



# SPECT AND PET: HOW USEFUL FOR PCI PATIENTS

Won Jun Kang, MD, PhD.

Division of Nuclear Medicine  
Department of Radiology  
Yonsei University College of Medicine

# Nuclear Imaging: SPECT and PET

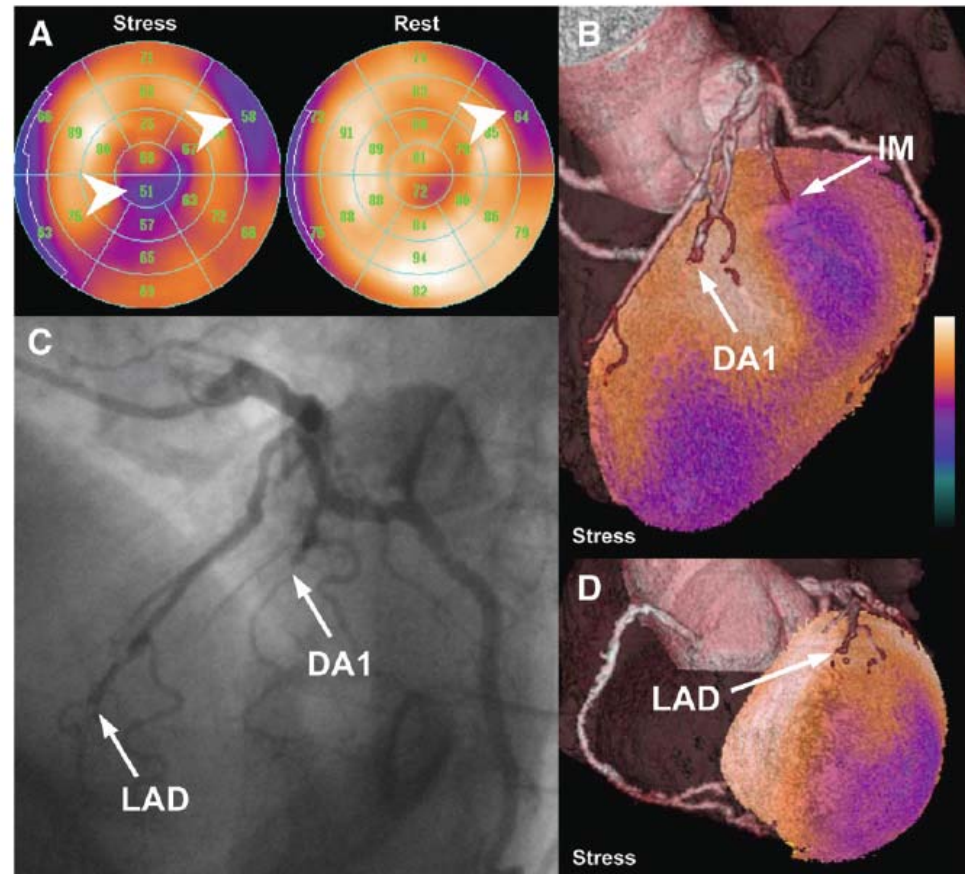
- Before PCI
  - Culprit vessel identification
  - Viability study
- After PCI
  - Restenosis detection

# Role of SPECT before PCI

- Documenting ischemia
- Determining the **functional impact** of the lesion
  - Stratify risk in 25-75% stenosis
- Identifying the lesion responsible for the ischemic symptom: '**Culprit lesion**' (>80%)
- Selection of patients to perform PCI
  - Based on CAG alone vs. considering of ischemia
  - No ischemia → low risk

# Cardiac Image Fusion from Stand-Alone SPECT and CT: Clinical Experience

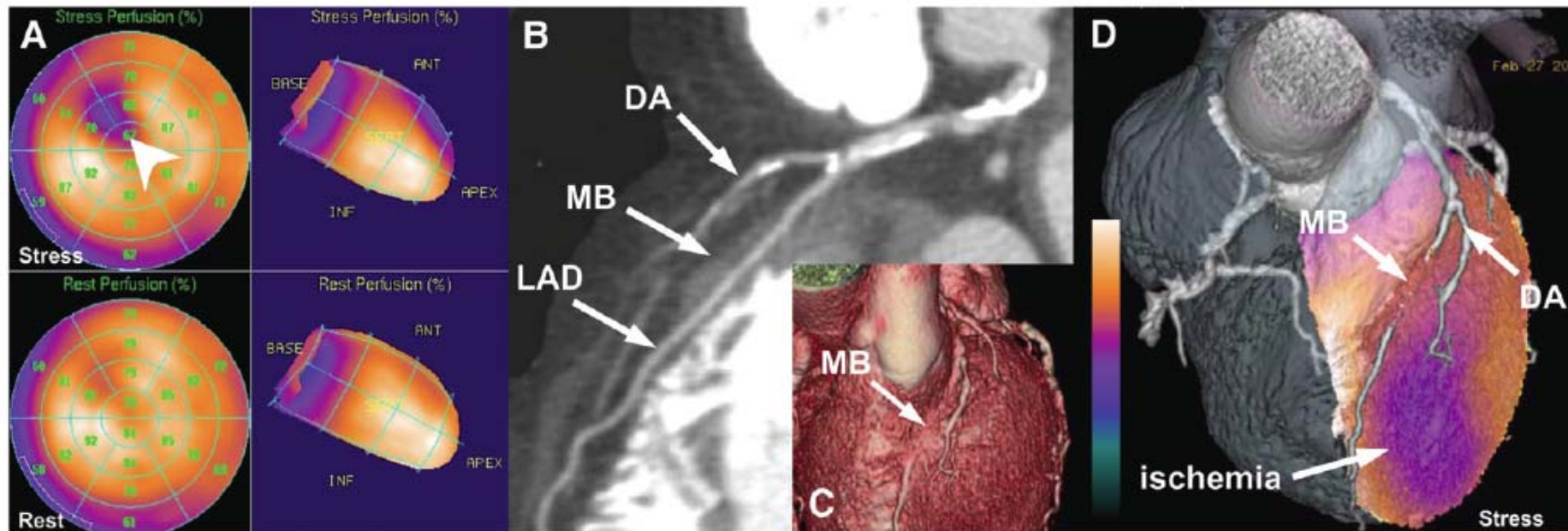
Supplying vessel  
Perfusion defect



**FIGURE 2.** (A) Stress and rest perfusion polar maps of SPECT-MPI study show mixed basal anterolateral defect and reversible inferoapical perfusion defect (arrowheads). (B and D) Fused SPECT/CT images reveal total occlusion of LAD and subtotal occlusion of first diagonal branch (DA1), which are confirmed by conventional CA (C). Anterolateral perfusion defect is caused by lesion of partially calcified small intermediary branch (IM); however, this vessel is not well visualized by CA.

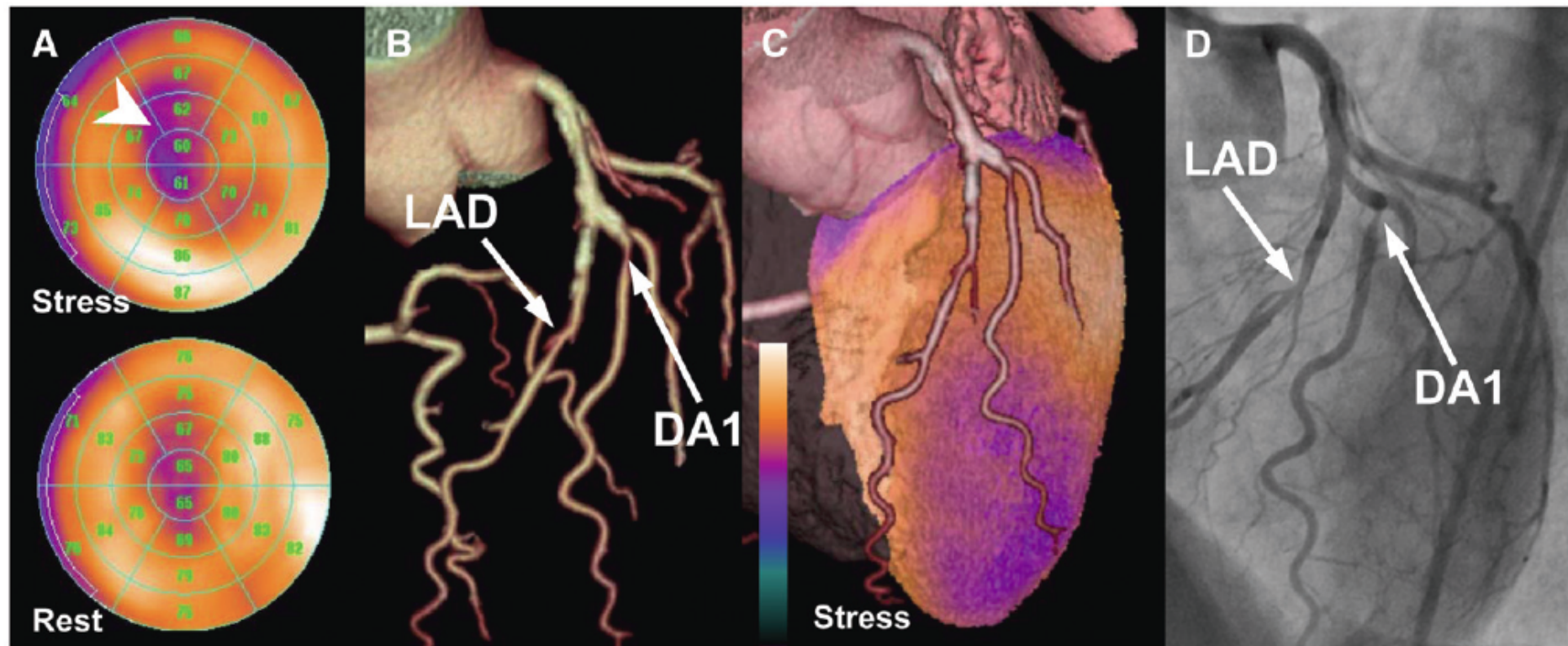
Anterolateral defect by intermediary a. stenosis

J Nucl Med 2007; 48:696-703



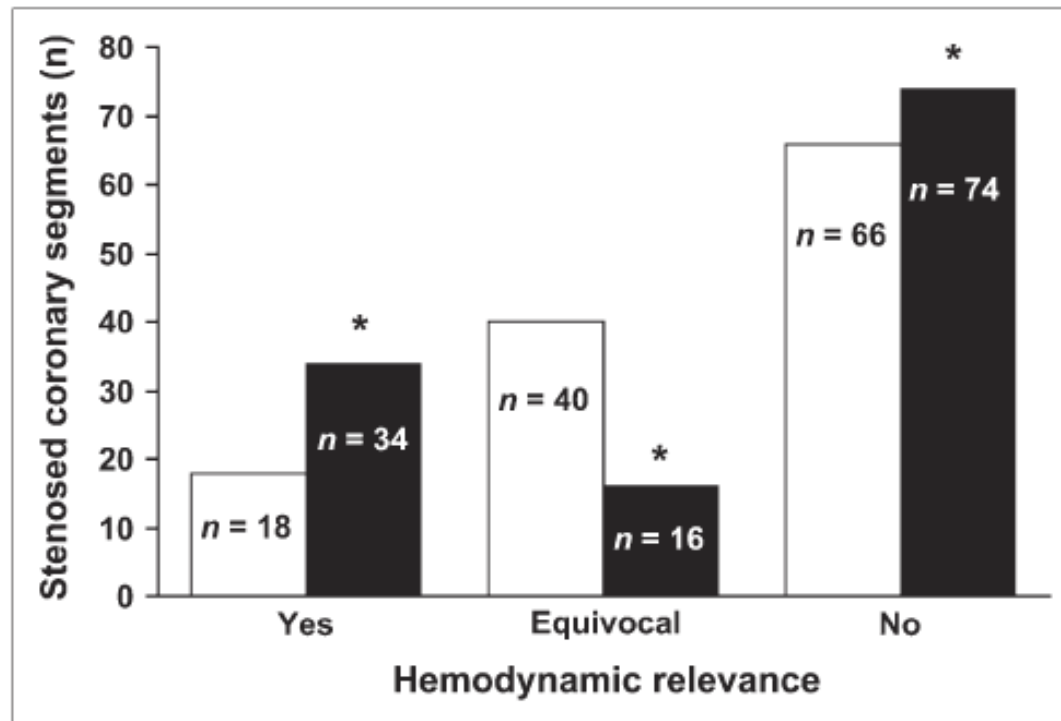
**FIGURE 3.** (A) Perfusion polar maps at stress (dobutamine stress) and rest show reversible anteroseptal perfusion defect. (B and C) 64-slice CTA revealed myocardial bridging (MB) of mid LAD of >2-cm length and calcified plaque at origin of first diagonal branch (DA). (D) Fused 3D SPECT/CT images could allocate reversible perfusion defect to DA, whereas MB seemed to be hemodynamically insignificant.

Anteroseptum defect—diagonal branch  
Myocardial bridging—insignificant



**FIGURE 4.** (A) Perfusion polar maps of SPECT-MPI at stress and rest show largely reversible anteroapical perfusion defect (arrowhead). (B) 3D volume-rendered CTA images show coronary vessel tree with stenosis of mid LAD and proximal stenosis of first diagonal branch (DA1). (C) Fused 3D SPECT/CT images are able to identify DA1 stenosis as functionally relevant lesions. (D) Findings were confirmed by invasive CA.

Apical anterior wall defect  
 Stenosis in mid LAD and diagonal branch



**FIGURE 1.** Interpretation of stenosed coronary segments with regard to hemodynamic significance on side-by-side (white columns) or fused (black columns) analysis. \* $P < 0.001$  for comparison of fused vs. side-by-side analysis ( $\chi^2$  test).



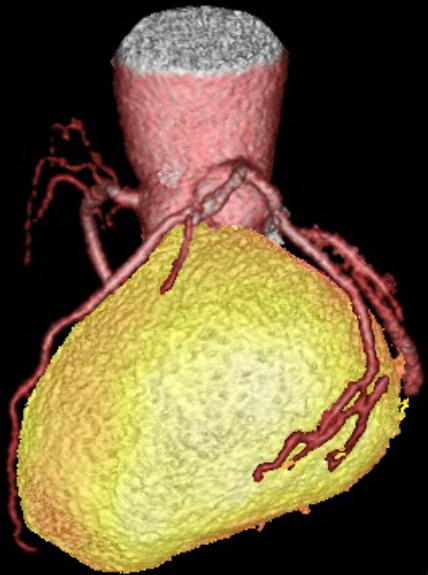
3D Volume 1  
 Ex: 801150601  
 Se: 5 +c  
 Volume Rendering No cut

DFOV 18.5cm  
 B25f



No VOl  
 kv 120  
 mA 266  
 0.2s  
 0.8mm /0.5sp  
 Tilt: 0.0  
 04:41:01 PM  
 W = 4095 L = 2048

SRA

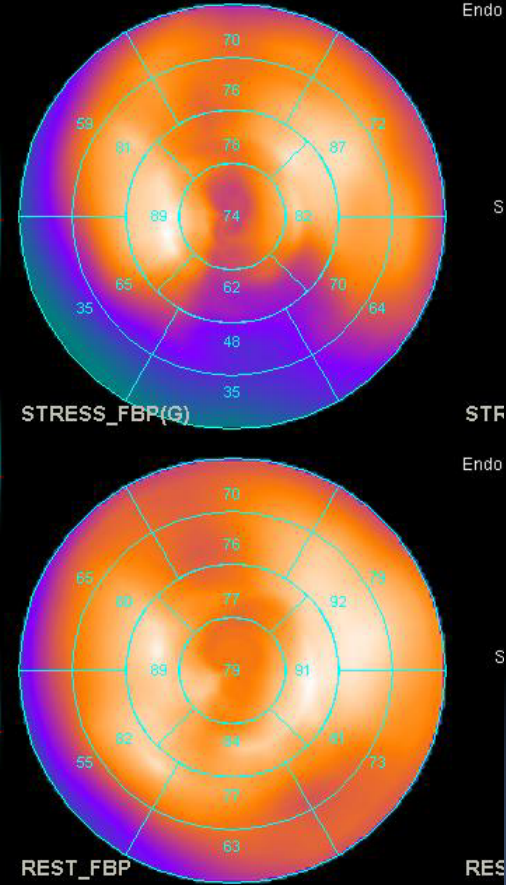
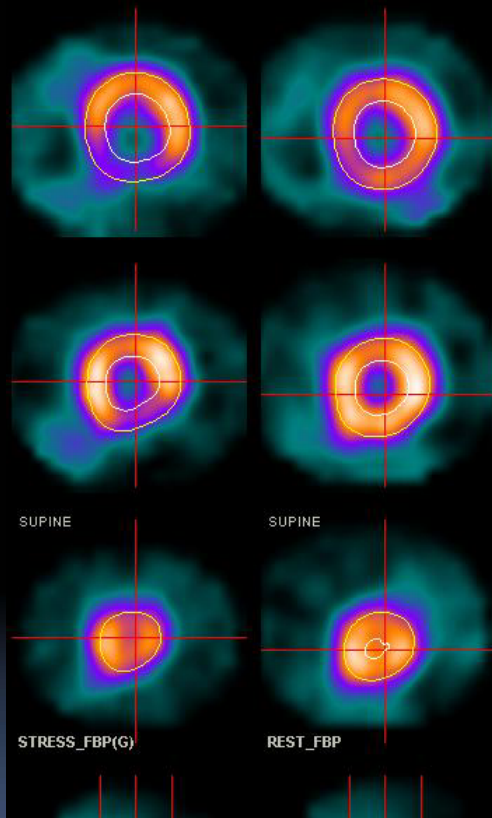


ILP

-999999.000000

TID: 1.1 (55/50)

P  
R  
I



Endo

S

STF

Endo

S

RES

RCA



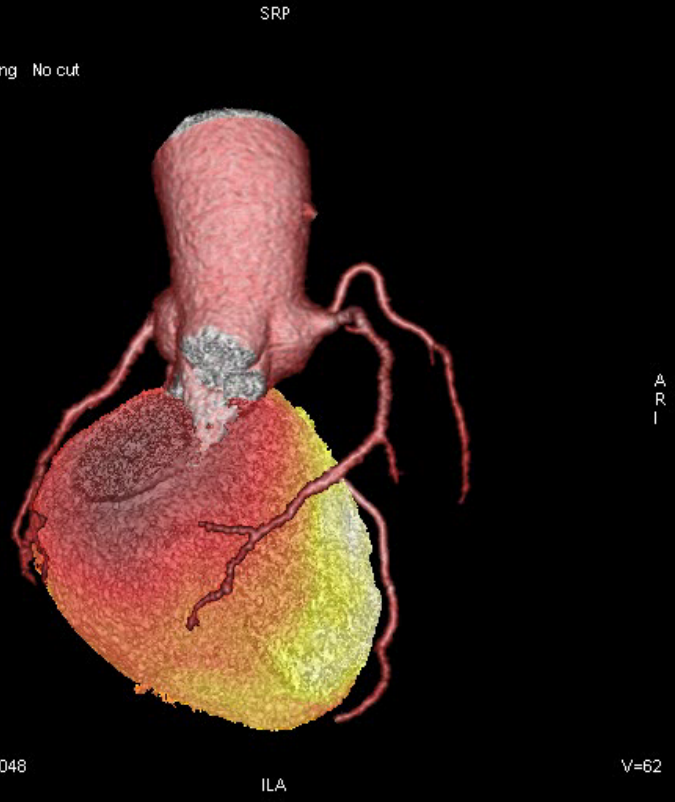


3D Volume 1  
Ex: 802026990  
Se: 5 +c  
Volume Rendering No cut

DFOV 19.0cm  
B25f

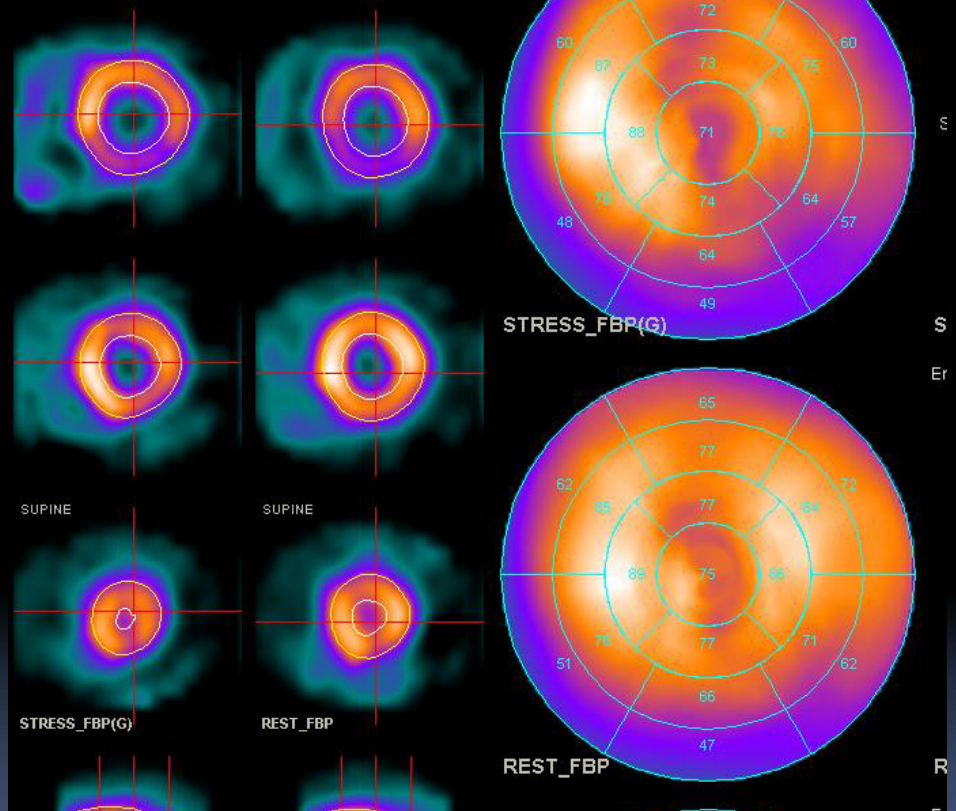


No VOI  
kv 120  
mA, 266  
0.2s  
0.8mm /0.5sp  
Tilt: 0.0  
05:32:25 PM  
W = 4095 L = 2048



Nuclear Imaging Environment

TID: 1.12 (73/66)



Anterior wall, inferolateral wall  
LAD diagonal, LCX



# Nuclear imaging for viability

- Preserved cell membrane integrity
  - Thallium
  - MIBI, tetrofosmin
- Preserved cell metabolism
  - FDG (glucose)

# Frequency of viable myocardium

**Table 1** Incidence of viable myocardium in patients with ischaemic left ventricular dysfunction

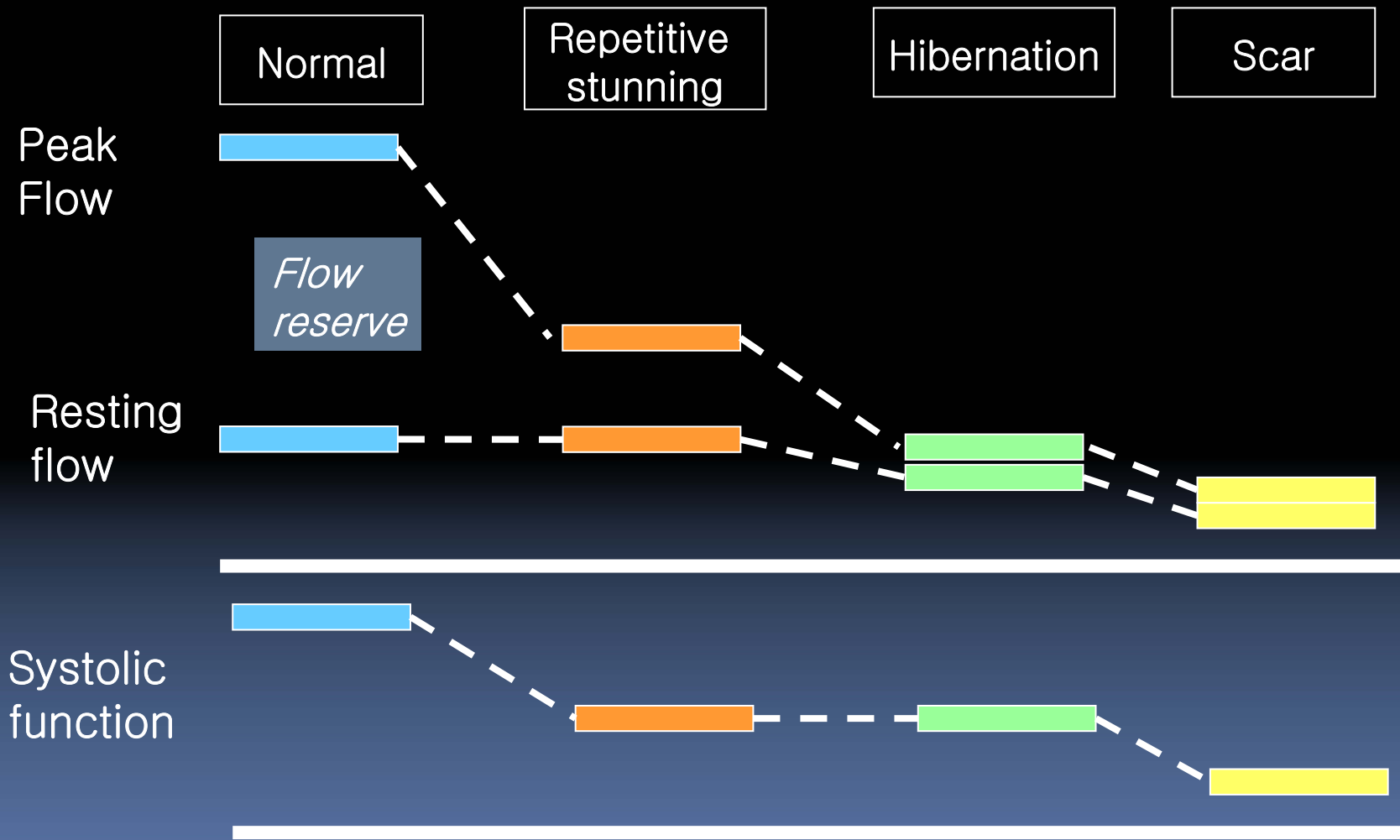
Author	Number of patients	LVEF	Viability technique	Incidence of viability
Auerbach <sup>77</sup>	283	26 (8%)	FDG/N13 ammonia PET	55%
Al-Mohammed <sup>78</sup>	27	19 (6%)	FDG/N13 ammonia PET	52%
Schinkel <sup>79</sup>	104	25 (7%)	FDG/Tc-99m TF SPECT	61%
Fox <sup>80</sup>	27	NA	Tc-99m MIBI/TF SPECT	37%

FDG, F18-fluorodeoxyglucose; LVEF, left ventricular ejection fraction; MIBI, sestamibi; PET, positron emission tomography; SPECT, single photon emission computed tomography; Tc-99m, technetium-99m; TF, tetrofosmin.

About 50%

Bax JJ, Heart 2004;90(Suppl V):v26–v33.

# Pathophysiology of ischemic LV dysfunction



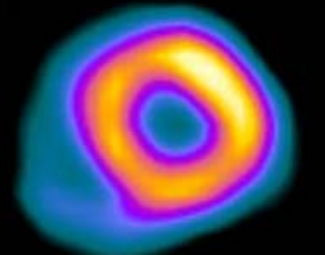
Pre-PCI

Post-PCI

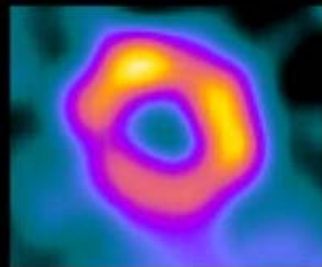
Rest



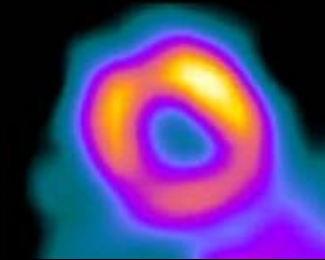
Stress



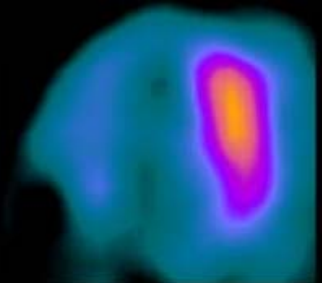
Delay



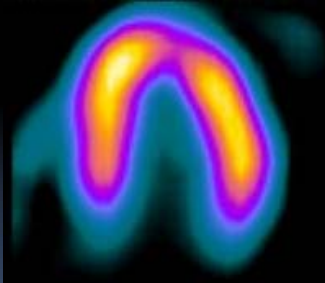
Rest



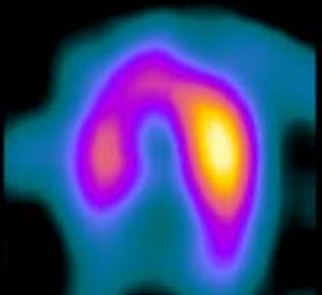
Rest



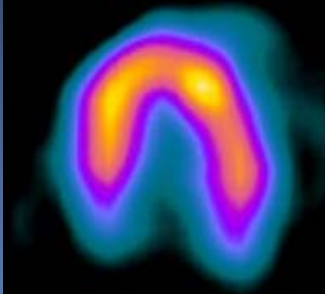
Stress



Delay



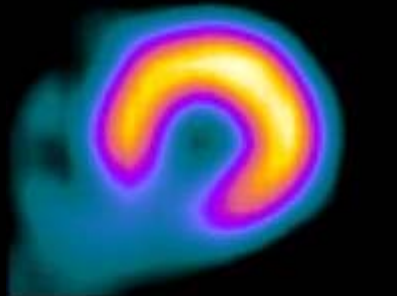
Rest



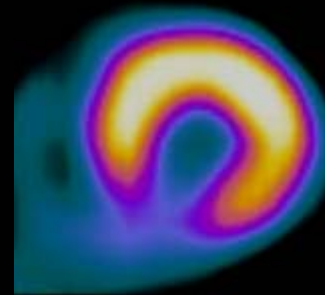
Pre-PCI

Post-PCI

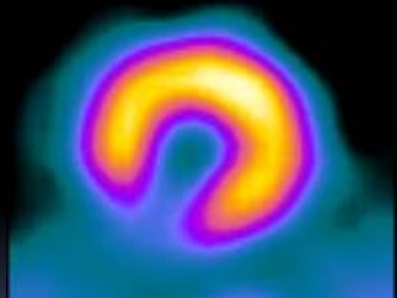
Stress



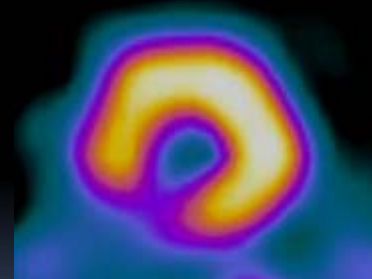
Stress



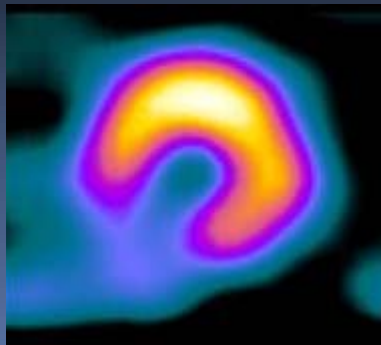
Rest



Rest



Delay



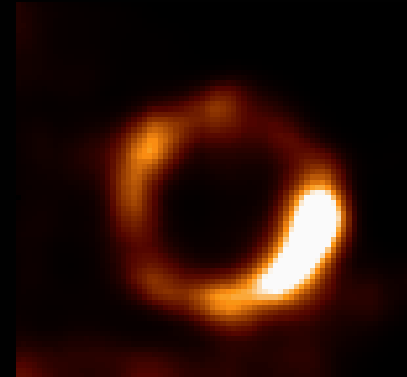
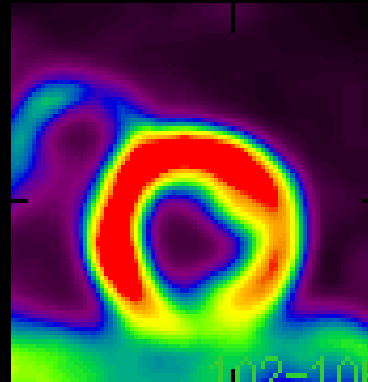
# PET patterns of myocardial viability assessment

	Rest perfusion	Stress perfusion	Rest FDG
Transmural MI	↓↓↓	↓↓↓	↓↓↓
Non-transmural MI	↓, ↓↓	No change or further decrease	↓, ↓↓
Hibernation	↓ or ↓↓↓	No change or further decrease	normal
Repetitive stunning	normal	Further decrease	normal

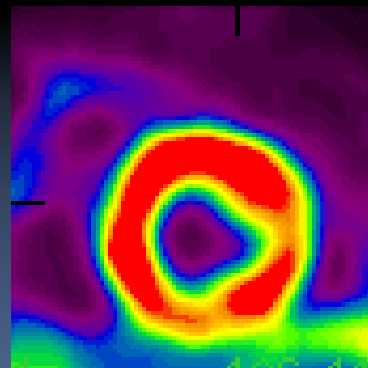
NH<sub>3</sub> PET

FDG PET

Stress



Rest





Pre-PCI

Post-PCI

Stress

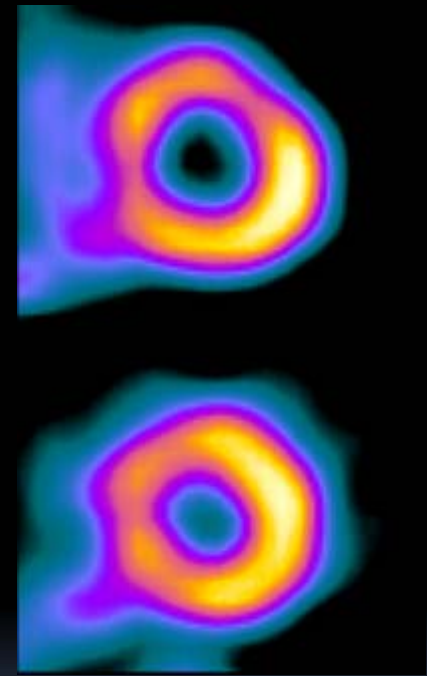
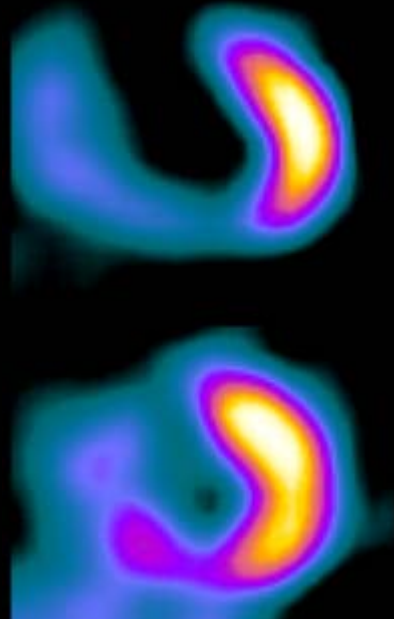
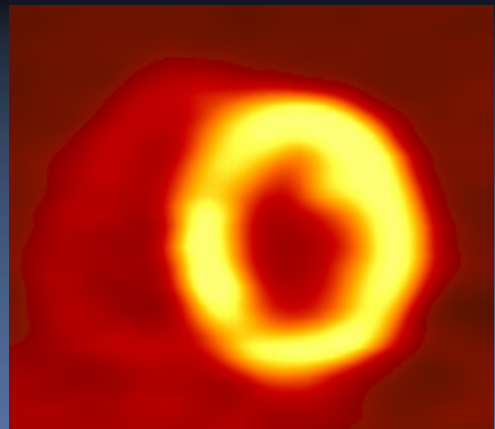
Stress

Rest

Rest



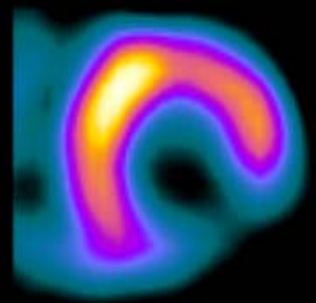
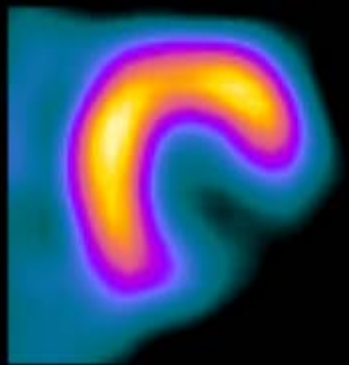
FDG



Pre-PCI

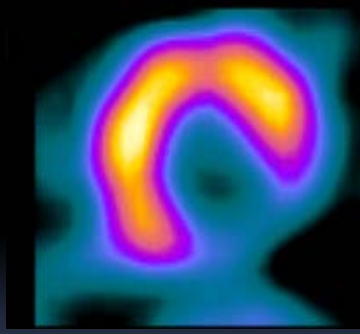
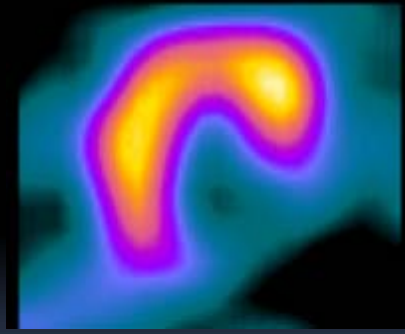
Post-PCI

Stress



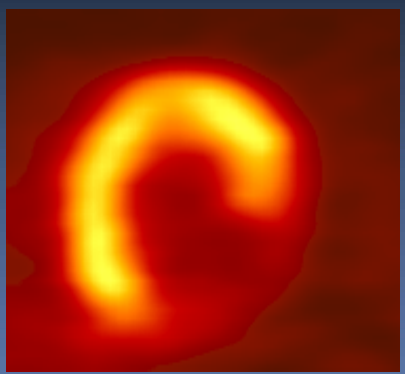
Stress

Rest



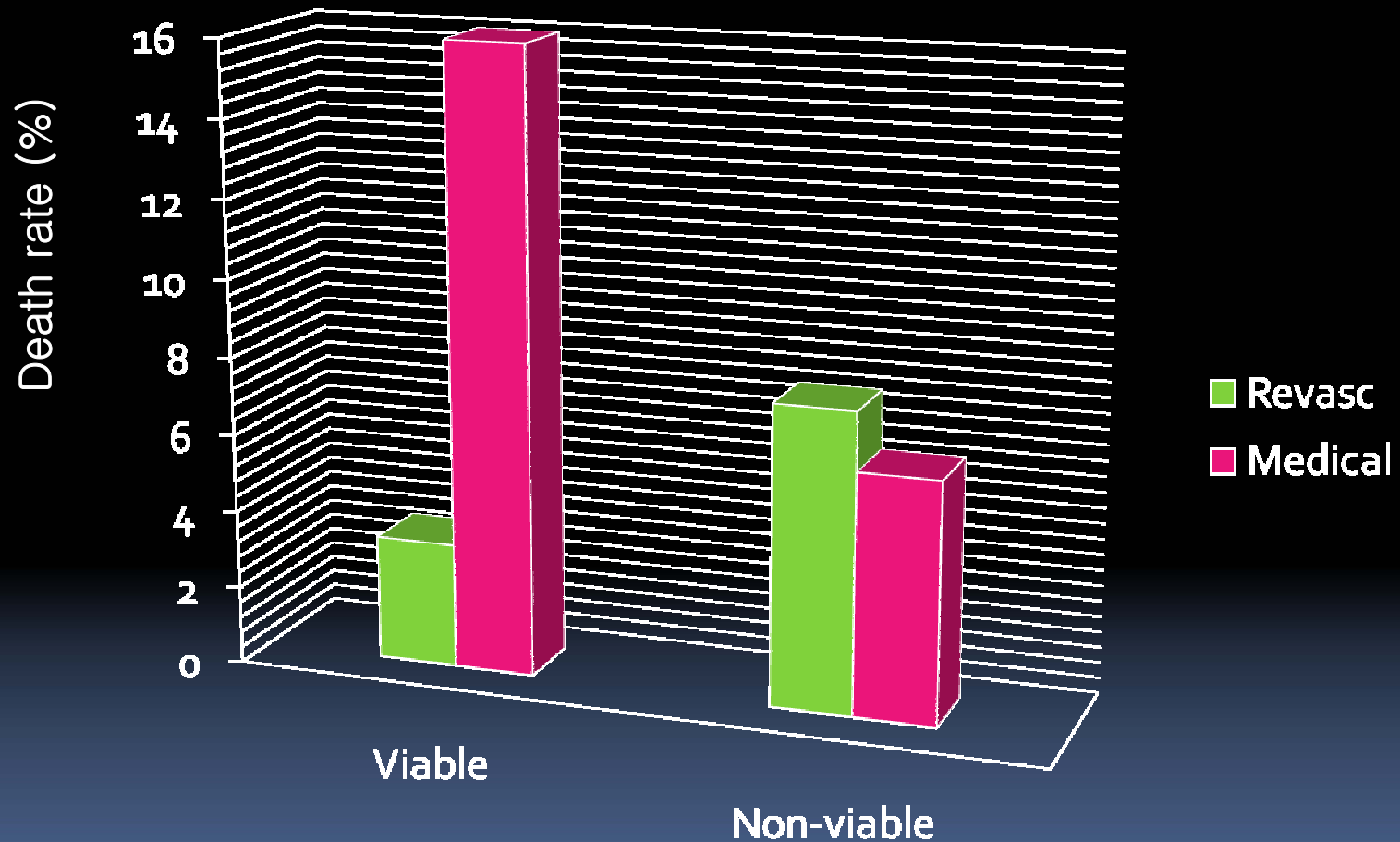
Rest

FDG



# Viability and outcome

Meta-analysis of 24 studies(3088 patients)



Allman KC, JACC 2002



# Following PCI

# MPI after PCI

- Symptom: unreliable index of restenosis
- MPI after PCI: 3-12 months after PCI
  - Sensitivity: 79-89%
- Prognostic value of MPI after PCI (Parisi et al, JACC)
  - Mortality
    - Reversible: 20%
    - Normal or persistent: 7%

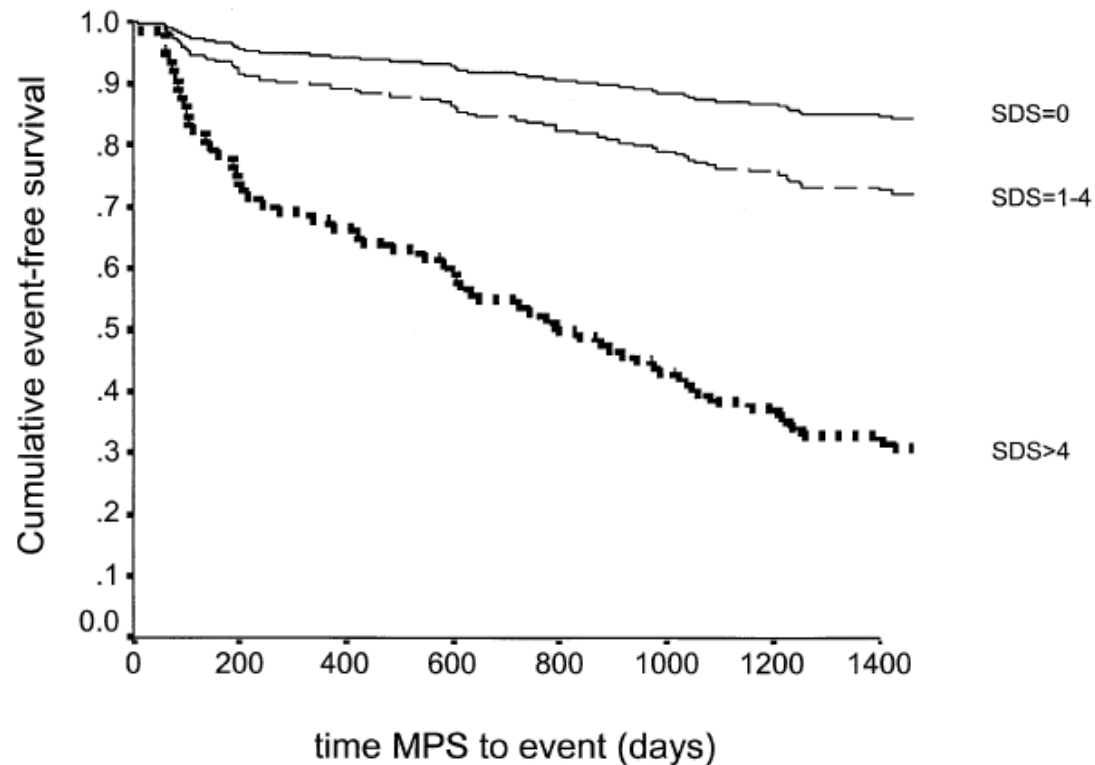
**Table 4.** Accuracy of Myocardial SPECT Imaging Following Percutaneous Coronary Intervention

Author (Ref.)	Year	No. of Patients	PCI Modality	% With Angina	Mean Time to SPECT	Mean Time to Angiogram	Sens %	Spec %	Acc %
Hecht et al. (15)	1991	116	PTCA	65	6 months	1 week	93	77	86
		41*		0			96*	75*	88*
		75†		100			91†	77†	85†
Marie et al. (37)	1993	62*	PTCA	0	6 months	3 days	94*	84*	87*
Milan et al. (54)‡	1996	37	PTCA	N/A	“late”	1 month	88	78	83
		20§					92§	67§	N/A
Kósa et al. (55)	1998	82 (99)	Stenting	N/A	7 months	1 month	79	78	79
		35 (52)§		N/A			100§	82§	85§
Milavetz et al. (56)	1998	33	Stenting	64	3 months	5 days	71	—	67
Caner et al. (57)#	1998	34 (37)	PTCA	N/A	2–48 months	1 month	76	79	78
		111 (138)					95¶	73¶	88¶
Beygui et al. (58)	2000	179 (208)*	PTCA	0	6 months	7 days	63*	77*	71*
		111 (138)					56§	81§	74§
Galassi et al. (59)	2000	97 (107)	Stenting	N/A	4 months	2 months	82**	84**	83**
		46 (56)§					76§	95§	89§
Overall performance of SPECT MPI							79††	79††	79††

Number of treated territories are indicated in parentheses; restenosis is defined as  $\geq 50\%$  diameter stenosis unless otherwise indicated; \*patients with “silent” ischemia; †patients with “symptomatic” ischemia; ‡all patients referred because of “equivocal” exercise stress tests, values reported are for qualitative analyses; §patients without prior infarction; ||calculations using  $\geq 50\%$  cross-sectional area narrowing definition; ¶calculations using  $\geq 70\%$  cross-sectional area narrowing definition; #dobutamine stress used for all patients; \*\*calculations based upon vascular territories; ††weighted average.

MPI = myocardial perfusion imaging; PCI = percutaneous coronary intervention; SPECT = single-photon emission computed tomography. Other abbreviations as in Tables 1 and 3.

## 6 month SPECT after PCI



**Figure 3.** Kaplan-Meier event-free survival curves by summed difference score (SDS) categories. Patients with an SDS of 0 ( $n = 242$ ) had significantly lower event rates than patients with an SDS of 1 to 4 ( $n = 49$ ;  $p = 0.03$ ); these patients had significantly lower event rates than patients with  $SDS > 4$  ( $n = 16$ ;  $p = 0.005$ ). Note the significant differences in outcome between patients with no, mild, and moderate to severe ischemia, as defined by SDS. MPS = myocardial perfusion SPECT.

Zellweger MJ et al. 2003 JACC

Residual ischemia (SDS) – prognostic value following PCI



# Conclusion

- Fusion software
  - Detecting ischemia
  - Culprit vessel
  - Restenosis
- Viability
  - Perfusion-metabolism mismatch

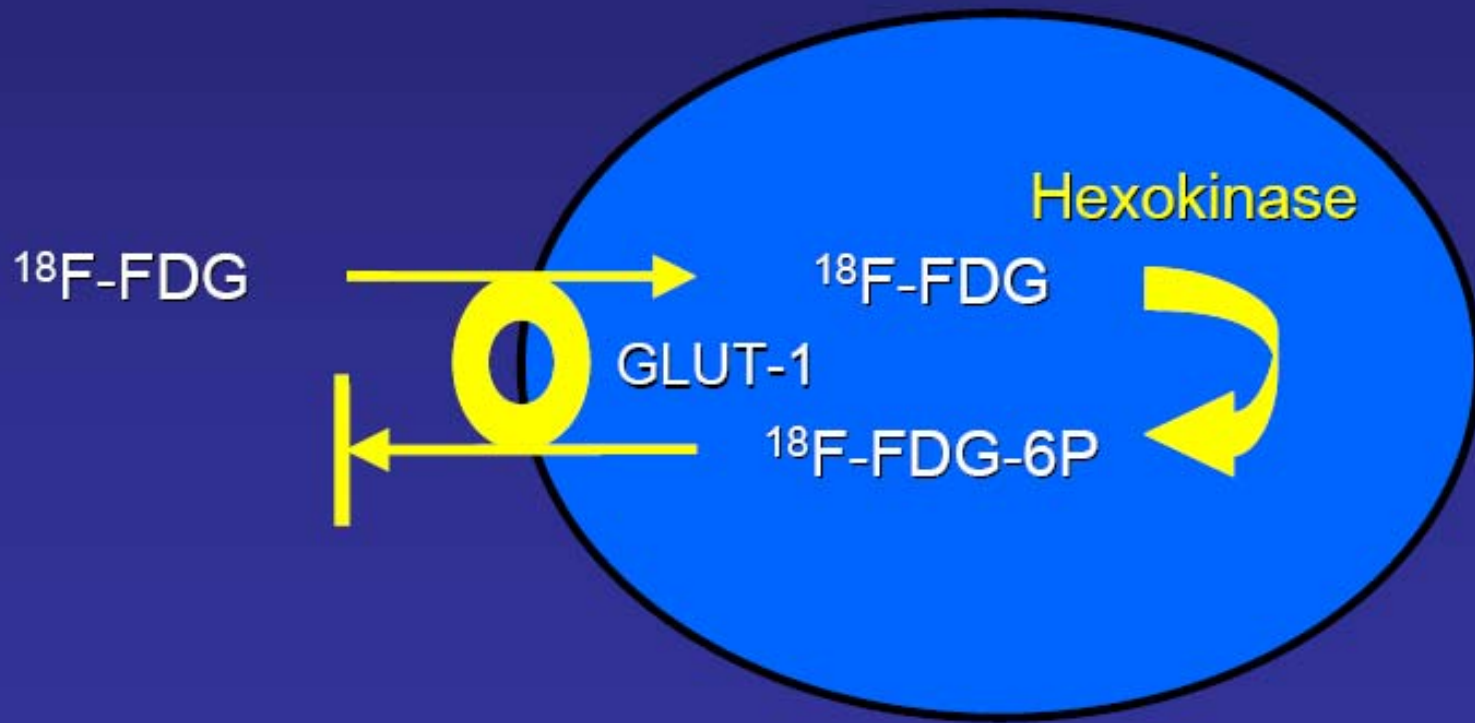


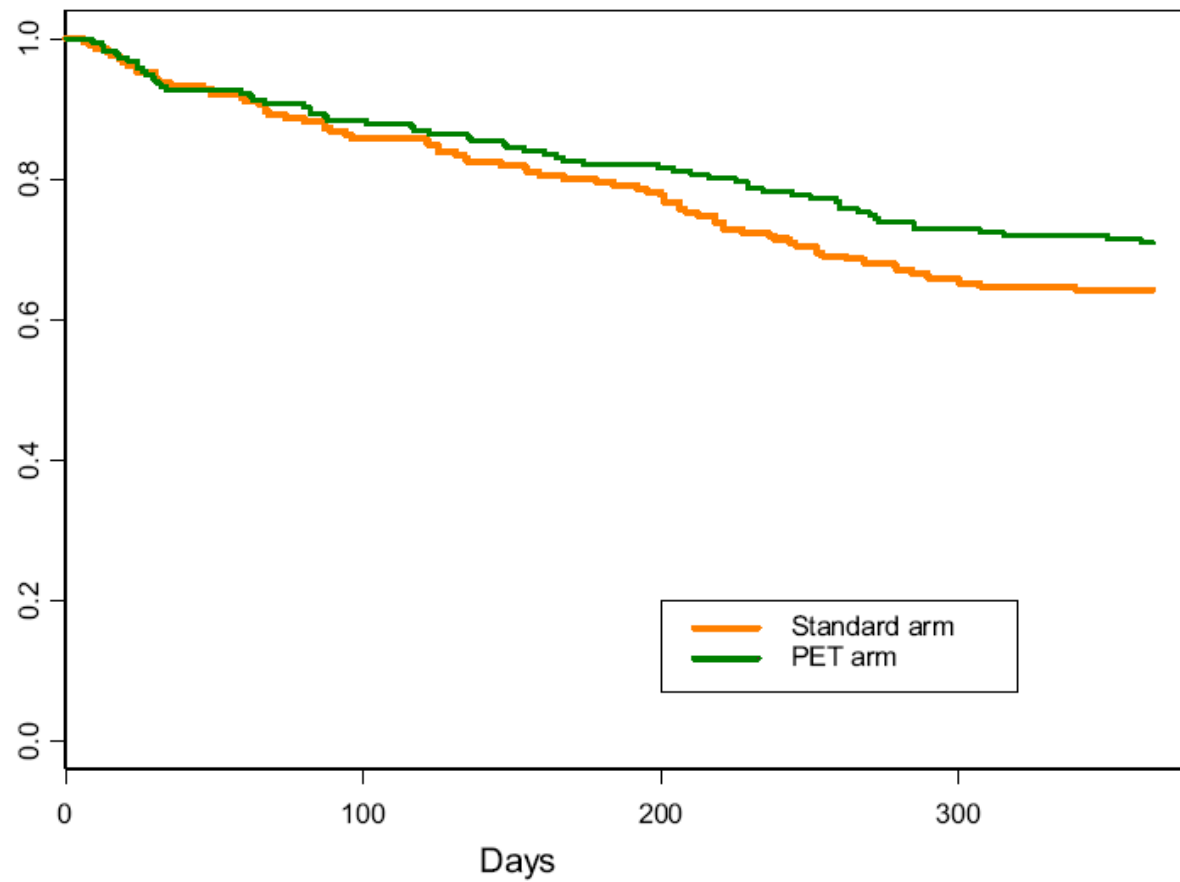
- 
- Thank you for your attention!

# Advantages of nuclear imaging

- Myocardial perfusion (stress/rest)
  - Quantitative analysis
  - Accurate
  - Reproducible
- Myocardial metabolism

# FDG





**Figure 3**

**“Survival Curves” (on the Basis of Time to First Occurring Outcome of the Composite Event)**

# Summary in following PCI

- < 3 months:
  - high false-positive
  - Limited role
- 3-6 months
  - Normal MPI exclude restenosis
  - Abnormal MPI → angiography
- > 6 months
  - Accurate to diagnose restenosis

# Why viability study?

- Morbidity and mortality of revascularization with low EF remain high
  - Need to determine risk vs. benefit
- Revascularization helps low EF with minimal or no angina?

# End points for clinical effectiveness

- Improvement of regional LV dysfunction
- Improvement of global LV dysfunction (LVEF)
- Improvement of heart failure symptoms
- Improvement of survival

# Characteristics of viable myocardium

**Table 2** Characteristics of dysfunctional but viable myocardium

Characteristic	Imaging modality	Markers of viability
Perfusion/intact cell membrane	Thallium-201 SPECT	Tracer activity >50%
Perfusion/intact mitochondria	Technetium-99m TF/MIBI SPECT	Redistribution >10% Tracer activity >50%
Glucose metabolism	FDG imaging (PET or SPECT)	Improved tracer uptake after nitrates Tracer activity >50% Preserved perfusion/FDG uptake Perfusion-metabolism mismatch
Free fatty acid metabolism	BMIPP SPECT	Tracer activity >50%
Contractile reserve	Dobutamine echo/MRI Dobutamine gated SPECT	Perfusion-BMIPP mismatch Improved contraction Infusion of low dose dobutamine

BMIPP,  $\beta$  methyliodophenyl pentadecanoic acid; FDG, F18-fluorodeoxyglucose; MIBI, sestamibi; MRI, magnetic resonance imaging; PET, positron emission tomography; SPECT, single photon emission computed tomography; TF, tetrofosmin.

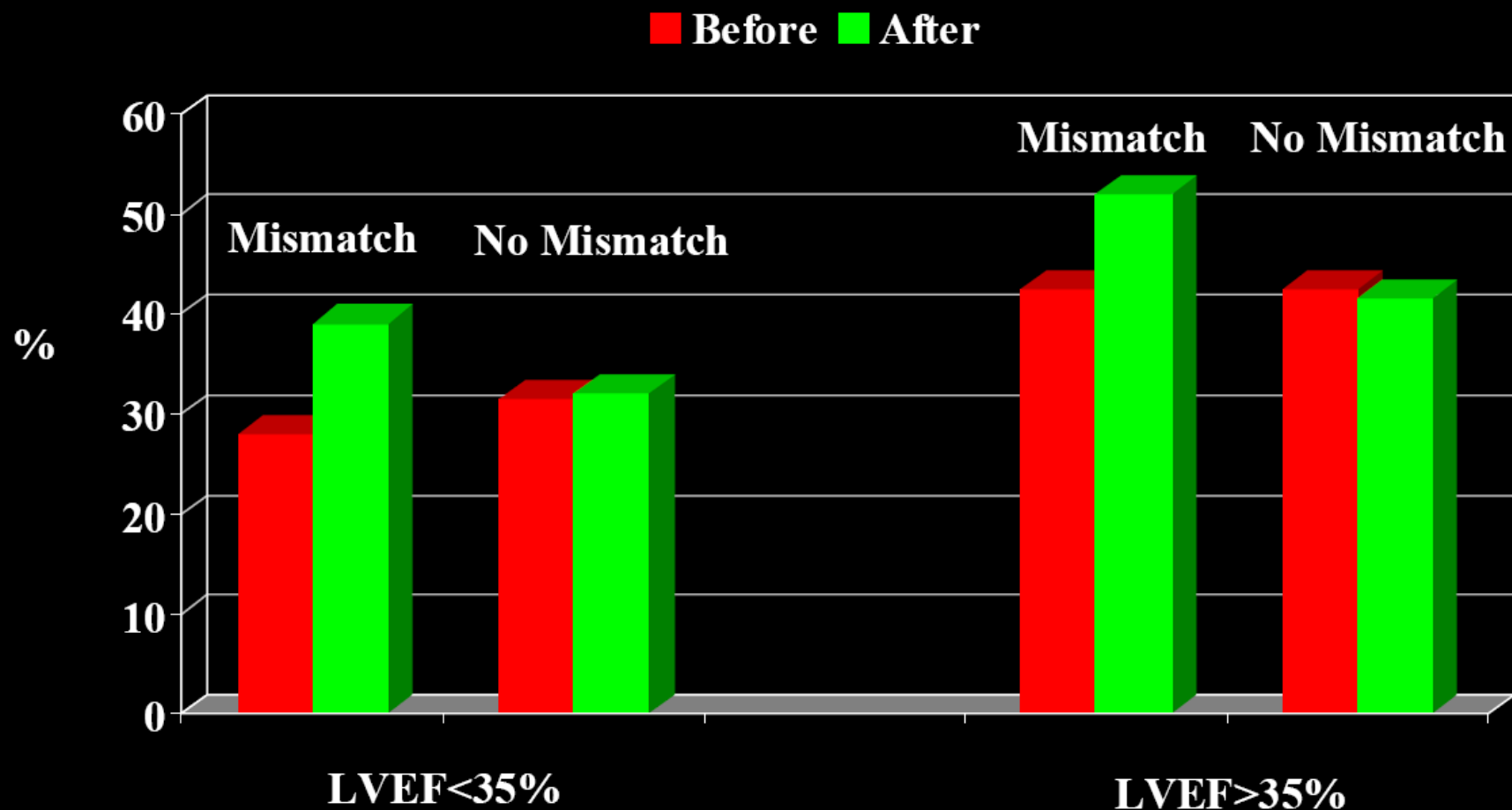


# Rest-redistribution Tl-201

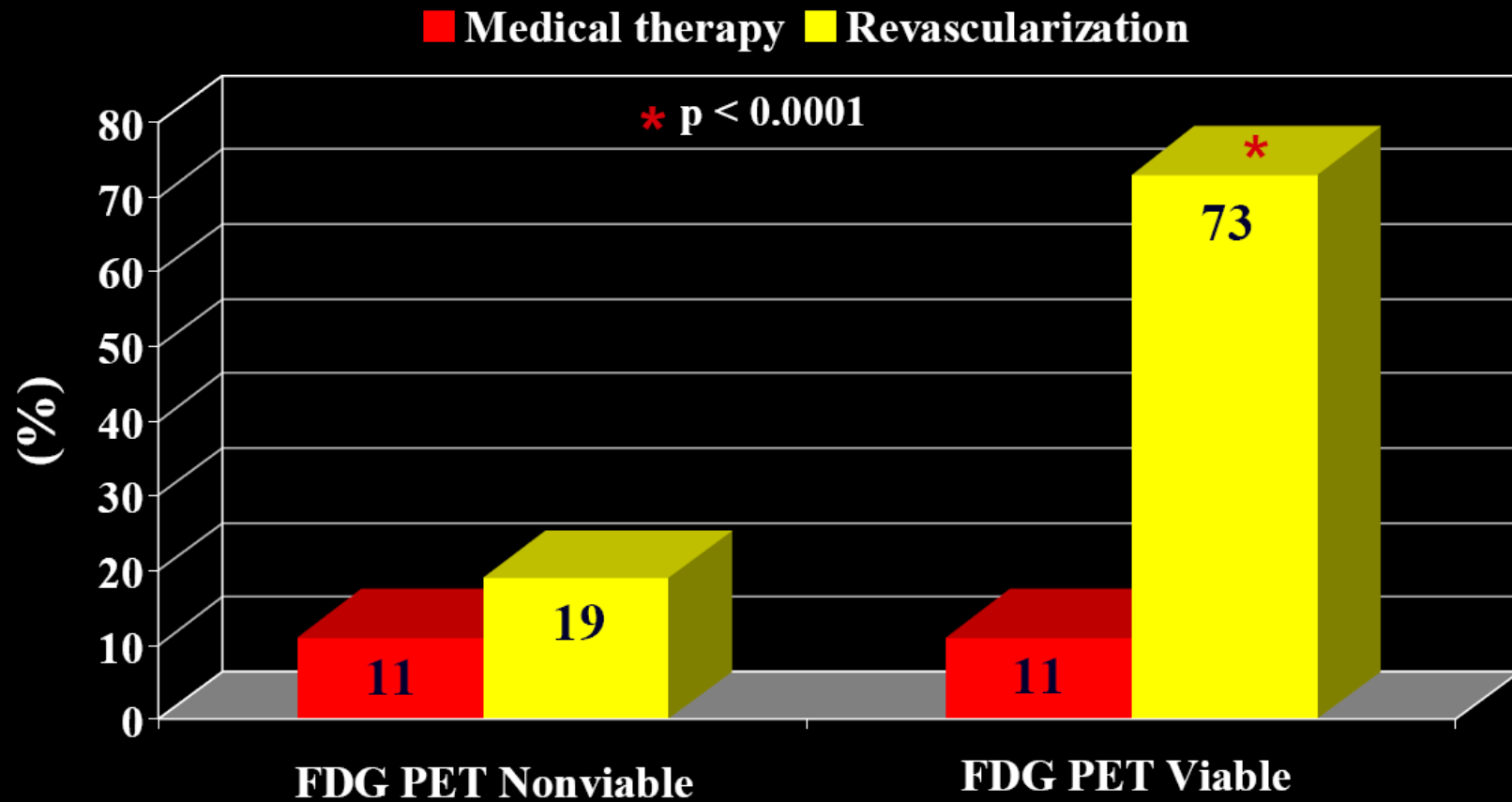


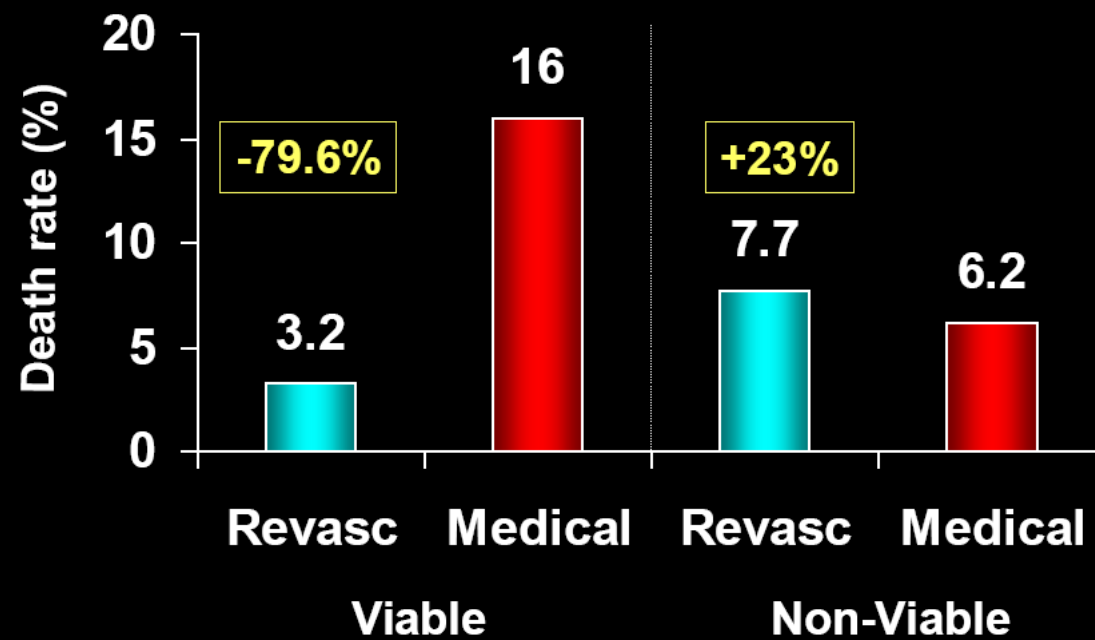
20% enhanced detection of viability by late imaging

# PET prediction of post-op LVEF improvement: 25 studies, 772 patients



# Improvement of Heart Failure Symptoms by treatment





Allman et al

# After PCI

- Restenosis
  - 1/3 of patients
  - 1/2 of restenosis: asymptomatic
  - 45% of chest pain: no stenosis
- Disease progression
  - 7% per year
- Annual risk of a major adverse cardiac events following PCI: 5-7%



3D Volume 1  
Ex: 801189503  
Se: 5 +c  
Volume Rendering No cut

RSP

Severance Hosp.  
KANG SIN YUNG  
M 69 5356237  
DoB: May 18 1938  
Ex: Jan 28 2008

DFOV 18.8cm  
B25f

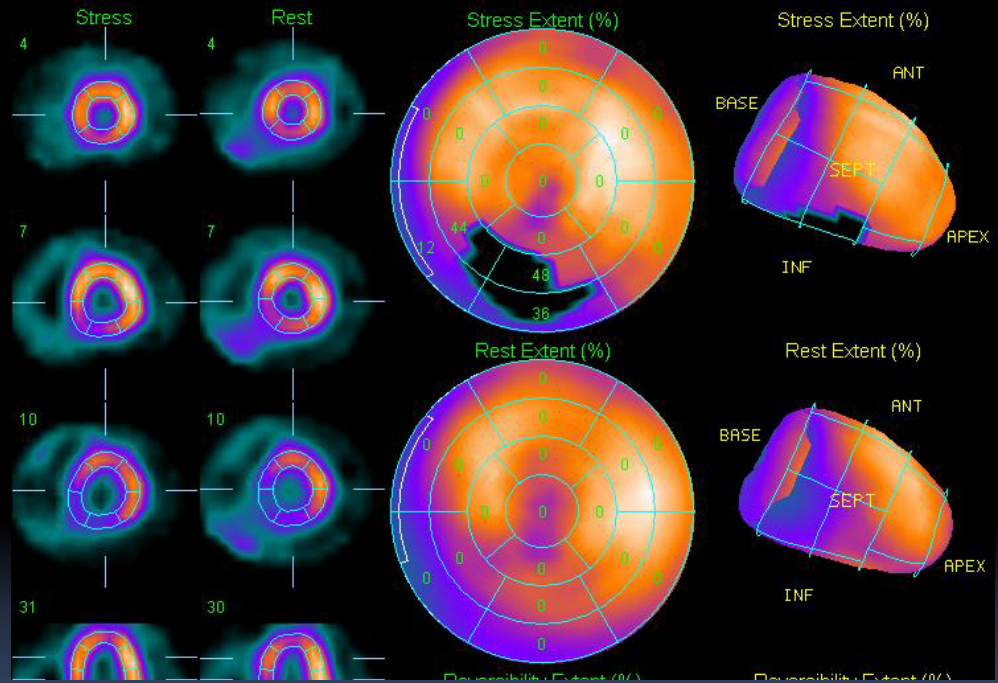


-16 L 169 RAO 55 CAU

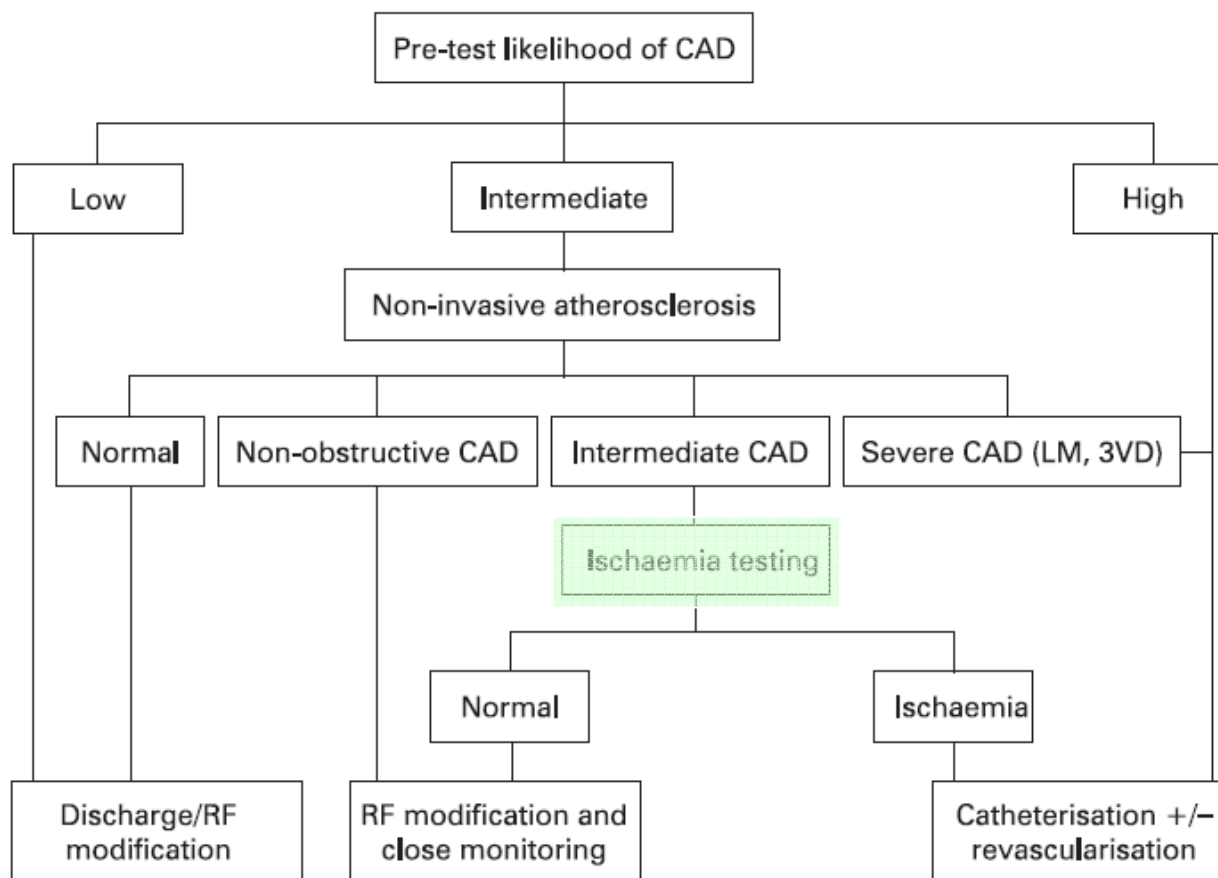
No VOI  
kv 120  
mA 261  
0.2s  
0.8mm /0.5sp  
Tilt: 0.0  
07:57:35 PM  
W = 4095 L = 2048

LIA

-999999.000000



RCA proximal



**Figure 3** Potential algorithm incorporating non-invasive anatomical and functional testing depending on pretest likelihood of CAD. In patients with low pretest likelihood, no imaging is performed. In patients with high pretest likelihood, referral for invasive coronary angiography is usually performed. In patients with an intermediate pretest likelihood, anatomy may be evaluated first, followed by functional testing or invasive coronary angiography based on the findings. CAD, coronary artery disease, LM, left main disease, 3VD, 3-vessel disease.