

Differential imaging: which imaging when?

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Lifescience

4 major imaging techniques:

- **Echocardiography**
- **Magnetic resonance imaging**
- **Multi-slice CT**
- **Nuclear imaging (PET and SPECT)**
- **Can provide all anatomical and functional information, but use should be clinically driven**

Based on the clinical presentation:

Ask yourself questions:

What information do I need to

- diagnose**

- treat**

this patient

Diagnosis is important

But the imaging results need
to have impact on choice of
therapy

Man 41 years old

Outpatient clinics:

No symptoms

Risk factors for CAD:

***Brother SCD age 43**

Asymptomatic individual,
low risk for atherosclerosis

The question is:

Risk stratification – early detection

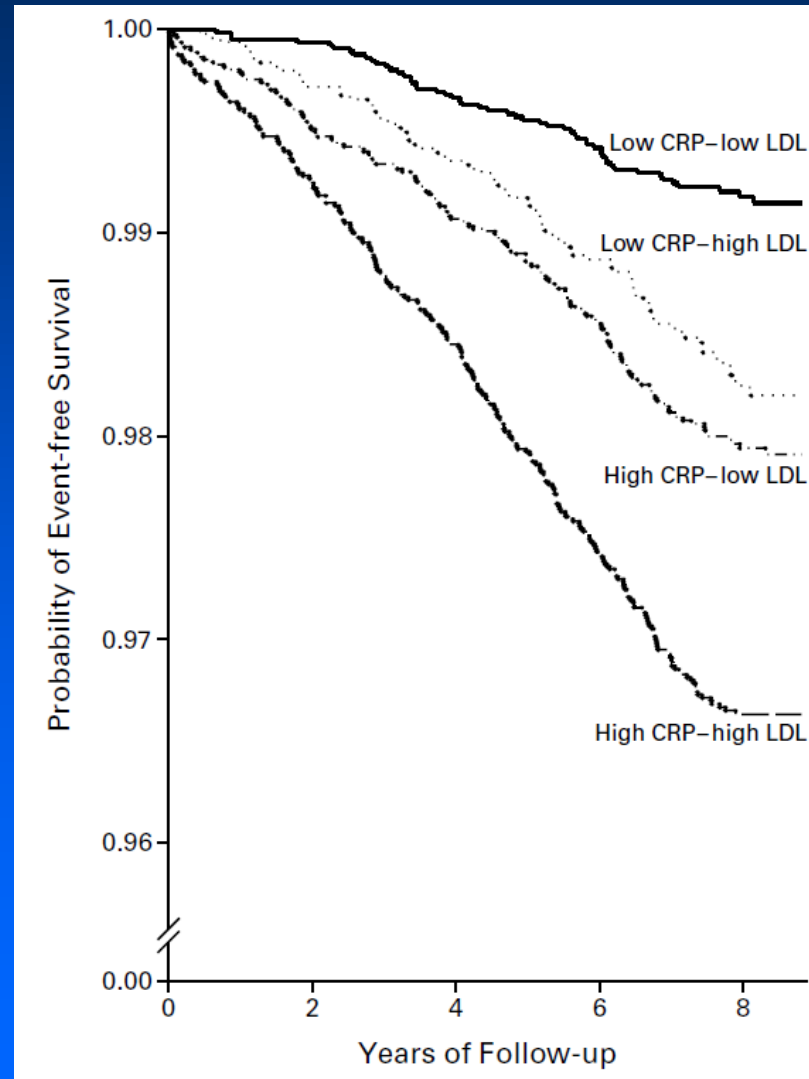
Blood:
biomarkers

Early
detection
of CVD

Large arteries:
Global: atherosclerosis

Coronary arteries:
Focal: lesion characteristics

Cardiovascular event-free survival, according to CRP and LDL



Blood:
biomarkers

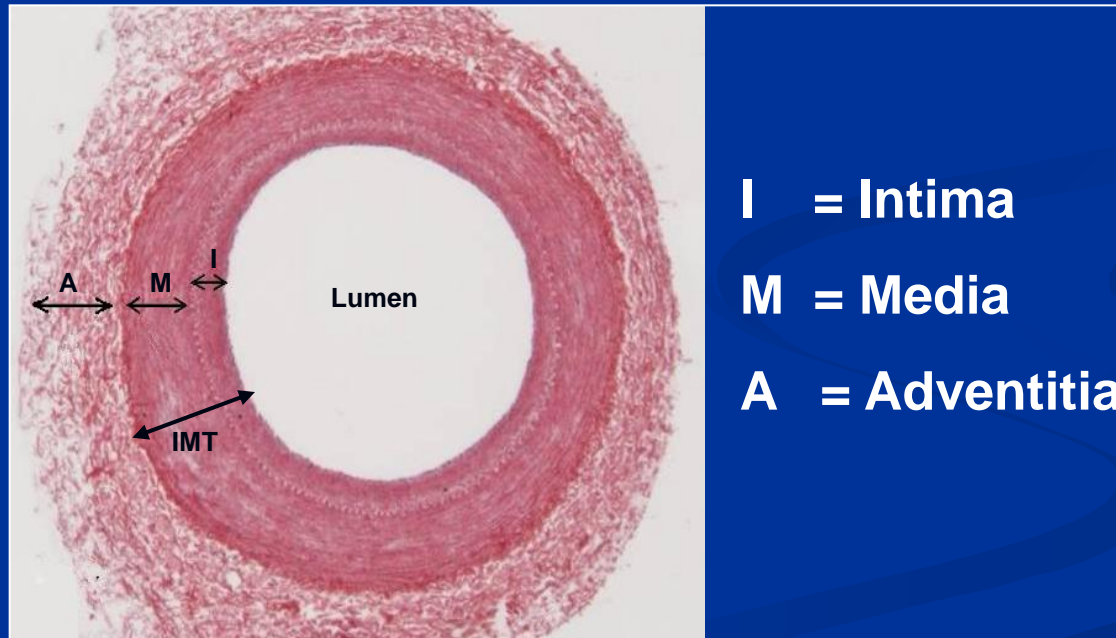
Early
detection
of CVD

Large arteries:
Global: atherosclerosis

Coronary arteries:
Focal: lesion characteristics

Carotid Intima Media Thickness (CIMT)

Tissue between luminal edge of the artery
and the boundary between
media and adventitia



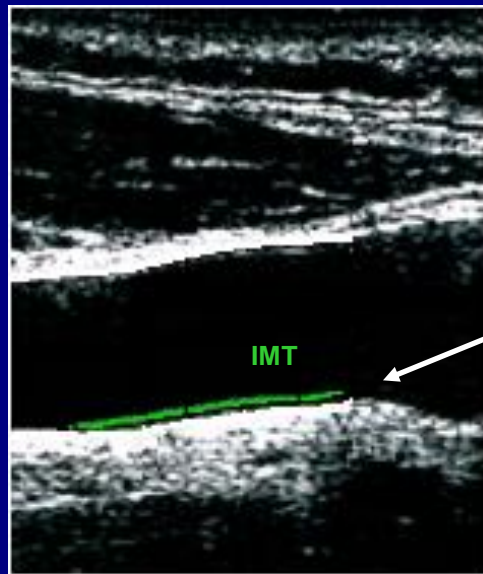
Assessment of CIMT

Semi-automatic B-mode ultrasound measurements

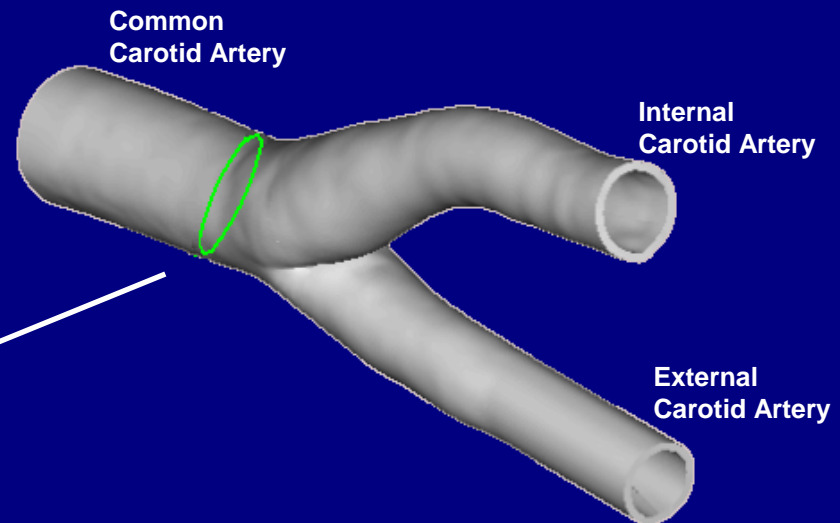
Left and right common carotid artery, directly proximal to the bifurcation

Mean CIMT measurements at four angles

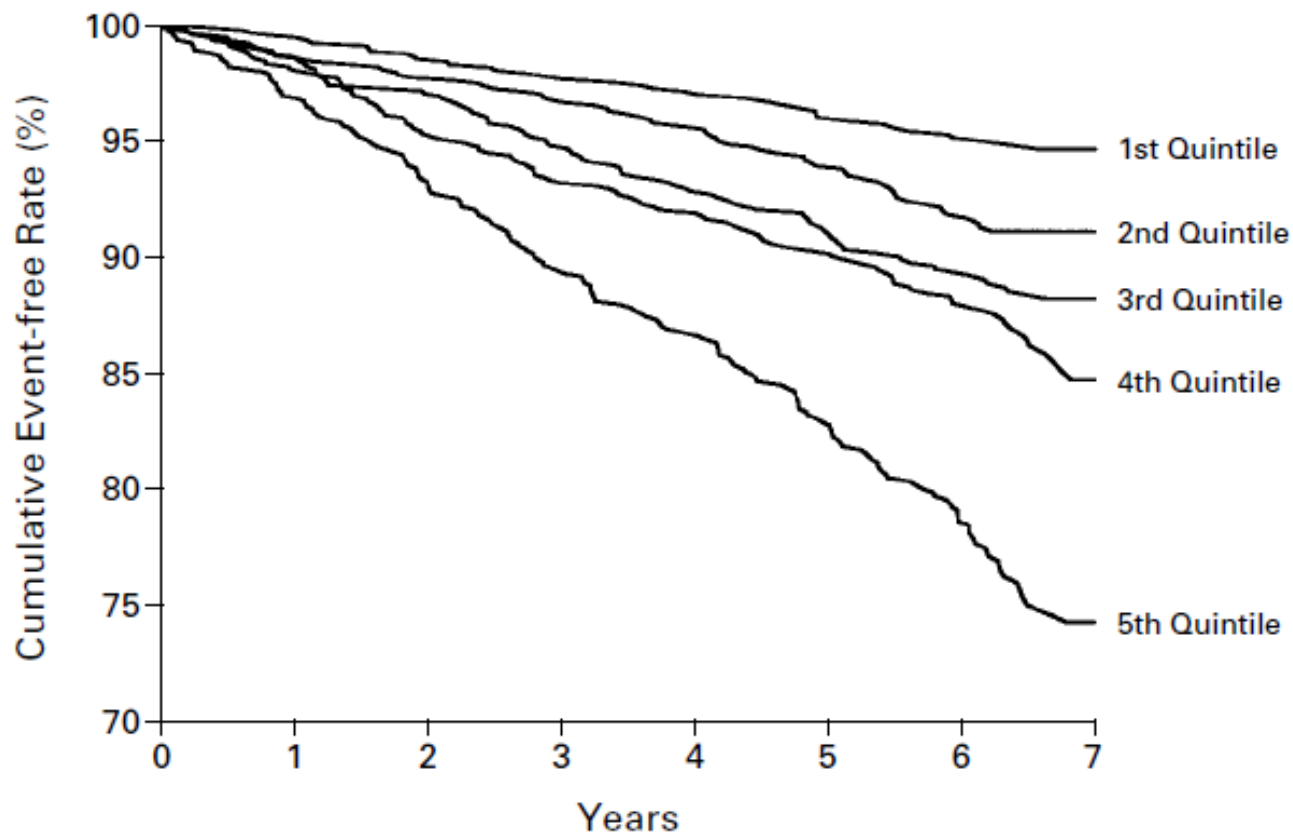
Calculation of the average of 8 mean CIMT per patient



Ultrasound measurement of CIMT



Cumulative event free rate (stroke or MI) according to IMT quintiles



Blood:
biomarkers

Early
detection
of CVD

Large arteries:
Global: atherosclerosis

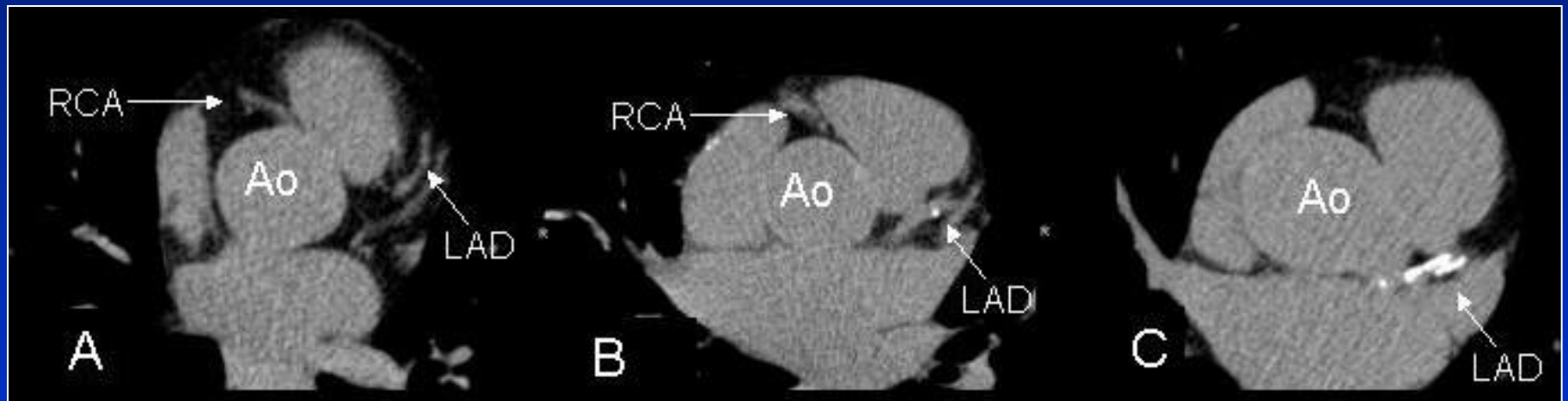
Coronary arteries:
Focal: lesion characteristics

Calcium Scoring (EBCT/MSCT)

No
calcification

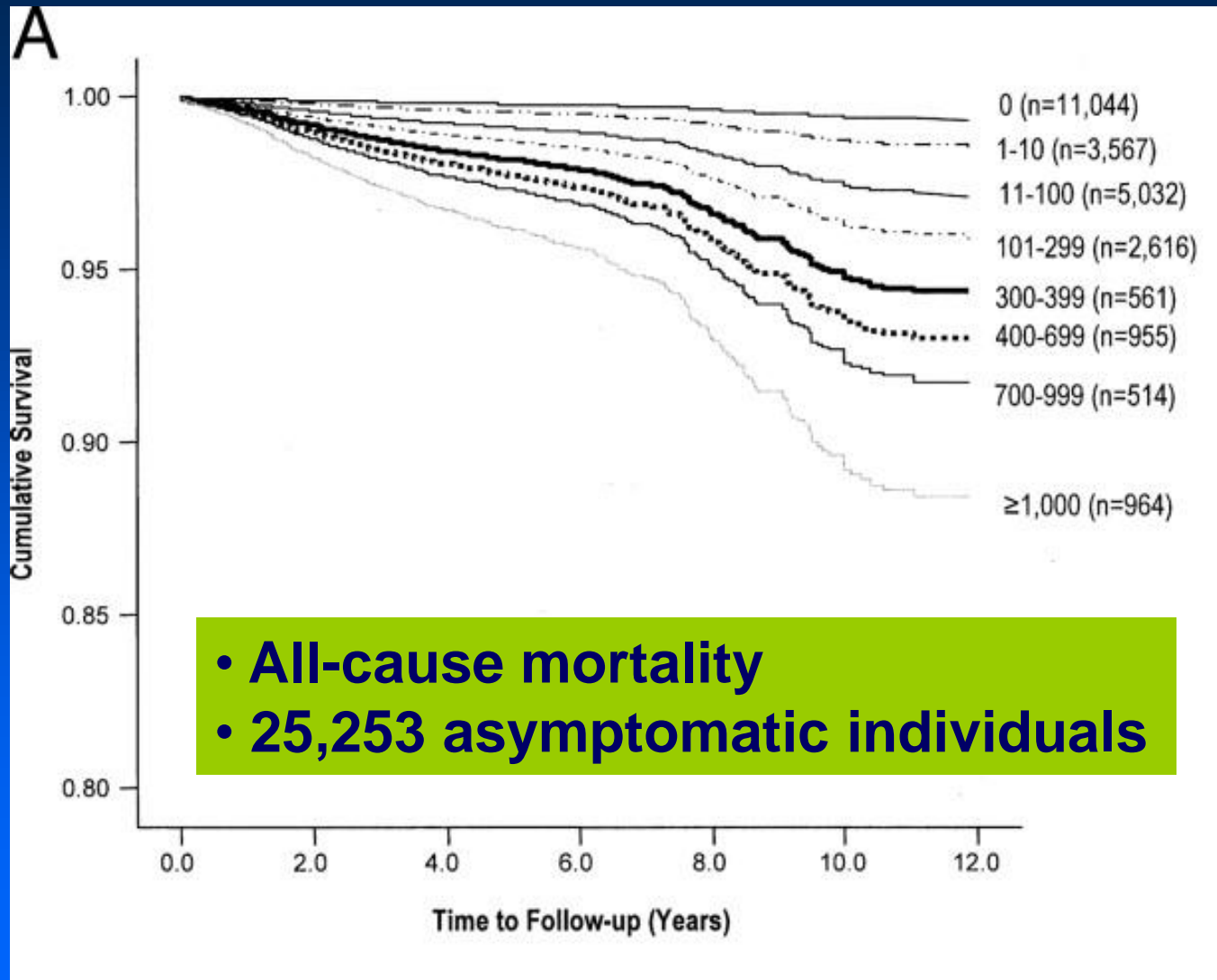
Moderate
calcification

Extensive
calcification



**Coronary calcifications provide a marker for
atherosclerotic disease burden**

Calcium score vs risk stratification



Man 54 years old

**We have screened some years ago:
nothing; now the symptoms change and
developed 1 RF**

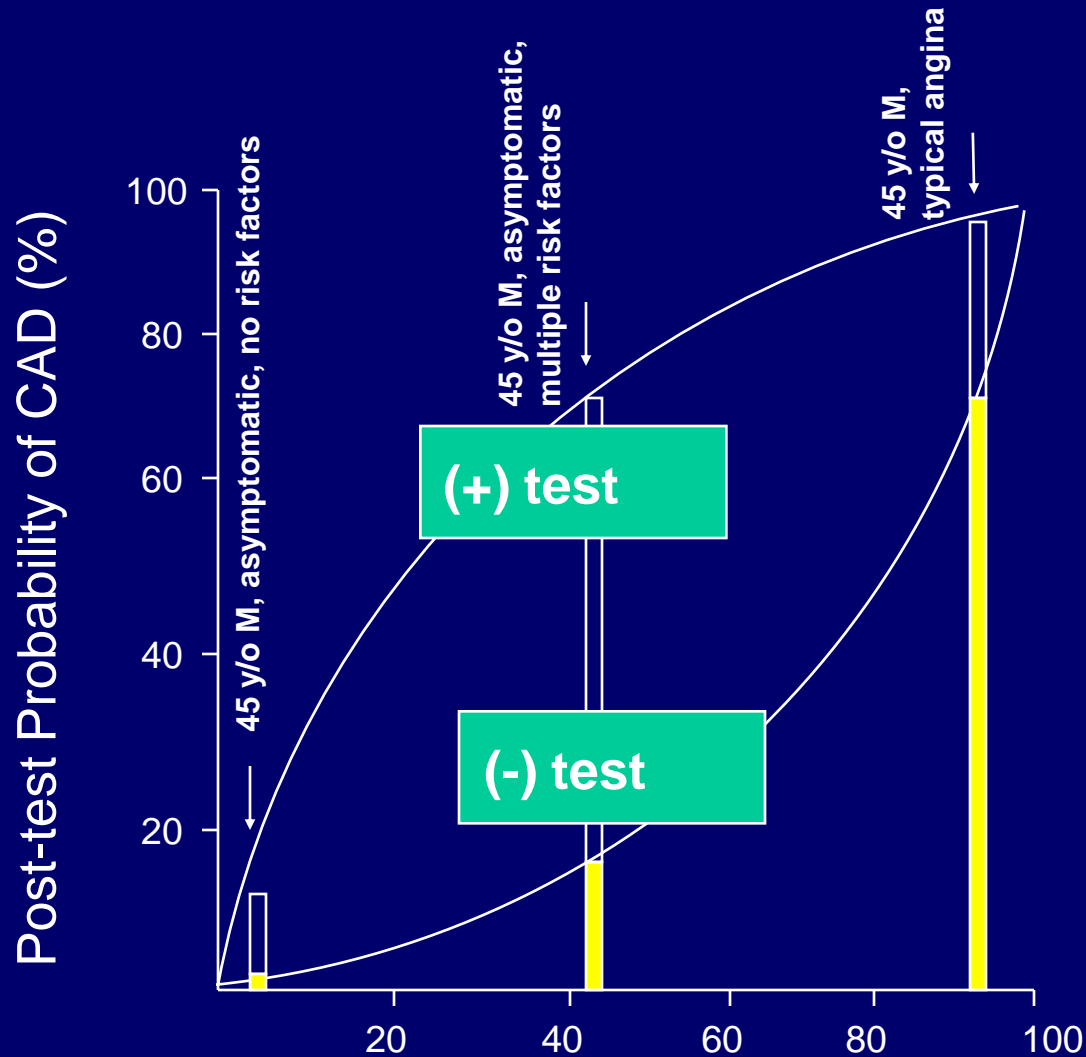
Outpatient clinics:

Dyspnea or atypical chest pain at exercise

Risk factors for CAD:

***Dyslipidemia**

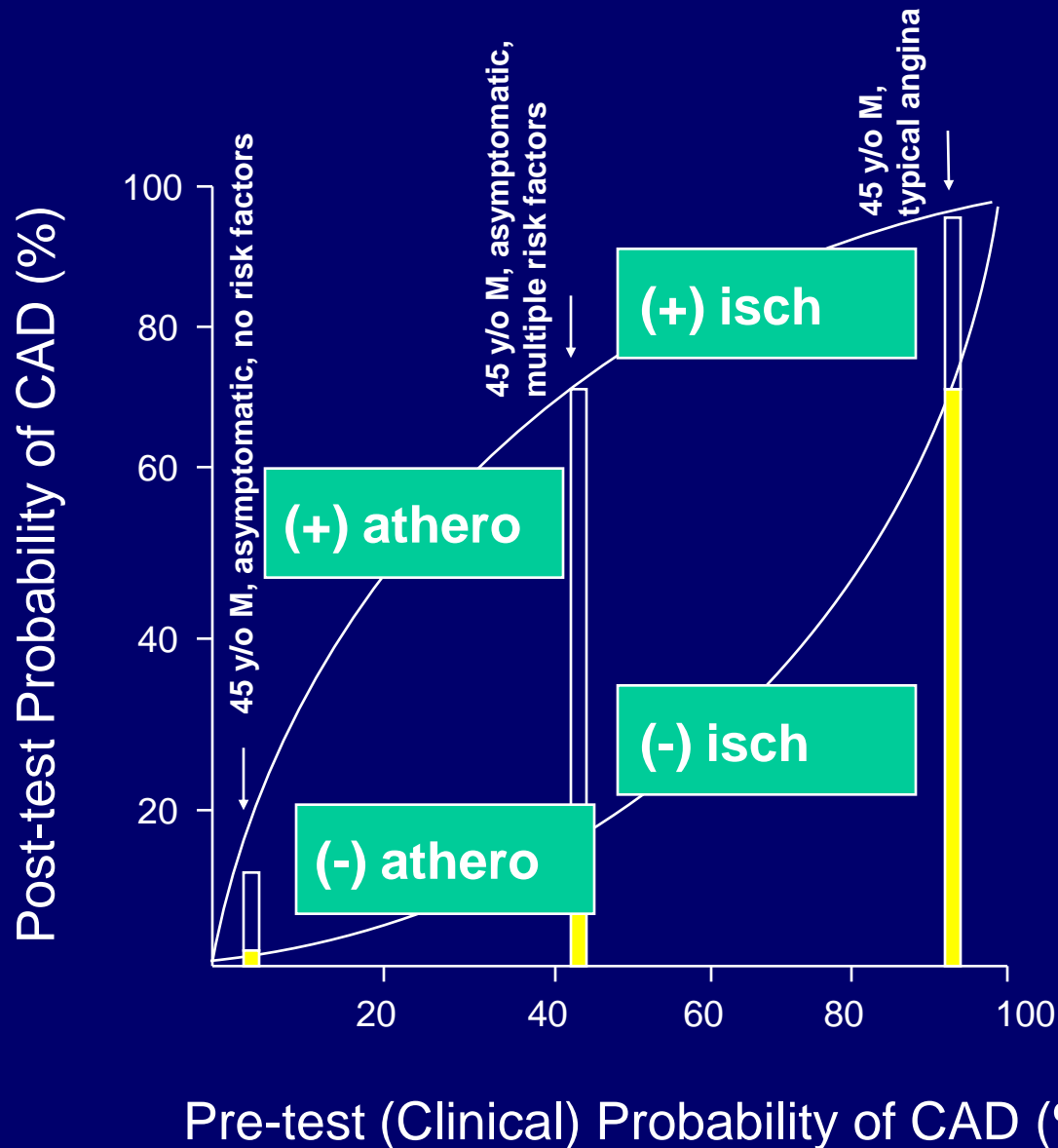
Non-invasive assessment of CAD: Which Patients?



Pre-test (Clinical) Probability of CAD (%)

Patterson et al. JACC 1989

Non-invasive assessment of CAD: Which Patients?



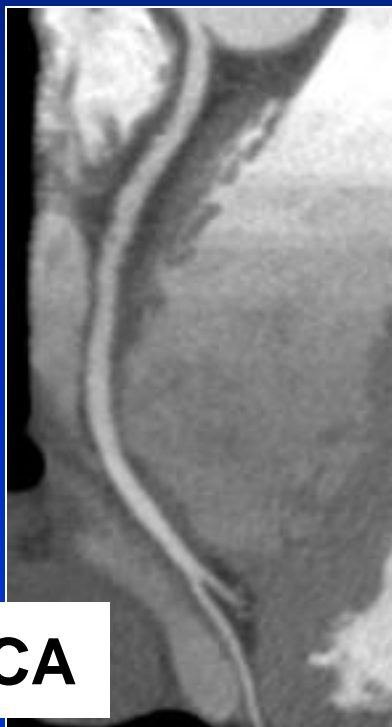
Symptomatic patient, low-intermed risk

The question is:

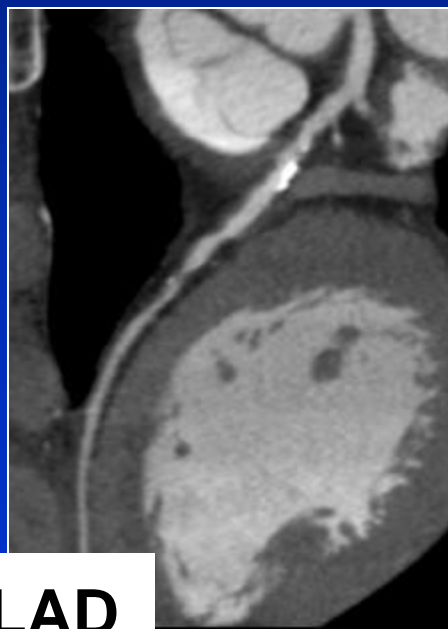
Atherosclerosis? (medical therapy needed and follow-up or discharge?)

☞ We order a non-invasive anatomical test to detect /exclude atherosclerosis

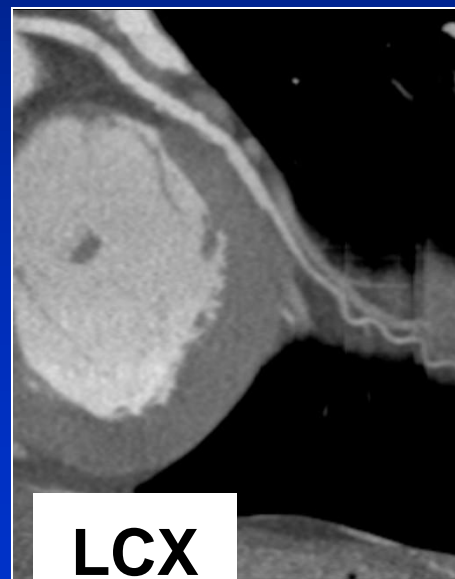
curved MPR



RCA



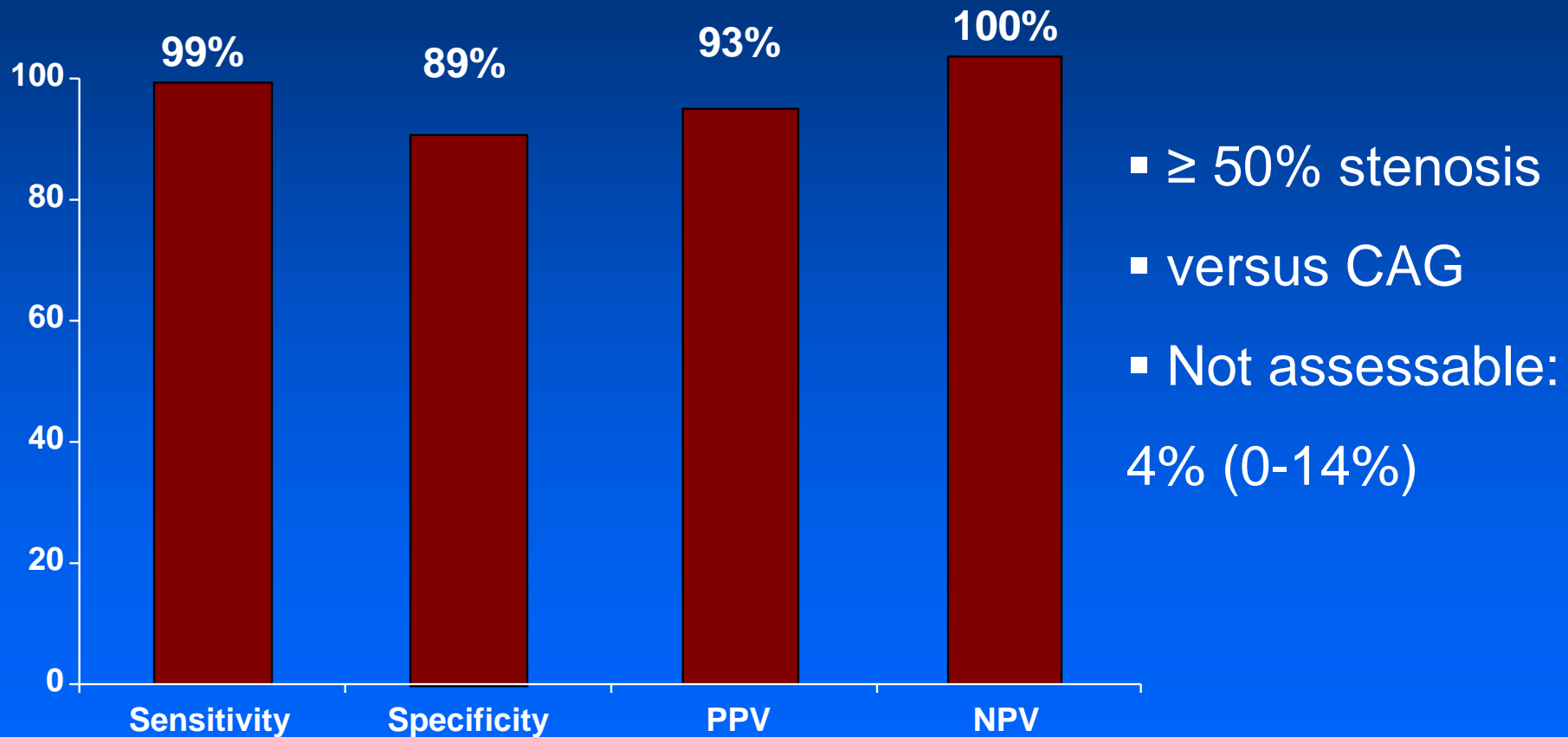
LAD



LCX

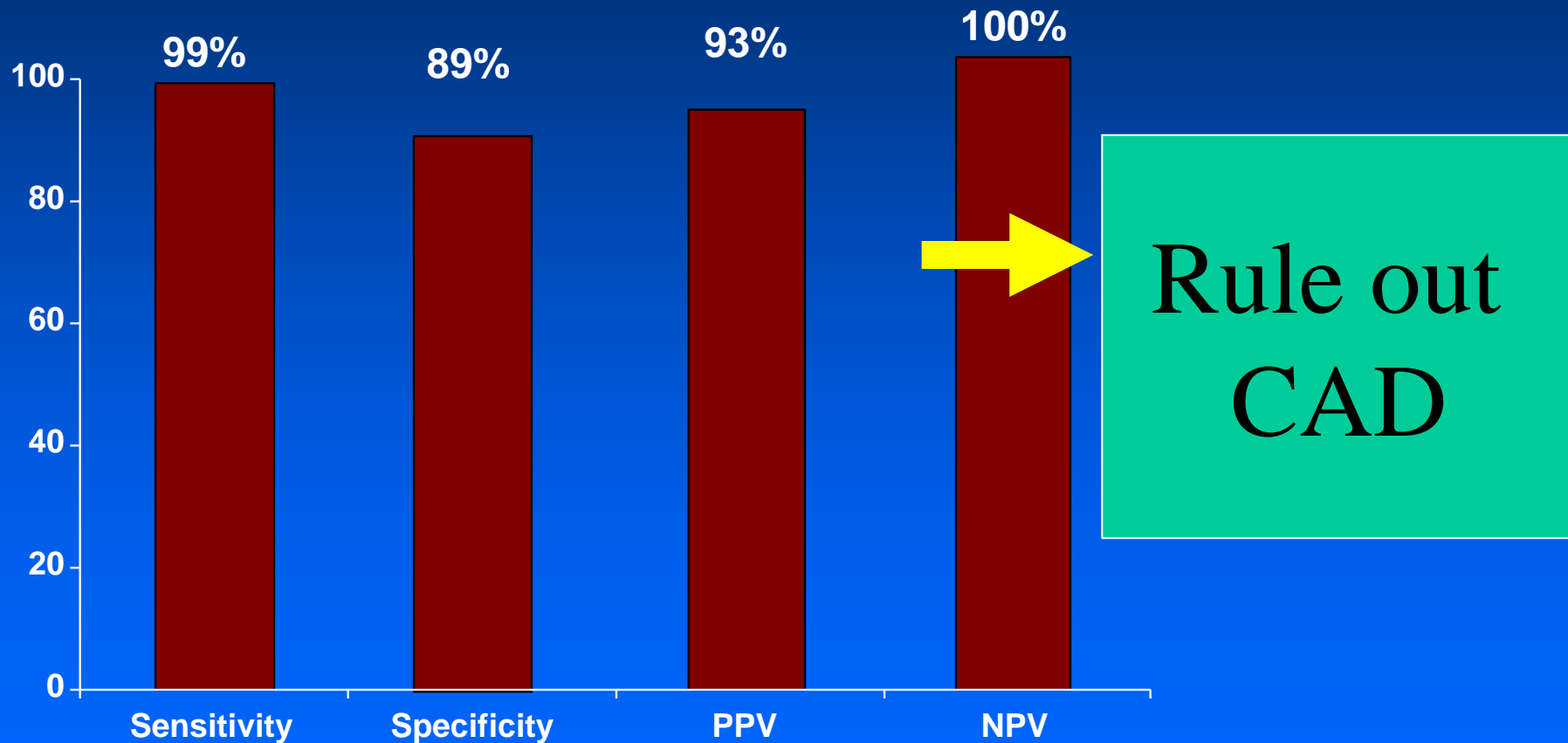
Meta-analysis 64-slice CT

Patient-based detection (n=1286)

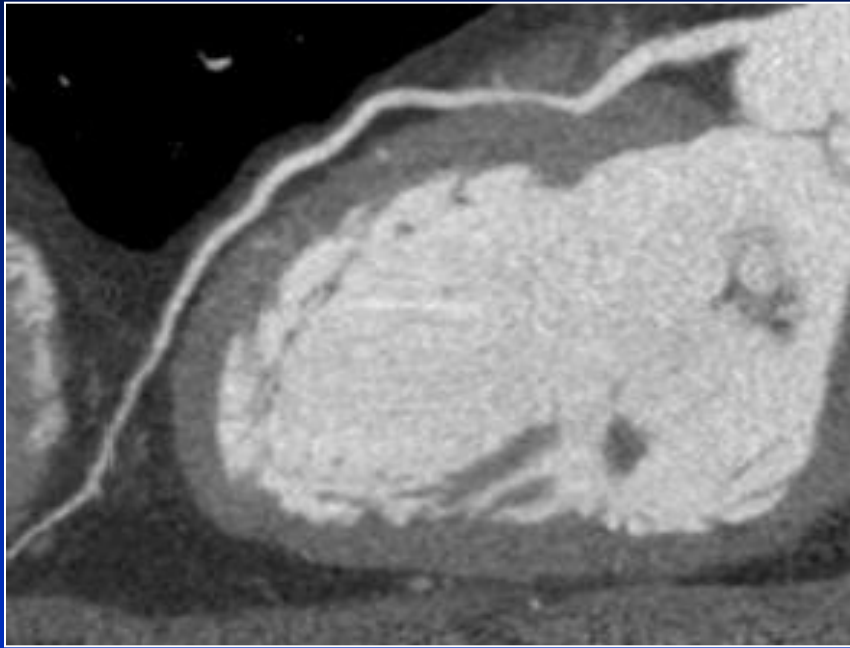


Meta-analysis 64-slice CT

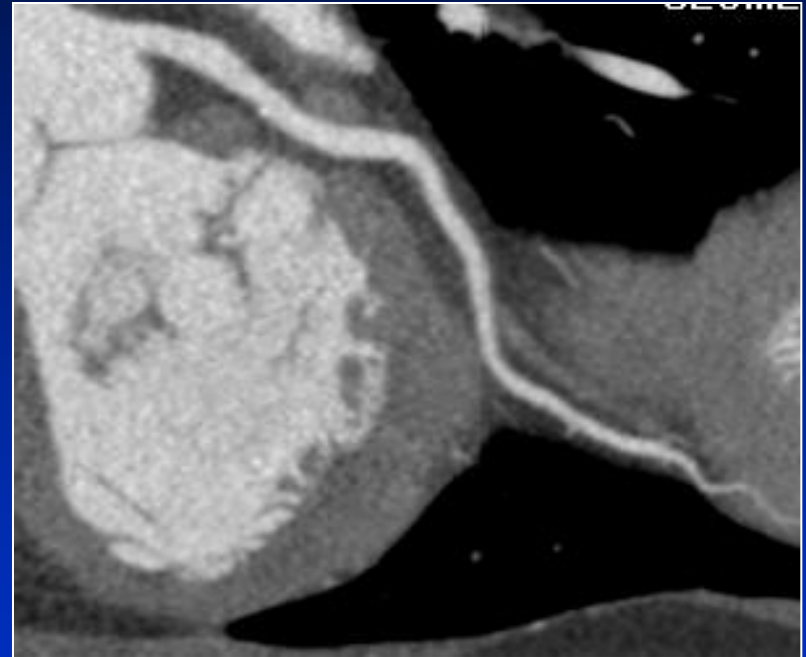
Patient-based detection (n=1286)



Non-invasive angiography - MSCT



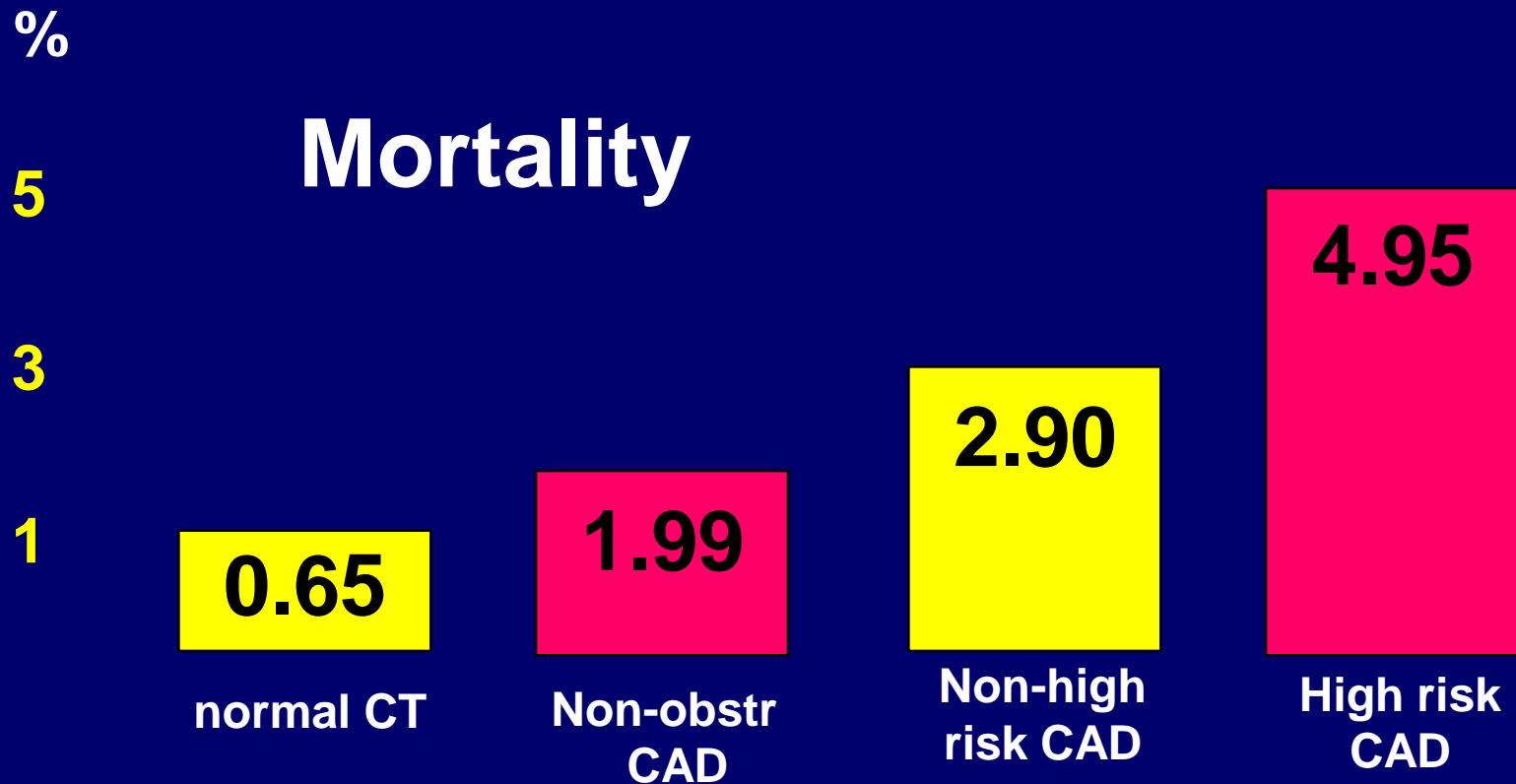
LAD: normal



LCx: normal

Prognosis MSCT

13,966 pts, mean F-up 22.5 months



Man 61 years old

Earlier on no atherosclerosis, but RFs have increased, symptoms have changed

Outpatient clinics:

chest pain at rest, sometimes stress

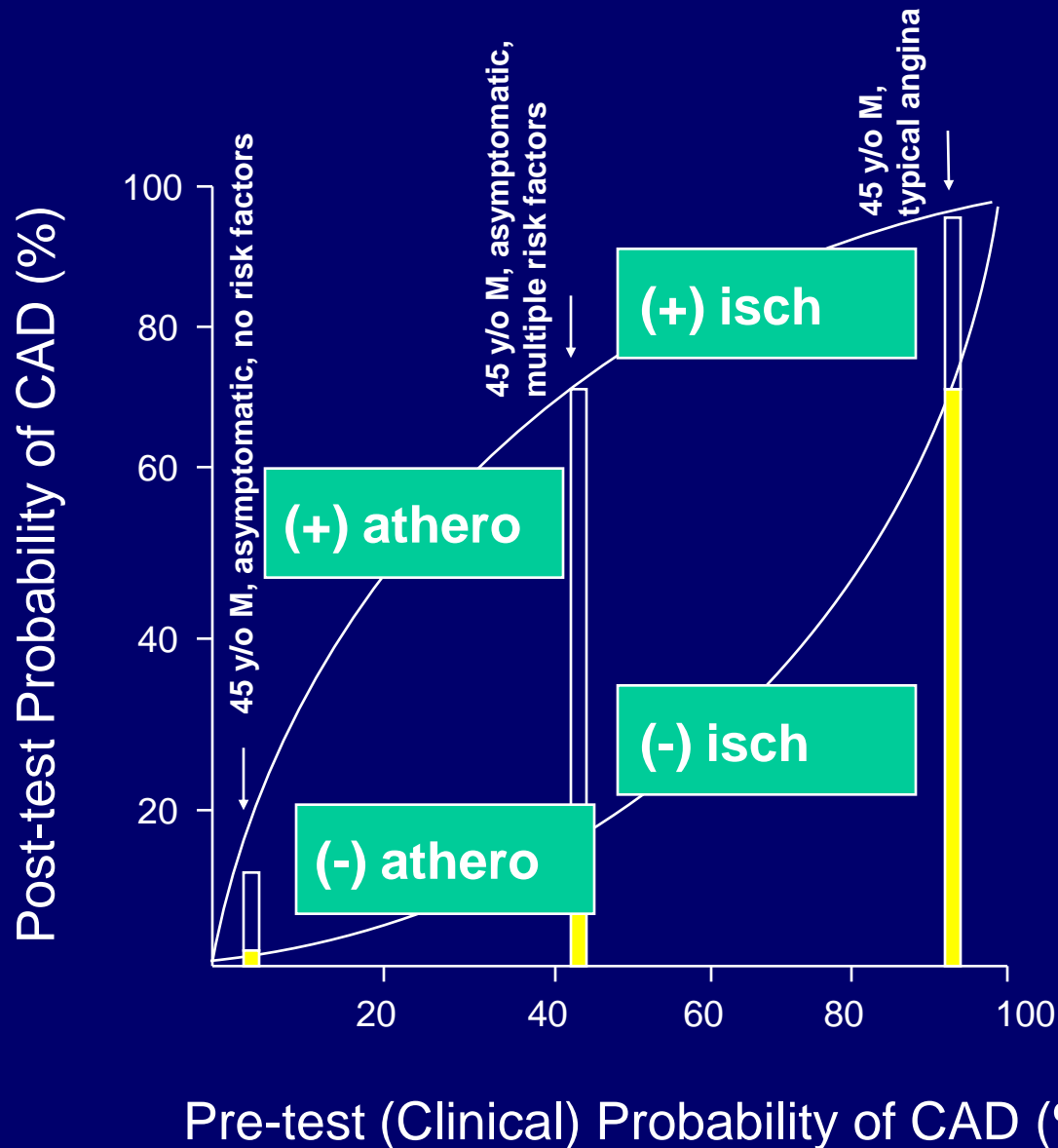
Risk factors for CAD:

- *Hypercholesterolemia

- *Hypertension

- *Smoking

Non-invasive assessment of CAD: Which Patients?



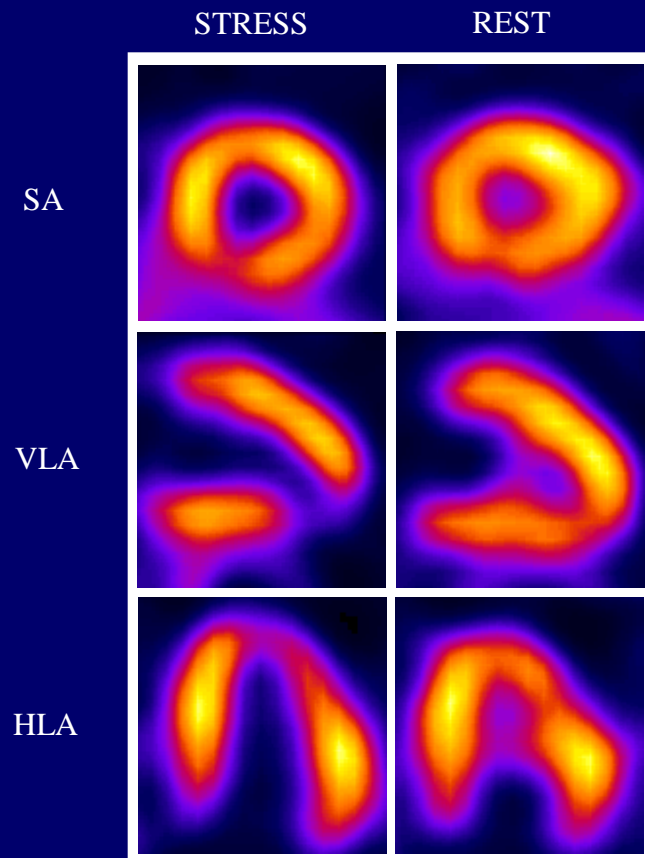
Symptomatic patient,
intermed – high pre-test
likelihood

The patient has high likelihood to have
atherosclerosis

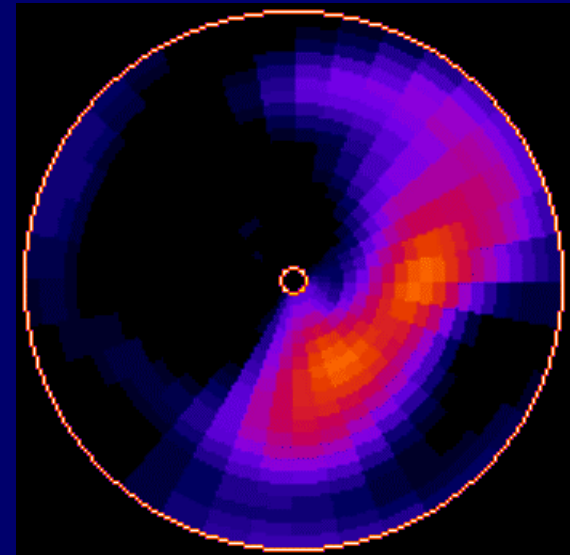
The question is: does he have ischemia?
(is intervention needed?)

☞ We order a non-invasive ischemia test

Nuclear perfusion imaging, SPECT

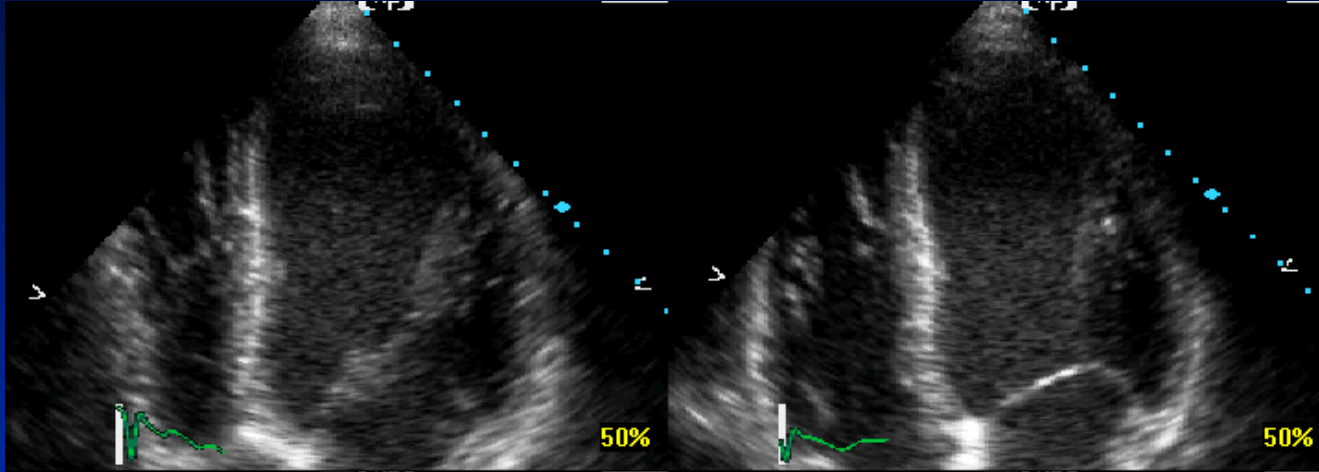


POLAR MAP TO QUANTIFY
EXTENT AND SEVERITY OF ISCHEMIA



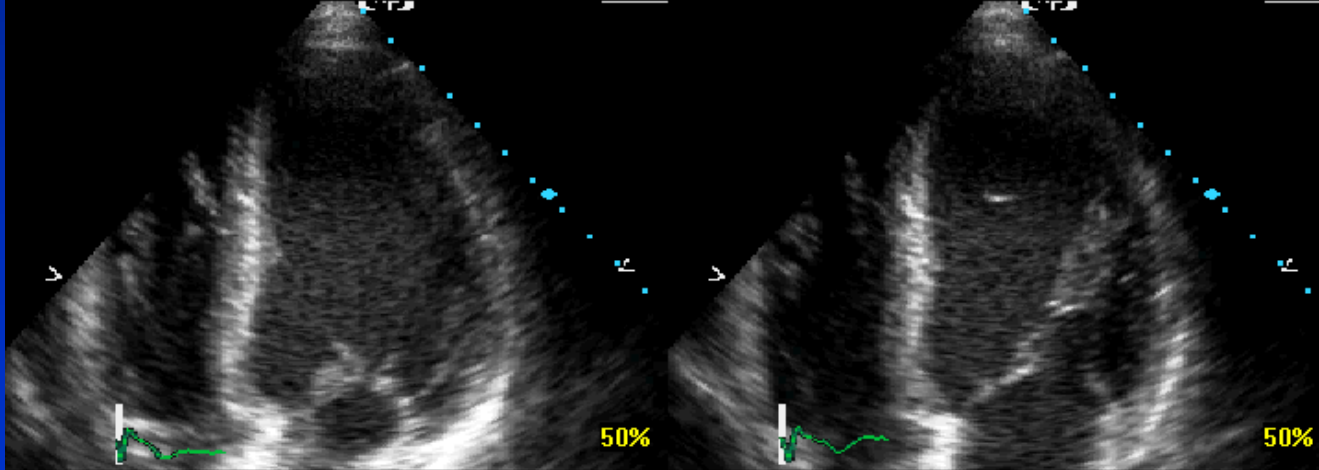
Stress echo to assess flow-limiting stenosis: wall motion

rest



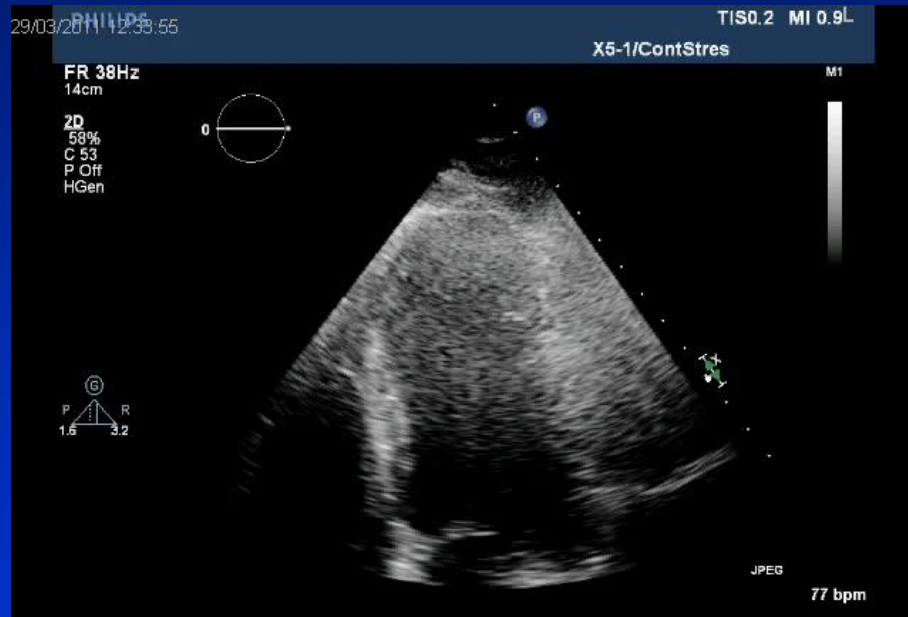
10 mcg

40

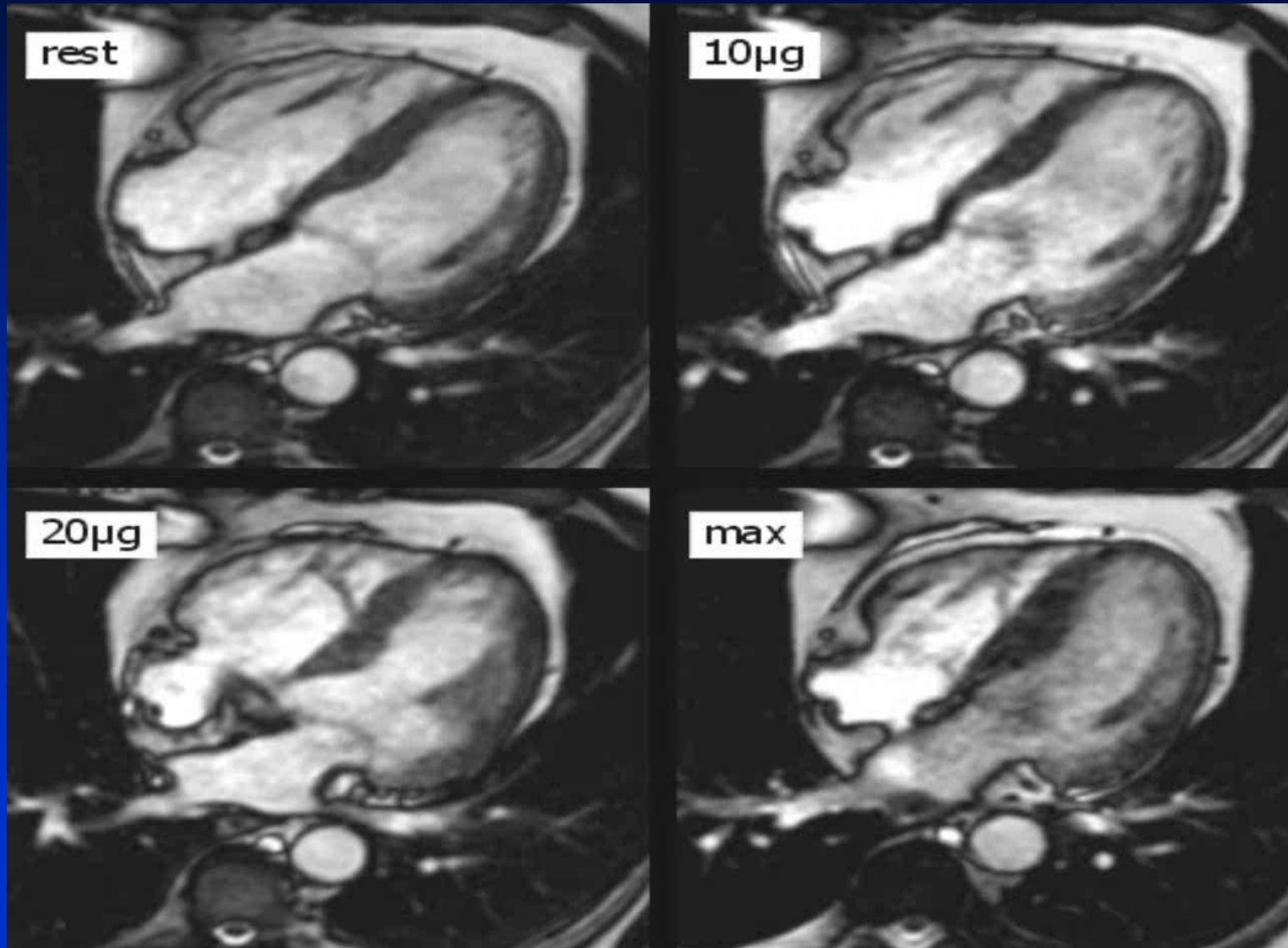


rest

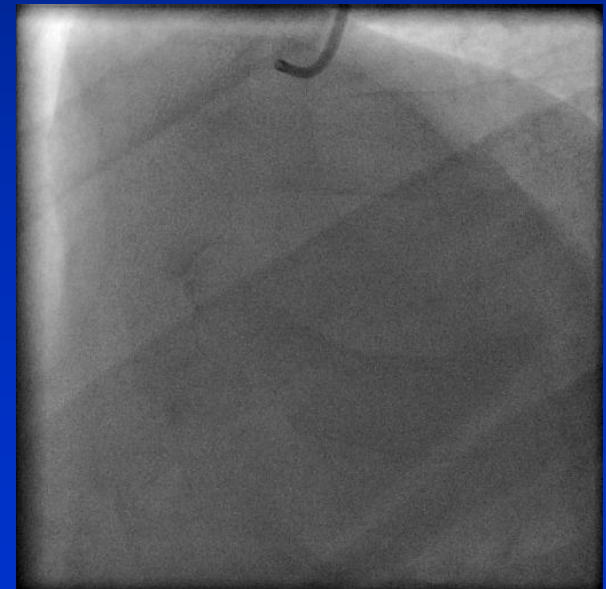
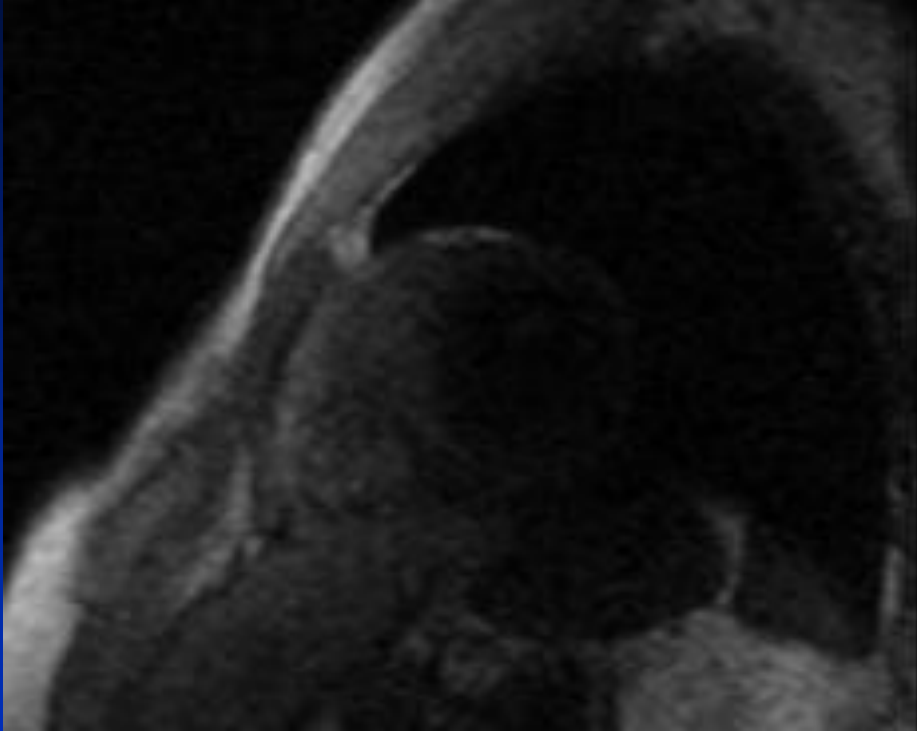
Addition on intravenous contrast to improve border opacification



Stress MRI to assess flow-limiting stenosis: wall motion



MRI – perfusion imaging



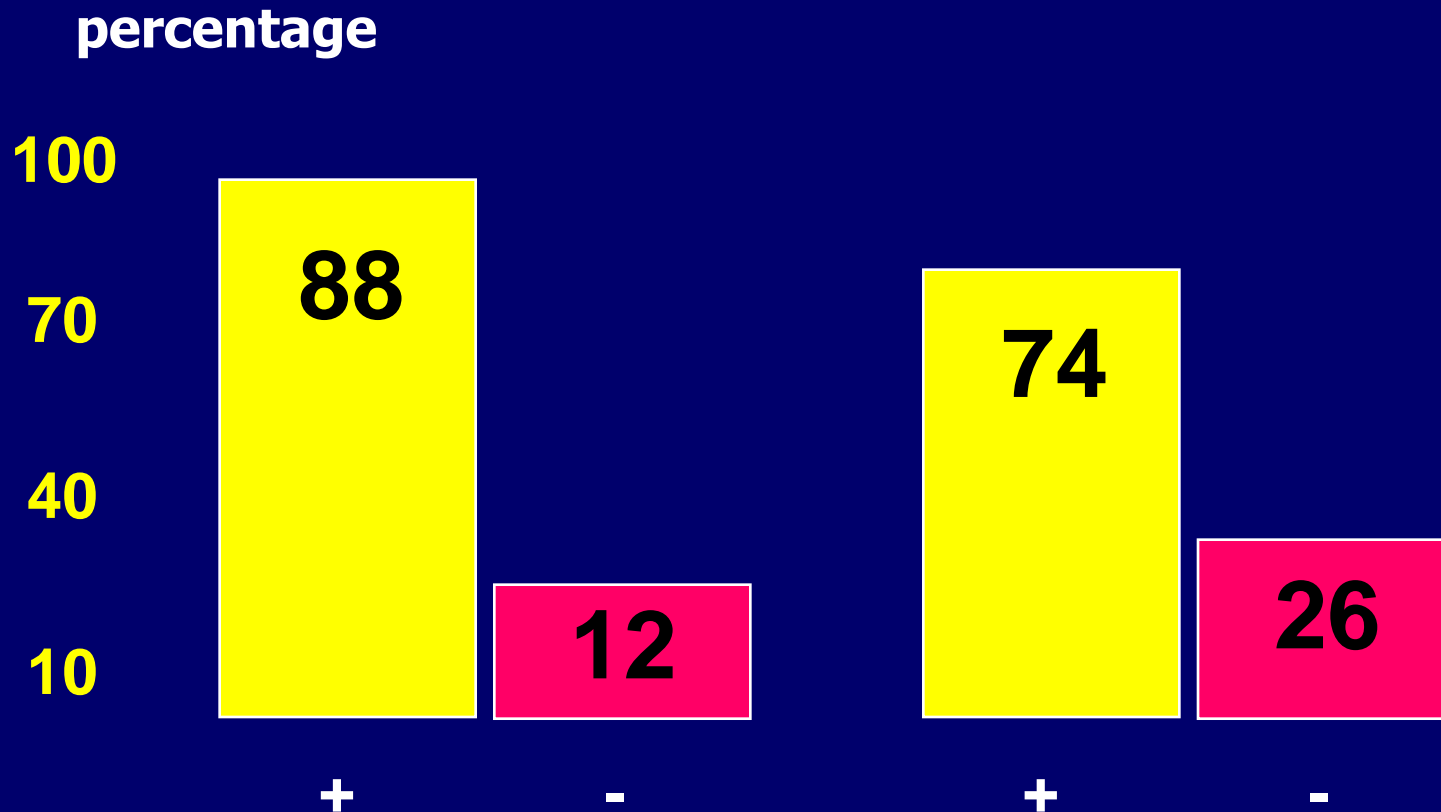
Man 61 years old

Outpatient clinics:

He has developed CAD, we treated based on ischemia

Now the patient developed PAF

AF ablation: success and failure



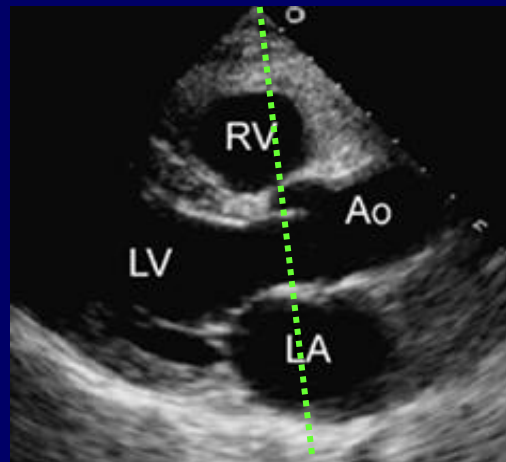
☞ Prediction of successful RFCA

Assessment of substrate for AF

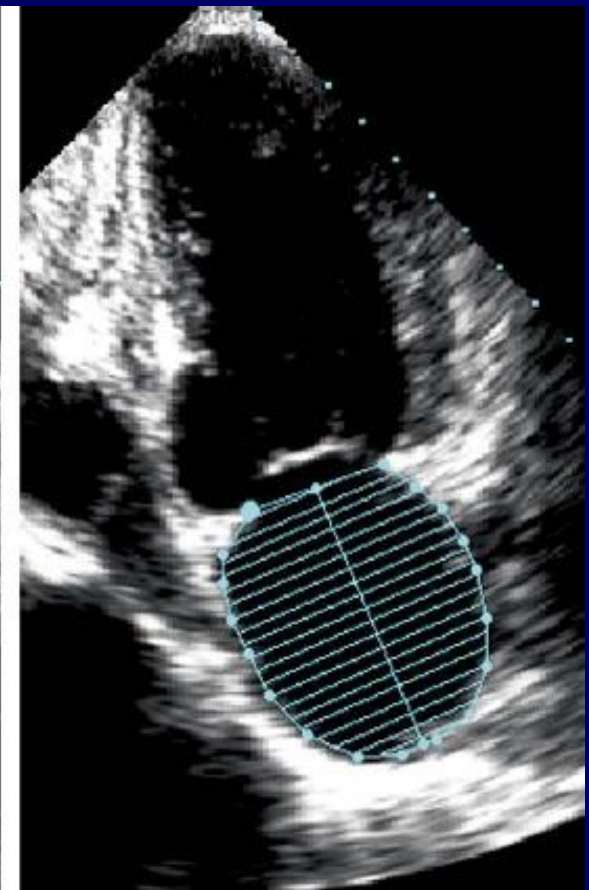
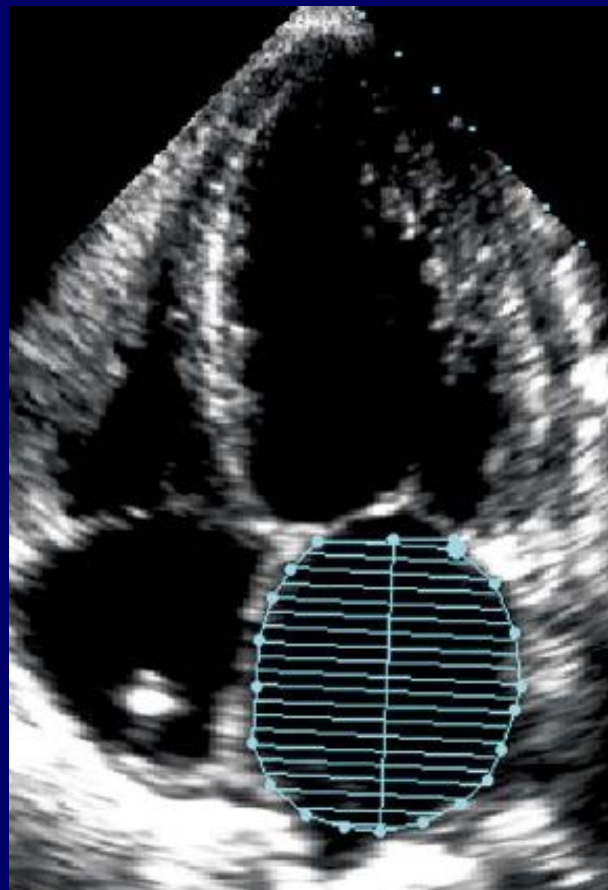
- LA enlargement
- LA fibrosis
 - Direct assessment
 - Indirect assessment
 - Mechanical consequence
 - Electrical conduction heterogeneities

Left atrial dimensions

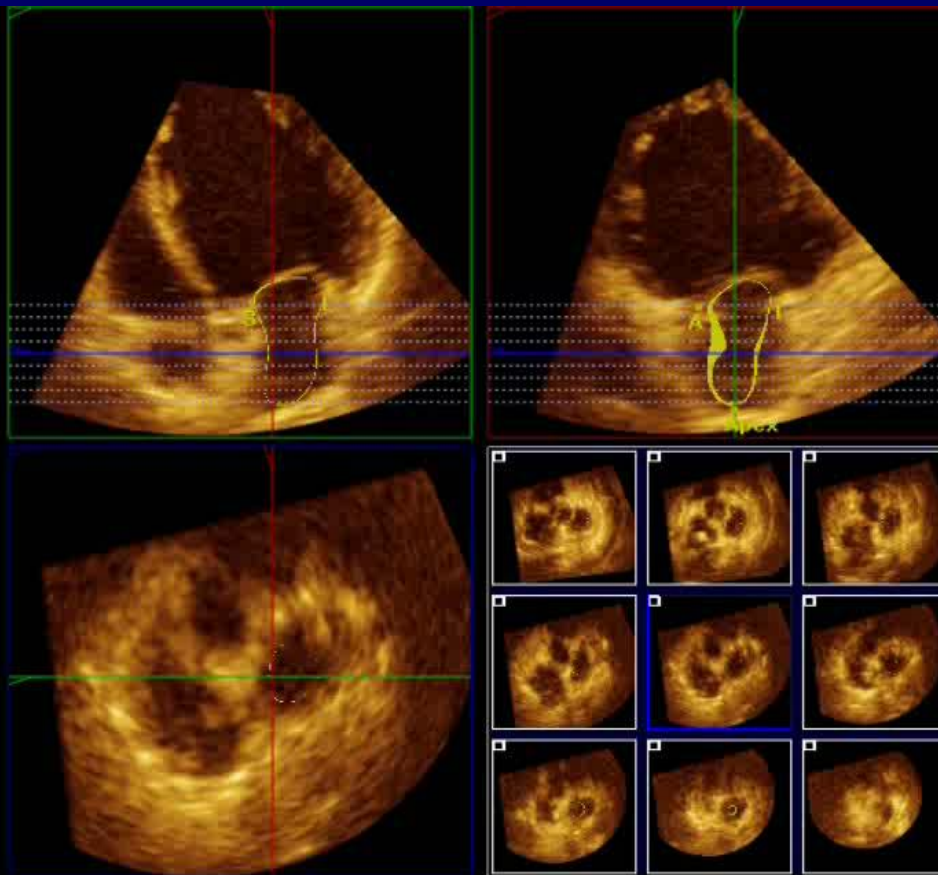
Linear dimensions
AP diameter



LA volume
Modified Simpson's rule



Real-time 3D echocardiography



Volume(s)

EDV = 58.0 ml

ESV = 24.2 ml

Calculation(s)

EF = 58.3 %

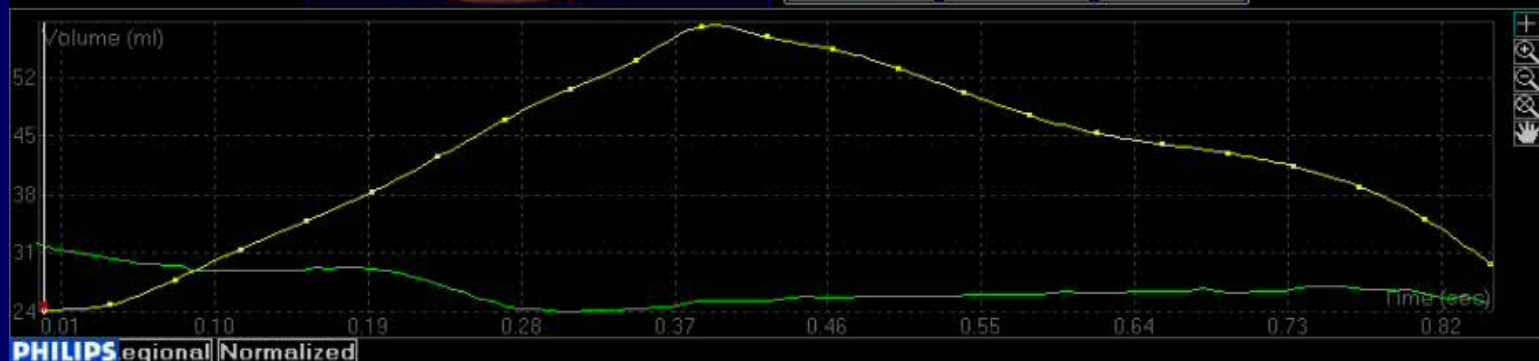
SV = 33.9 ml

Regional

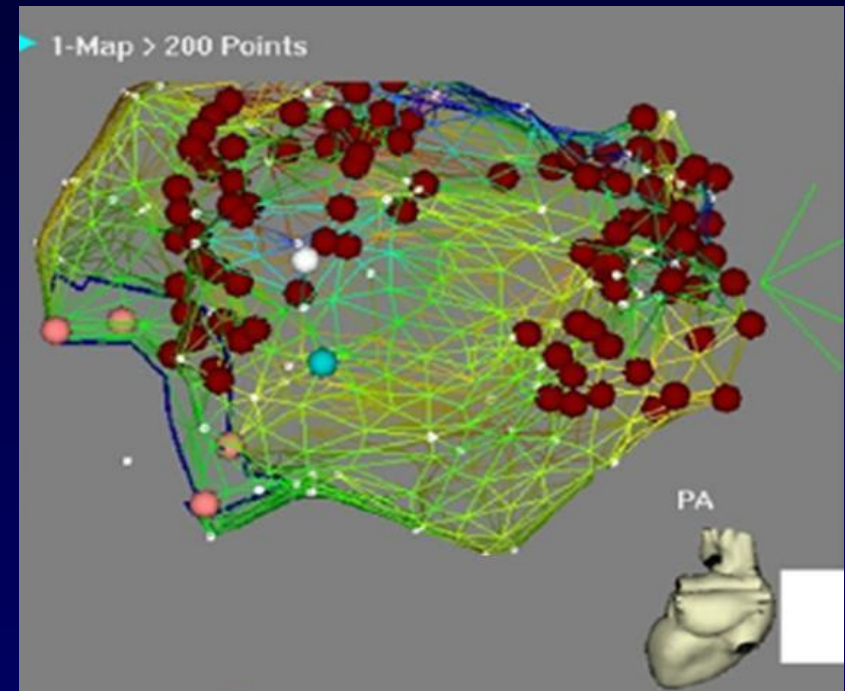
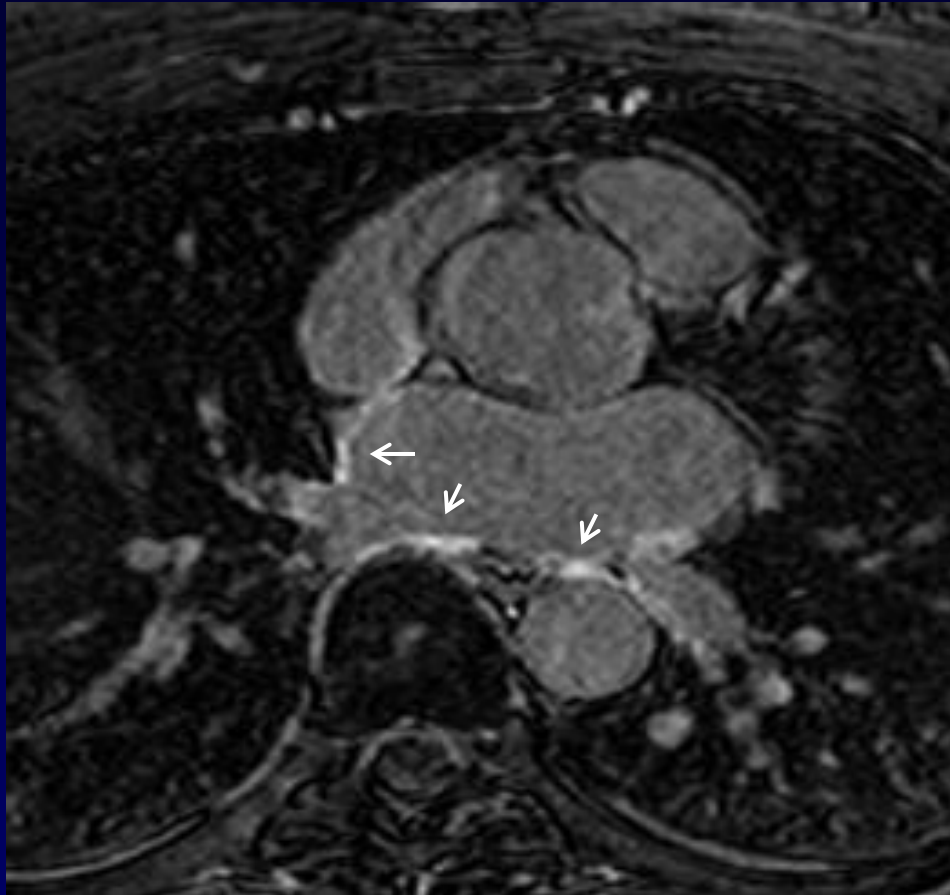
Tmsv Sel-SD = ☐

Tmsv Sel-Dif = ☐

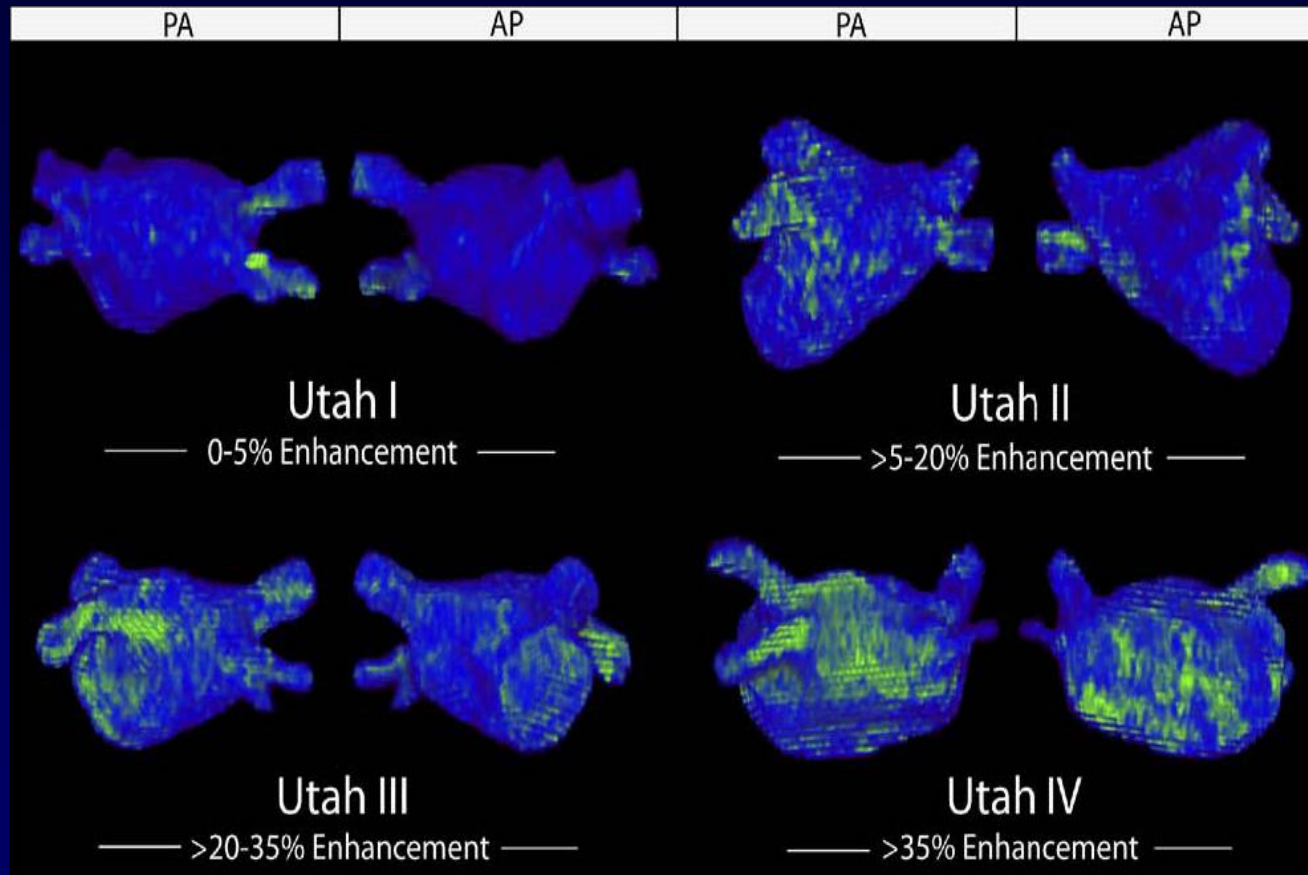
Tmsv Sel-SD = ☐



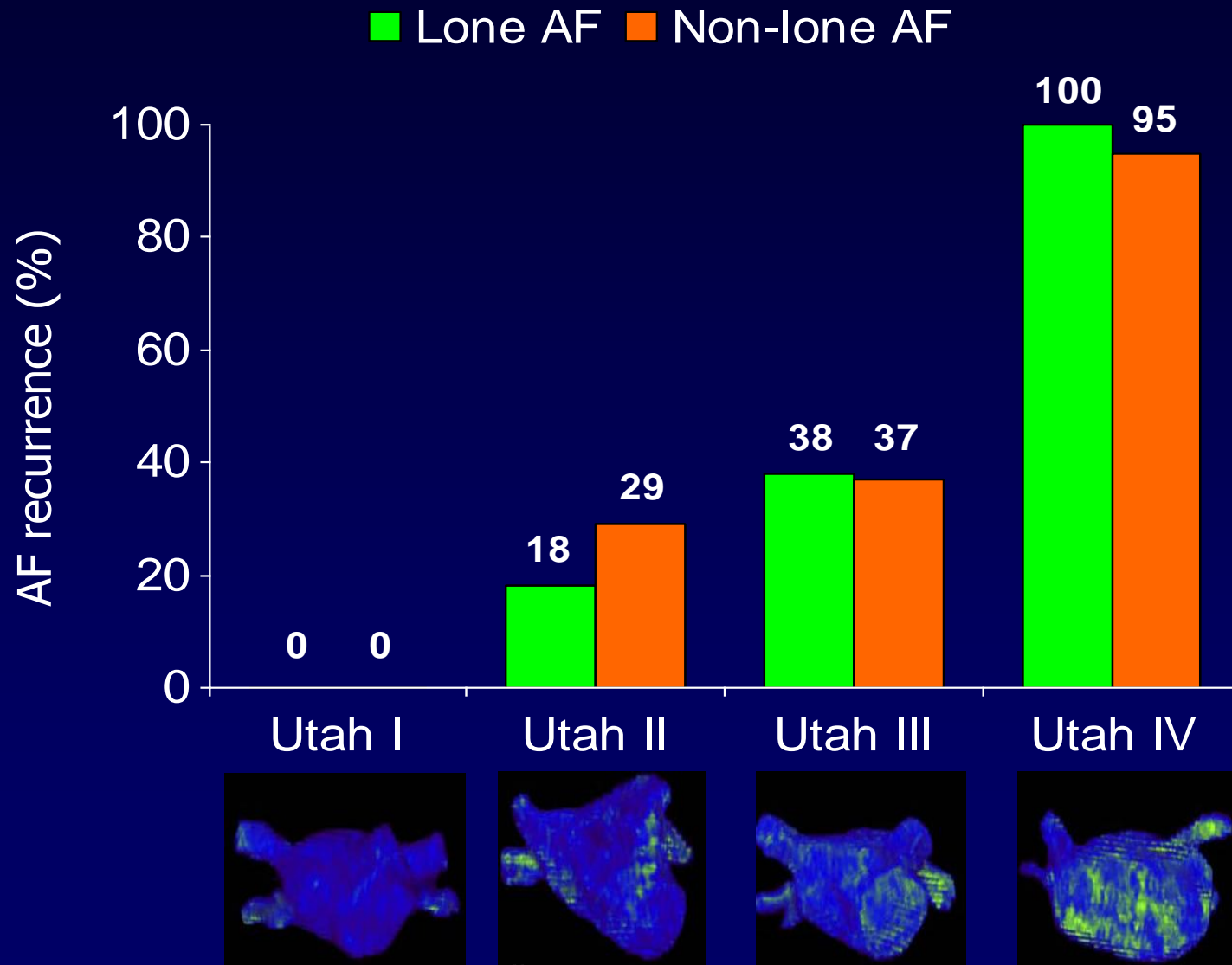
Left atrial fibrosis imaging - MRI



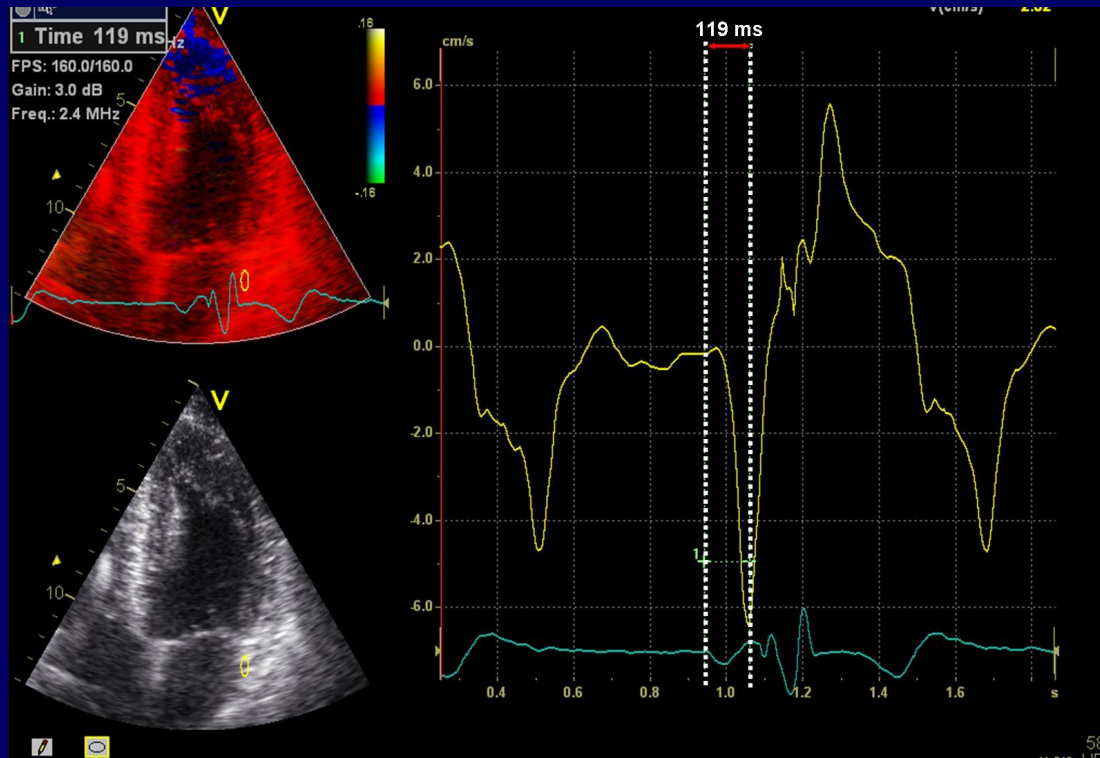
- **N = 333 AF patients**
- **LA fibrosis before RFCA: DE-MRI**



LA fibrosis vs. RFCA outcome



Left atrial electro-mechanical properties



TDI



**Total atrial conduction
time (PA-TDI)**



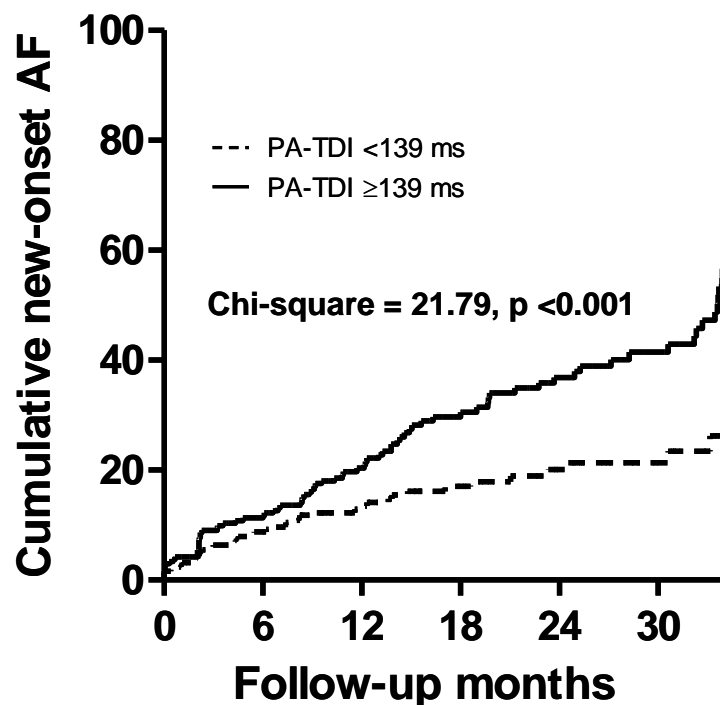
**Time interval from the
onset of the P-wave to
the A'-wave peak**

- PA-TDI as predictor of new onset AF in heart failure patients

N = 495

79% male

21% previous paroxysmal AF



Number of patients at risk:

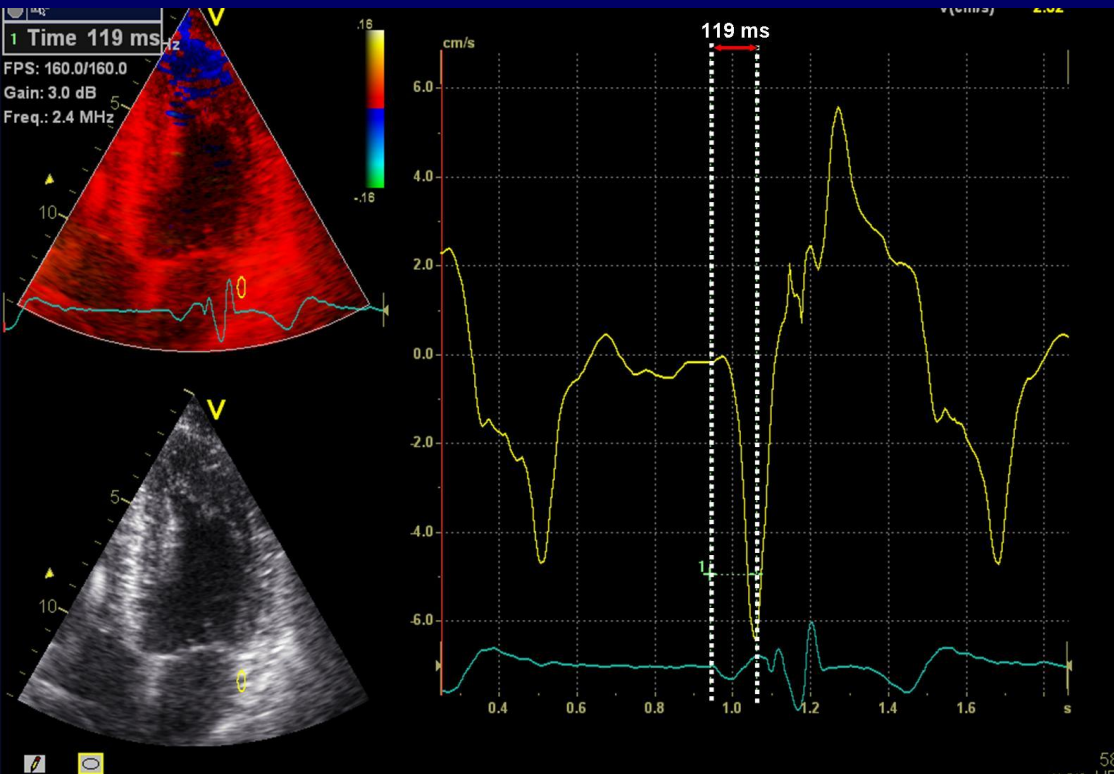
PA-TDI duration <139 ms	253	226	140	94	66	38
PA-TDI duration ≥139 ms	242	193	133	83	64	43

PA-TDI

HR: 1.01 (1.01-1.02)

P<0.001

Patient 1



No AF

Patient 2



New onset AF

Different patient: extensive CAD

Male, 72 yrs

- 2001: Infero-postero-lateral infarct – PCI
- 2002: 2003: Antero-septal infarct - PCI
- 2004: CABG: LIMA-graft LAD,
venous graft MO-LCX and RDP/RCA
- 2004: LV dilated, EF 28%

Co-morbidities

- Diabetes II

Man 72 years old

CAD has been treated

History of MI, EF is reduced

Outpatient clinics:

Does he need an ICD?

Does he need an ICD?

- Patients with:
previous infarction
LVEF <30-35%
- Benefit from ICD:
- MADIT II: improved survival

ICD needed?

ICD shocks in primary prevention

percentage N=720 pts, MADIT II
Follow-up 21 months
Shocks:

100

70

40

10

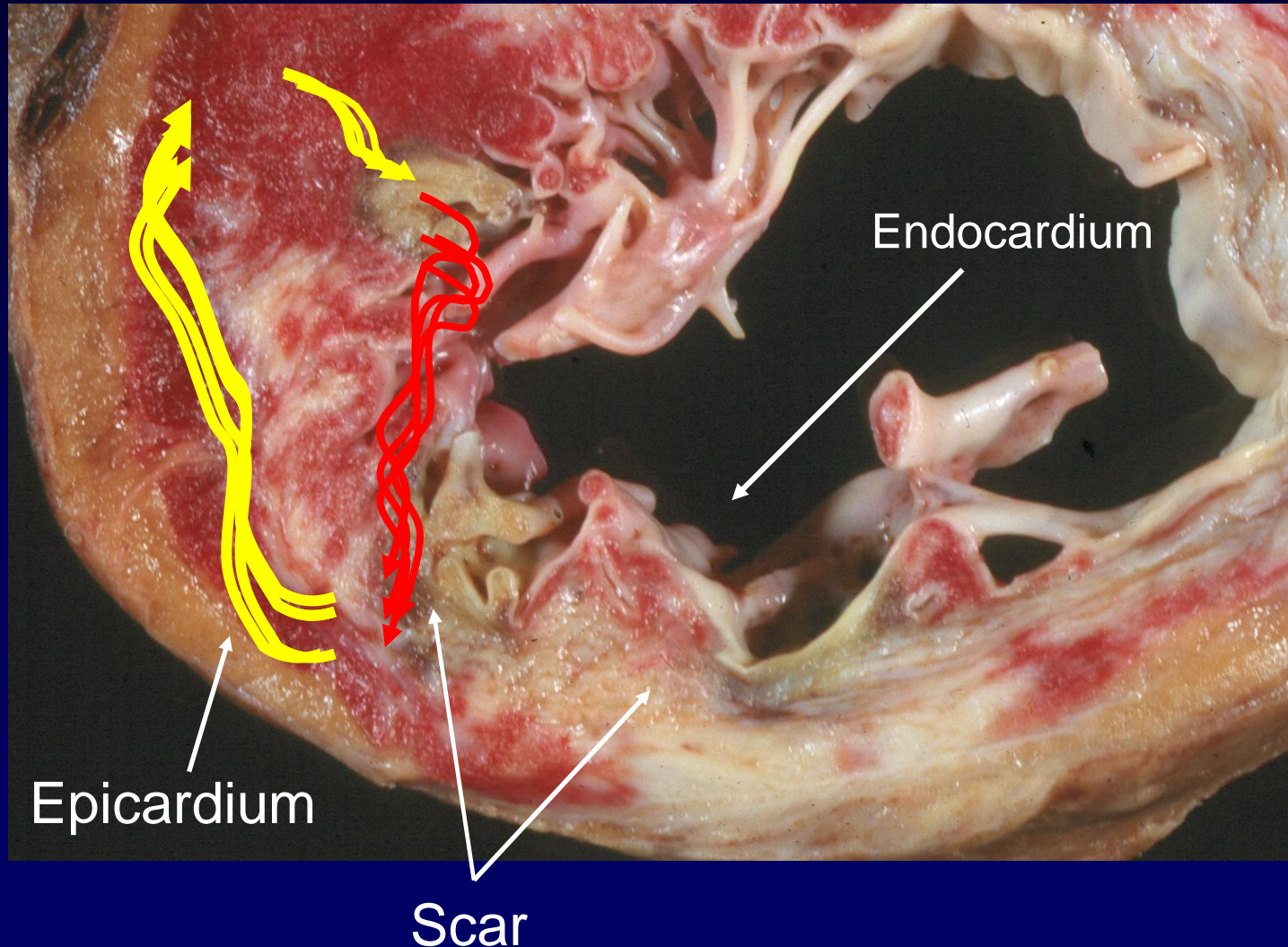
65

35

-

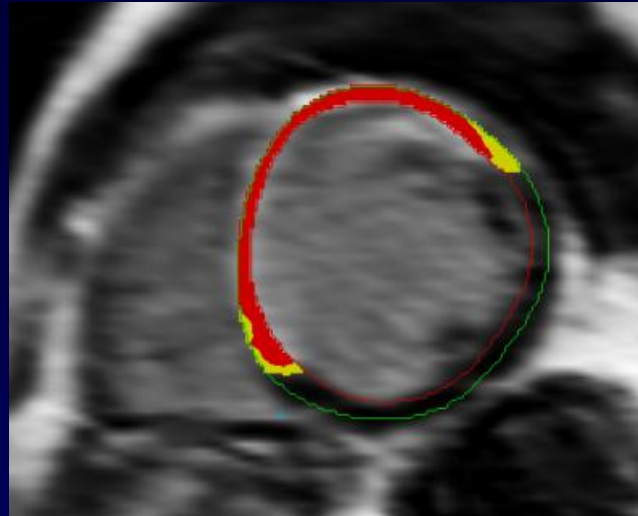
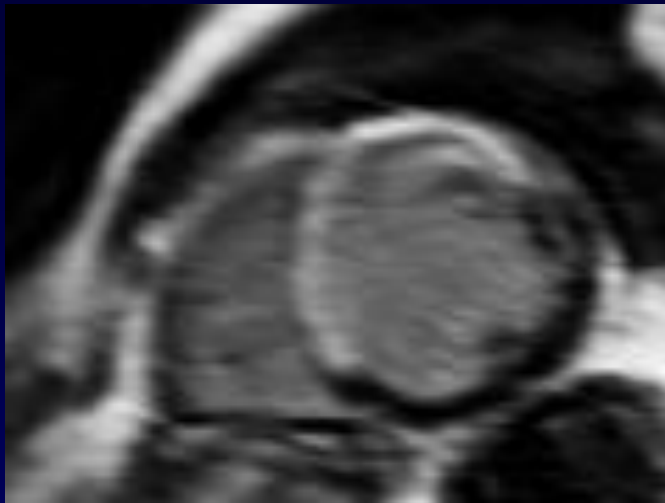
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What is the pathophysiological substrate for SCD in CAD?



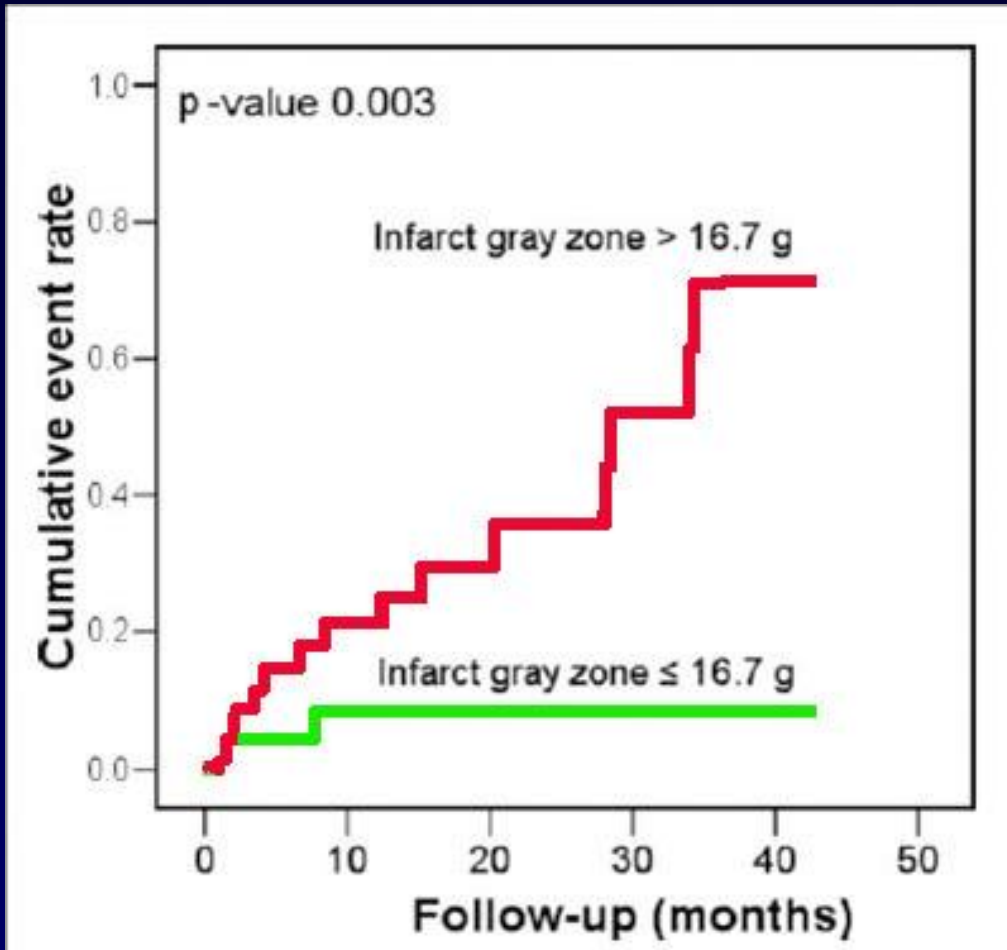
Courtesy W Stevenson

MRI to assess arrhythmogenic substrate:



- Late-gadolinium enhancement: scar area and peri-infarct zone

Value of border zone to predict VTs



HR (95%CI): 1.47 (1.04 to 2.08)
P = 0.003

Conclusions

- **Virtually all anatomical and functional information can be obtained by (a combination) of the available imaging techniques**
- **The choice of techniques should be guided by the information needed (the questions we need answers to)**
- **The imaging results must affect treatment**