

FFR in Left Main Disease

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Why FFR instead of IVUS?

- Physiologic versus anatomic information
- Limitations of IVUS for assessing Left Main (LM)
- Data supporting FFR assessment of LM
- Limitations/Practical Aspects of FFR of LM

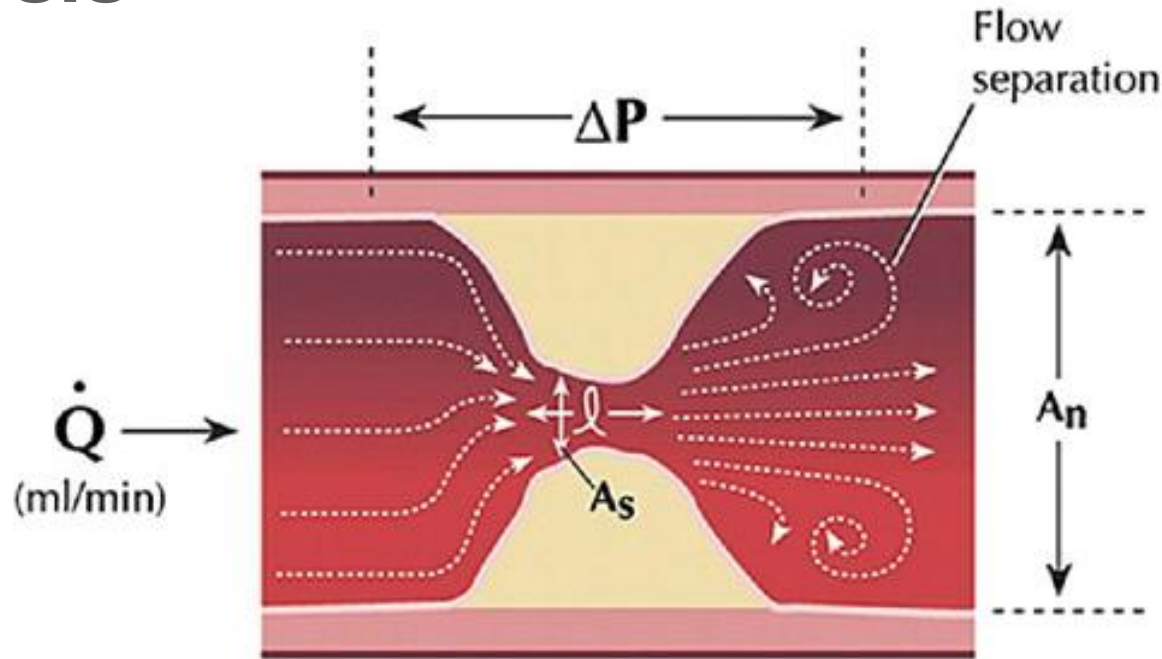


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Factors impacting ischemic potential of a stenosis



$$\Delta P = \underbrace{f_1(1/A_s^2, \ell, \dot{Q})}_{\text{Viscous}} + \underbrace{f_2(1/A_s^2, 1/A_n^2, \dot{Q}^2)}_{\text{Separation}}$$



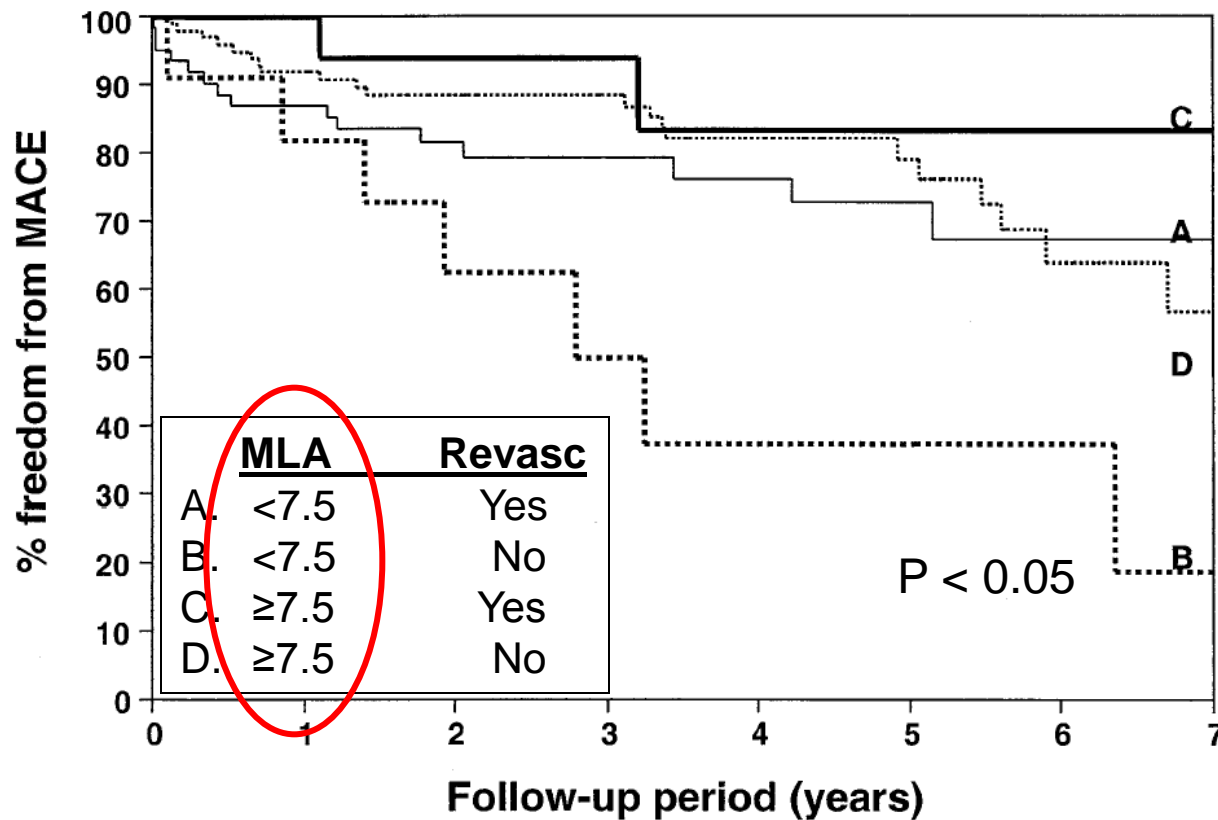
Variability of IVUS Assessment of the LM

- 73 patients with distal left main disease underwent IVUS pullback from the LAD and from the L Cx.
- The average MLA of the LM from the LAD pullback was $6.7 \pm 3.1 \text{ mm}^2$ and from the L Cx pullback was $6.8 \pm 3.3 \text{ mm}^2$
- However, in $\frac{1}{2}$ the patients the L Cx measurement was smaller and in 11% the difference was $> 1 \text{ mm}^2$.
- In the other $\frac{1}{2}$ of the patients the LAD measurement was smaller and in 16% the difference was $> 1 \text{ mm}^2$



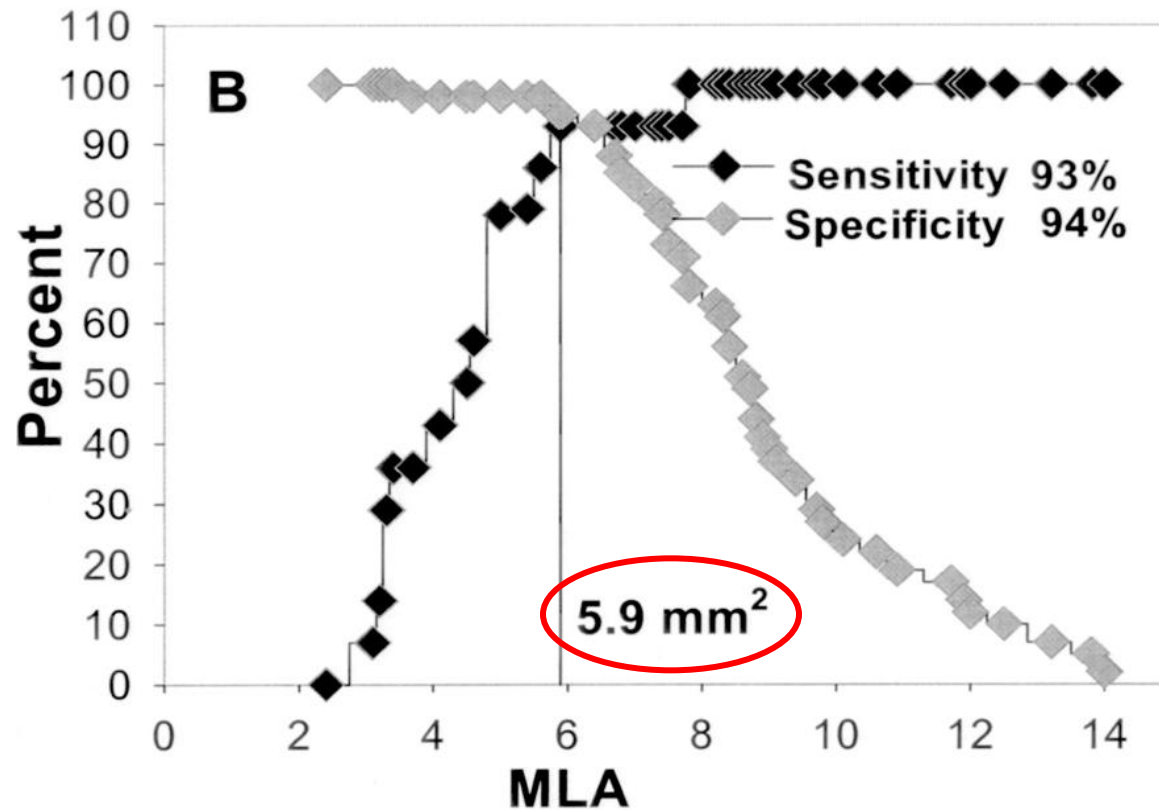
Variability of IVUS Cutoff Values

3 Yr Follow-up in 214 Intermediate Left Mains Assessed by IVUS



Variability of IVUS Cutoff Values

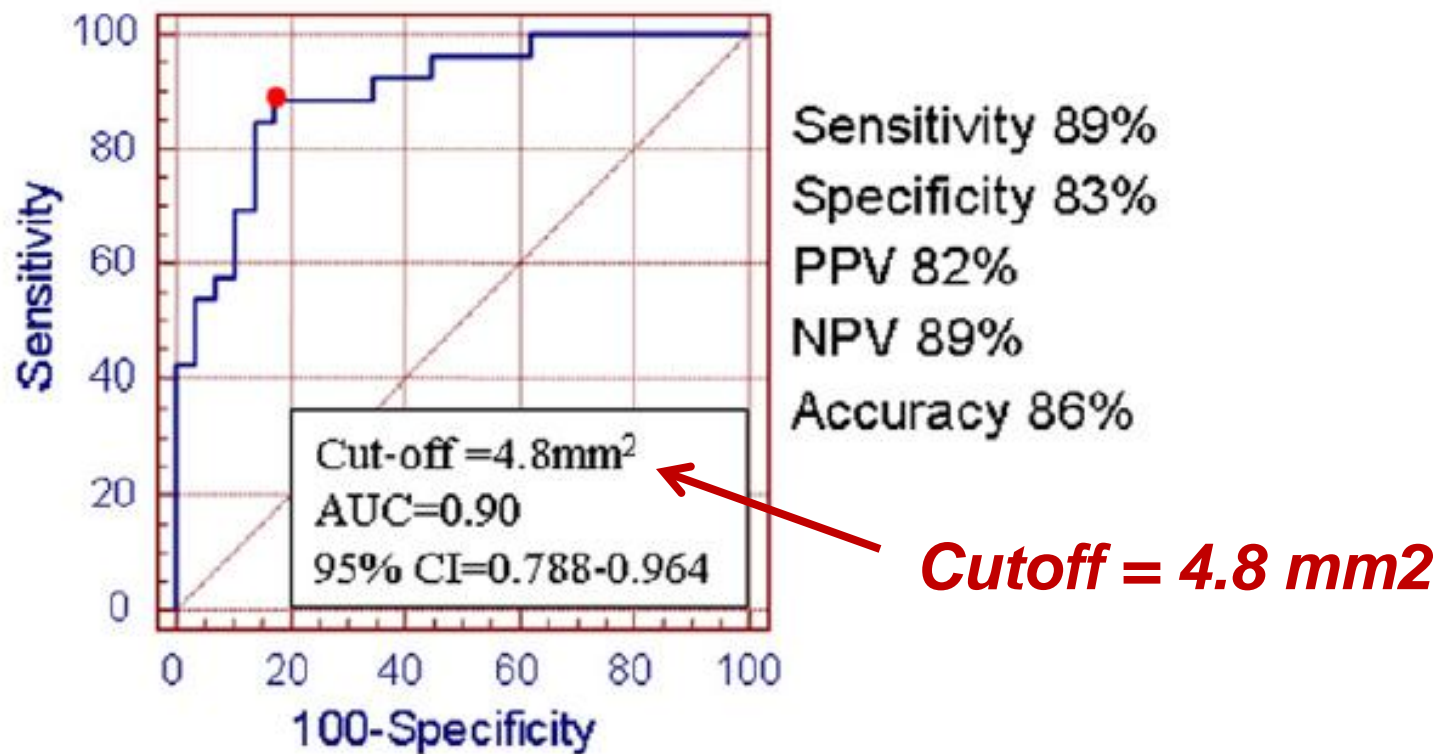
55 patients with ambiguous left main disease



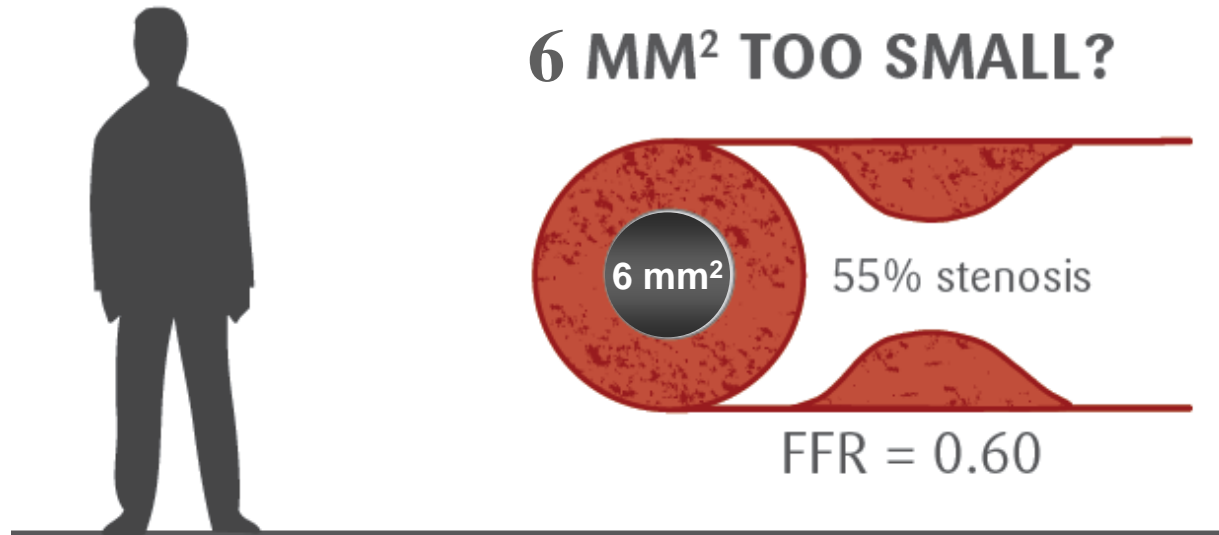
Variability of IVUS Cutoff Values

55 patients with 30-80% LM and FFR and IVUS

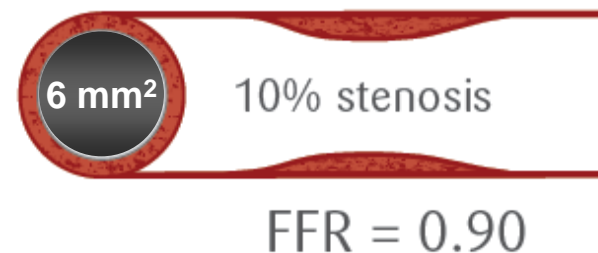
A. MLA predicting FFR<0.80



Variability of IVUS Cutoff Values



6 MM² SUFFICIENT?



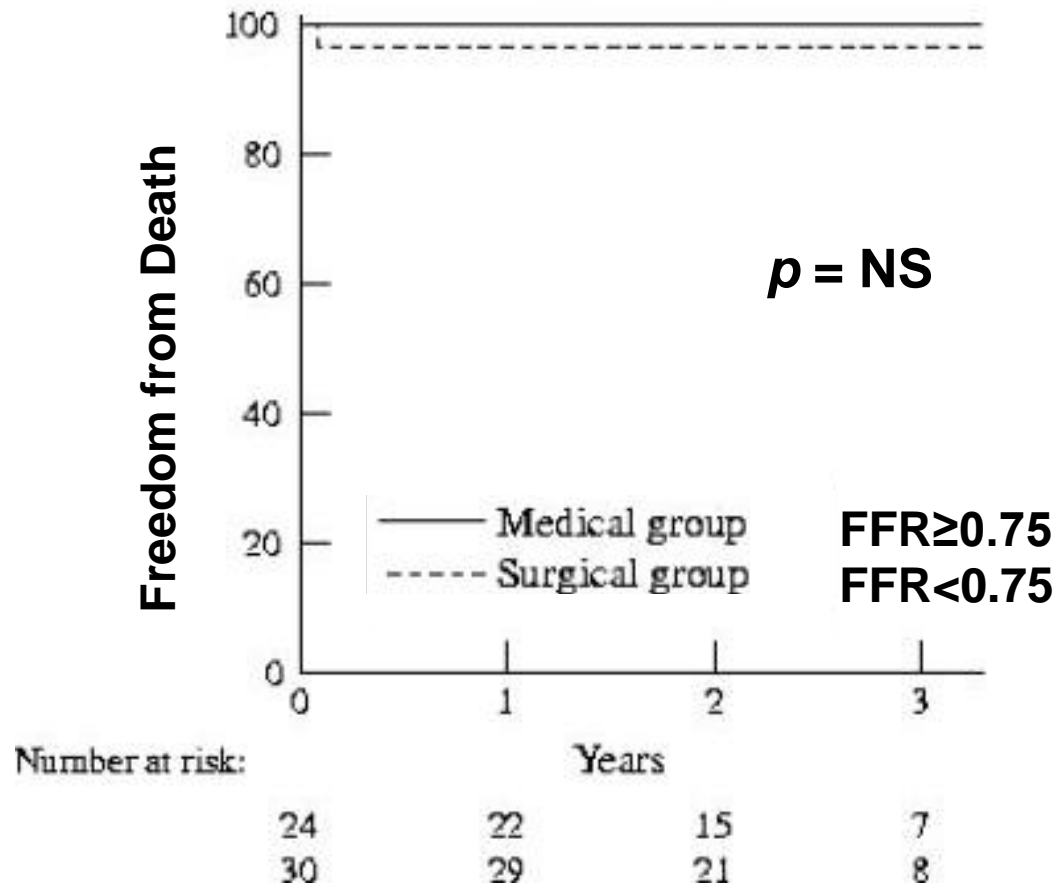
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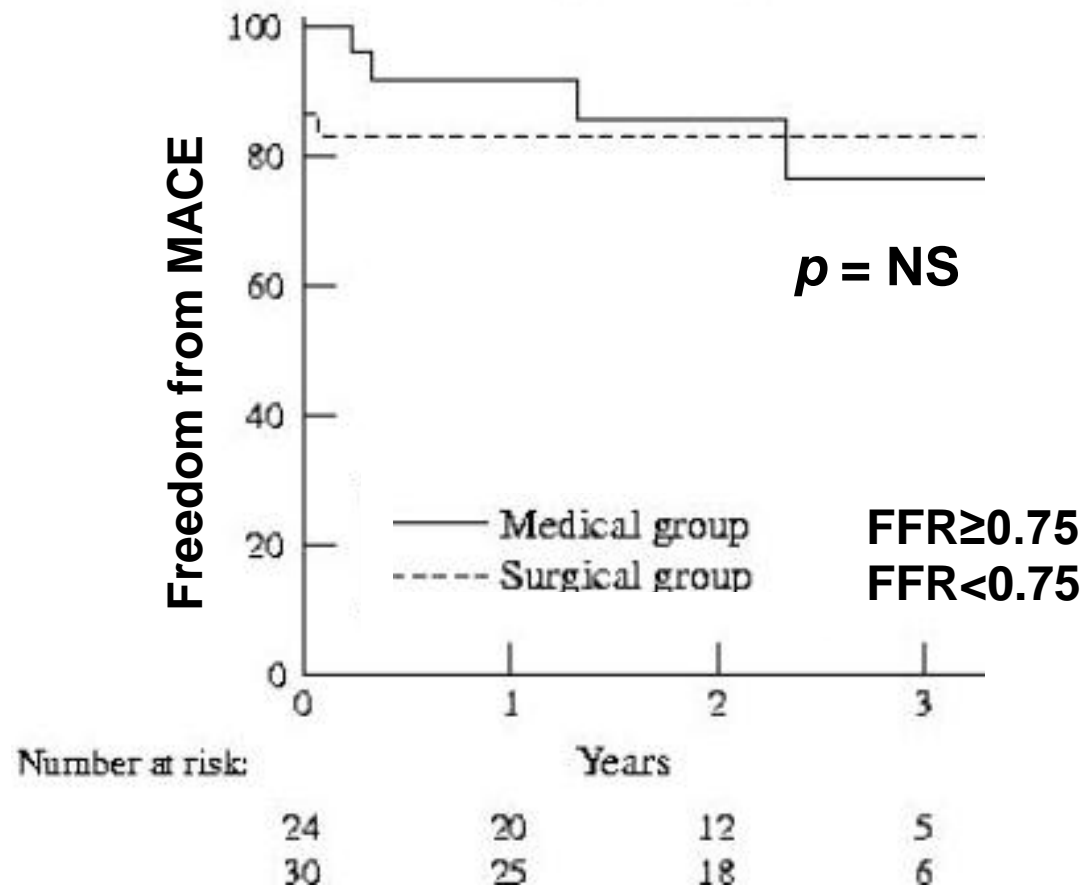
Is it safe to defer LM Rx based on FFR?

FFR measured in 54 patients with equivocal left main



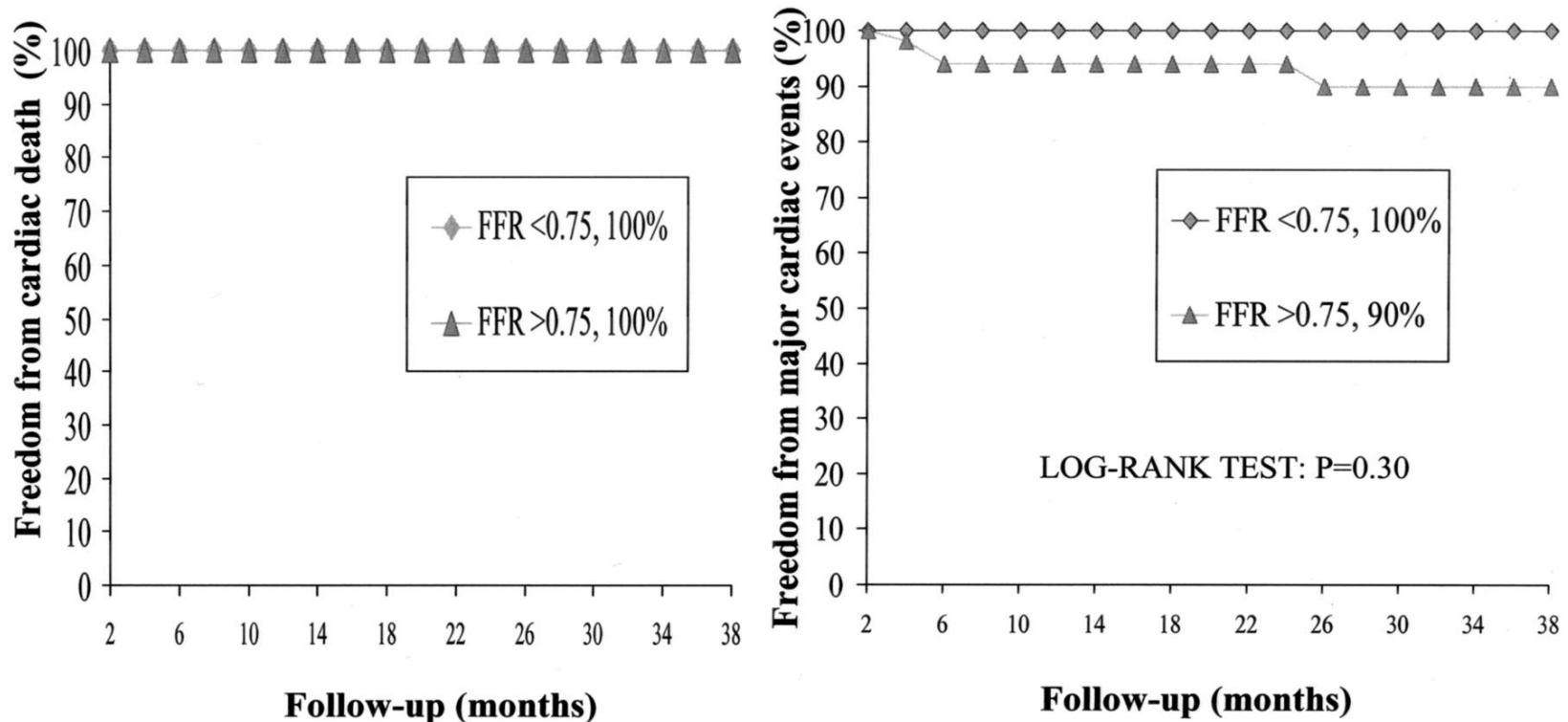
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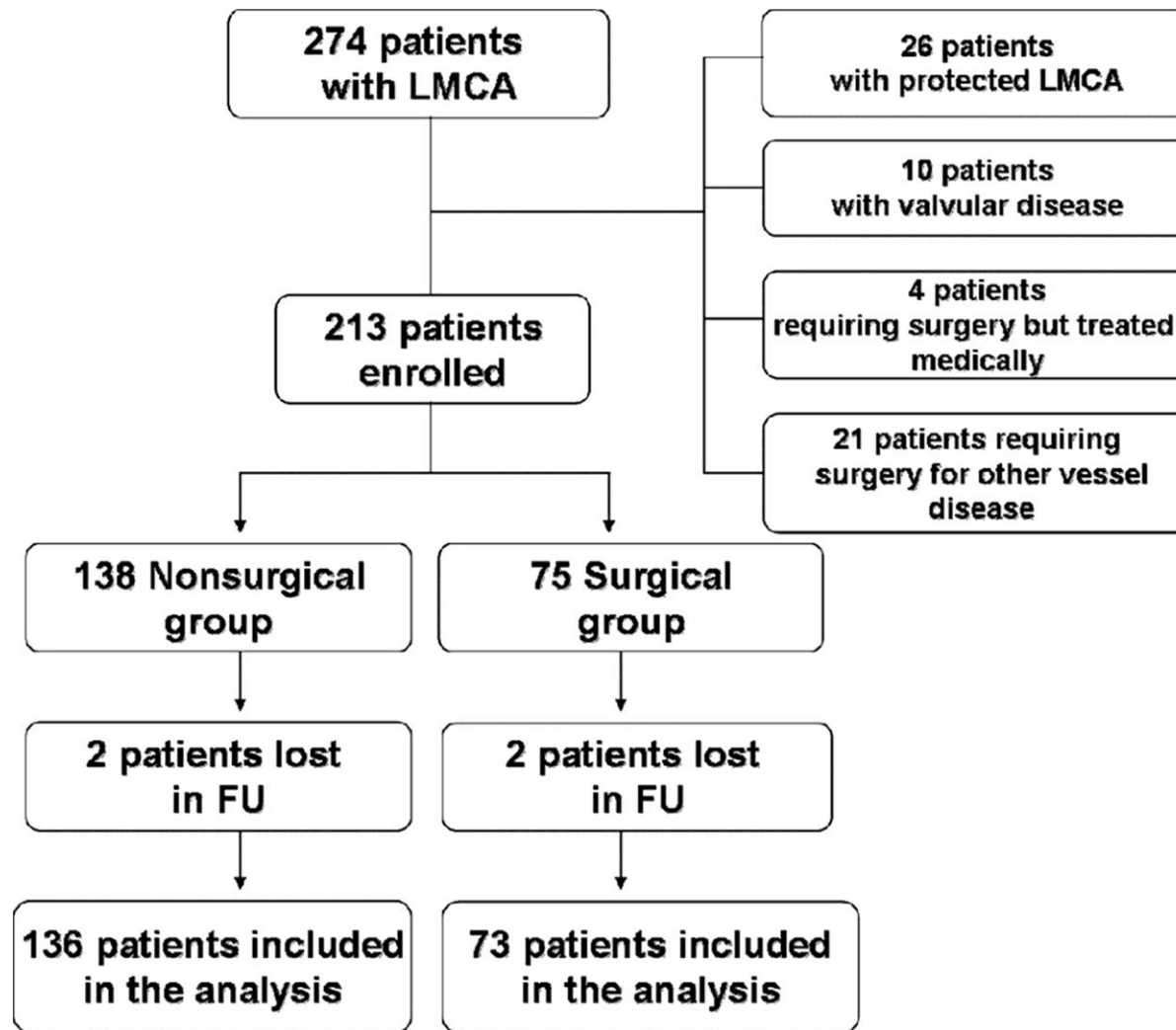
FFR for Assessing LM Significance

Summary of Published Studies

Patient #			FFR	FU	EFS		Survival	
Total population	Defer group*	Surgical group	Cut-off value	Mean (months)	Defer group* (%)	Surgical group (%)	Defer group* (%)	Surgical group (%)
54	24	30	0.75	29±15	76	83	100	97
51	37	14	0.75	25±11	90	100	100	100
27	20	7	0.75	26±12	90	86	100	86
38	20	18	0.75	24±12	90	89	100	89
15	8	7	0.75	33±10	100	71	100	100
51	24	27	0.75	29±16	69	66	100	81
(236)	(133)	(103)						

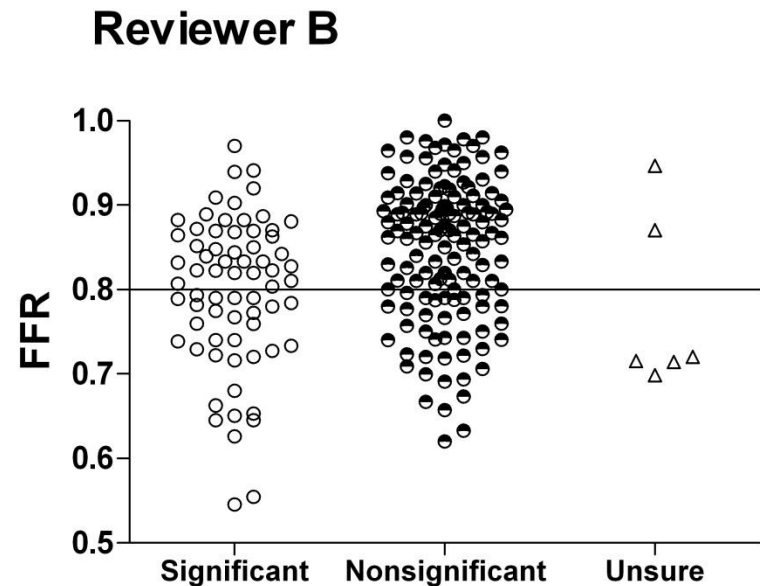
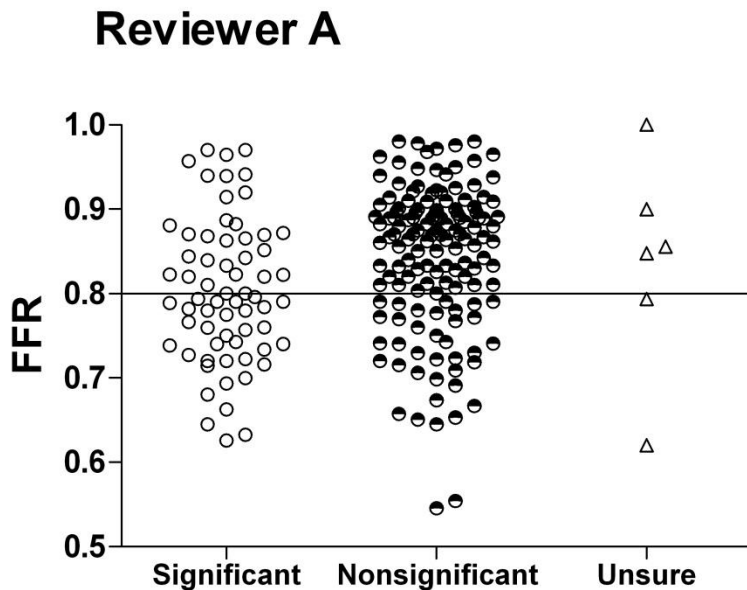


FFR and Intermediate Left Main



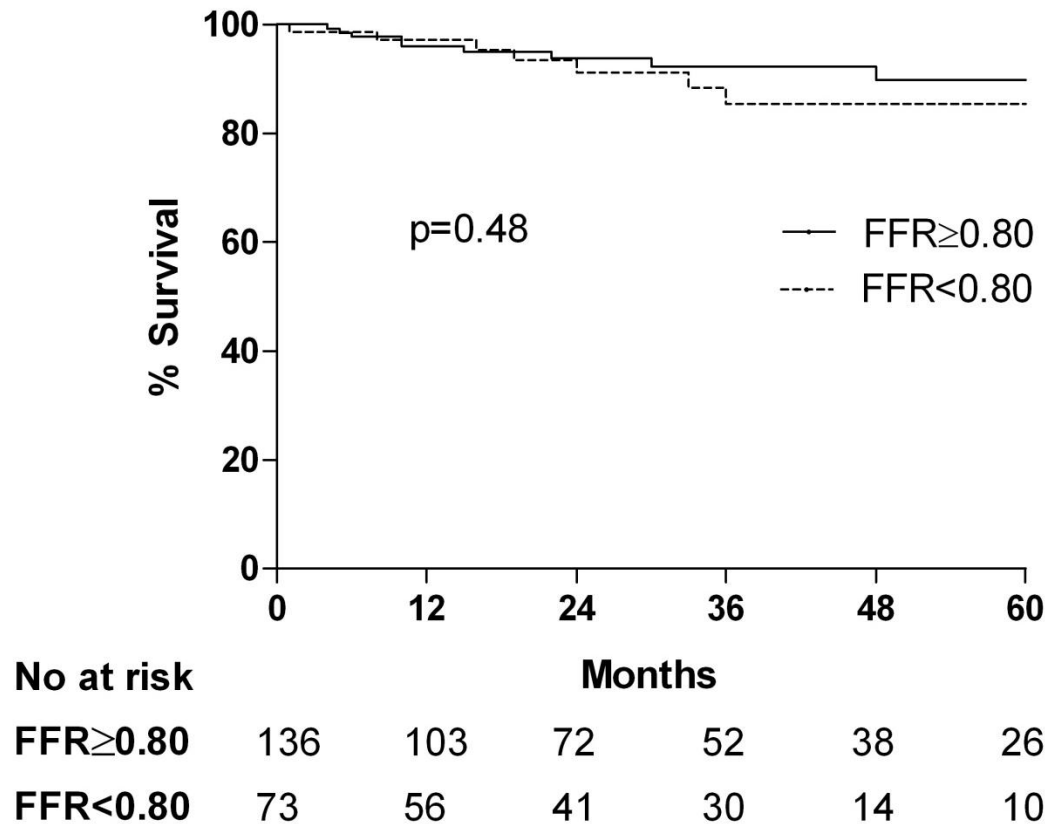
FFR for Assessing LM Significance

Poor correlation between “eyeball” and FFR



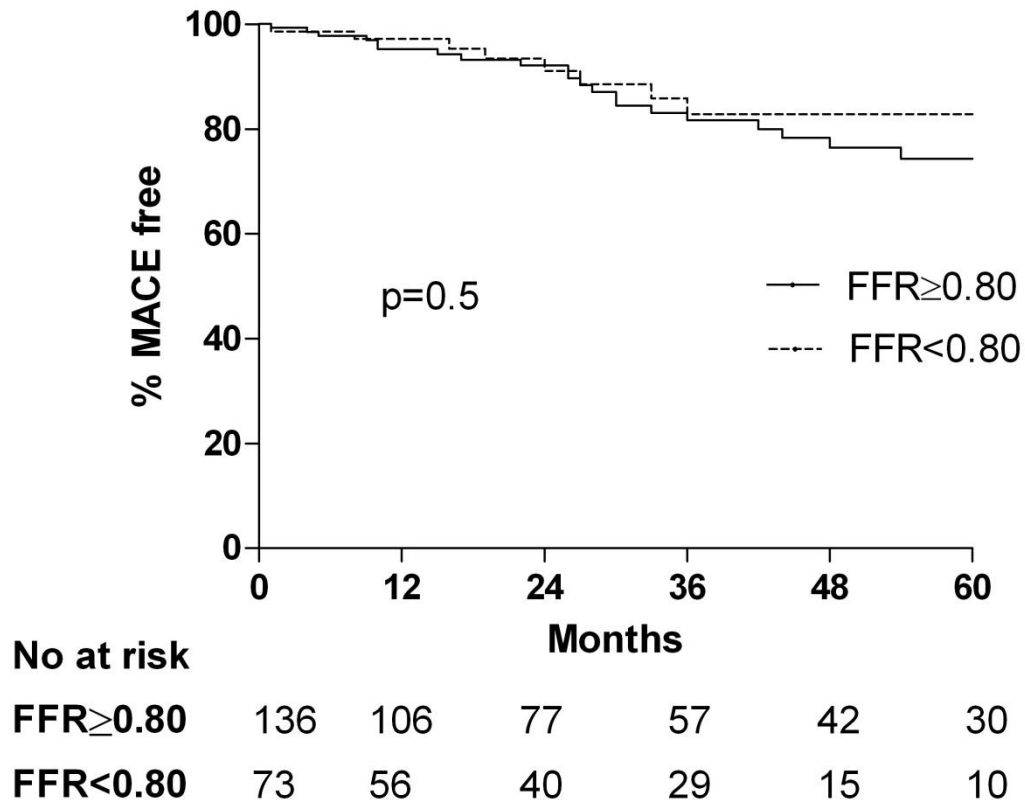
FFR for Assessing LM Significance

Survival Rate



FFR for Assessing LM Significance

MACE Rate



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Filter: Enhance 2

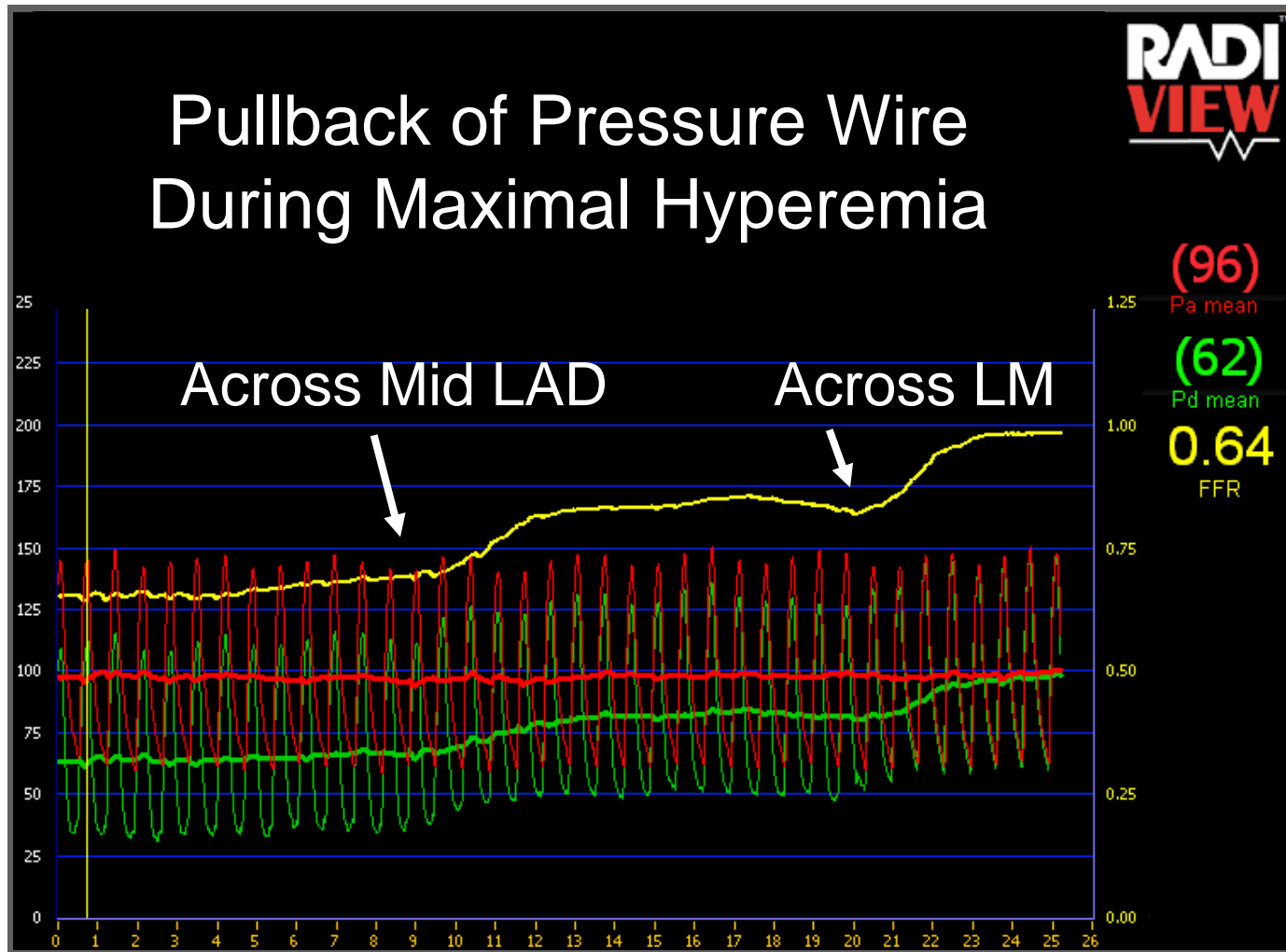
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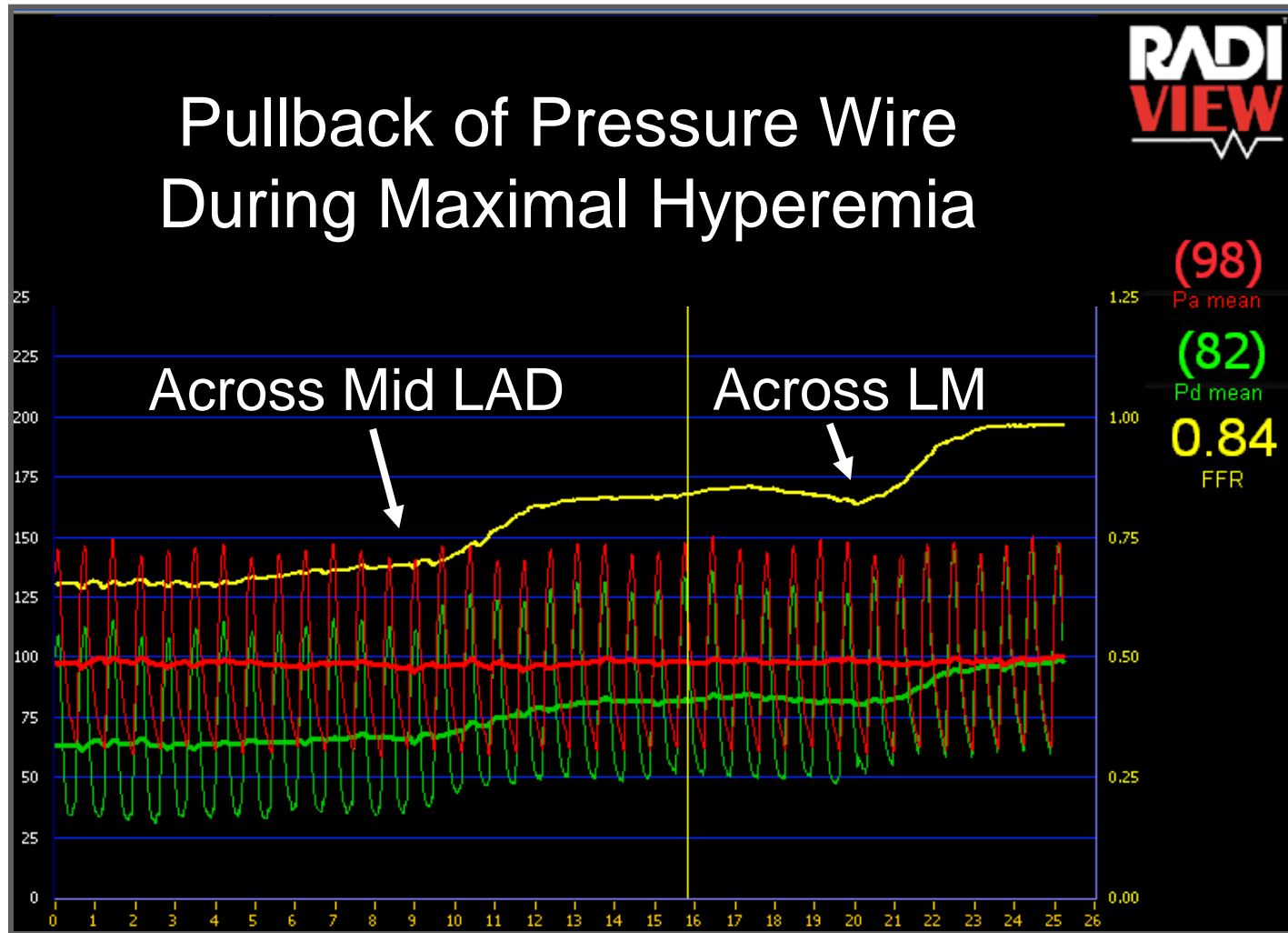
13.60 RAO
30.90 CRA



FFR of Left Main



FFR of Left Main

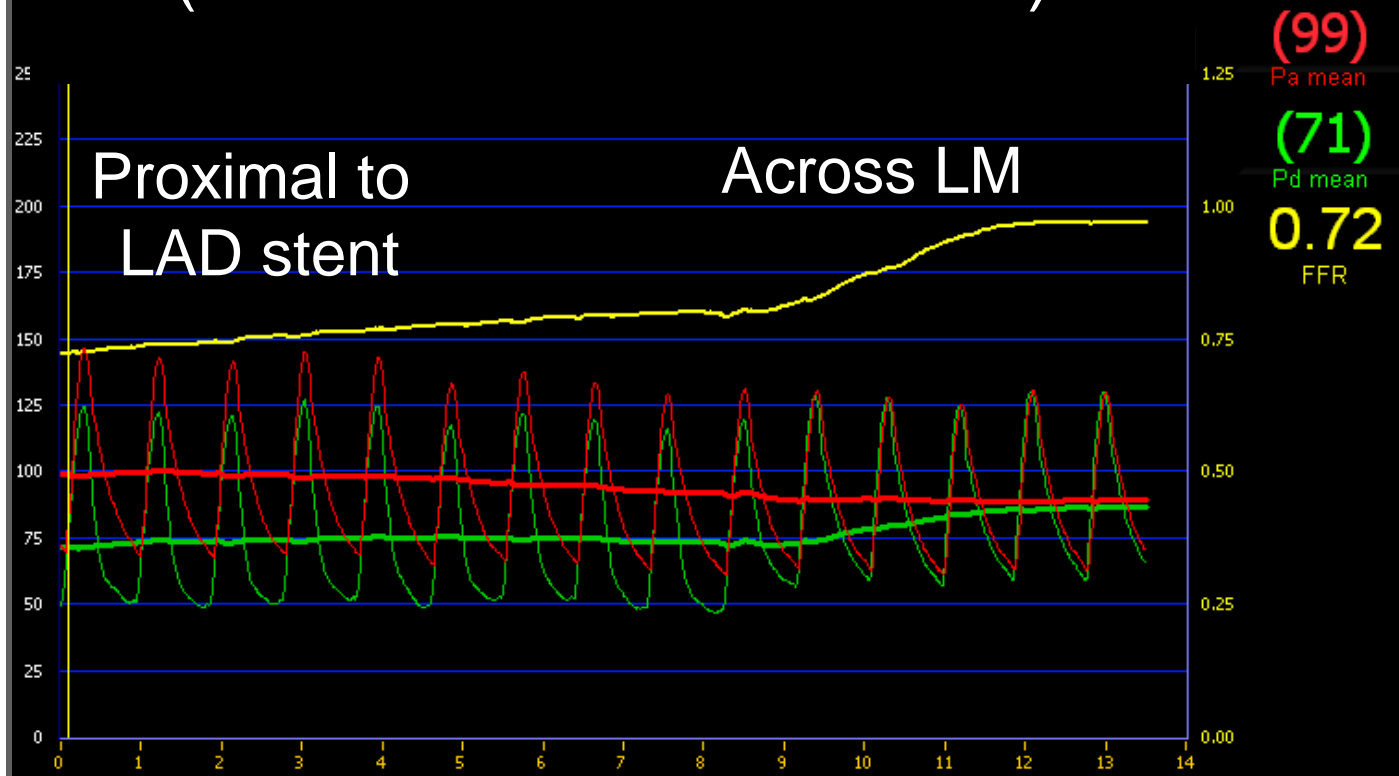


After rotational atherectomy and 2.5x28 mm DES, post-dilated to 3.0 mm

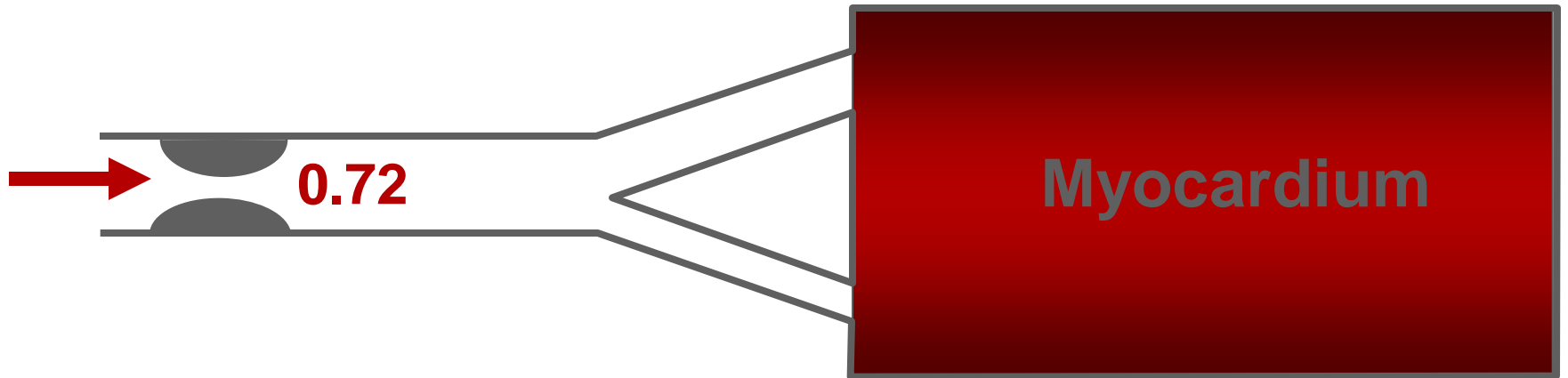
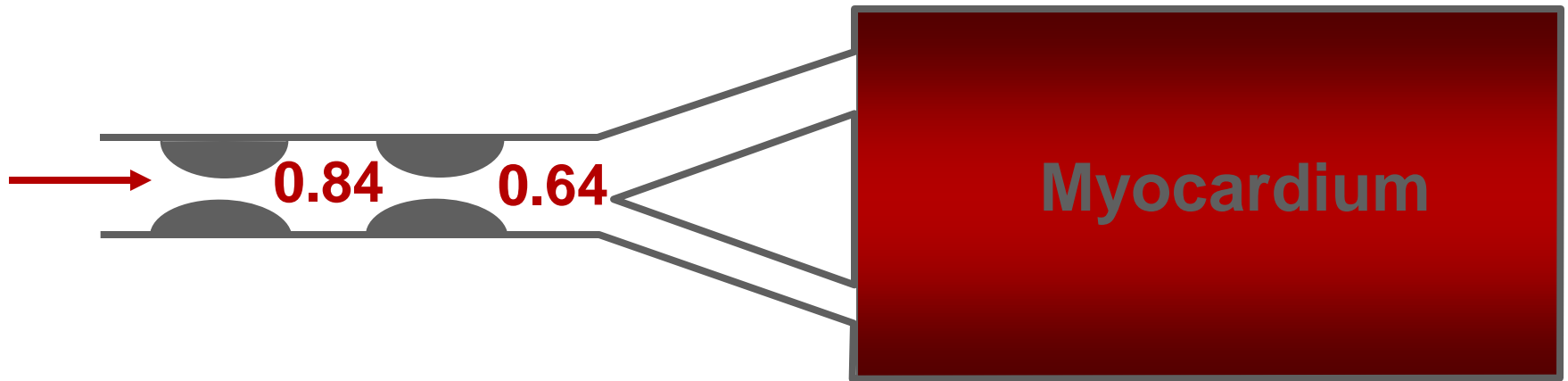


FFR of Left Main

FFR of Left Main = 0.72
(In absence of LAD lesion)



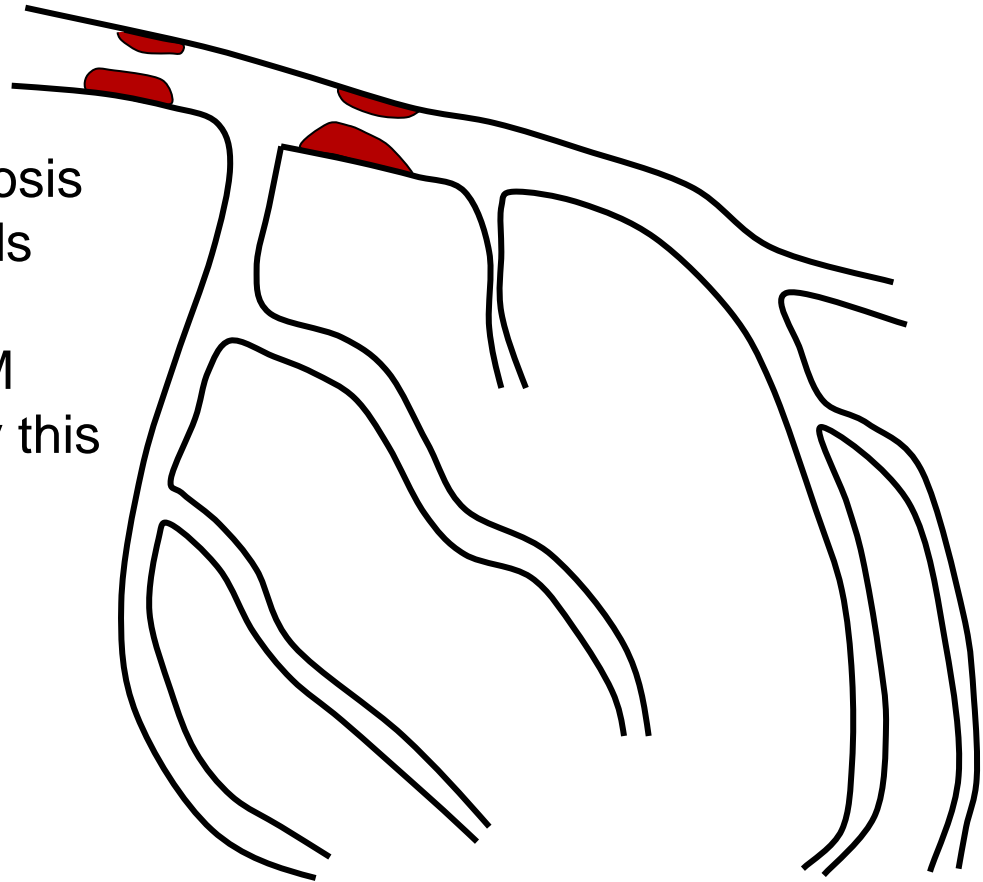
Effect of Tandem Lesions



Left Main Stem Stenoses are Rarely Isolated

The influence of a distal stenosis on the FFR of the LM depends on the extent to which hyperemic flow across the LM stenosis will be decreased by this distal lesion

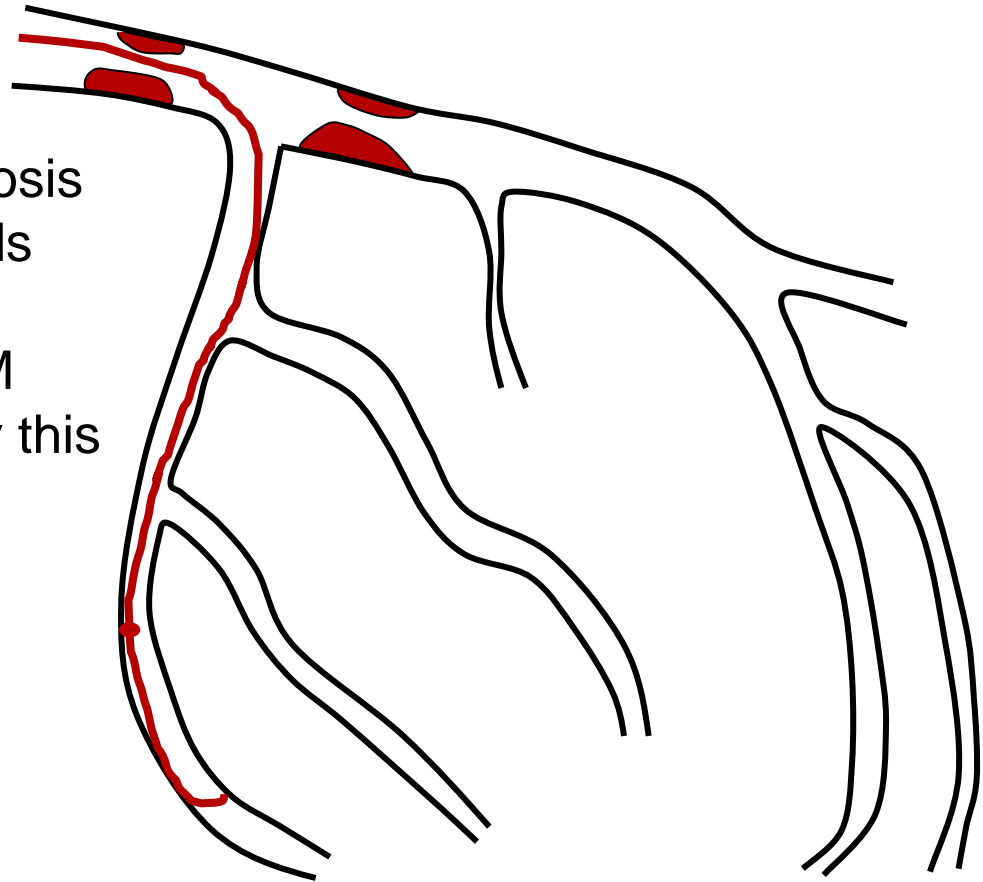
- Severity
- Myocardial mass



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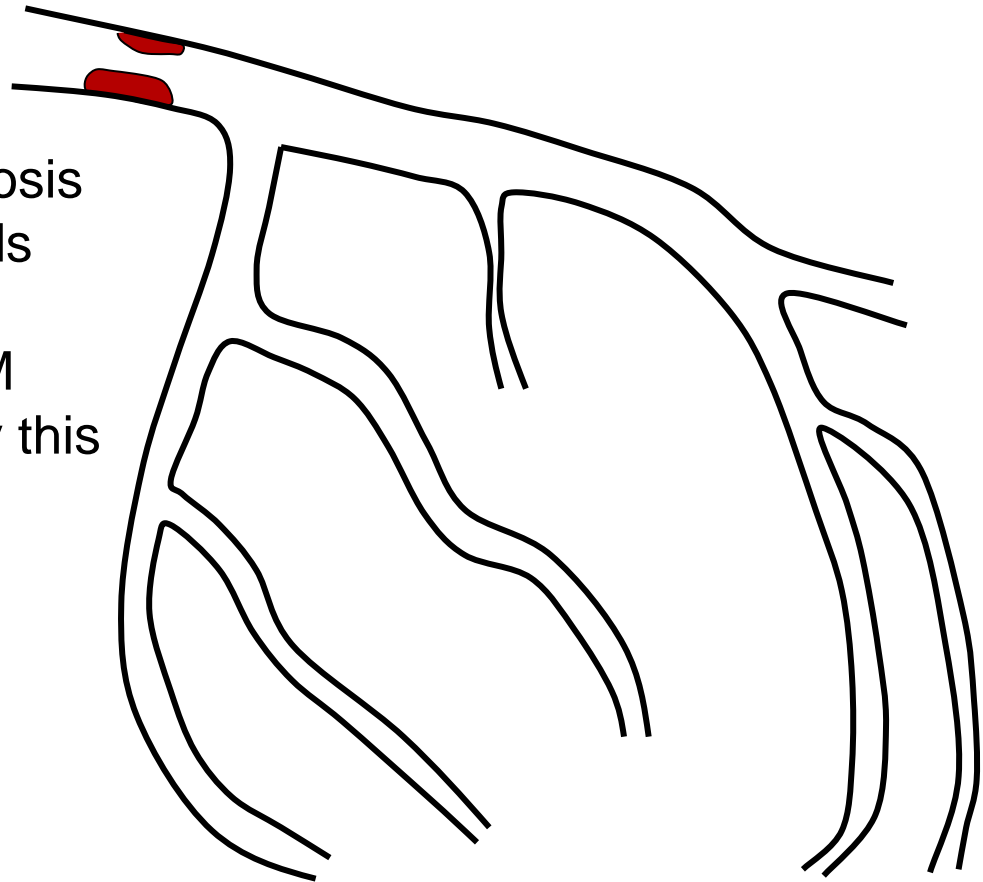
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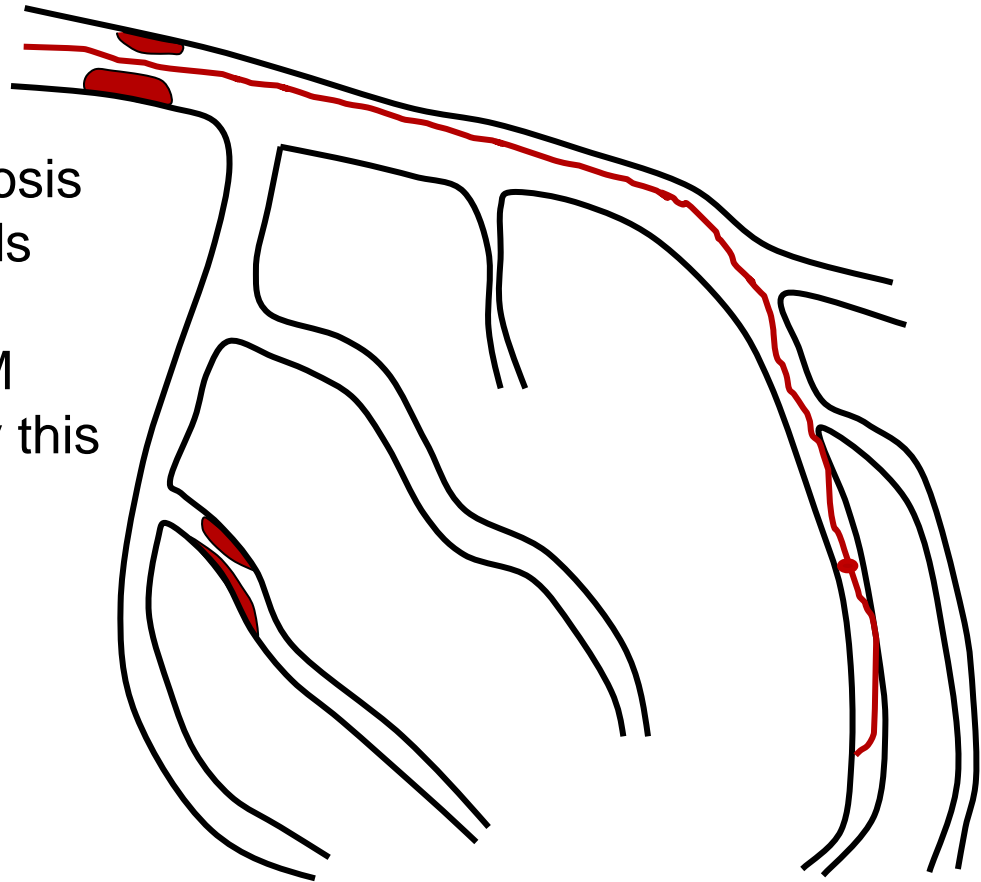
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- Severity
- Myocardial mass



Effect of Epicardial Lesions on FFR

Assessment of Intermediate LM Disease

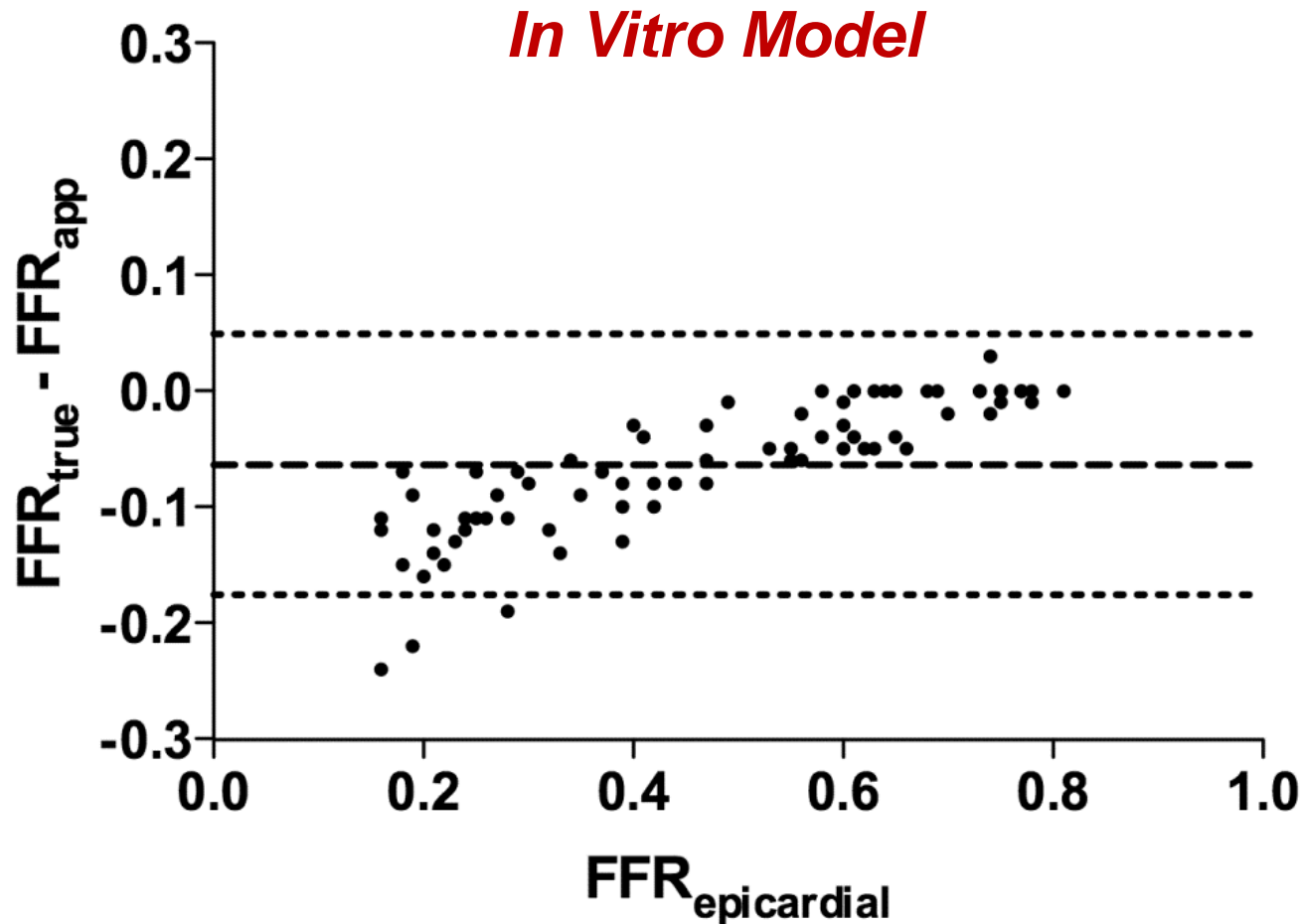


In Vitro Model



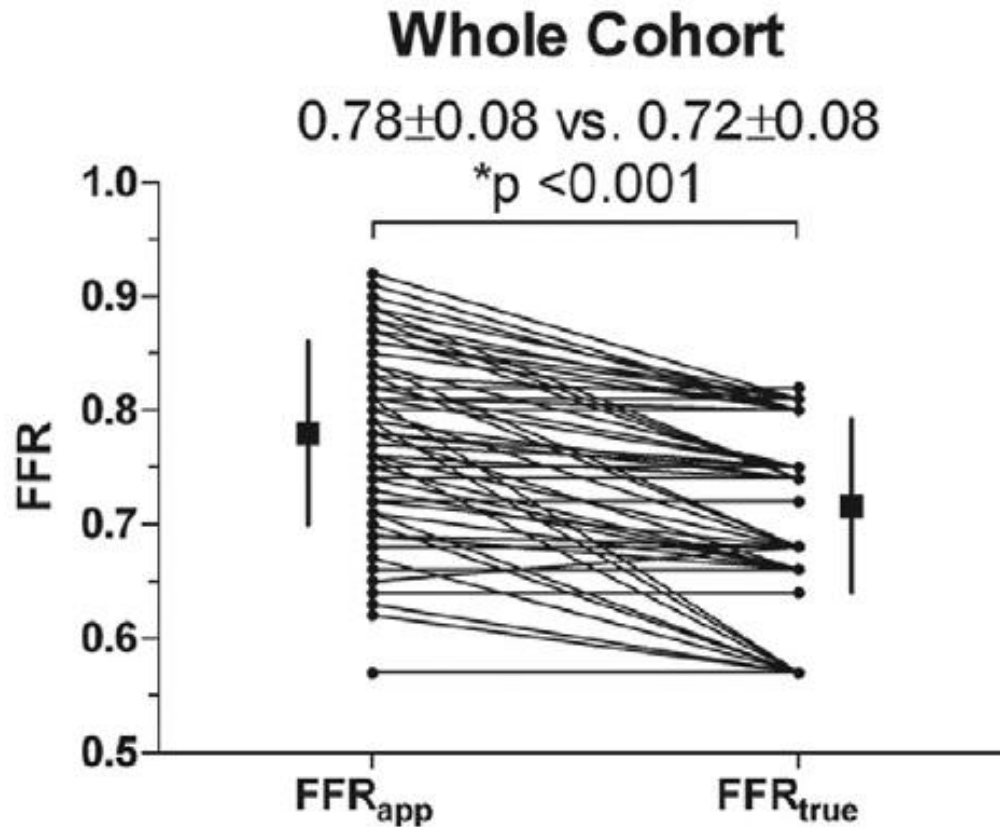
Effect of Epicardial Lesions on FFR

Assessment of Intermediate LM Disease



Effect of Epicardial Lesions on FFR Assessment of Intermediate LM Disease

In Vitro Model



Effect of Epicardial Lesions on FFR

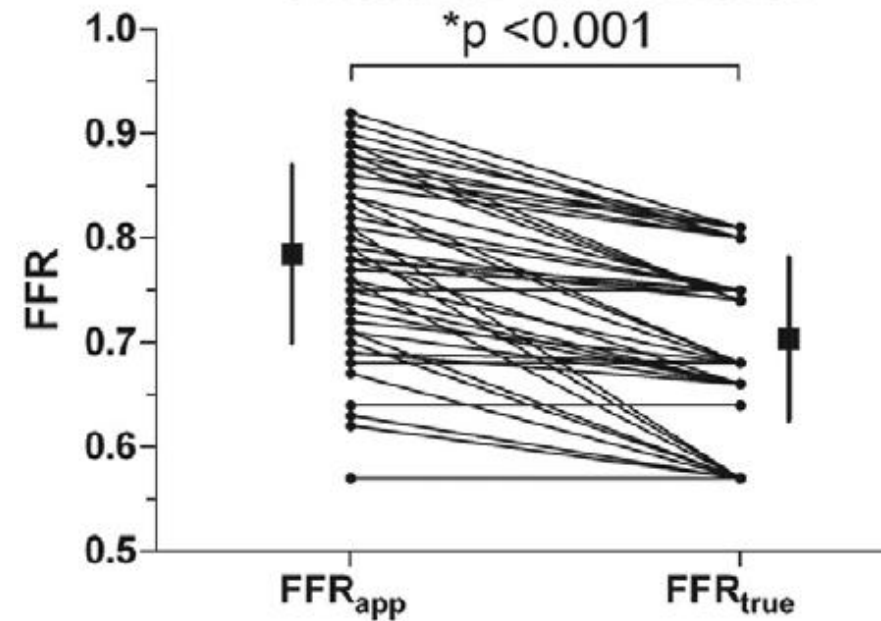
Assessment of Intermediate LM Disease

In Vitro Model

Composite FFR <0.65

0.79 ± 0.09 vs. 0.70 ± 0.08

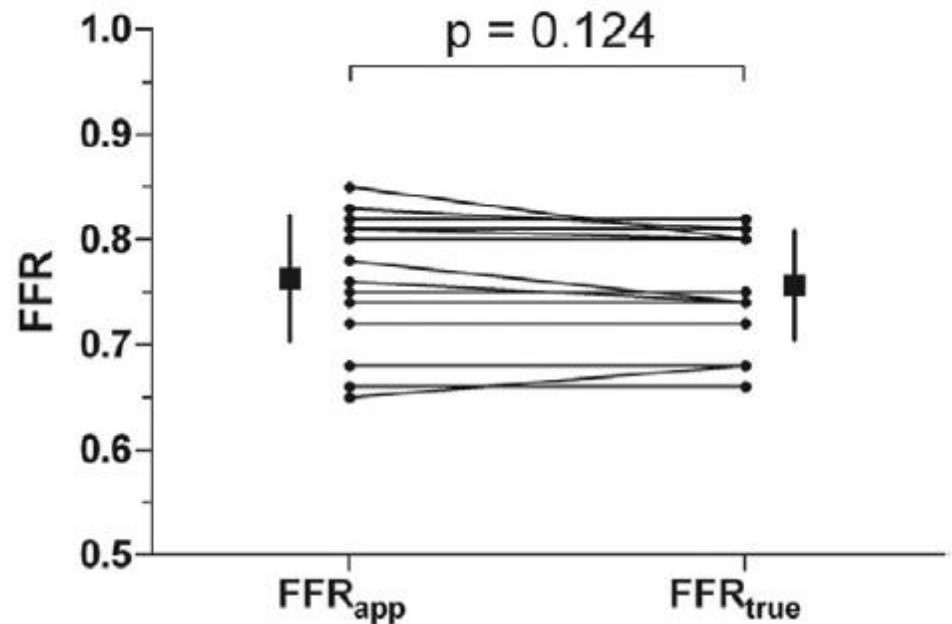
* $p < 0.001$



Composite FFR ≥ 0.65

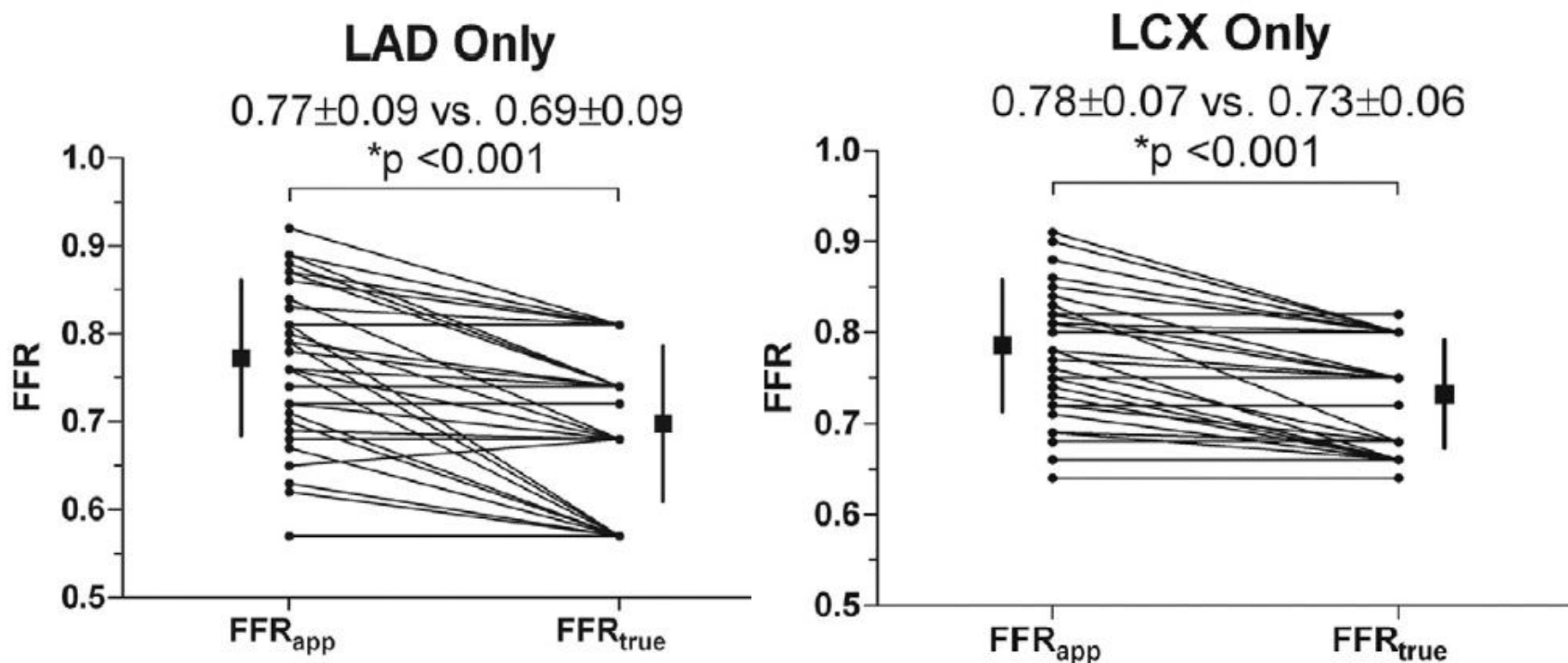
0.76 ± 0.06 vs. 0.76 ± 0.05

$p = 0.124$



Effect of Epicardial Lesions on FFR Assessment of Intermediate LM Disease

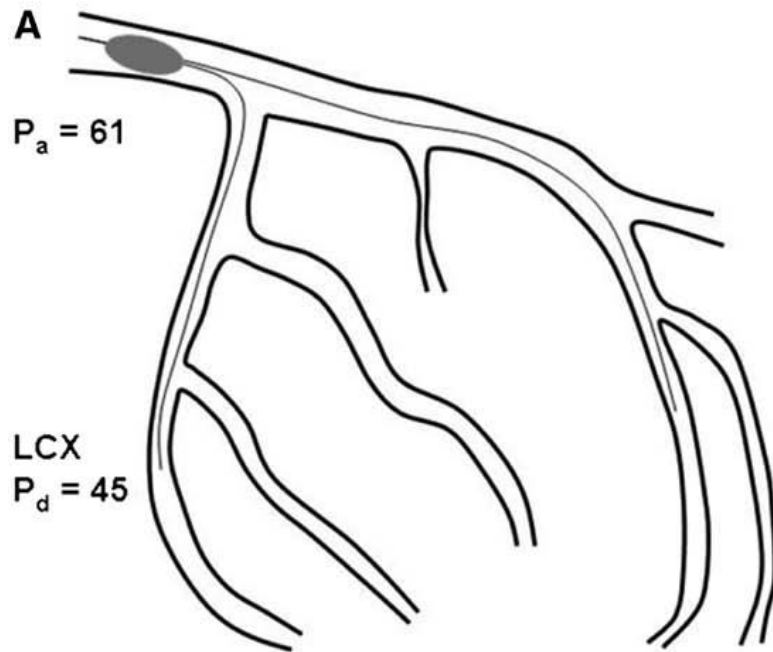
In Vitro Model



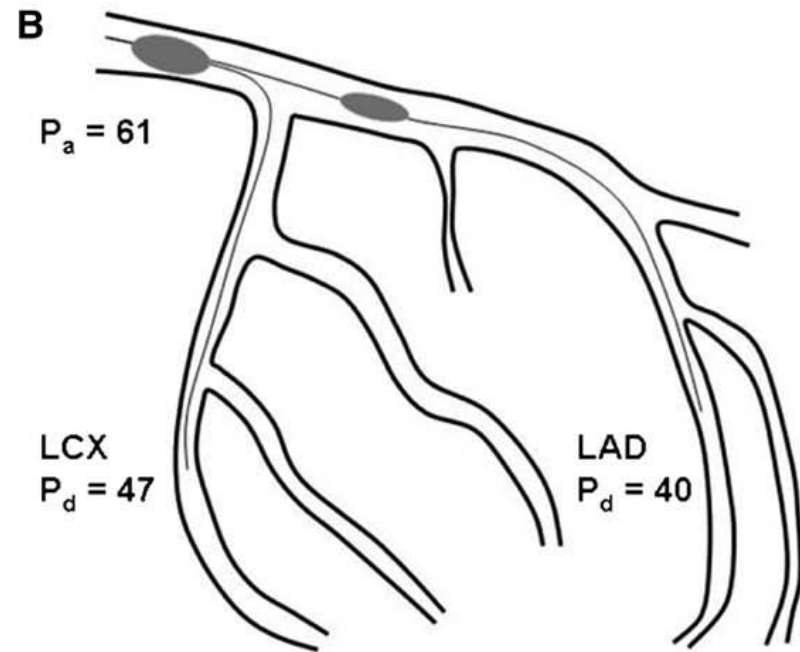
Effect of Epicardial Lesions on FFR

Assessment of Intermediate LM Disease

Animal Model



$$FFR_{\text{true}} = 45/61 = 0.74$$



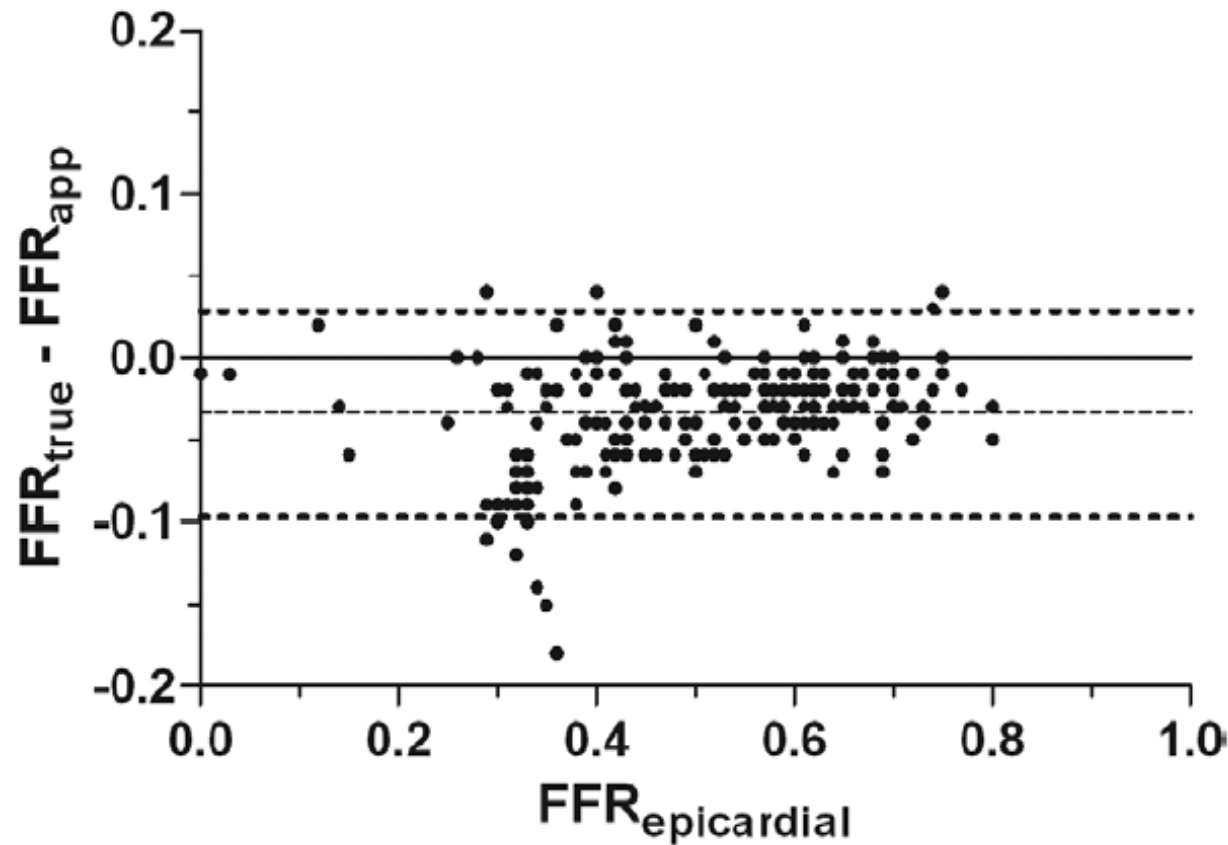
$$FFR_{\text{app}} = 47/61 = 0.77$$

$$FFR_{\text{epicardial}} = 40/61 = 0.66$$



Effect of Epicardial Lesions on FFR Assessment of Intermediate LM Disease

Animal Model

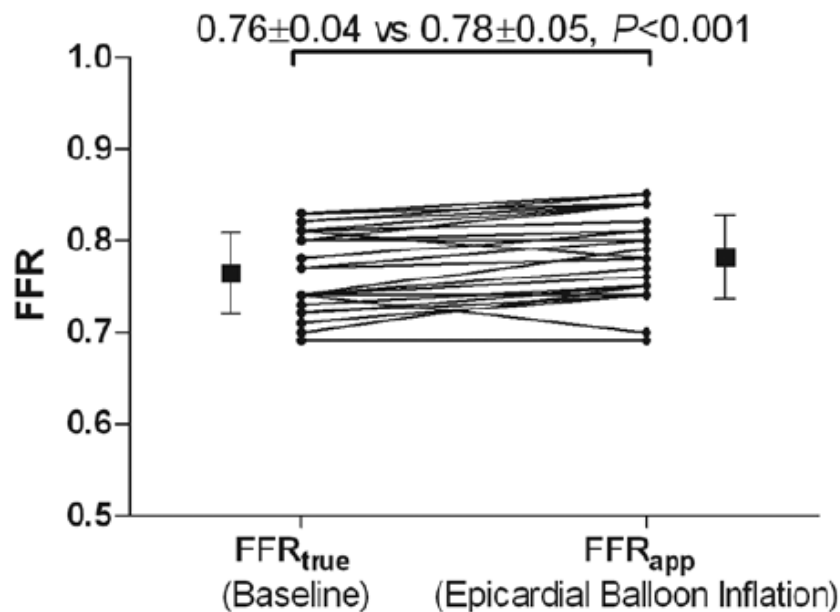


Effect of Epicardial Lesions on FFR

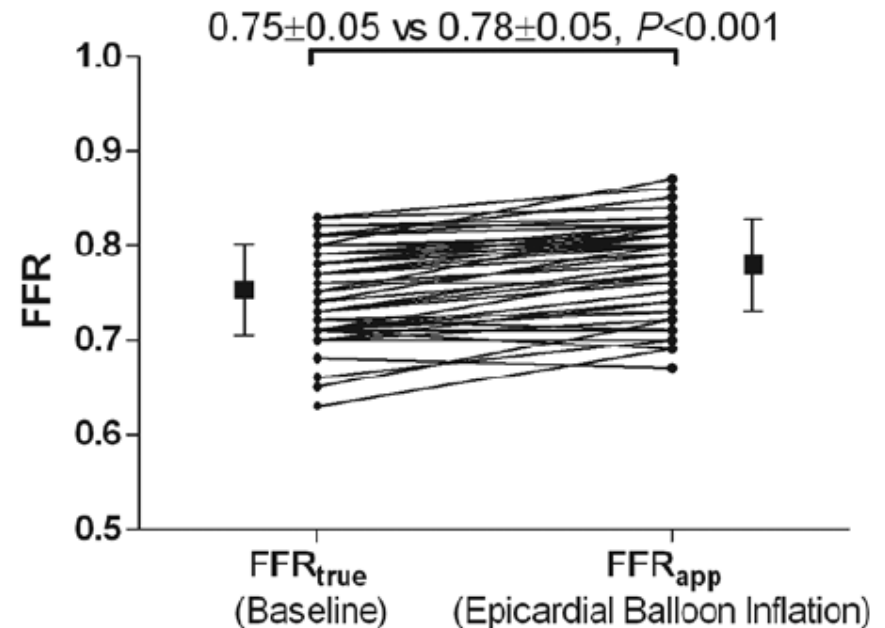
Assessment of Intermediate LM Disease

Animal Model

Mild Epicardial Disease
(FFR_{epicardial} 0.70-0.80)



Moderate Epicardial Disease
(FFR_{epicardial} 0.60-0.69)

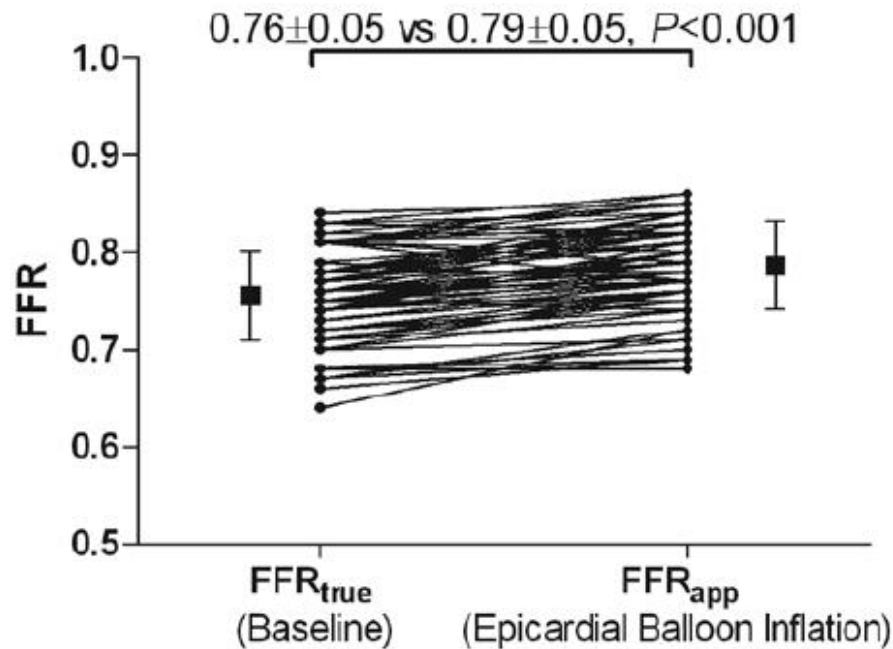


Effect of Epicardial Lesions on FFR

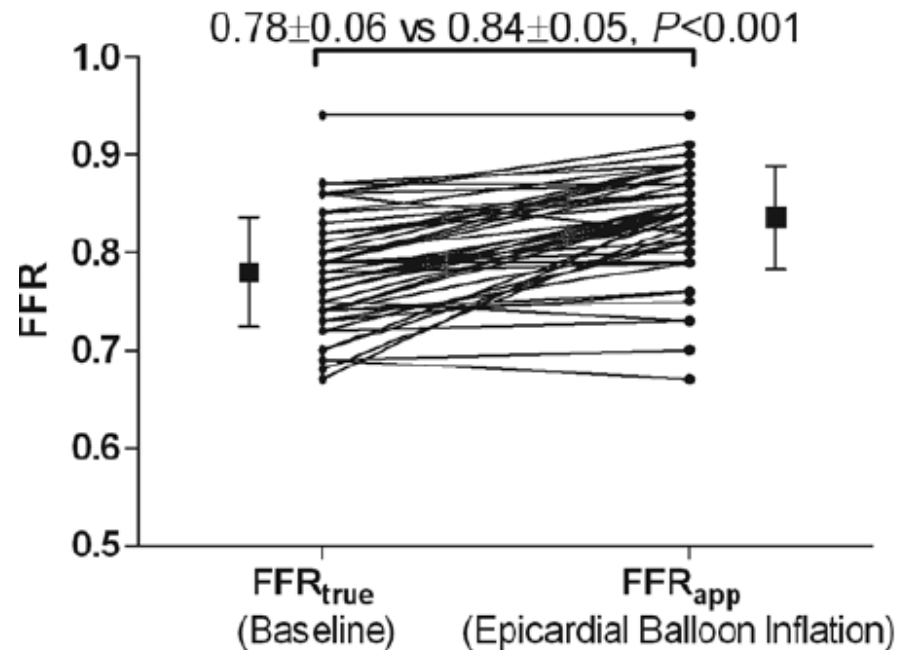
Assessment of Intermediate LM Disease

Animal Model

Severe Epicardial Disease ($\text{FFR}_{\text{epicardial}} 0.40-0.59$)

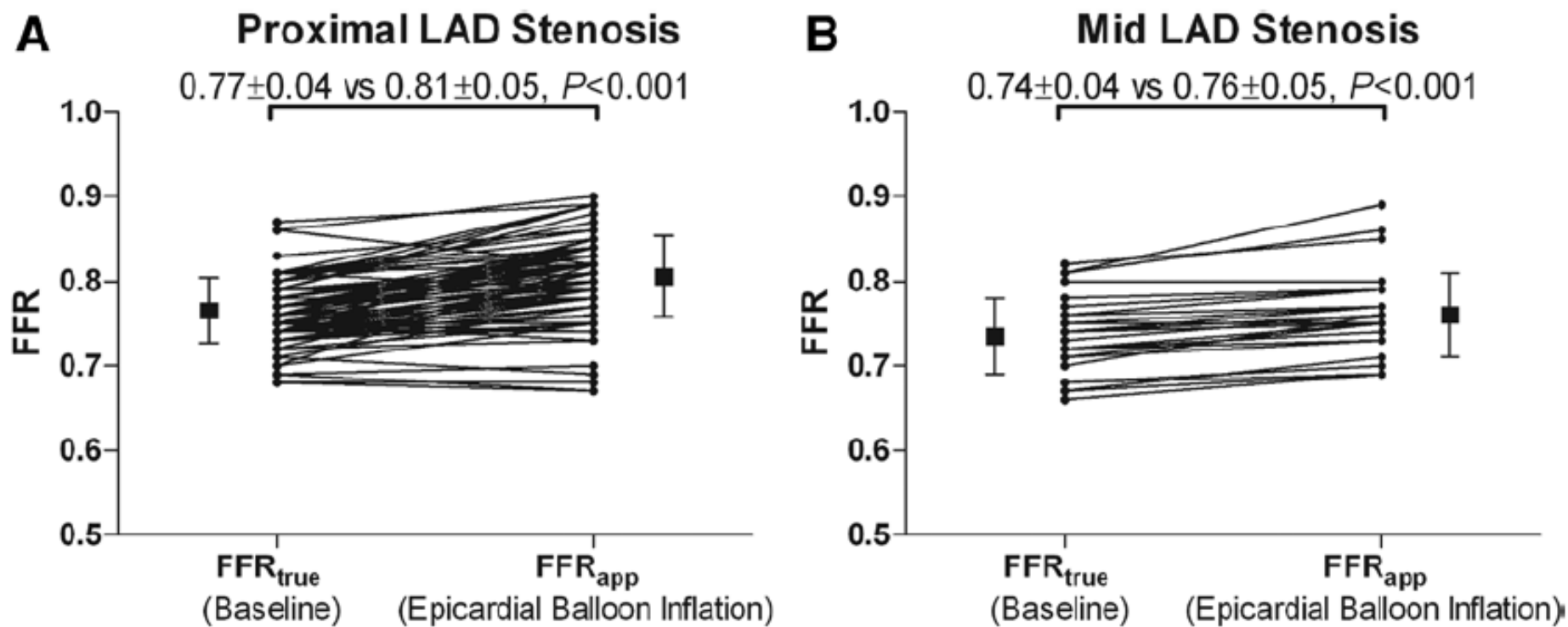


Complete Epicardial Occlusion ($\text{FFR}_{\text{epicardial}} < 0.40$)



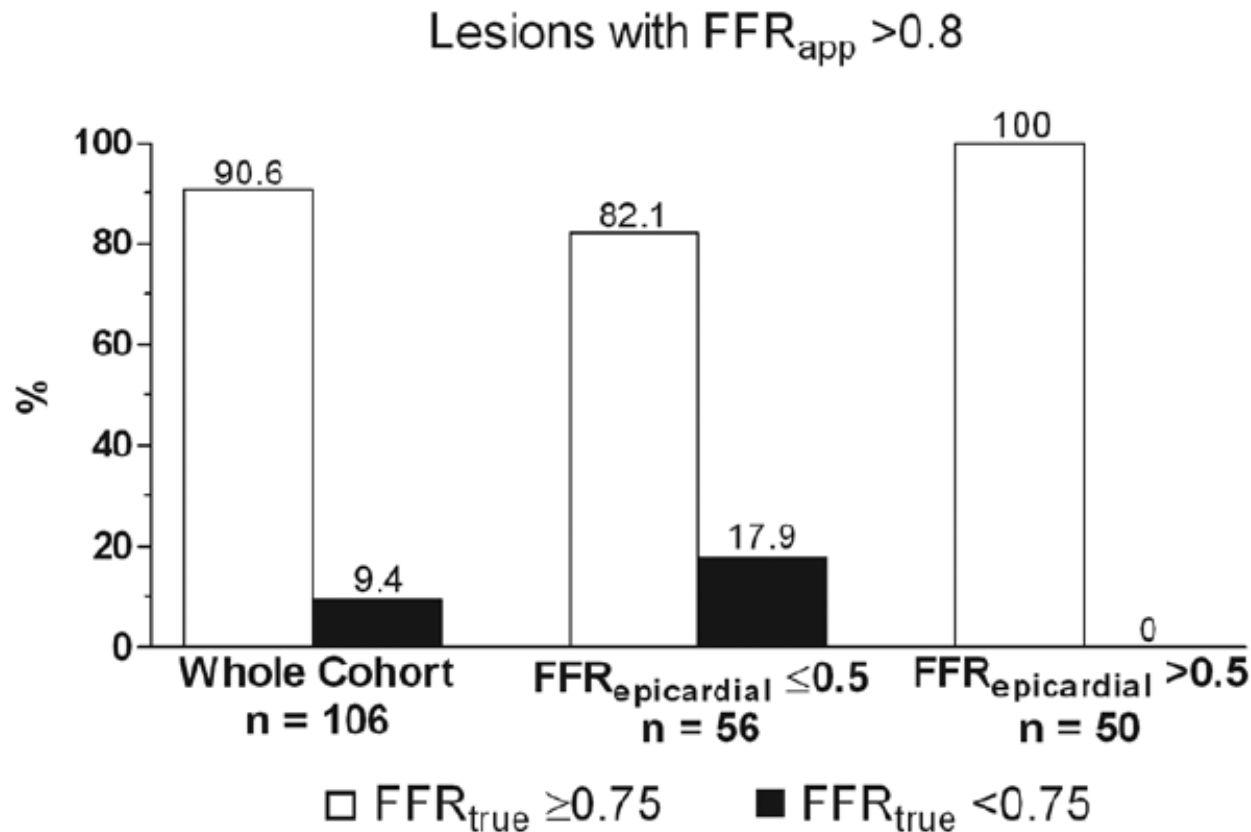
Effect of Epicardial Lesions on FFR Assessment of Intermediate LM Disease

Animal Model



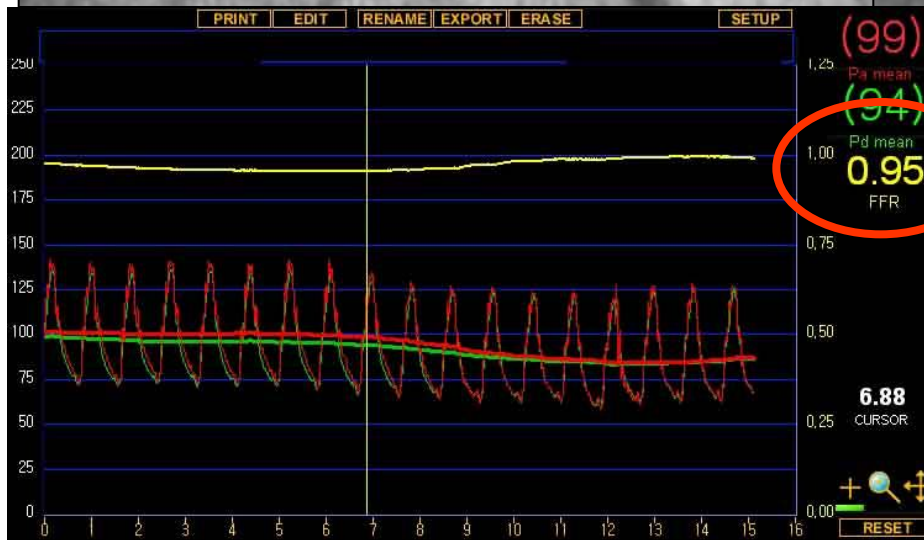
Effect of Epicardial Lesions on FFR Assessment of Intermediate LM Disease

Animal Model



Pre Stent

Post Stent

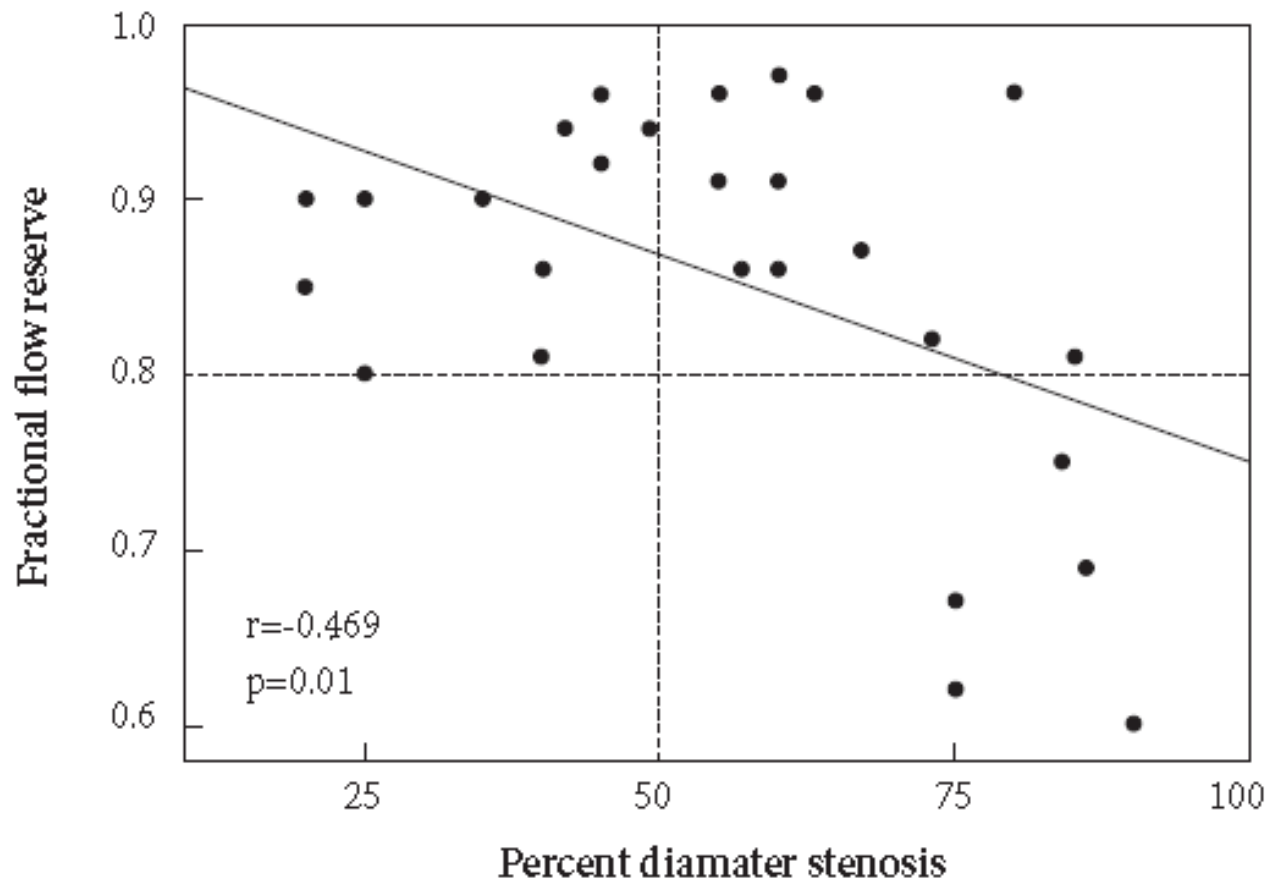


FU @ 8 mo



FFR of “Jailed” Left Circumflex

29 patients with LM/LAD crossover stenting with FFR of “jailed” Cx



FFR of “jailed” Circumflex

Mean 20 month follow-up	Defer group n = 24	PCI group n = 5
Death, n	0	1
Myocardial Infarction, n	0	0
TLR, n	3	1
Stent Thrombosis, n	0	0
Total Events, n	3	2



An Approach to the Equivocal LM

- First measure FFR in the least diseased vessel, preferably the LAD, with a pullback
 - If FFR < 0.80 , then revascularize
 - If FFR > 0.85 , then treat medically
 - If FFR between 0.80 and 0.85 and there is significant downstream epicardial disease in the other epicardial vessel, then consider IVUS
- Never forget the patient and the clinical scenario



Practical Aspects

Intravenous adenosine is the ideal hyperemic agent because it allows time to pull the guide catheter out of the ostium.

If possible, confirm pressure gradient across left main by checking FFR in both the LAD and Circumflex and by performing a pullback of the pressure wire.

A physiologic evaluation of left main disease, compared to an anatomic evaluation alone, is safe and appropriate, just as it is in non-left main CAD.

