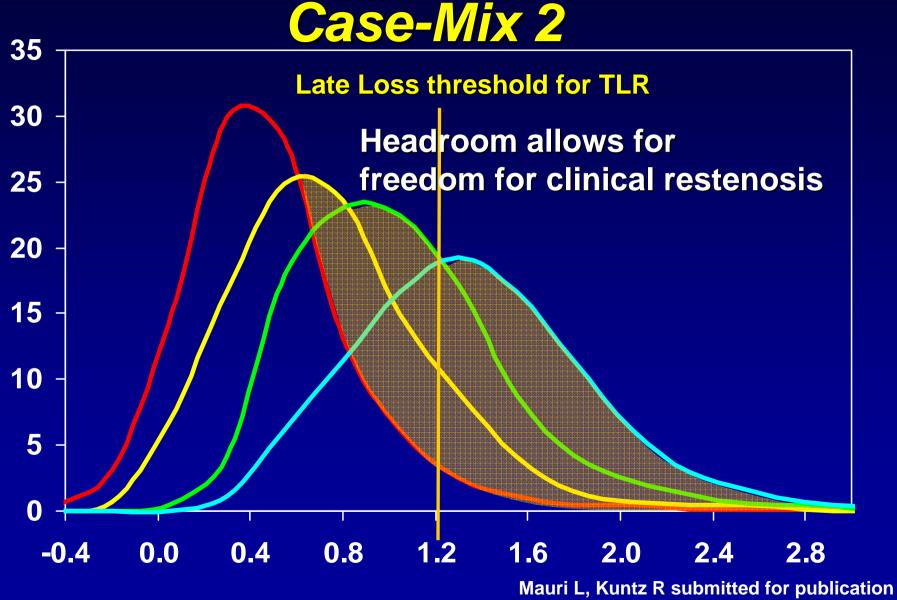
Late Loss Headroom



Late Loss Headroom Case-Mix 1



Late Loss Headroom

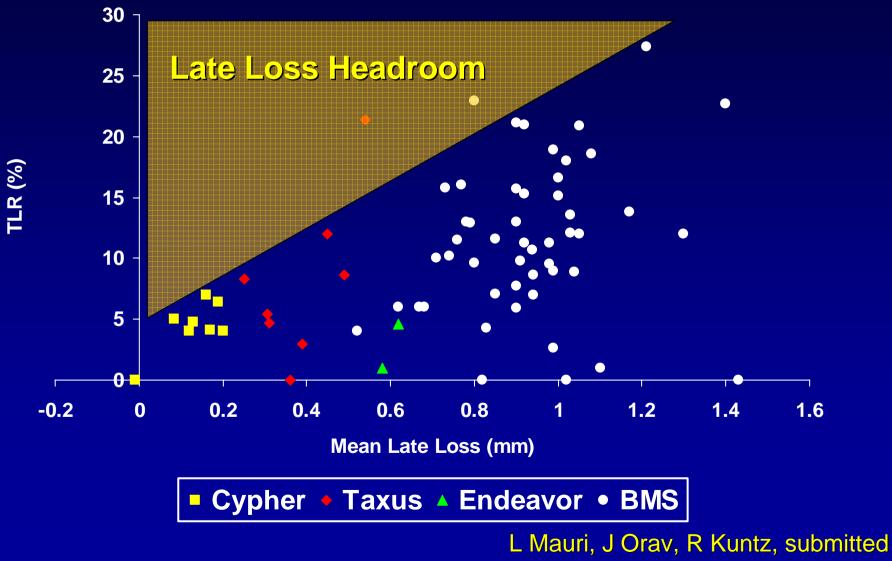


Late Loss Headroom

- Late Loss headroom is the space of extra late loss available for high risk restenosis case-mix cohorts
 - Headroom highest for low in-stent late loss stent systems
- For low Late Loss stent systems, the headroom concept reduces the chance of high TLR over the wide range of case-mix risk

- Evident in real data from clinical trials

In-Stent Late Loss and TLR Late Loss Headroom



Clinical Results to 9 months

	Endeavor N = 582	Control N = 585	P value
Composite MACE (%)	7.4	14.7	<0.0001
Death	1.2	0.5	ns
Q-Wave MI	0.3	0.9	ns
Non Q-Wave MI	2.4	3.1	ns
CABG	0.0	0.0	ns
TLR	4.6	12.1	<0.0001
CABG	0.3	0.5	ns
PCI	4.3	11.6	<0.0001
TVR (%)	5.7	12.8	<0.0001
TVF (%) (Primary endpoint)	8.1	15.4	<0.0005

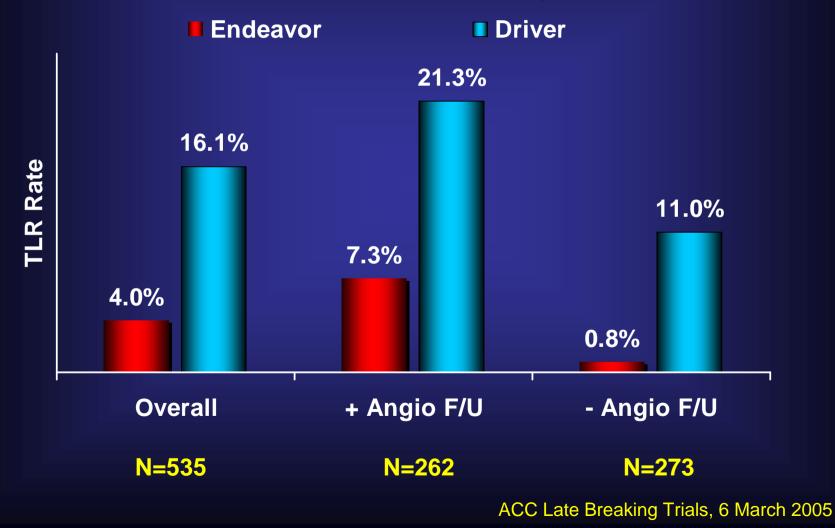
Endeavor II

ACC Late Breaking Trials, 6 March 2005

Endeavor II

TLR by Angiographic Follow-up

LAD Subset Analysis

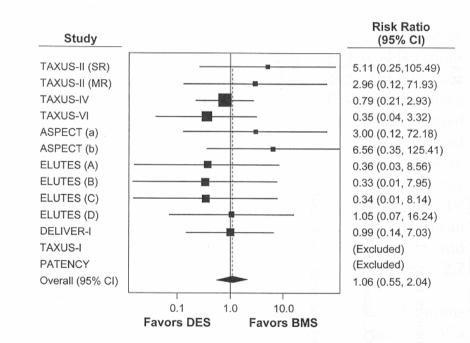


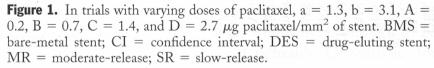
Drug Eluting Stent Late Stent Thrombosis

- EP McFadden, E Stabile, E Regar, et. al. Research Letter, *Lancet* 2004: 364:1519
 - 4 cases of angiographically documented late stent thrombosis, accompanied by acute MI
 - SES (335, 375 days)
 - PES (343, 442 days)
 - All cases occurred soon after clopidogrel cessation

Paclitaxel Stent Thrombosis Bavry et al, J Am Coll Cardiol 2005

 Meta-anlysis of 8 PES/BMS trials





PES, SES, and BMS Thrombosis Moreno et al, J Am Coll Cardiol 2005

Β

10 RCT DES studies of 5030 patients pooled

Study	DES n / N	BMS n/N	OR (95% CI Fixed)	Weight %	OR (95% CI Fixed)
RAVEL	0 / 120	0/118	225 m Fach Stud	0.0	Not estimable
SIRIUS	2 / 533	4 / 525 🔶		27.6	0.49 (0.09,2.69)
E-SIRIUS	2/175	0/177	e suchas	■→ 3.4	5.12 (0.24,107.32
C-SIRIUS	1/50	1/50 🔶	LOAD - NO.	→ 6.7	1.00 (0.06,16.44)
ASPECT	0 / 90	0 / 48	+ 0.483 +	0.0	Not estimable
ELUTES	1 / 153	1/39 🔶		- 10.9	0.25 (0.02,4.09)
TAXUS-I	0/31	0/30	-1.681	0.0	Not estimable
TAXUS-II	3 / 266	0 / 270	272	→ 3.4	7.19 (0.37,139.80
TAXUS-IV	4 / 662	5 / 652		34.4	0.79 (0.21,2.94)
DELIVER	2 / 522	2/519 -	102 - SLI = -1.765	13.7	0.99 (0.14,7.09)
Total (95% CI)	15 / 2602	13 / 2428	bb 224 -m	100.0	1.05 (0.51, 2.15)
۸		-1 -1	2 1	5 10	
A	DES	Favo	rs DES F	avors BMS	OR
A Study	DES n/N	Favo	rs DES F	avors BMS	OR (95% CI Fixed)
		Favo	rs DES F	avors BMS	
Study	n / N	Favo BMS n / N	rs DES F	Favors BMS Weight %	(95% CI Fixed)
Study RAVEL	n / N 0 / 120	Favo BMS n / N 0 / 118	rs DES F	Favors BMS Weight %	(95% CI Fixed) Not estimable
Study RAVEL SIRIUS	n / N 0 / 120 1 / 533	Favo BMS n/N 0/118 3/525 ←	rs DES F	Favors BMS Weight % 0.0 43.1	(95% Cl Fixed) Not estimable 0.39 (0.09,3.15)
Study RAVEL SIRIUS E-SIRIUS	n / N 0 / 120 1 / 533 0 / 175	Favo BMS n/N 0/118 3/525 0/177 ←	rs DES F	Favors BMS Weight % 0.0 43.1 0.0	(95% CI Fixed) Not estimable 0.39 (0.09,3.15) Not estimable 0.33 (0.01,8.22) Not estimable
Study RAVEL SIRIUS E-SIRIUS C-SIRIUS	n / N 0 / 120 1 / 533 0 / 175 0 / 50	Favo BMS n/N 0/118 3/525 0/177 1/50 ◀	rs DES F	avors BMS Weight 0.0 43.1 0.0 21.2 0.0 0.0 0.0	(95% CI Fixed) Not estimable 0.39 (0.09,3.15) Not estimable 0.33 (0.01,8.22) Not estimable Not estimable
Study RAVEL SIRIUS E-SIRIUS C-SIRIUS ASPECT	n / N 0 / 120 1 / 533 0 / 175 0 / 50 0 / 90	Favo BMS n/N 0/118 3/525 0/177 1/50 0/48	rs DES F	avors BMS Weight % 0.0 43.1 0.0 21.2 0.0 0.0 0.0 0.0 0.0	(95% CI Fixed) Not estimable 0.39 (0.09,3.15) Not estimable 0.33 (0.01,8.22) Not estimable
Study RAVEL SIRIUS E-SIRIUS C-SIRIUS ASPECT ELUTES	n/N 0/120 1/533 0/175 0/50 0/90 0/153	Favo BMS n/N 0/118 3/525 0/177 1/50 0/48 0/39	rs DES F	avors BMS Weight 0.0 43.1 0.0 21.2 0.0 0.0 0.0	(95% CI Fixed) Not estimable 0.39 (0.09,3.15) Not estimable 0.33 (0.01,8.22) Not estimable Not estimable
Study RAVEL SIRIUS E-SIRIUS C-SIRIUS ASPECT ELUTES TAXUS-I	n/N 0/120 1/533 0/175 0/50 0/90 0/153 0/31	Favo BMS n/N 0/118 3/525 0/177 1/50 0/48 0/39 0/30	rs DES F	Eavors BMS Weight % 0.0 43.1 0.0 21.2 0.0 0.0 0.0 0.0 0.0 0.0 14.4	(95% CI Fixed) Not estimable 0.39 (0.09,3.15) Not estimable 0.33 (0.01,8.22) Not estimable Not estimable Not estimable 5.11 (0.24,107.02 1.97 (0.18,21.81)
Study RAVEL SIRIUS E-SIRIUS C-SIRIUS ASPECT ELUTES TAXUS-I TAXUS-II	n/N 0/120 1/533 0/175 0/50 0/90 0/153 0/31 2/266	Favo BMS n/N 0/118 3/525 0/177 1/50 0/48 0/39 0/30 0/270	rs DES F	avors BMS Weight % 0.0 43.1 0.0 21.2 0.0 0.0 0.0 0.0 0.0 0.0 7.0	(95% CI Fixed) Not estimable 0.39 (0.09,3.15) Not estimable 0.33 (0.01,8.22) Not estimable Not estimable 5.11 (0.24,107.02
Study RAVEL SIRIUS E-SIRIUS C-SIRIUS ASPECT ELUTES TAXUS-I TAXUS-I TAXUS-II TAXUS-II	n/N 0/120 1/533 0/175 0/50 0/90 0/153 0/31 2/266 2/662	Favo BMS n/N 0/118 3/525 0/177 1/50 0/48 0/39 0/30 0/270 1/652	rs DES F	Eavors BMS Weight % 0.0 43.1 0.0 21.2 0.0 0.0 0.0 0.0 0.0 0.0 14.4	(95% CI Fixed) Not estimable 0.39 (0.09,3.15) Not estimable 0.33 (0.01,8.22) Not estimable Not estimable Not estimable 5.11 (0.24,107.02 1.97 (0.18,21.81)

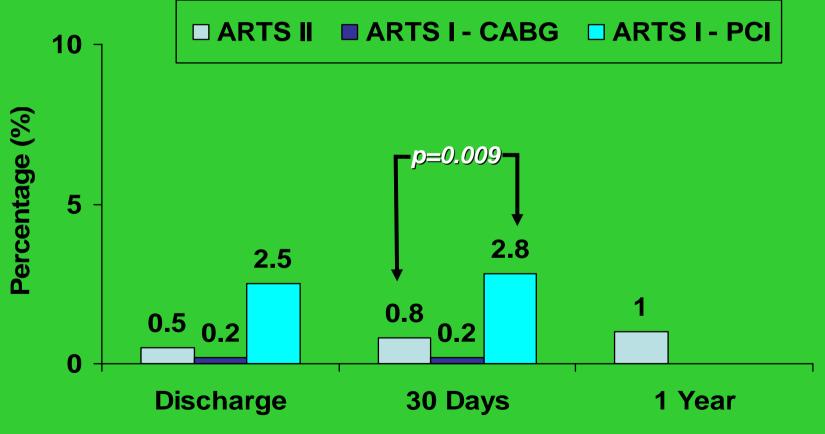
Figure 1. (A) Comparison between the rate of stent thrombosis in patients allocated to drug-eluting stents (DES) or bare-metal stents (BMS) in the randomized studies and in the pooled population. (B) Comparison between the rate of late stent thrombosis in patients allocated to DES or BMS in the randomized studies and in the pooled population. CI = confidence interval; OR = odds ratio.

Favors DES

Favors BMS

ARTS II Angiographic Occlusions

* Definition of thrombotic occlusion: Angiographically proven occlusion (TIMI 0 or 1) or flow limiting thrombus (TIMI 1 or 2)



ARTS II up to 1 year: 5 TLR (1 Q wave MI, 4 with substantial cardiac enzyme release)

Safety Results

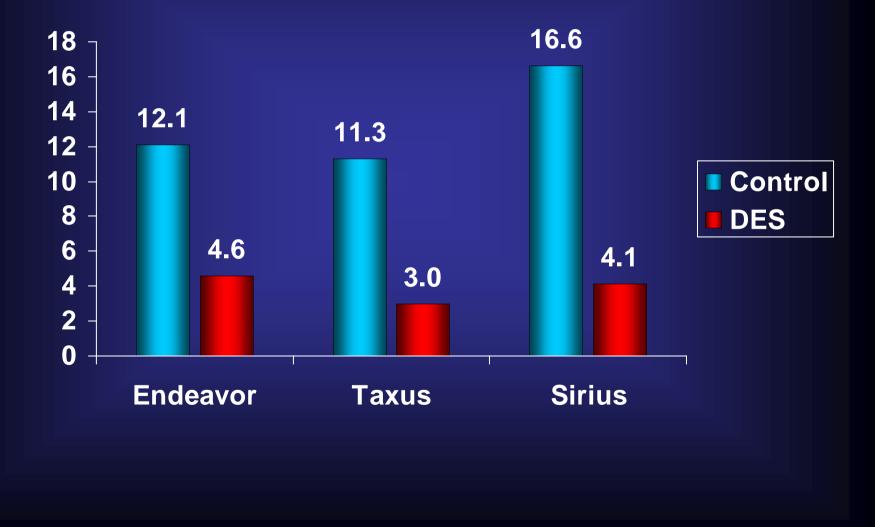
Stent Thrombosis	Endeavor N = 582	Driver N = 585	P value
In-hospital	0.3% (2)	0.3% (2)	
Discharge to 30 days	0.2% (1)	0.9% (5)*	
>30 – 270 days	0	0	
Total at 270 days	0.5% (3)	1.2% (7)	0.34
IVUS Results	Endeavor N = 100	Driver N = 83	P value
Late Acquired Stent Malapposition	0%	0%	ns
Late Aneurysm	0%	0%	ns

Stent thrombosis defined as angiographic thrombus or subacute closure in the stented vessel or any death not attributed to a non-cardiac cause within the 1st 30 days <u>*3/6 post-discharge stent thrombosis cases occurred in Driver arm when Plavix was stopped prematurely</u>

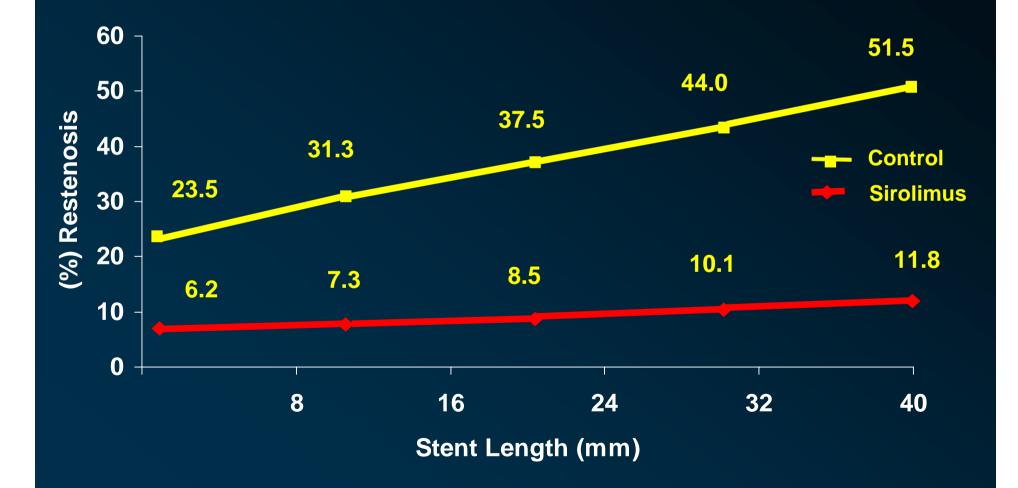
Endeavor II

ACC Late Breaking Trials, 6 March 2005

Pivotal Trial Comparisons TLR to 9 Months



SIRIUS: Restenosis vs. Stent Length In-Segment



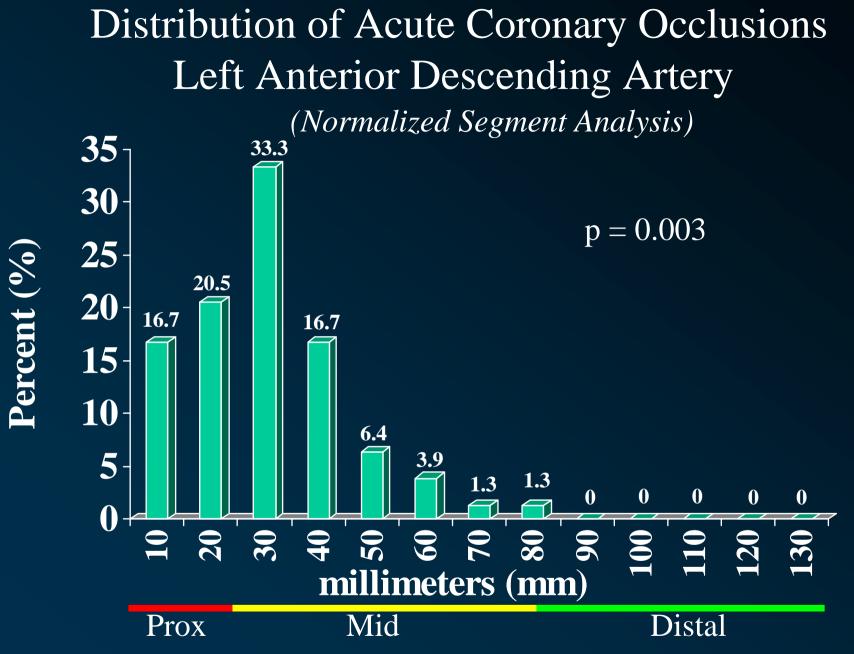
Now That We've Conquered Restenosis Can We Prevent Plaque Rupture?

The Stented Coronary Segement

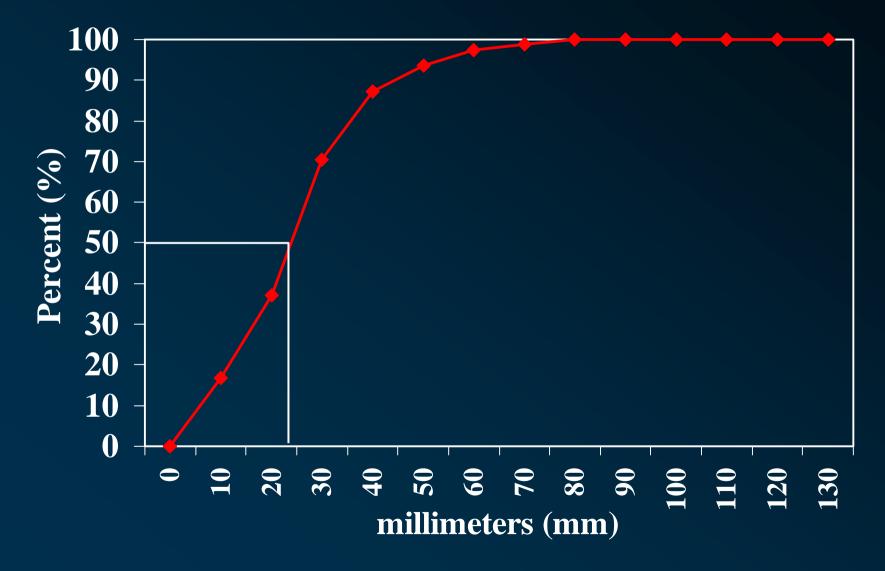
Low incidence of ACS in the segment <0.005 over 4 years

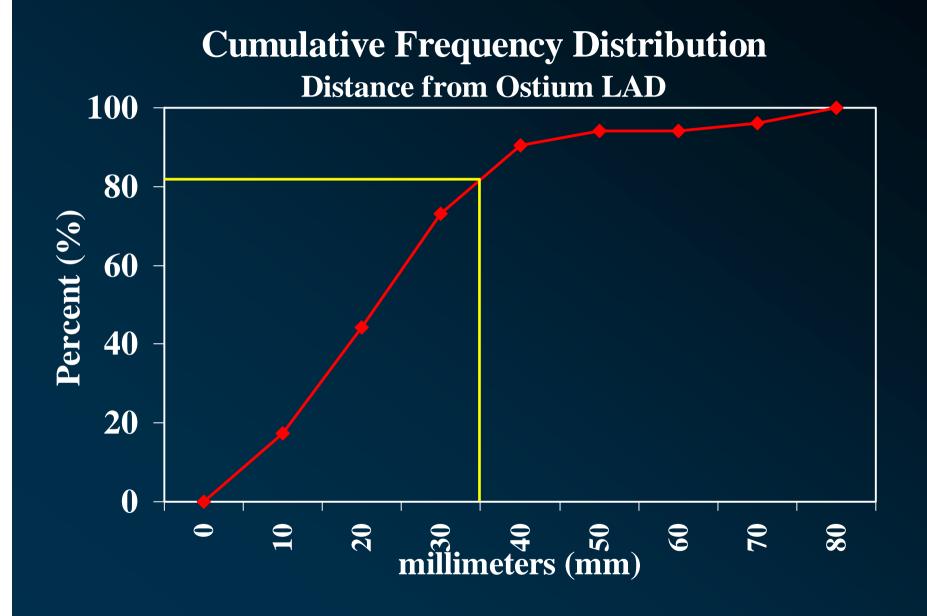
Can't grow atherosclerosis here!





Cumulative Frequency Distribution Curve of Acute Coronary Occlusions by Distance from the Ostium Left Anterior Descending Artery

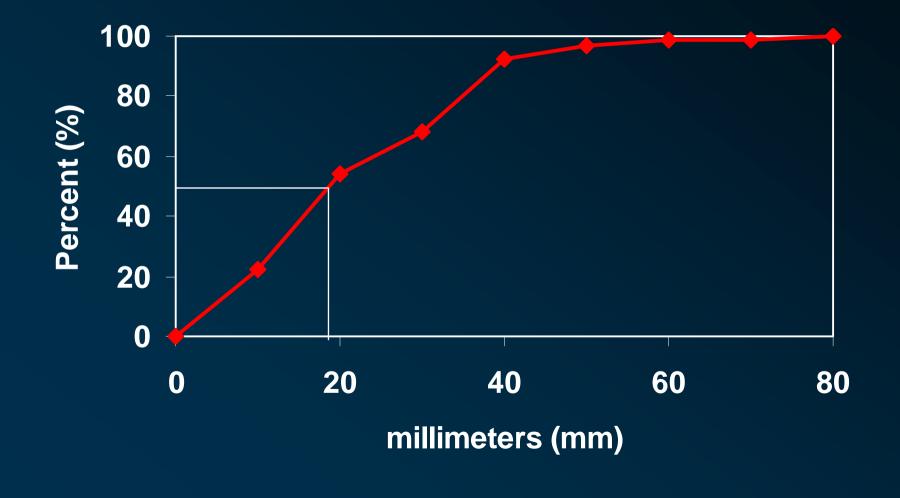


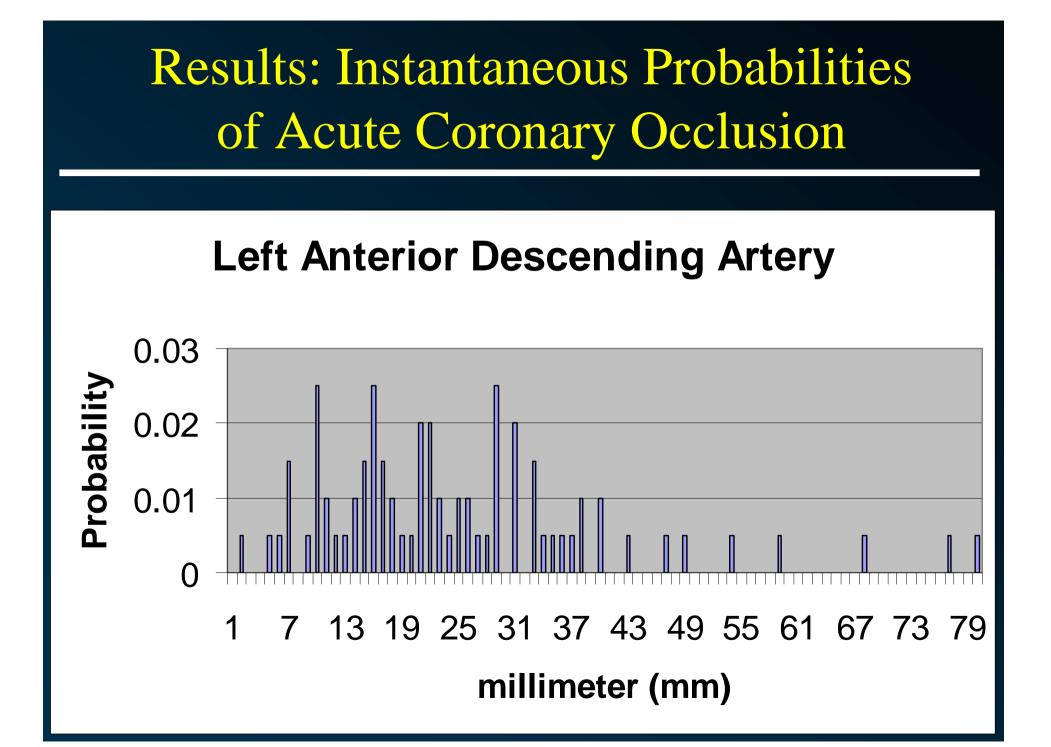


Vulnerable Hot Spots How About A Few DESs



Cumulative Frequency Curve for LAD Non-Q MI

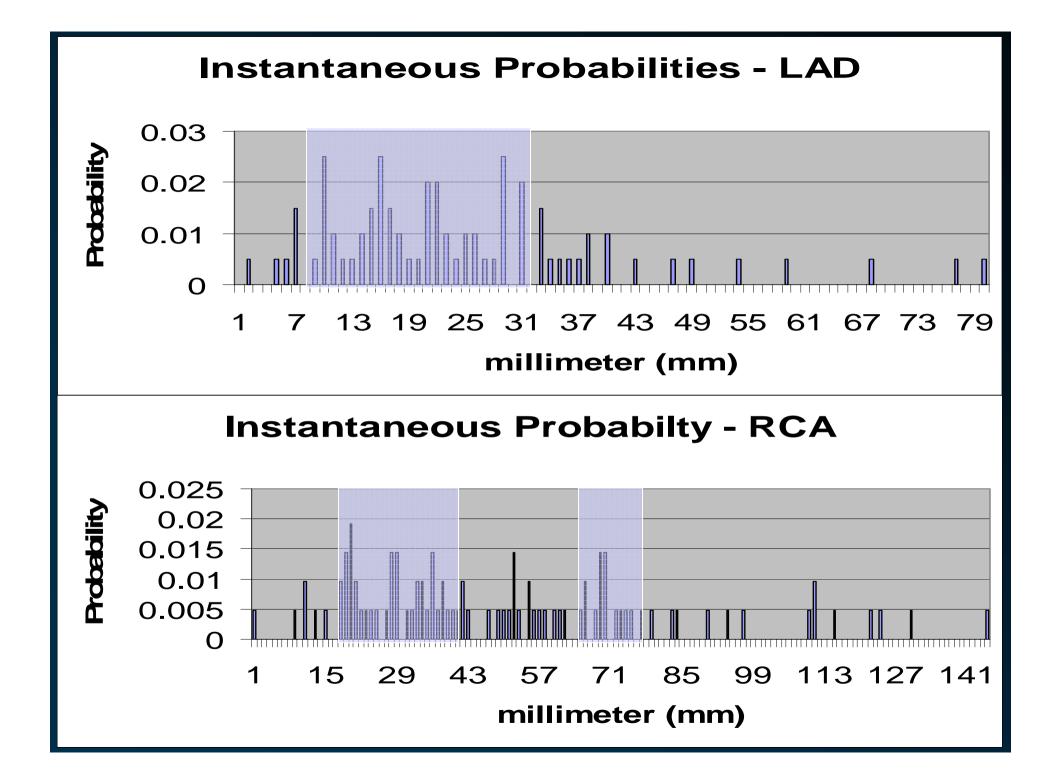


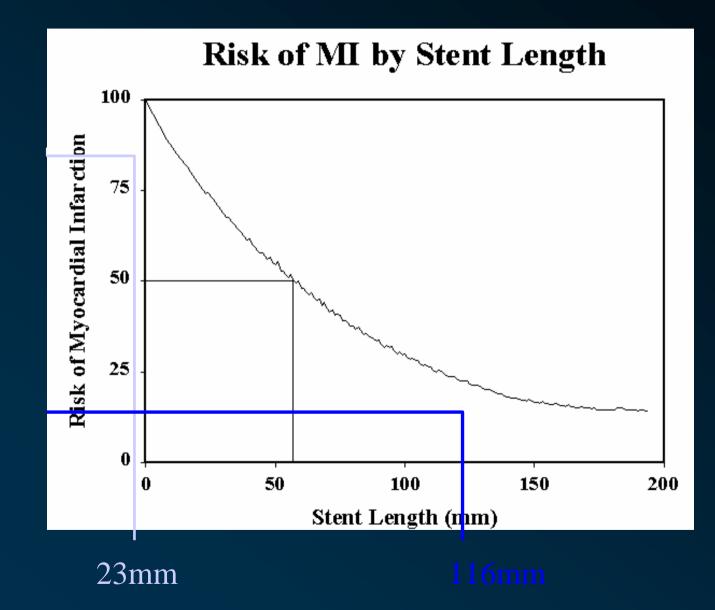


Simulation Optimum Combination

Vessel	Stent Length	Starting Location *
LAD	23mm	9mm
RCA	23mm	18mm
RCA	13mm	65mm

* Absolute distance from ostium





DES, Late Loss, Preventing MIs

- The introduction of DES has been a remarkable advancement for the Interventional Cardiology community and patients who suffer from coronary disease
 - Already three products have demonstrated breakthrough antiresults in the prevention of restenosis
- Late Loss "Head Room" is the extra space available for higher risk lesions to provide freedom from repeat revascularization
 - Choice of stent will incorporate patient risk of restenosis and ease-of-use of the stent

DES, Late Loss, Preventing MIs

- All three DES products appear at lest as safe, in terms of stent thrombosis, as bare metal stents
- The use of stents to prevent MIs may be the first step in prophylactic therapy, but medical therapy must be optimized
 - The use of geographic spatial maps and imaging techniques will guide placement
- It's good to be an Interventional Cardiologist at this time!