



# Advances in Cardiology and Cardiac Interventions

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# Cardiology is never static...

▼ “what we can do.”

▼ “what we should do”



# The future appears very promising...

✓Diagnostics

✓Therapeutics



# DIAGNOSTICS

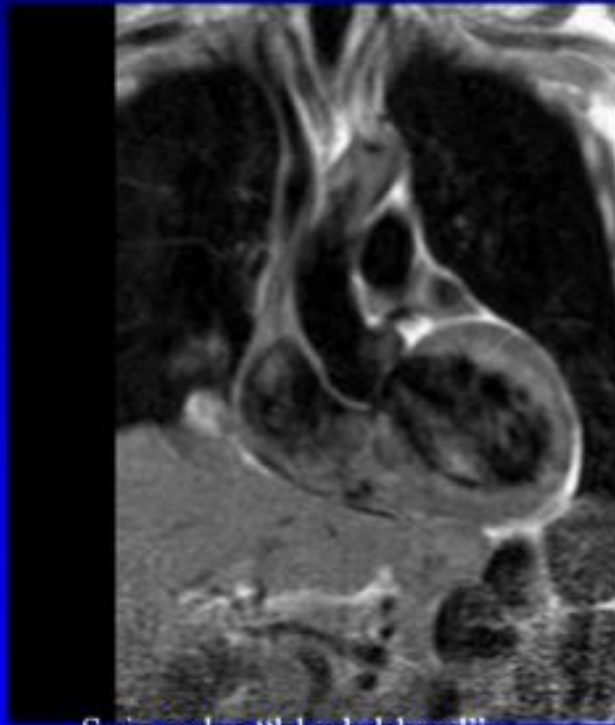


# Cardiac MRI

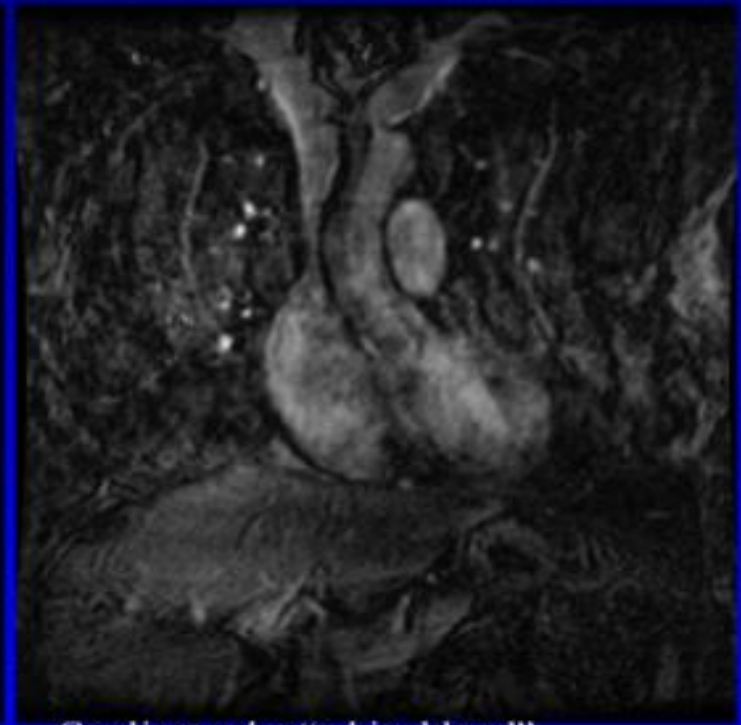
- Cardiac and great vessel anatomy
- Cardiac volumes and mass
- Global and regional contractile function
- Regional myocardial tissue perfusion
- Regional myocardial tissue characteristics:  
Viability, oedema, inflammation, fibrosis,  
metabolism
- Coronary artery lumen, wall anatomy, blood flow



# Coronal MRI shows aorta, av, lv (can eval for stenosis and regurg)



Spin echo "black blood"  
anatomy



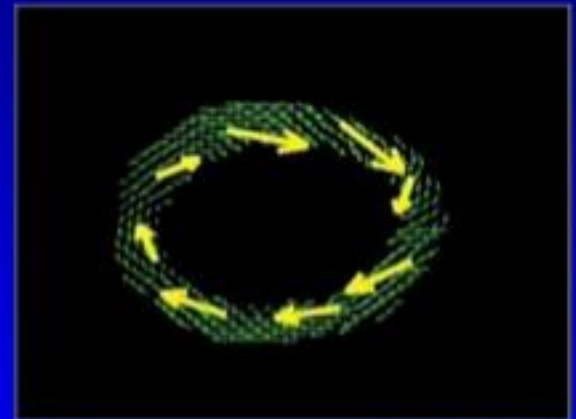
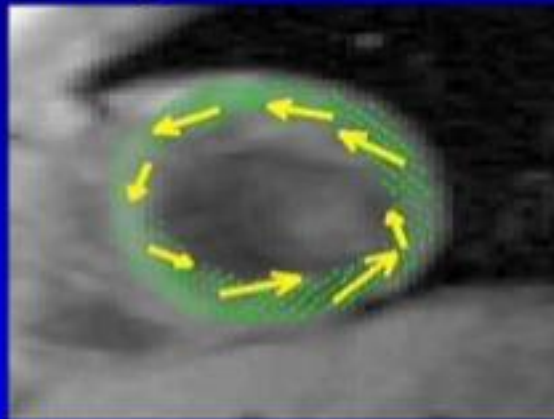
Gradient echo "white blood"  
function & flow



# Regional Tissue Contractility

## *Tissue Phase Mapping*

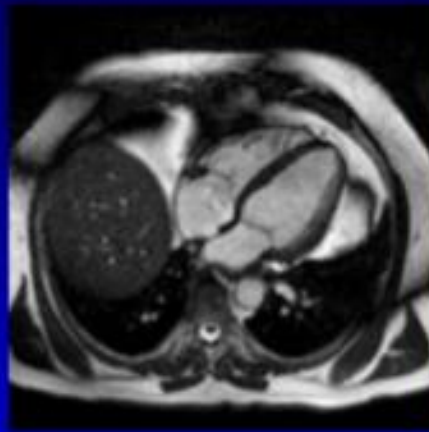
3D Velocities: Radial, circumferential,  
longitudinal



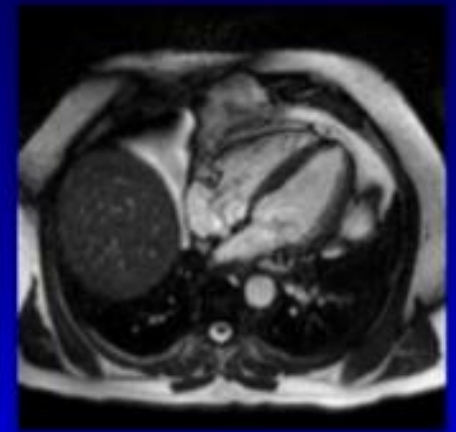


## Dobutamine-Stress MR: 4-Chamber

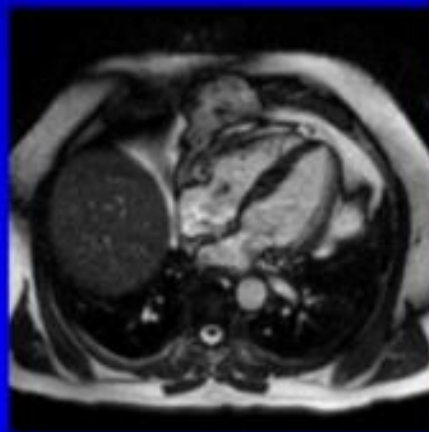
rest



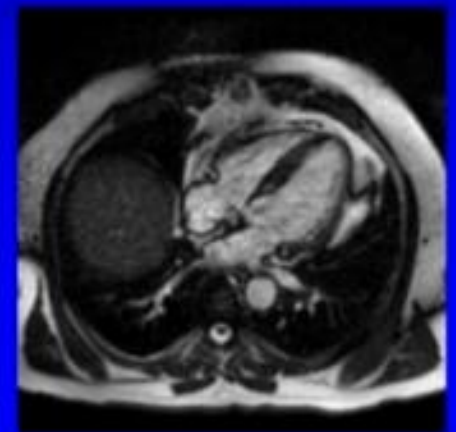
20  $\mu$ g



30  $\mu$ g



40  $\mu$ g



# Delayed Enhancement MRI

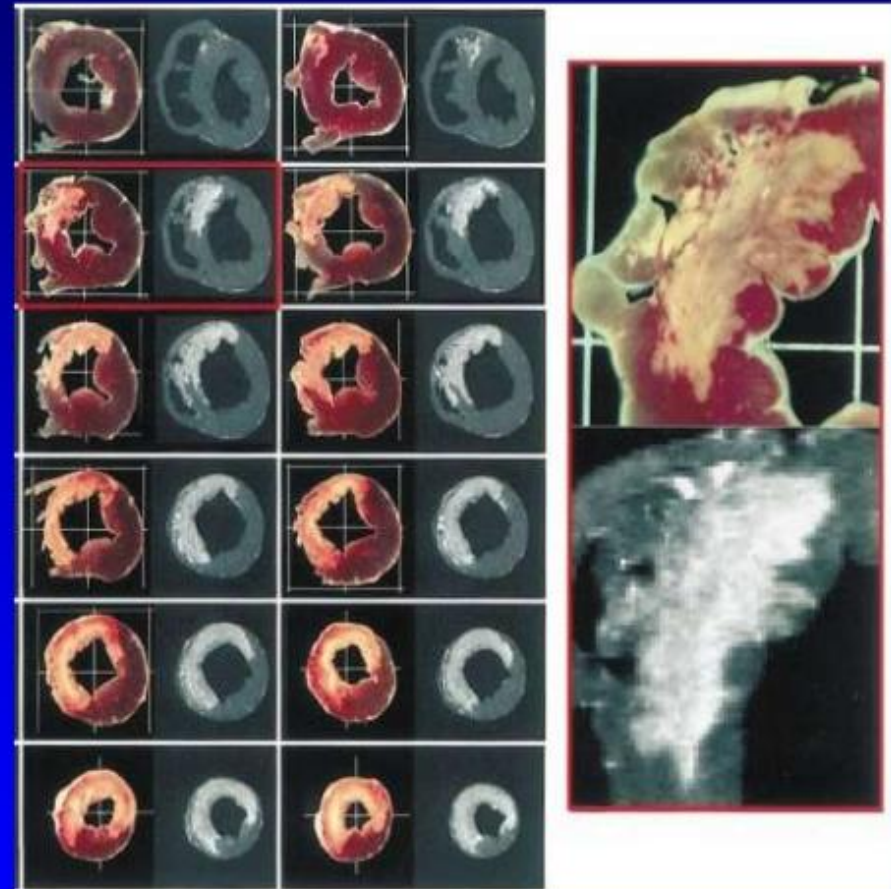
10 – 20 min post Gd DTPA

Inversion recovery

FLASH or True FISP

“Bright is dead”

Normal, stunned, hibernating  
myocardium is dark

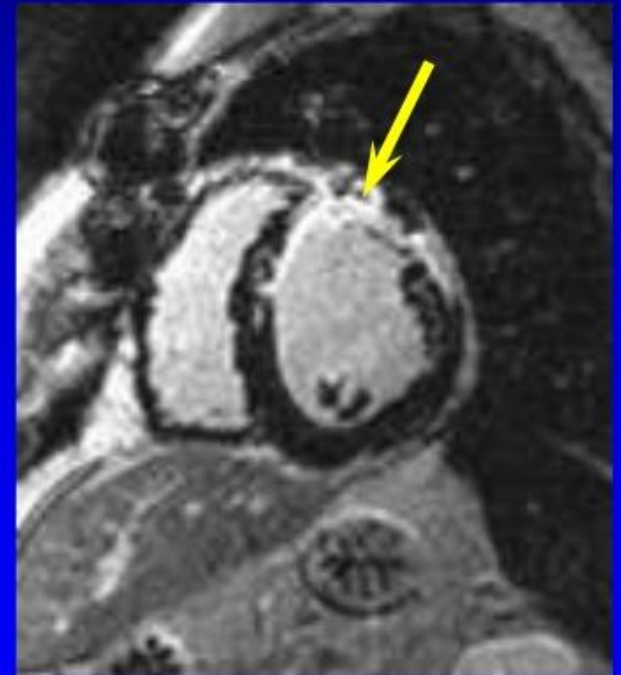
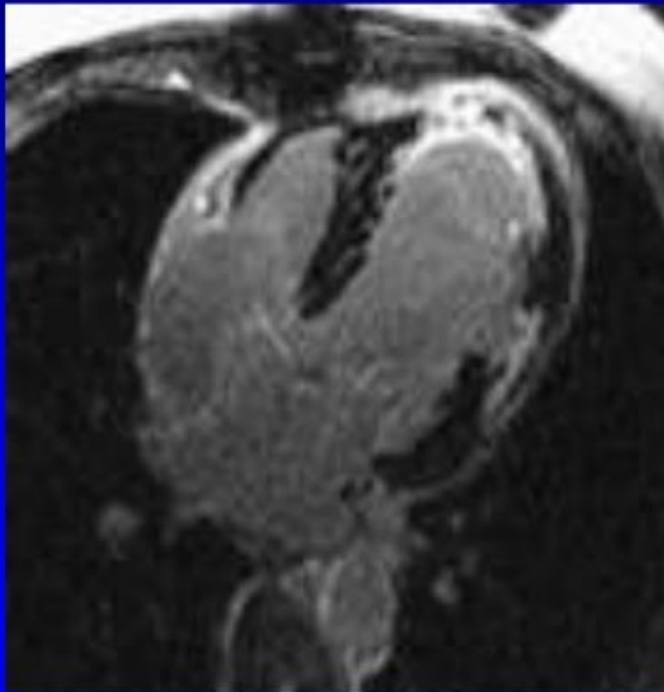




# Delayed Enhancement MRI

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*In vivo infarct imaging*





FFR, IVUS, OCT



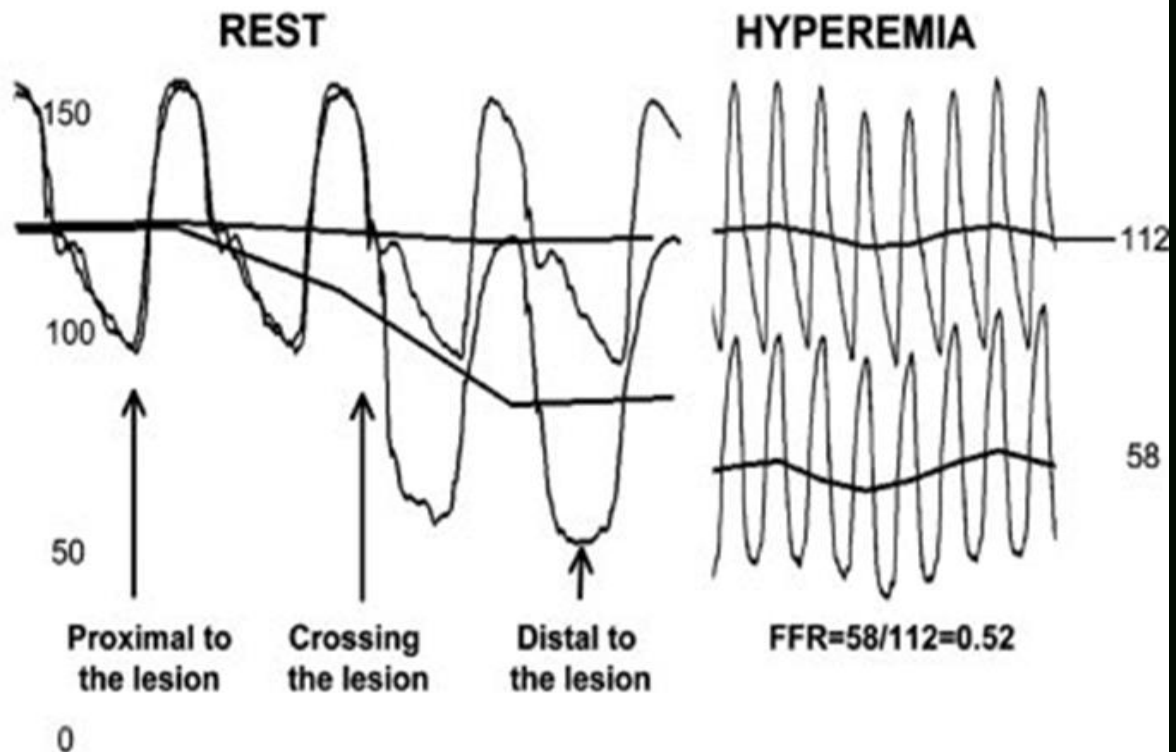
# Limitations of CAG ....

- Intermediate lesion (40-70%) significance cannot be accurately determined
- Won't provide intraluminal detail ; vessel wall details ;
- diffuse disease, angiographic artifacts of contrast streaming, image foreshortening and calcification often complicates the interpretation ;
- Bifurcation or ostial lesion locations may be obscured by overlapping branch



# FFR(Fractional Flow Reserve)

- Gold standard for invasive assessment of physiologic stenosis significance.
- Ratio of the coronary pressure distal to the stenosis to aortic pressure as the normal perfusion pressure (distal coronary pressure/aortic pressure) when the microvascular resistance was minimal and assumed to be constant (ie at maximal hyperemia).
- Normal value of FFR - 1
- $FFR < 0.75$  .... inducible myocardial ischemia [ 88% sensitivity, 100% specificity, 100% positive predictive value and overall accuracy 93%]



- In this example....During maximal hyperemia (at the right side of the pressure tracing), the hyperemic distal pressure decreased to 58 mm Hg with aortic pressure of 112 mm Hg for an FFR of 0.52 (58/112).
- Maximal blood flow to the myocardium of the anterior wall of this patient is decreased to 52% of expected normal flow.

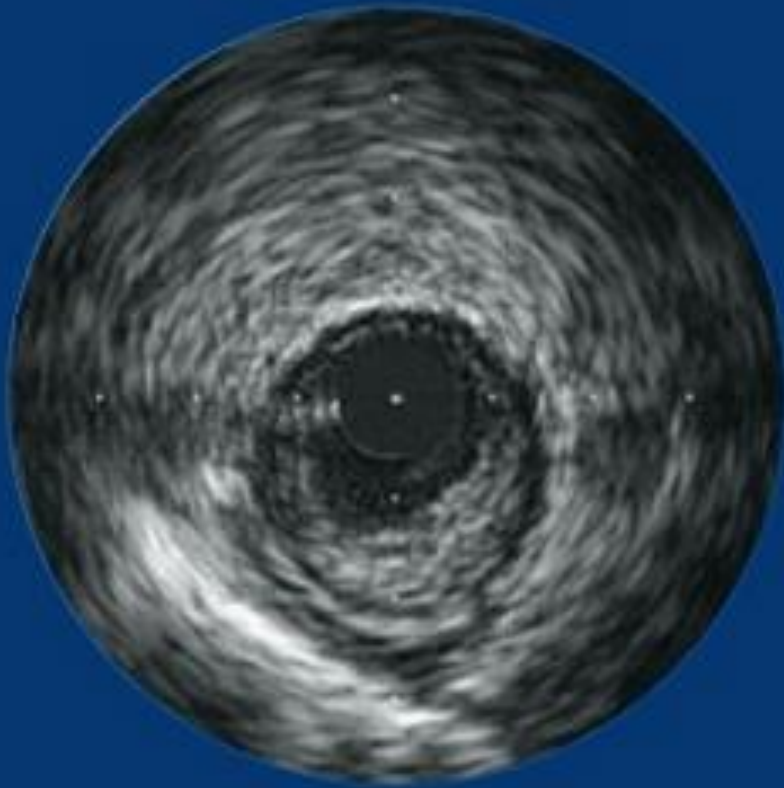


# IVUS to guide PCI :

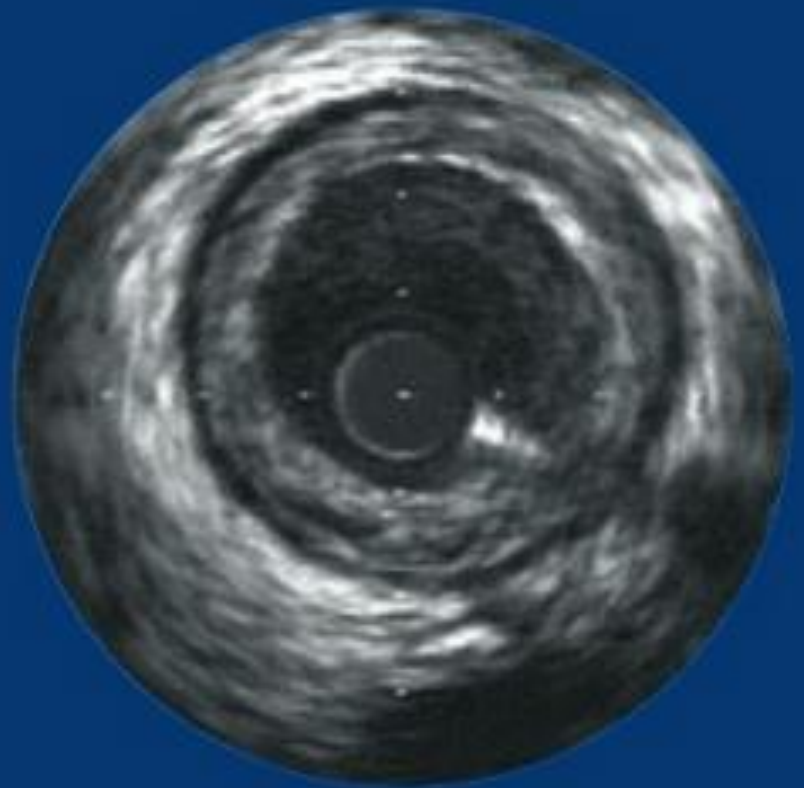
- ✓ plan interv strategy & to optimise stent deployment.....
- ✓ find reference lumen dimensions and lesion length for appropriate stent sizing;
- ✓ identification of superficial calcium
- ✓ Assess thrombus burden
- ✓ Post-stent IVUS assessment may detect complications of PCI and suboptimal stent deployment
- ✓ Addition of radio frequency backscatter signal analysis allows for improved characterization of plaque composition1)virtual histology IVUS 2) iMAP 3) Integrated Backscatter IVUS



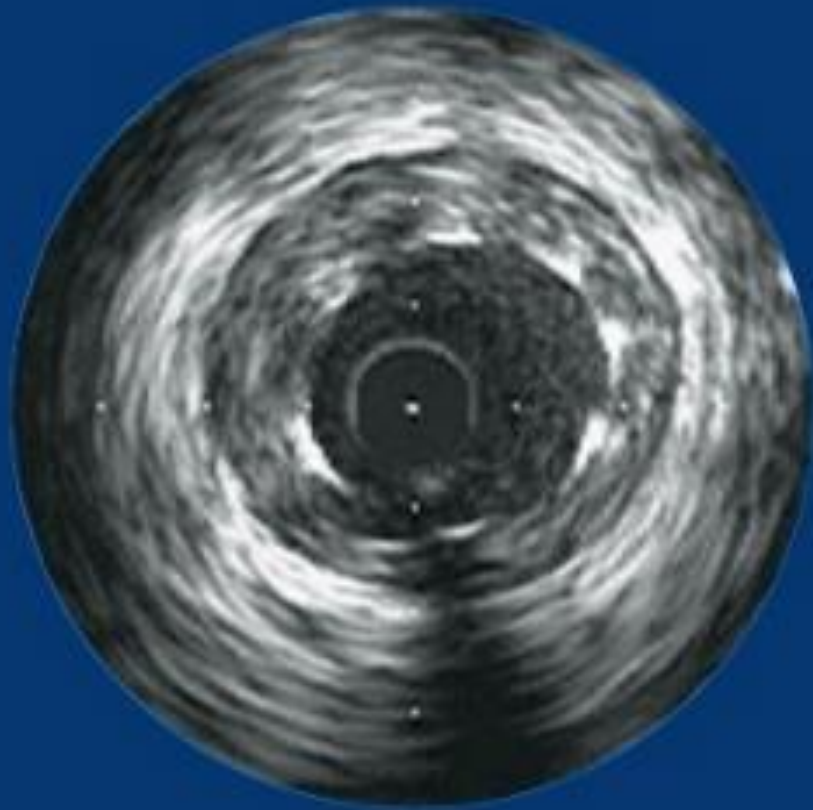
# IVUS images



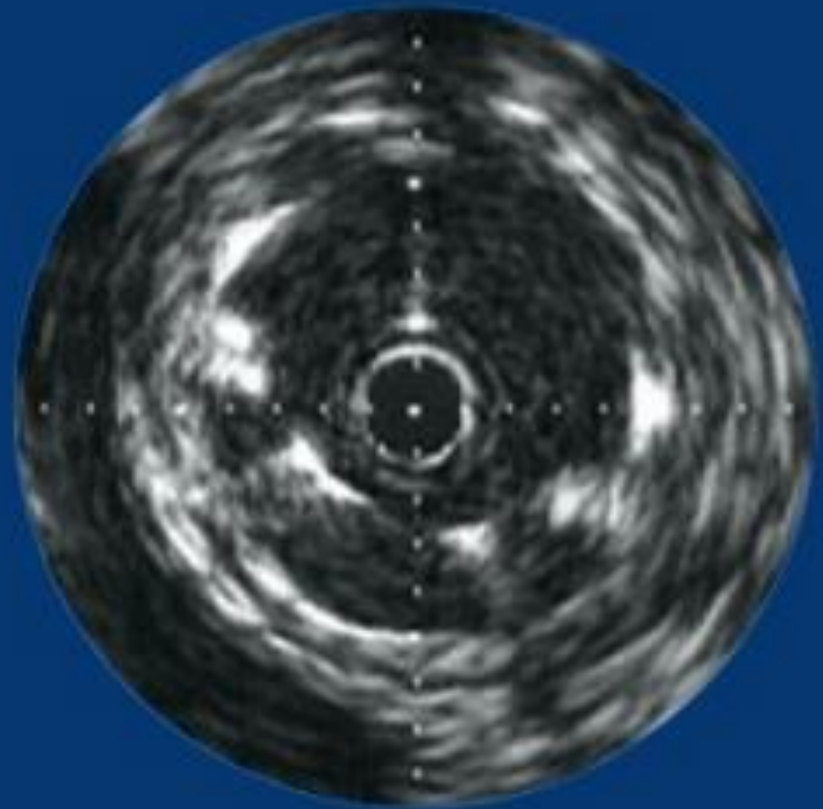
Eccentric fibrous plaque



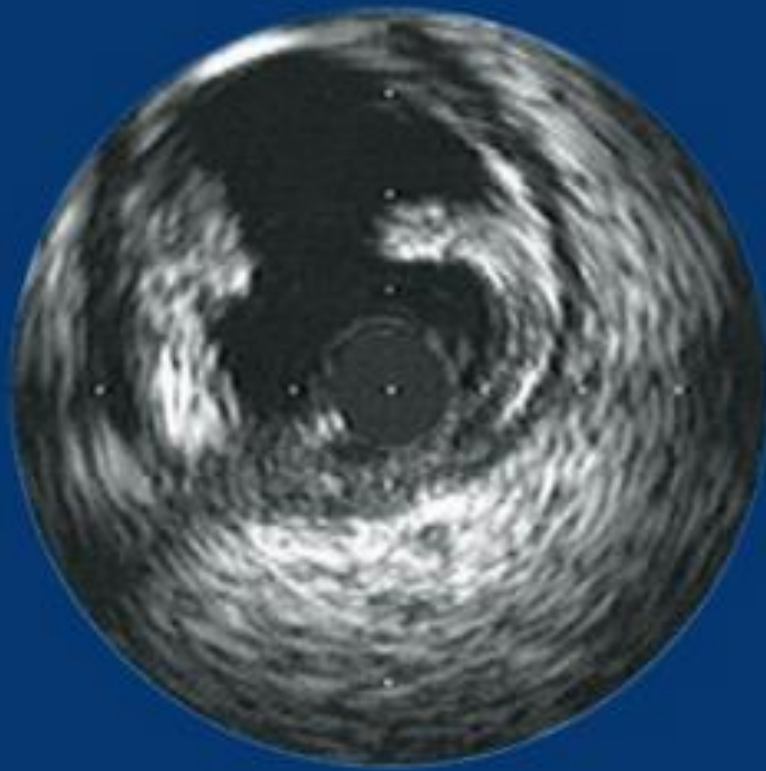
Concentric fibro-fatty plaque



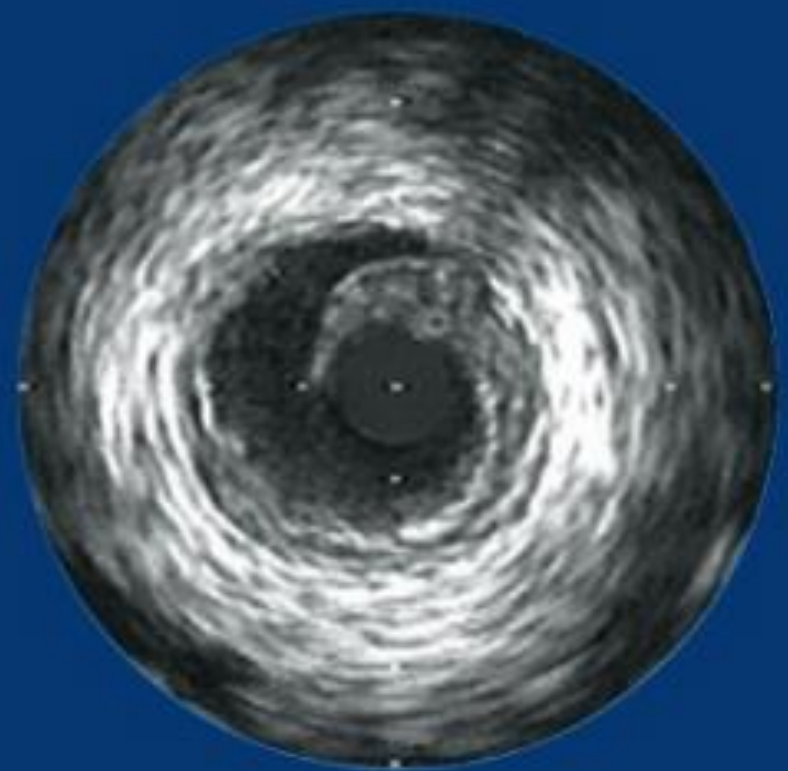
Fully apposed stent



Incompletely apposed stent



Fibrous plaque dissection  
extending into the intima

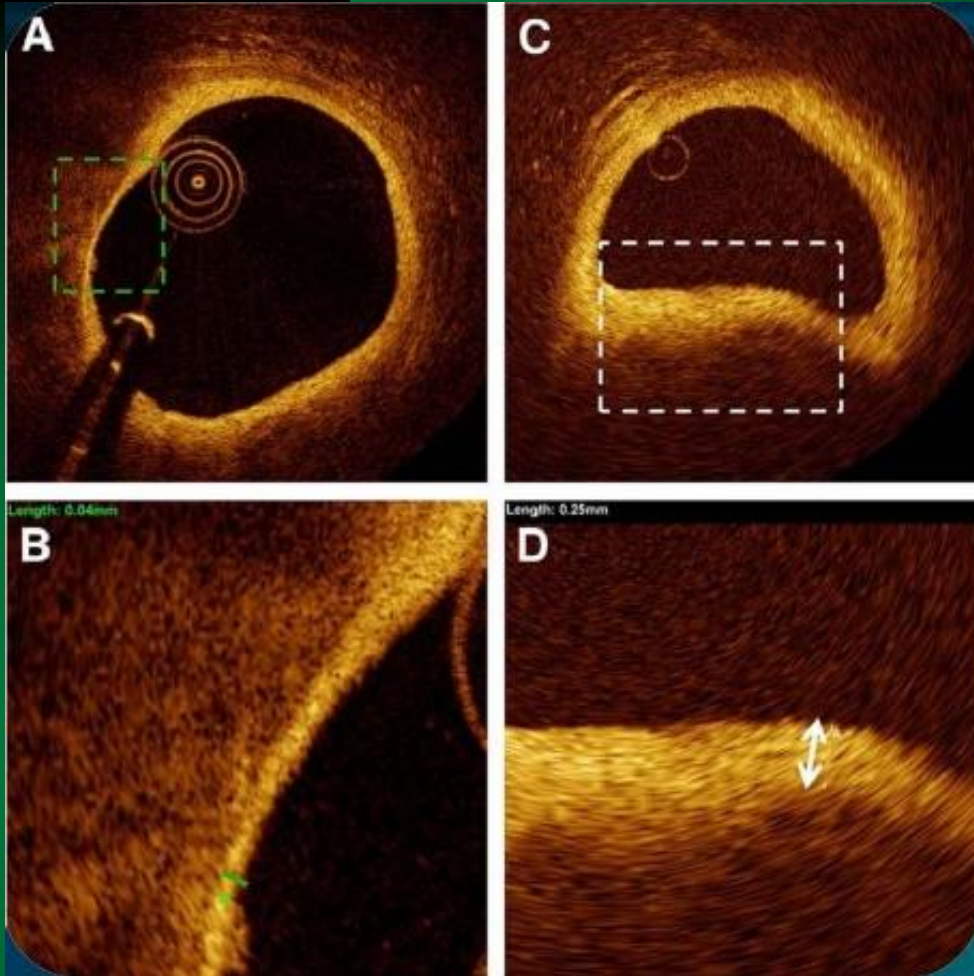


Deep fibro-fatty plaque dissection  
extending into the media



# OCT

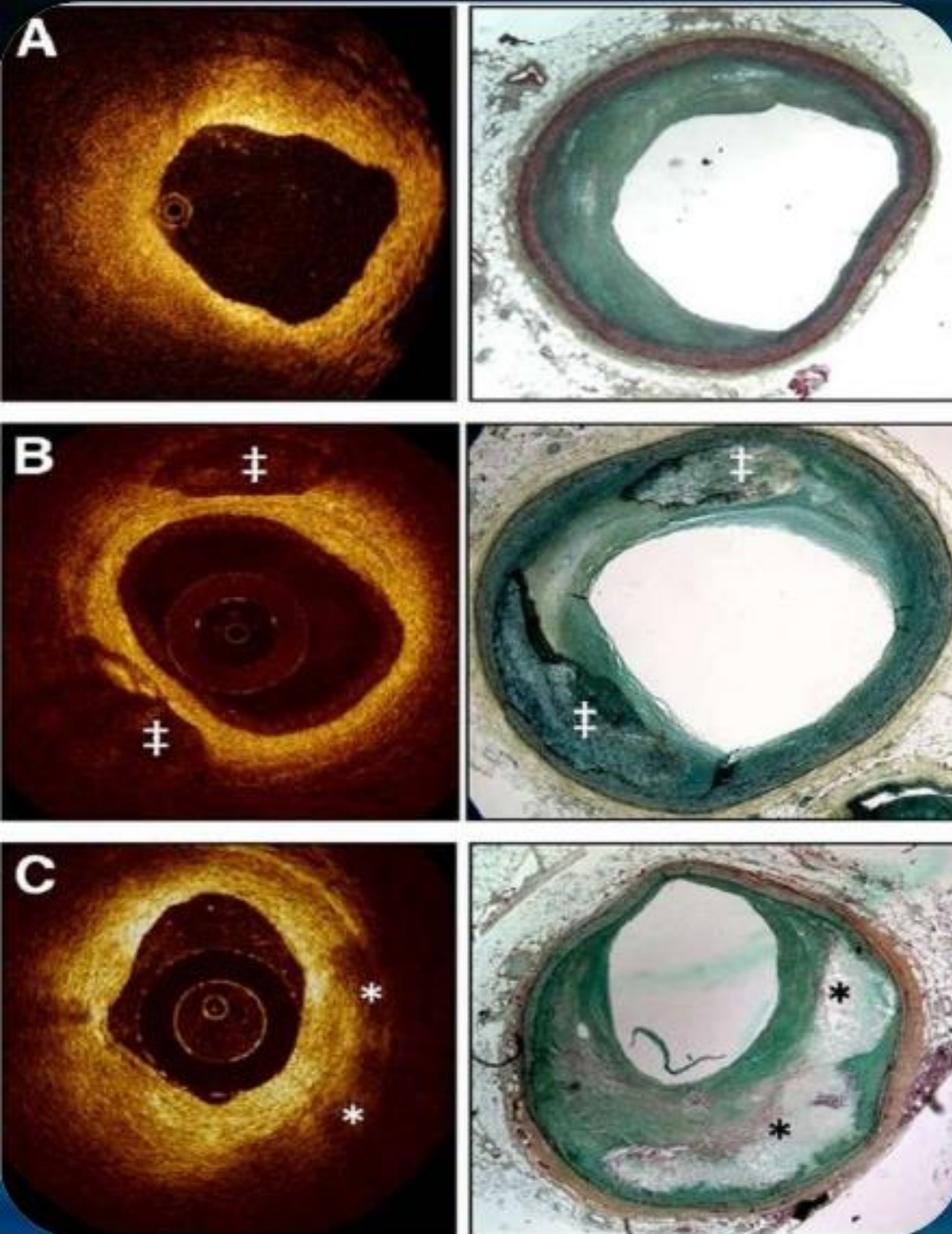
- ACS ----- Rupture of a coronary plaque .
  - Detection of lesions at high risk of rupture (so called “vulnerable plaques”) is of major importance for the prevention of future ACS.
  - OCT --- one of the most promising --- provides unique information about the plaque composition, the presence of macrophages, and the thickness of the fibrous cap
- 
- **Fibrous plaques** -- homogeneous, signal-rich regions
  - **Fibro - calcific plaques** --- signal-poor regions/ sharp borders
  - **Lipid-rich plaques** ---signal-poor regions with diffuse borders



# OCT images

- Thin capped fibroatheromas (TCFA) - defined pathologically by the triad of:

- ☐ **Lipid core.**
- ☐ **Fibrous cap with a thickness  $< 65$  micron m.**
- ☐ **Cell infiltration of the fibrous cap.**



## OCT- Histology co-relation



# NUCLEAR CARDIAC IMAGING

Available for the evaluation of -

- ✓ Myocardial Perfusion
- ✓ Ventricular Function
- ✓ Cardiac Metabolism
- ✓ Myocardial Viability
- ✓ Myocardial Innervation
- ✓ Infarct Imaging



- It involves the injection of a radiolabelled substance which is extracted by the myocardium and accumulates in proportion to myocardial blood flow.
- Such radiopharmaceuticals are injected under stress as well as resting conditions,
- **Myocardial ischaemia** is defined as a perfusion defect present during stress but not resting conditions.
- **Scar** tissue is associated with a relative perfusion defect at rest as well as under stress.



## ✓ **Exercise Stress**

## ✓ **Pharmacologic Stress**

- \* vasodilators : adenosine  
dipyridamole
- \* inotropic : dobutamine



SIEMENS

1

Symbia  
TruePoint SPECT-C

STRESS [fbp]

21-Nov-2007 11:40:20

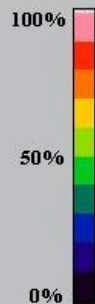
Intervals: 1

Pharma: Tc-99m MIBI

Ug Vol: 62 ml, TID: 1.72

SSS: 26

V2-GSRD/TC/NC/M



REST [fbp]

28-Nov-2007 14:00:05

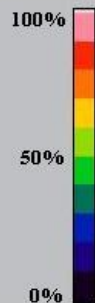
Intervals: 1

Pharma: Tc-99m MIBI

Ug Vol: 36 ml, TID: N/A

SRS: 7

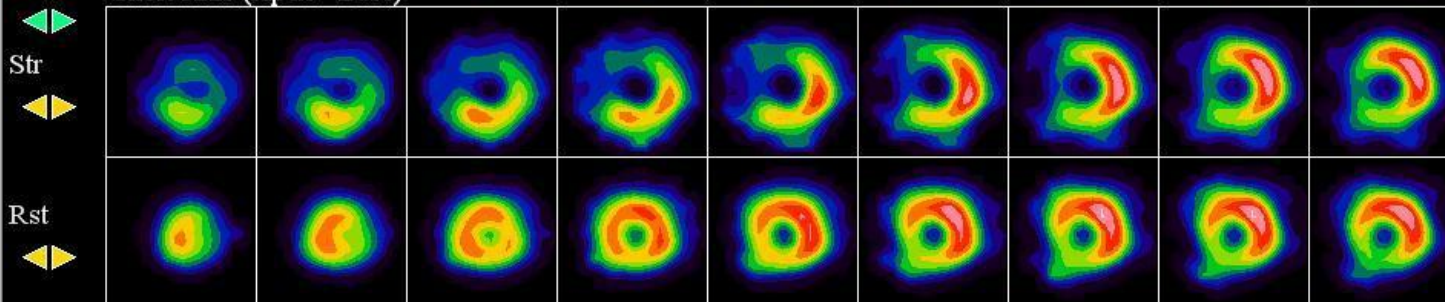
V2-GSRD/TC/NC/M



Study Info

# REVERSIBLE DEFECT

Short Axis (Apex->Base)

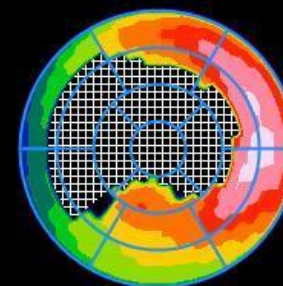
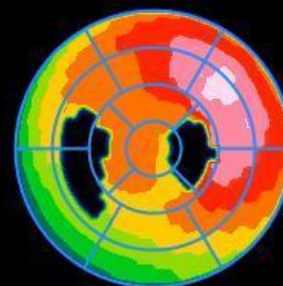
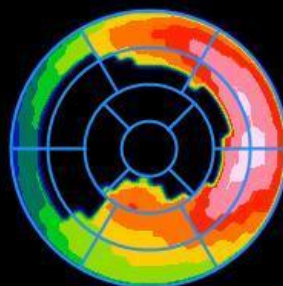
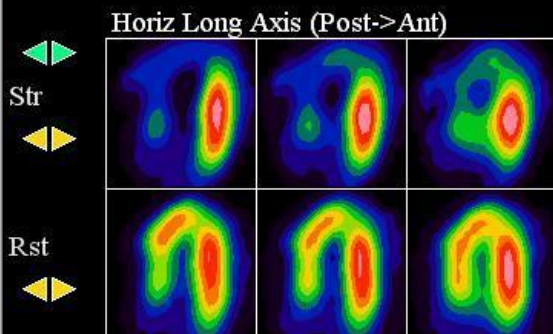


Defect Blackout Map

STRESS [fbp]

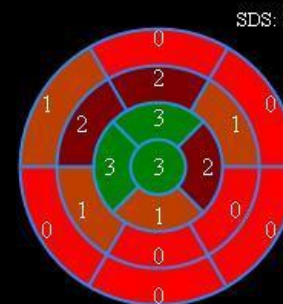
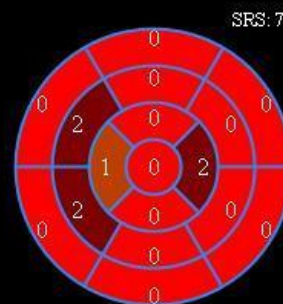
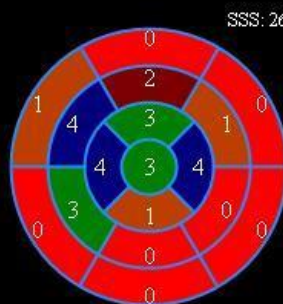
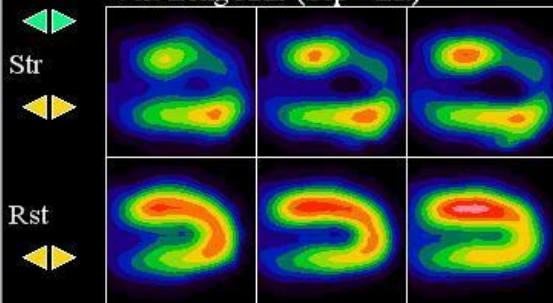
REST [fbp]

Reversibility



Horiz Long Axis (Post->Ant)

Vert Long Axis (Sep->Lat)



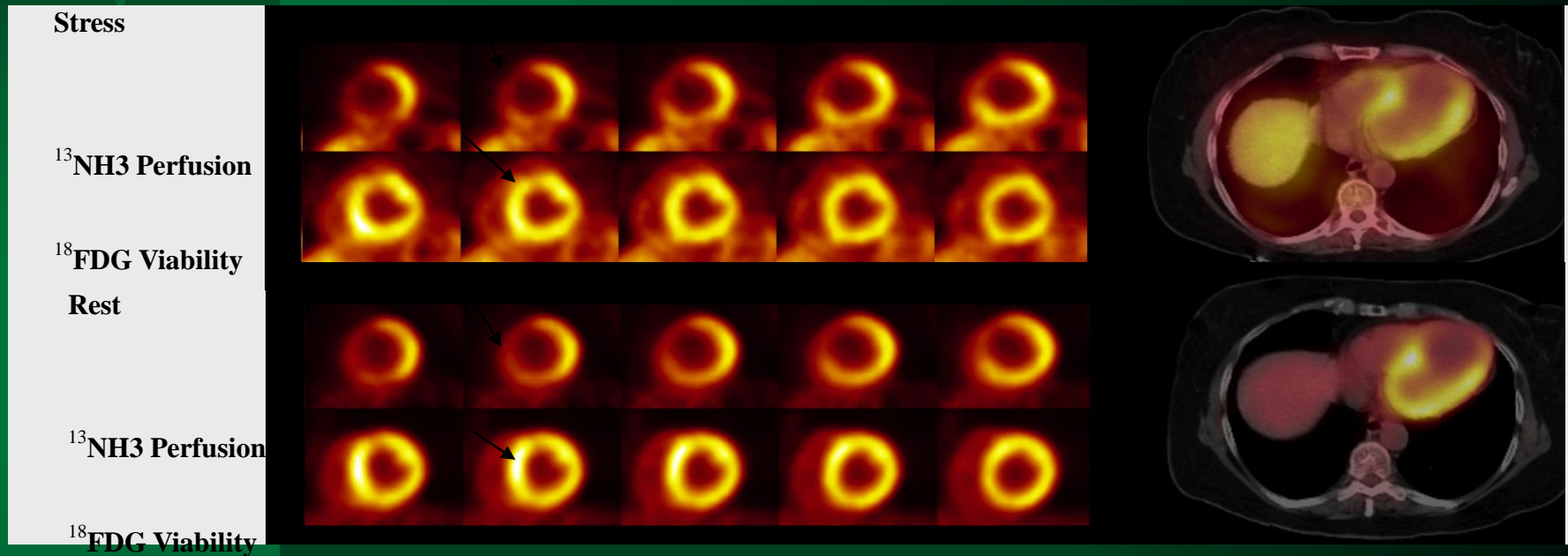
Perf. 0: Normal 1: Equivocal 2: Abnormal 3: Severe 4: Absent



# PET - Scan

- ✓ **The gold standard method for myocardial viability-evaluation of myocardial glucose utilisation with fluorine-18 fluorodeoxyglucose (FDG)**
- ✓ **Principle** -Under fasting conditions the normal myocardium primarily utilises free fatty acids. In ischaemic myocardium glucose becomes an important energy substrate, FDG uptake will be enhanced.
- ✓  **$^{13}\text{NH}_3$  for Perfusion**
- ✓  **$^{18}\text{FDG}$  for Metabolism**

# PET•CT cardiac perfusion and viability mismatch study



Perfusion-viability mismatch in  $^{13}\text{N}$ - $\text{NH}_3$  PET perfusion and  $^{18}\text{F}$ -FDG PET viability study

Mismatch indicative of ischemic but viable myocardium - capable of return to normal contractility with revascularization

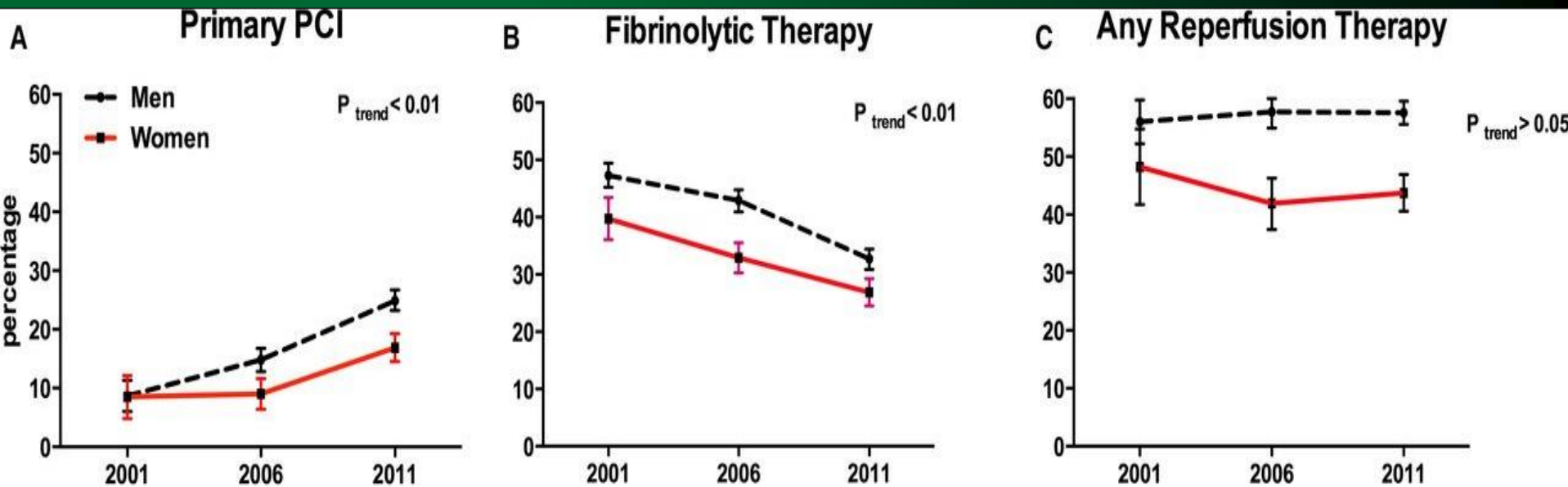
Patient identified as a candidate for revascularization.



# THERAPEUTICS



# Primary PCI ... on the rise



# Culprit Artery – Only Versus Multivessel PCI

- Previous clinical practice guidelines recommended against PCI of nonculprit artery stenoses at the time of primary PCI in hemodynamically stable patients with STEMI, based primarily on the results of nonrandomized studies and meta-analyses and safety concerns.
- Four RCTs (PRAMI, CvLPRIT, DANAMI 3 PRIMULTI, PRAGUE-13) have since suggested that a strategy of multivessel PCI, either at the time of primary PCI or as a planned, staged procedure, may be safe and beneficial in **selected patients** with STEMI.
- On the basis of these findings, the prior Class III-harm recommendation with regard to multivessel primary PCI in hemodynamically stable patients with STEMI has been upgraded and modified to a Class IIb recommendation to include consideration of multivessel PCI, either at the time of primary PCI or as a planned, staged procedure.
- The writing committee emphasizes that this change should not be interpreted as endorsing the routine performance of multivessel PCI in all patients with STEMI and multivessel disease. Rather, when considering the indications for and timing of multivessel PCI, physicians should integrate clinical data, lesion severity/complexity, and risk of contrast nephropathy to determine the optimal strategy.



## WHY TO OPEN UP A CTO ?

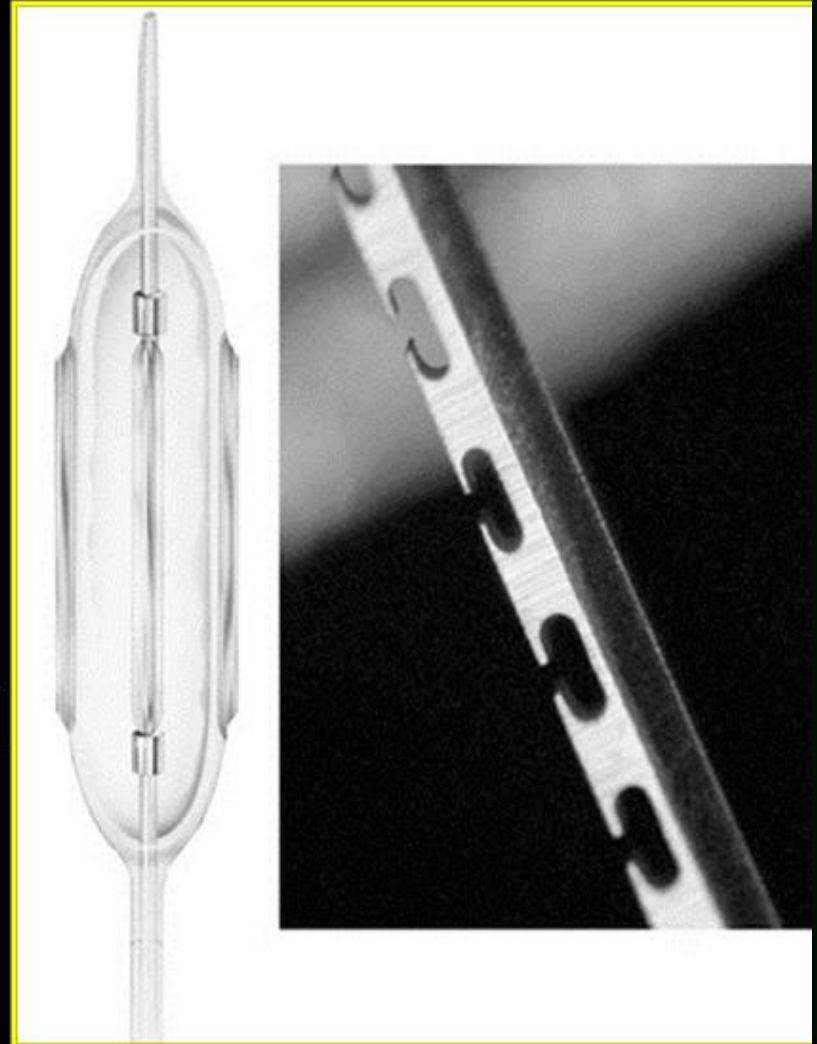
Significant clinical problem	JACC intvn 2009;2:489 –97
Similar risk to non CTO PCI	JACC intvn 2009;2:489 –97
Angina relief	(FACTOR TRIAL-2010)
Improved L V function	JACC 2006;47:721–5
Improved tolerance of a future ACS	JACC intvn 2009;2:1128 –34
Potentially better survival with successful PCI	AmHeart J 2010;160:179-87
Avoidance of CABG	AmHeart J 2010;160:179-87



- Cutting Balloon Angioplasty (CBA)
- Directional Coronary Atherectomy (DCA)
- Laser angioplasty
- **Percutaneous Transluminal Rotational Atherectomy (PTRA)**

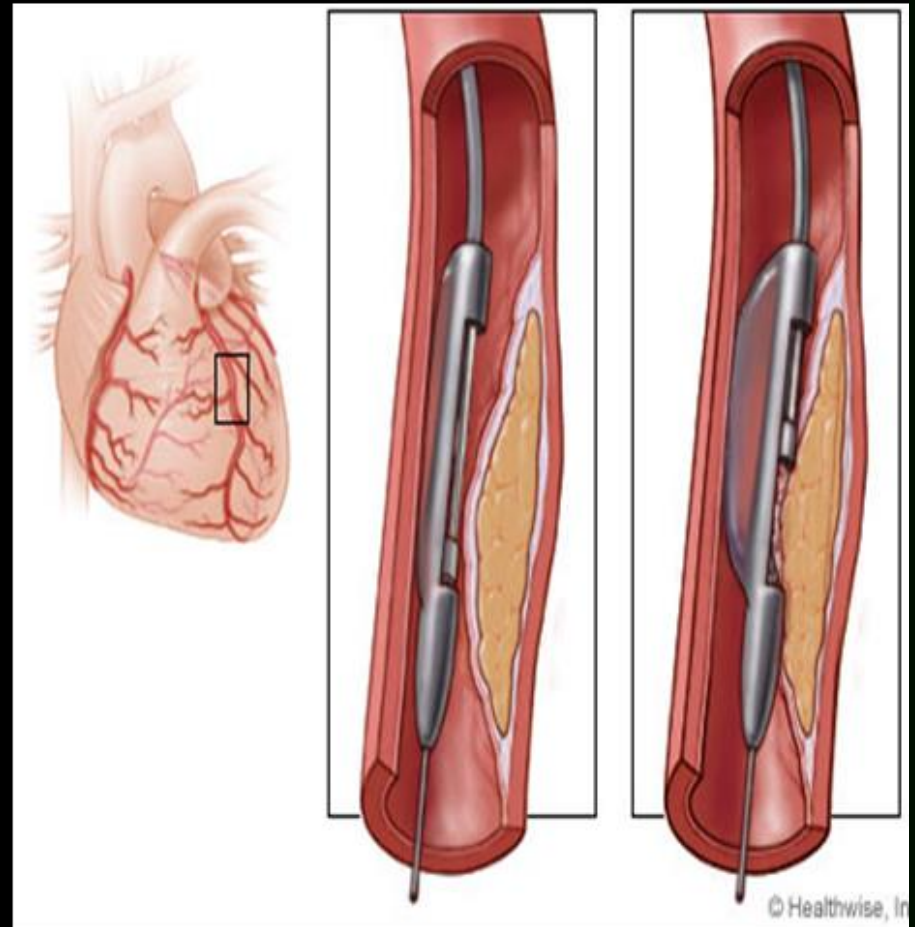
# CUTTING BALLOON ANGIOPLASTY

- Makes controlled microincisions into the atheromatous plaque (radial atherectomy)
- Lesser barotrauma to vessel



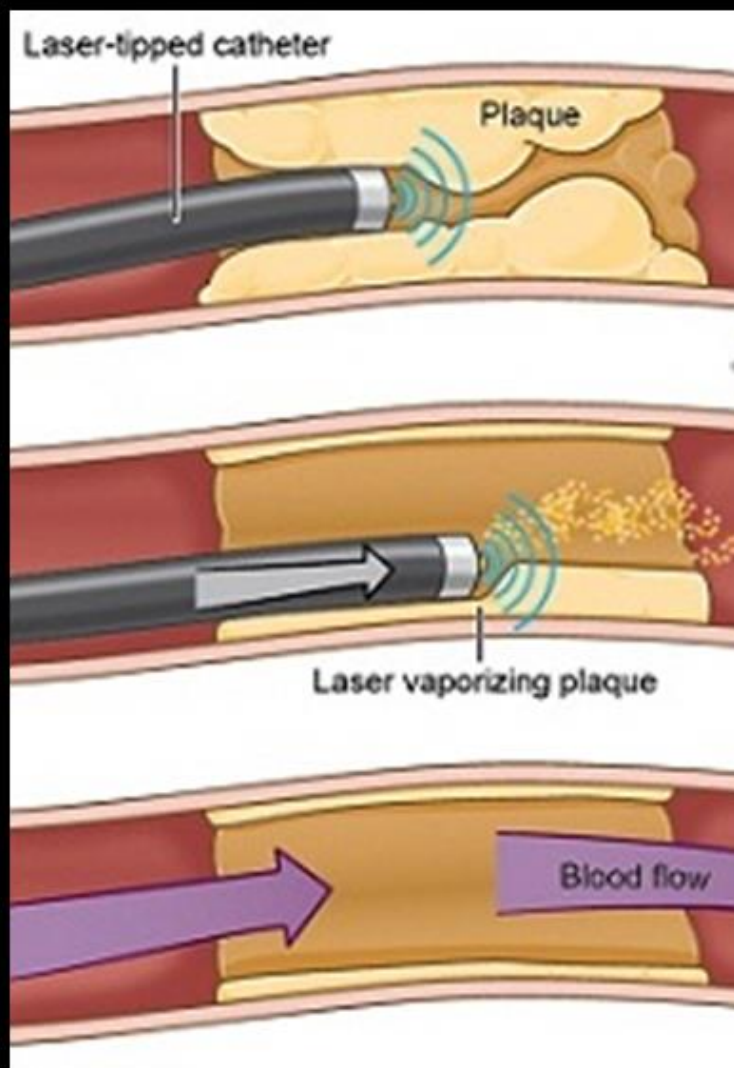
# DIRECTIONAL ATHERECTOMY

- Apposes a cutting device to a section of the plaque.
- Manually advances the cutting cup
- Debulking procedure



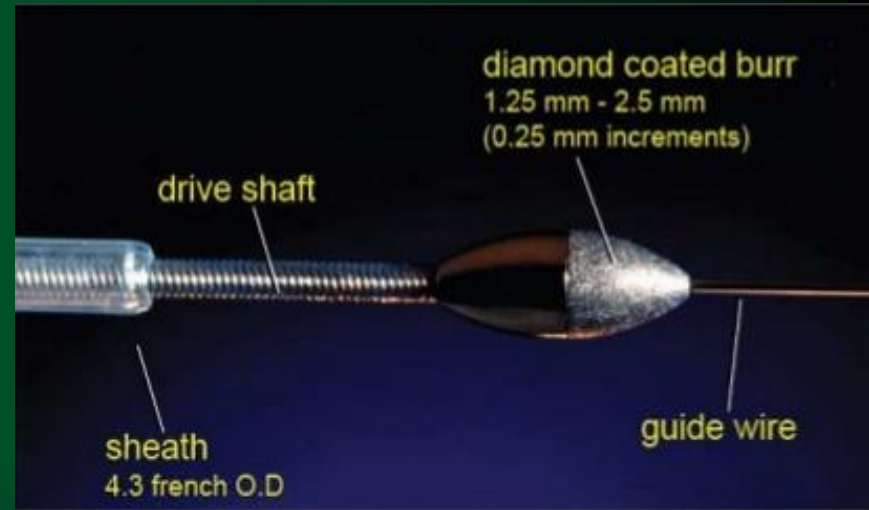
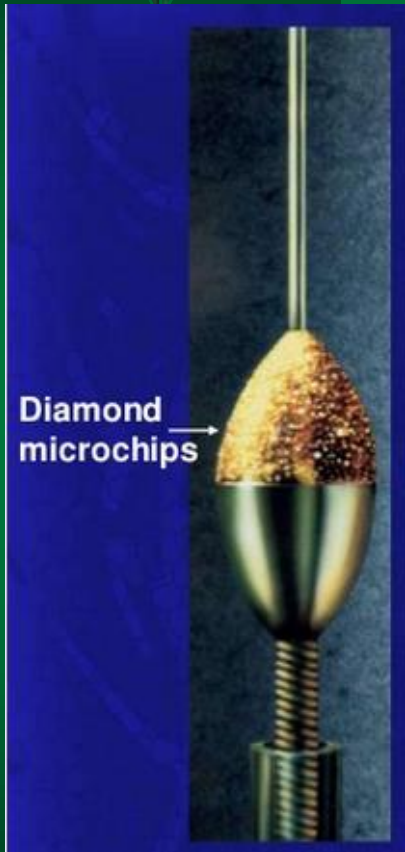
# LASER ANGIOPLASTY

- Laser beam is channelled in pulse/continuous from to help in debulking.
- Photo-thermal effect, photo-acoustic effect, photo-chemical dissociation





# ROTABLATOR



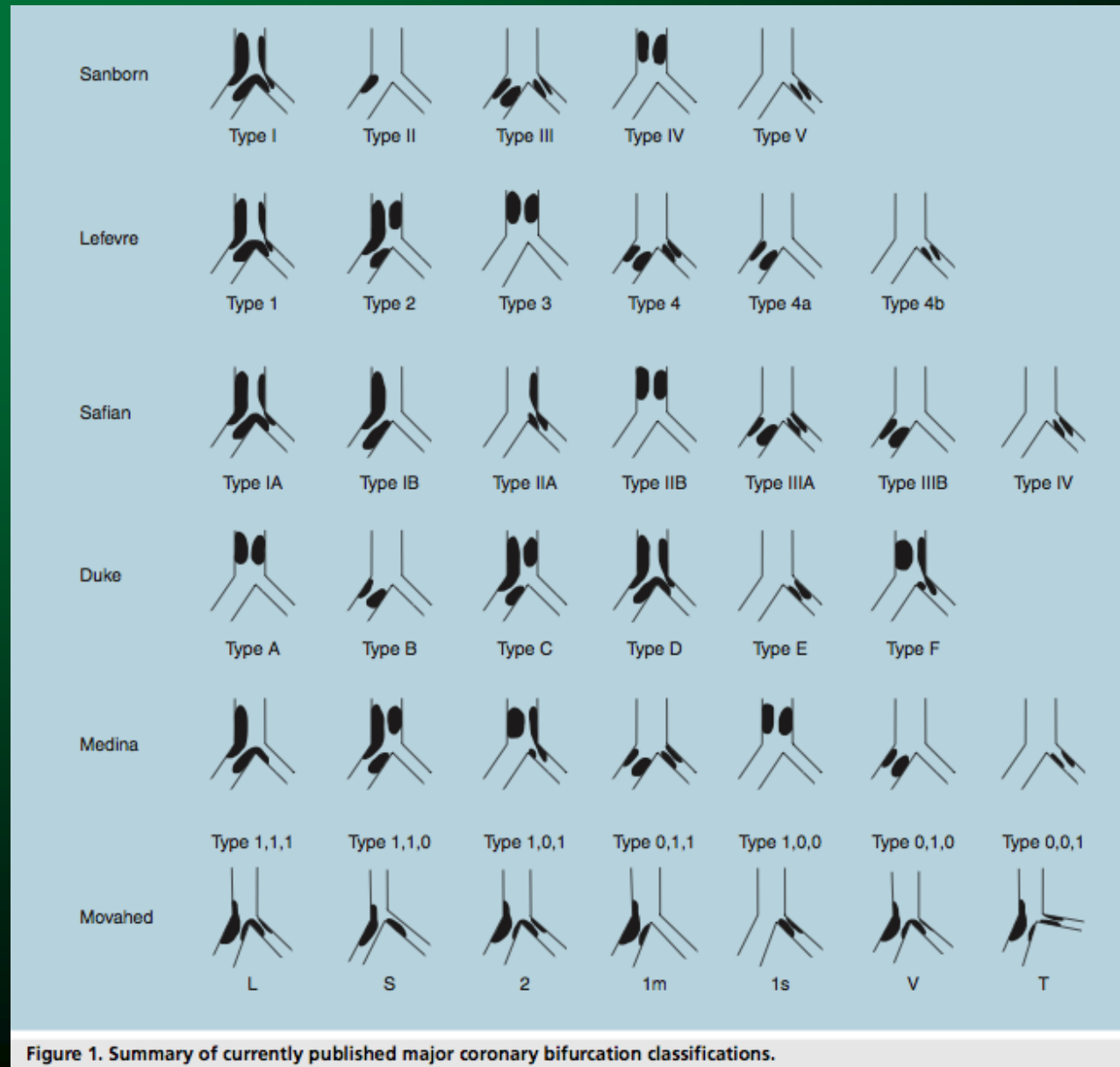


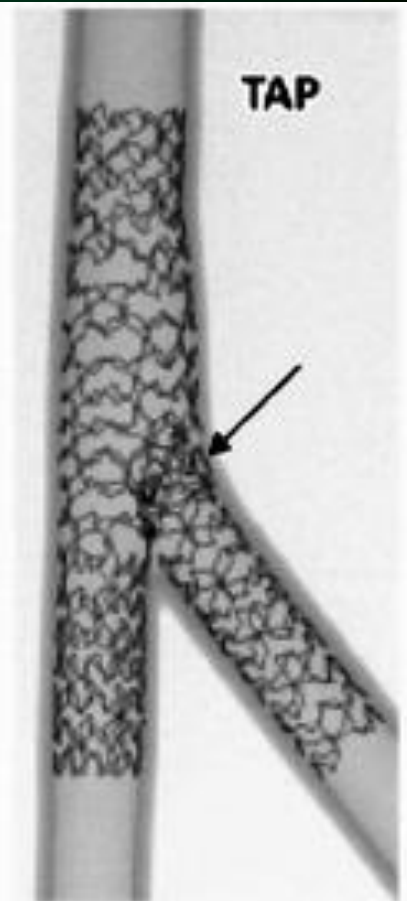
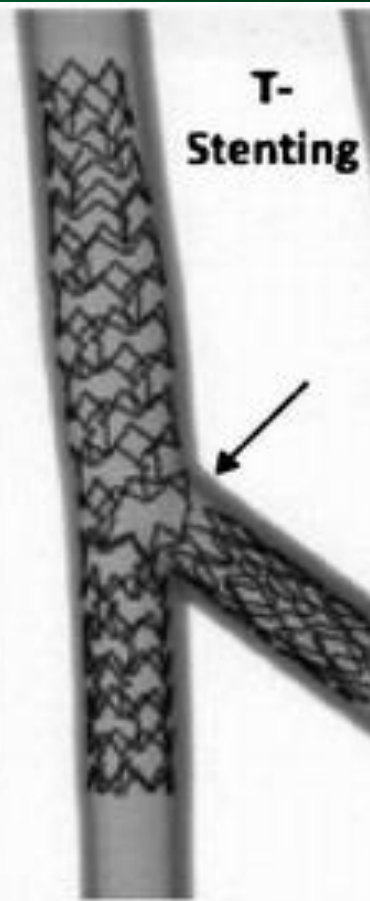
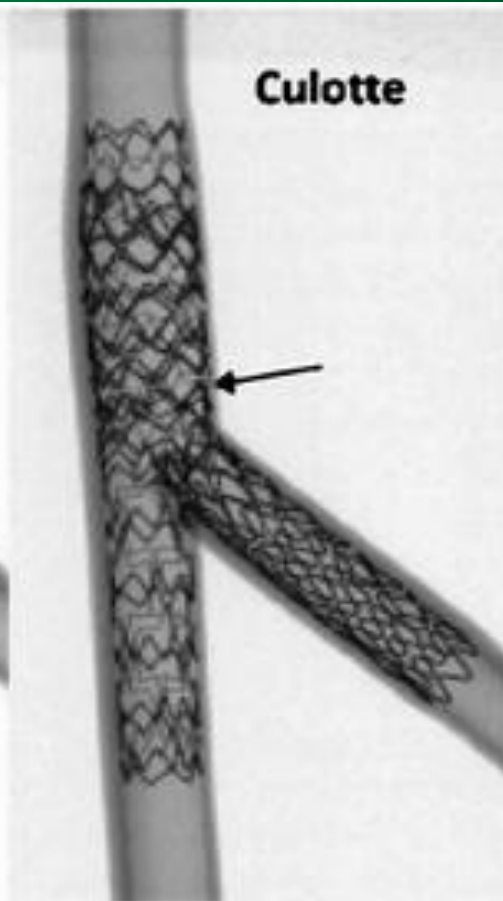
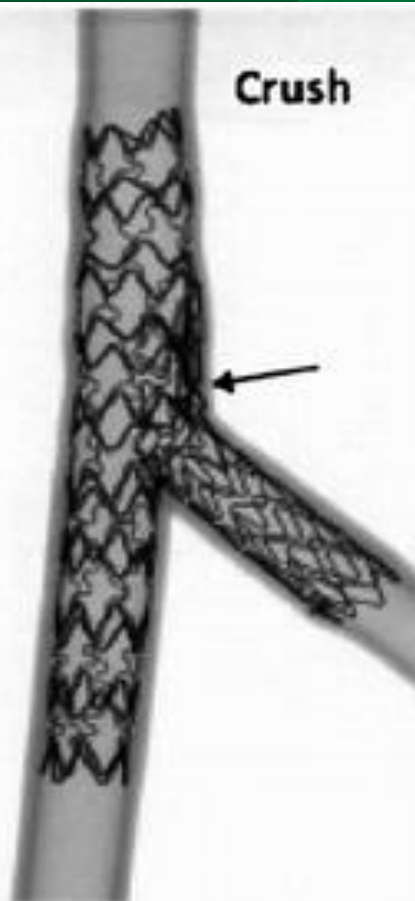
Encouraging results were reported from the ‘Synergy Between Percutaneous Coronary Intervention with Taxus and Cardiac Surgery’ (SYNTAX) trial

	PCI (n = 358)	CABG (n = 357)	P-value
1-year clinical outcomes			
Death (%)	4.2	4.4	0.88
Stroke (%)	0.3	2.7	0.0009
MI (%)	4.3	4.1	0.97
Revascularization (%)	12	6.7	0.02
ST or graft occlusion (%)	2.7	3.7	0.49
Overall MACCE (%)	15.8	13.6	0.44
MACCE low SYNTAX score (0–17)	7.7	13	0.19
MACCE intermediate SYNTAX score (23–32)	12.6	15.5	0.54
MACCE high SYNTAX score (≥ 33)	25.3	12.9	0.008
3-year clinical outcomes			
Death (%)	7.3	8.4	0.64
Stroke (%)	1.2	4	0.02
MI (%)	6.9	4.1	0.14
Revascularization (%)	20	11.7	0.004
ST or graft occlusion (%)	4.1	3.7	0.80
Overall MACCE (%)	26.8	22.3	0.20
MACCE low SYNTAX score (0–17)	18	23	0.33
MACCE intermediate SYNTAX score (23–32)	23.4	23.4	0.90
MACCE high SYNTAX score (≥ 33)	37.3	21.2	0.003



# Coronary Bifurcation Stenosis



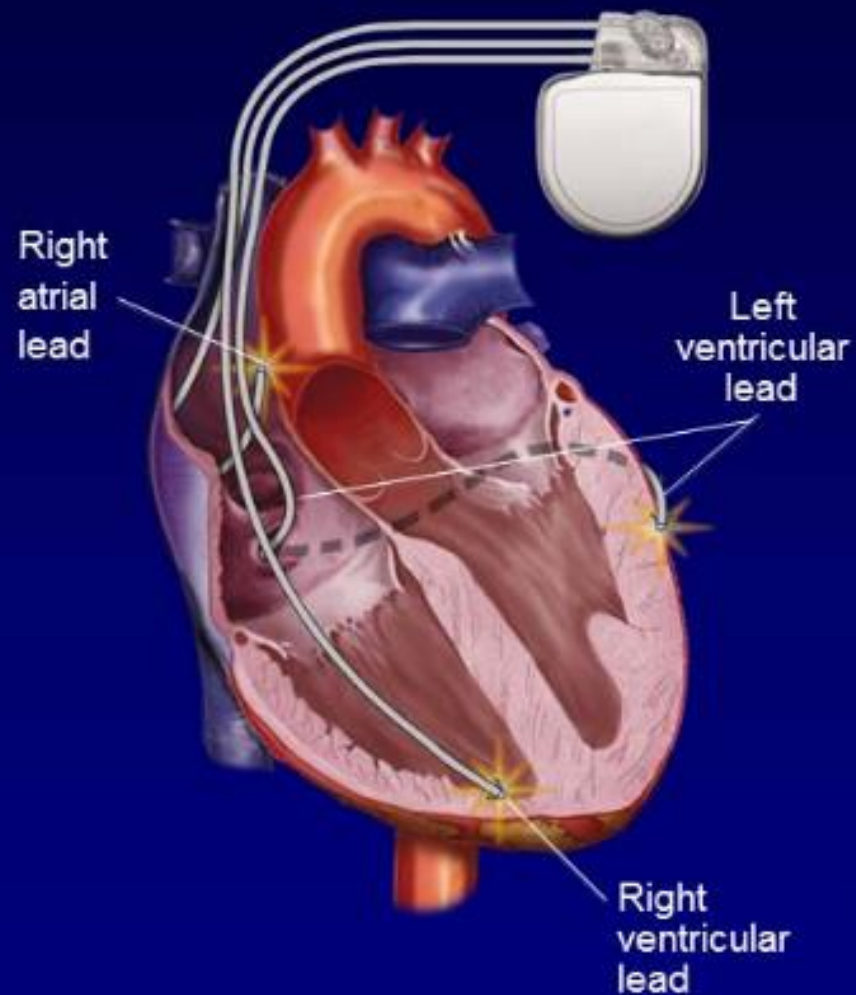




# CARDIAC RESYNCHRONIZA TION THERAPY (CRT-D)

# CRT-D System Adds LV Lead to an ICD System

- Provides Biventricular Pacing With ICD Capability
- Composed of:
  - Pulse generator
  - Right atrial pacing and sensing lead
  - Right ventricular pacing plus defibrillation lead
  - Left ventricular pacing lead
- Implant of right atrial and right ventricular leads is the same as a dual chamber ICD





# CRT Indications

- ✓ LBBB with QRS duration  $>150$  ms.
- ✓ chronic HF patients and LVEF  $\leq 35\%$
- ✓ who remain in NYHA functional class II, III and ambulatory IV despite adequate medical treatment.



# LEADLESS PACEMAKERS



## EVOLUTION OF PACEMAKER TECHNOLOGY



**1958**

Weight: 73.4g  
Size: 35cc



**1981**

Weight: 55g  
Size: 25cc



**1995**

Weight: 14g  
Size: 6cc



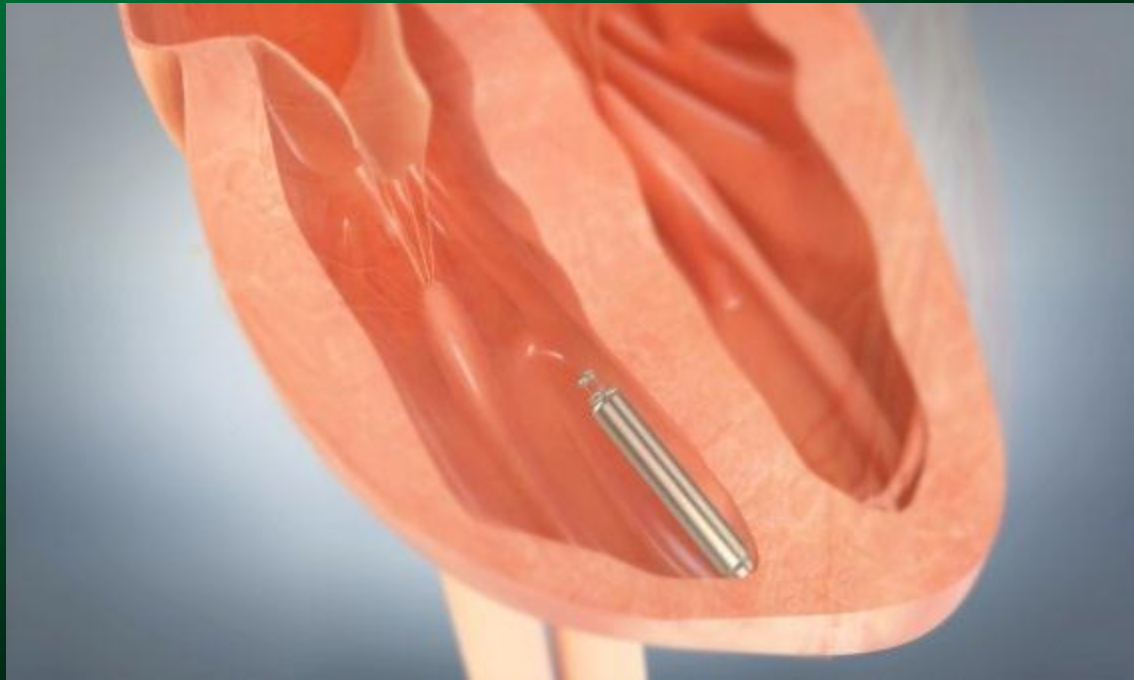
**2009**

Weight: 23g  
Size: 12.8cc



**2013**

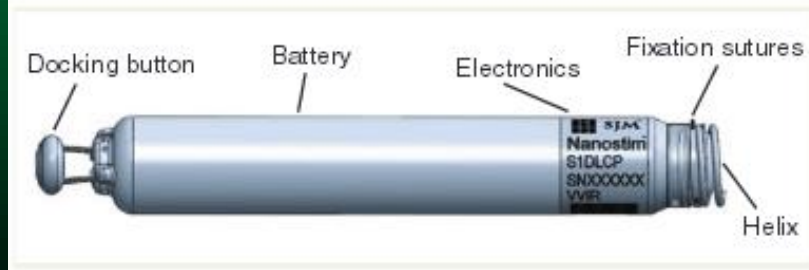
Weight: 2g  
Size: 1cc



# Leadless systems for right ventricle pacing

## St. Jude Medical Nanostim™ leadless pacemaker (LCP)

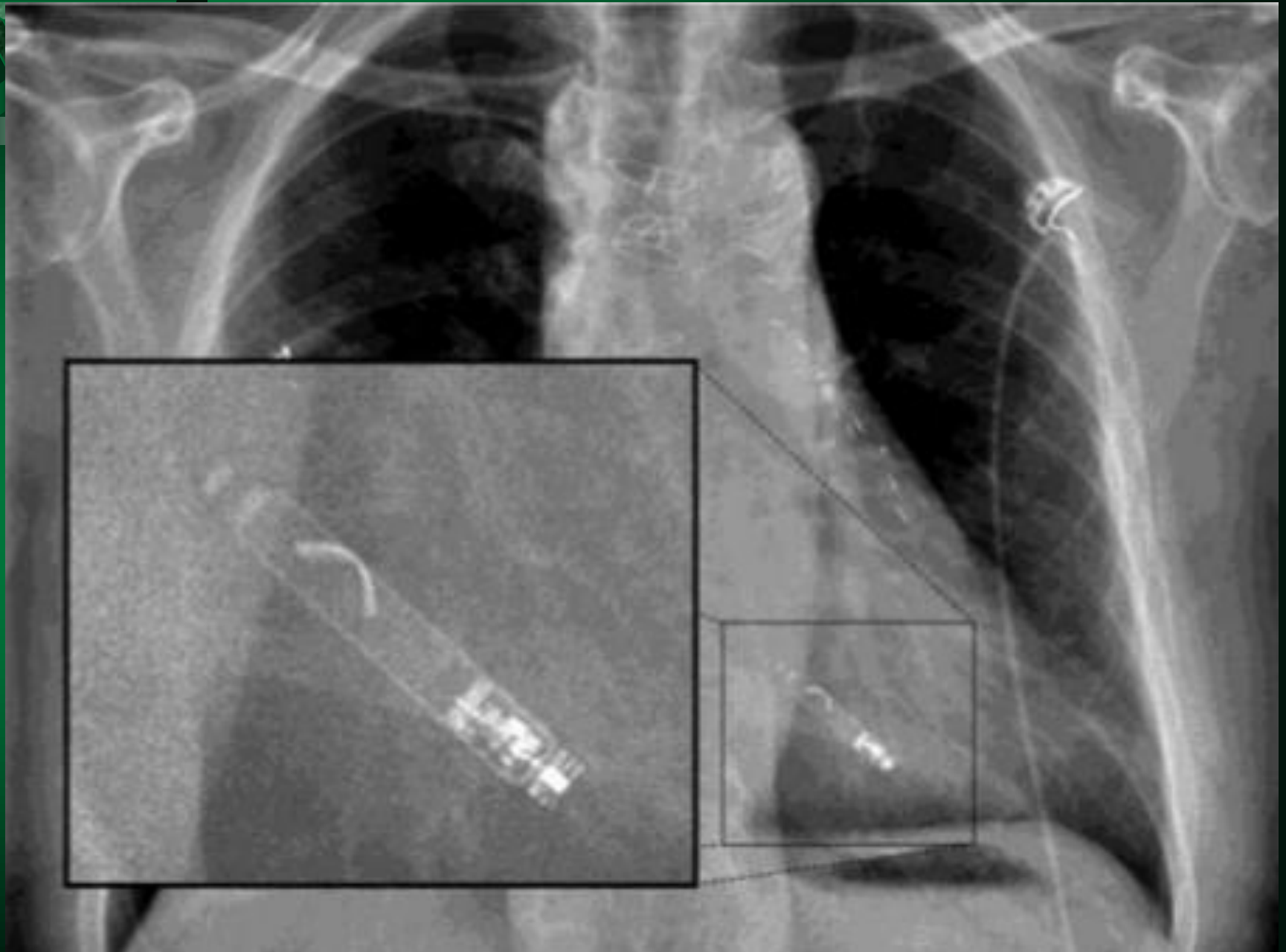
- Available in 2012
- VVI(R)
- Screw-in helix



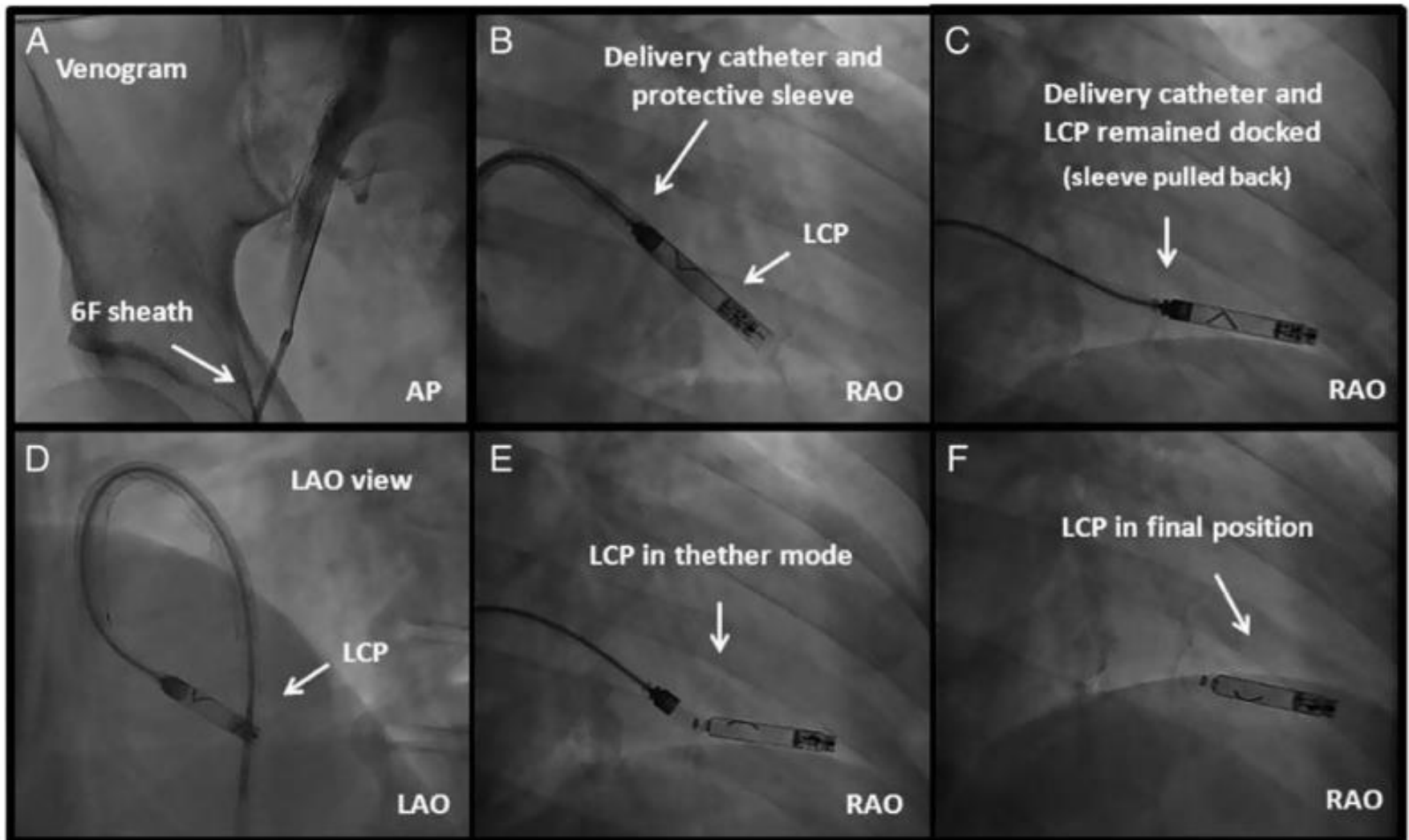
## Medtronic Micra™ Transcatheter Pacing System (TPS)

- Available in 2013
- VVI(R)
- Self-expanding nitinol tines





# Implantation (LCP)





# **Leadless Pacemaker: Potential Advantages**

**Less Invasive**

**More cosmetic for patient (“invisible”)**

**More readily MRI conditional**

**Reduced length of hospital stay (same day)**

**Fewer acute and chronic complications (infection, erosion)**



# Leadless Pacemaker: Potential Disadvantages

Only VVIR pacing

External communication interference

Memory capacity and download limitations

Large diameter sheaths

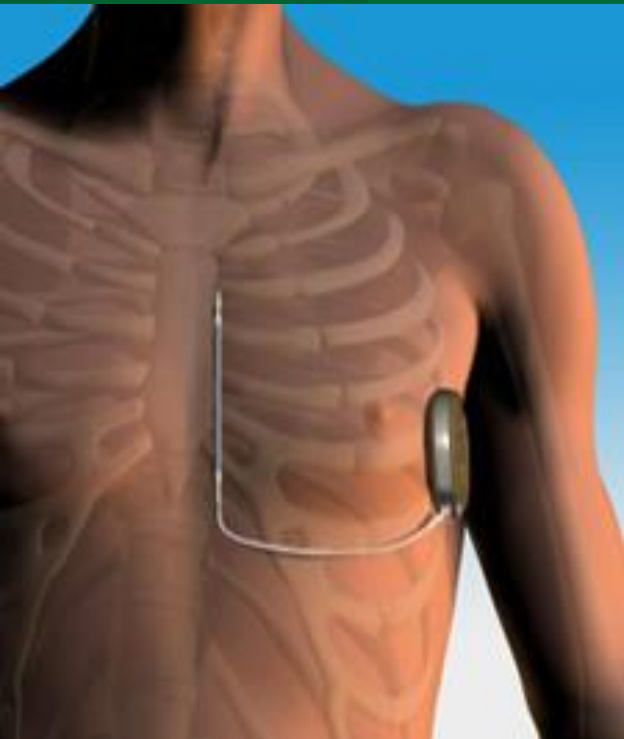
Embolization / retrieval

Repositioning difficulty for high threshold

Longevity limitations ( Less battery life )



# Subcutaneous AICD





# AICD- Indications Class I

- Structural heart disease, sustained VT
- Syncope of undetermined origin, inducible VT or VF at electrophysiologic study (EPS)
- Left ventricular ejection fraction (LVEF)  $\leq 35\%$  due to prior MI, at least 40 days post-MI, NYHA class II or III
- LVEF  $\leq 35\%$ , NYHA class II or III
- LVEF  $\leq 30\%$  due to prior MI, at least 40 days post-MI
- LVEF  $\leq 40\%$  due to prior MI, inducible VT or VF at EPS



# AICD- Indications

## Class IIa

- Unexplained syncope, significant LV dysfunction, nonischemic cardiomyopathy
- Sustained VT, normal or near-normal ventricular function
- Hypertrophic cardiomyopathy with 1 or more major risk factors
- Arrhythmogenic right ventricular dysplasia/cardiomyopathy (ARVD/C) with 1 or more risk factors for sudden cardiac death (SCD)
- Long QT syndrome, syncope or VT while receiving beta-blockers
- Nonhospitalized patients awaiting heart transplant
- Brugada syndrome, syncope or VT
- Catecholaminergic polymorphic VT, syncope or VT while receiving beta-blockers
- Cardiac sarcoidosis, giant cell myocarditis, or Chagas disease



# Advantages:

- ✓ Less invasive surgery
- ✓ Eliminates potential for infection and damage to venous system
- ✓ May be implanted using anatomical landmarks
- ✓ Potential for less inappropriate shocks in children



# Disadvantages:

## ✦ Size

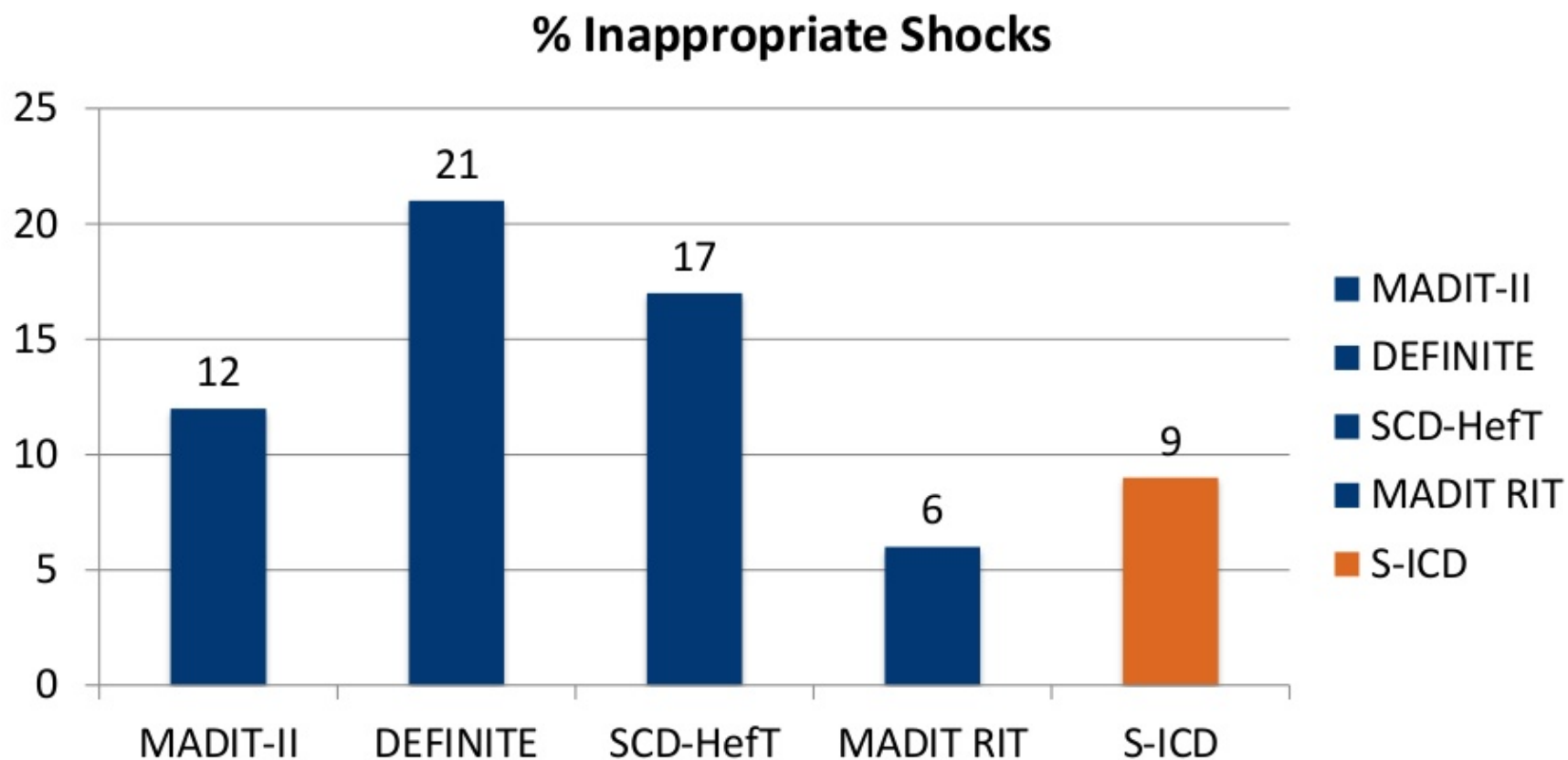
- Twice that of current T-ICD

## ✦ Battery life

- 5 years as opposed to upwards of 10 with T-ICD

## ✦ Does not provide anti-tachycardia pacing (ATP) or bradycardia pacing

# Inappropriate Shocks





# Transcatheter aortic valve implantation

TAVI is indicated in patients with  
severe symptomatic AS  
who are not suitable for AVR,  
have a  $>1$ -year life expectancy  
and are likely to gain improvement  
in quality of life

# From PVT to Edwards balloon expandable Valves

## Edwards Valves

2000: PVT Valve

2003-2004

2005-2009

2009

Percutaneous Heart Valve

Cribier Edwards

Edwards Sapien

Edwards Sapien XT



Bovine pericardium Equine pericardium Treated bovine pericardium

Stainl. steel frame Stainl. steel frame Stainl. steel frame

23mm

23mm

23 and 26mm

Next to come  
20mm / 29mm

Next generation

24F

22F

22F, 24F

18F, 19F

TF sheath sizes

# Self expandable Medtronic CoreValve

Generation 1  
25F

Generation 2  
21F

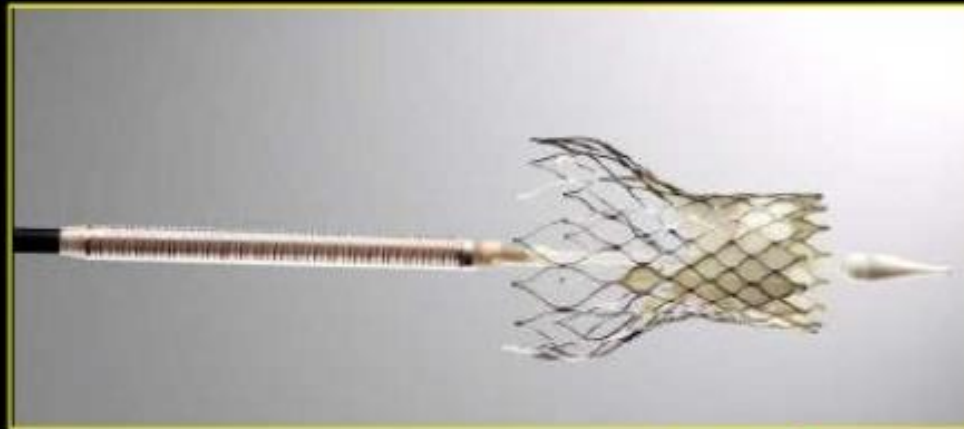
Generation 3  
18F

Generation 4  
18F

2004-2005

From 2006

2010



Porcine pericardium valve  
Nitinol stent

Improved  
delivery ?

### Antegrade technique

The catheter is advanced via the femoral vein, traversing the interatrial septum and the mitral valve, and is positioned within the diseased aortic valve.

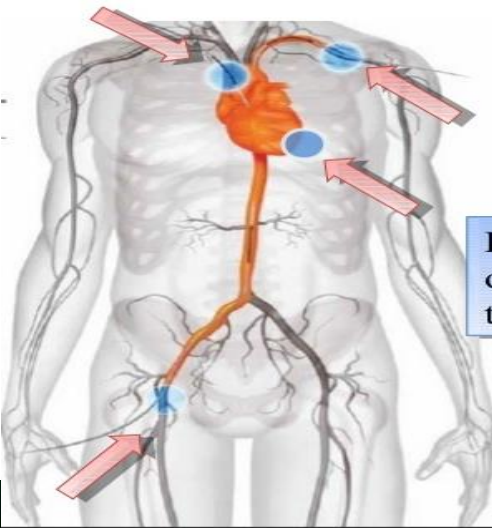
#### Advantages

- Femoral vein accommodates the large catheter sheath
- Easy management of peripheral access site

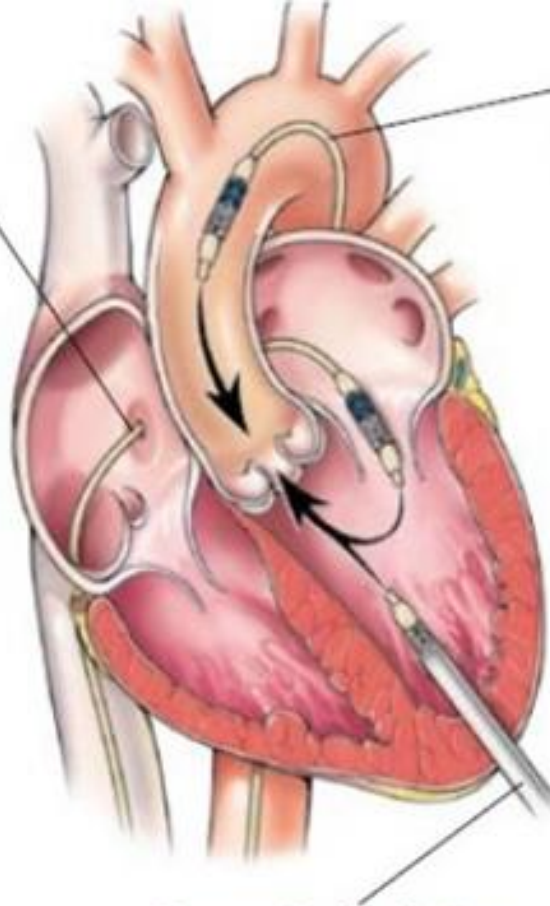
#### Disadvantages

- Risk of mitral valve injury and severe mitral valve regurgitation
- Correctly positioning the prosthetic valve can be challenging

This technique is no longer in use.



Percutaneous  
or Cut-down  
technique



### Retrograde or transfemoral technique

The catheter is advanced to the stenotic aortic valve via the femoral artery.

#### Advantages

- Faster, technically easier than antegrade approach

#### Disadvantages

- Potential for injury to the aortofemoral vessels
- Crossing the stenotic aortic valve can be challenging

### Transapical technique

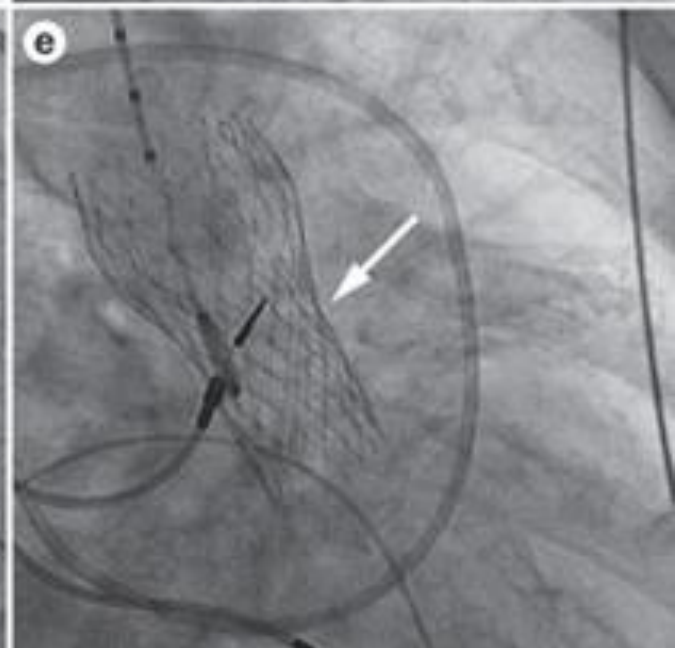
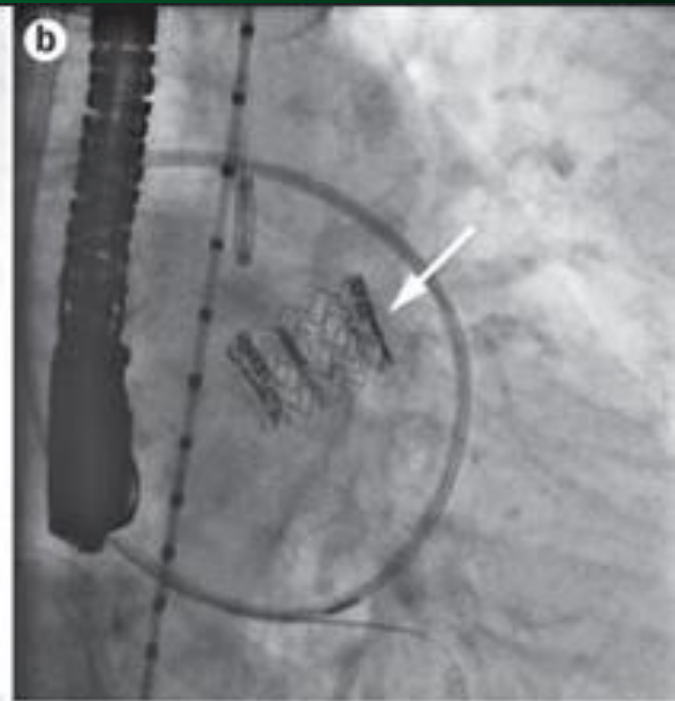
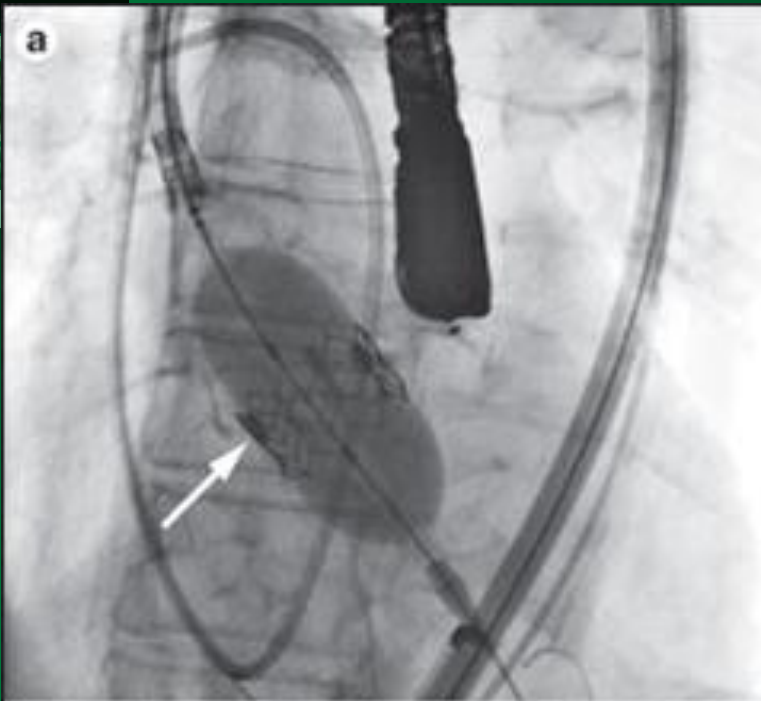
A valve delivery system is inserted via a small intercostal incision. The apex of the left ventricle is punctured, and the prosthetic valve is positioned within the stenotic aortic valve.

#### Advantages

- Access to the stenotic valve is more direct
- Avoids potential complications of a large peripheral access site

#### Disadvantages

- Potential for complications related to puncture of the left ventricle
- Requires general anesthesia and chest tubes





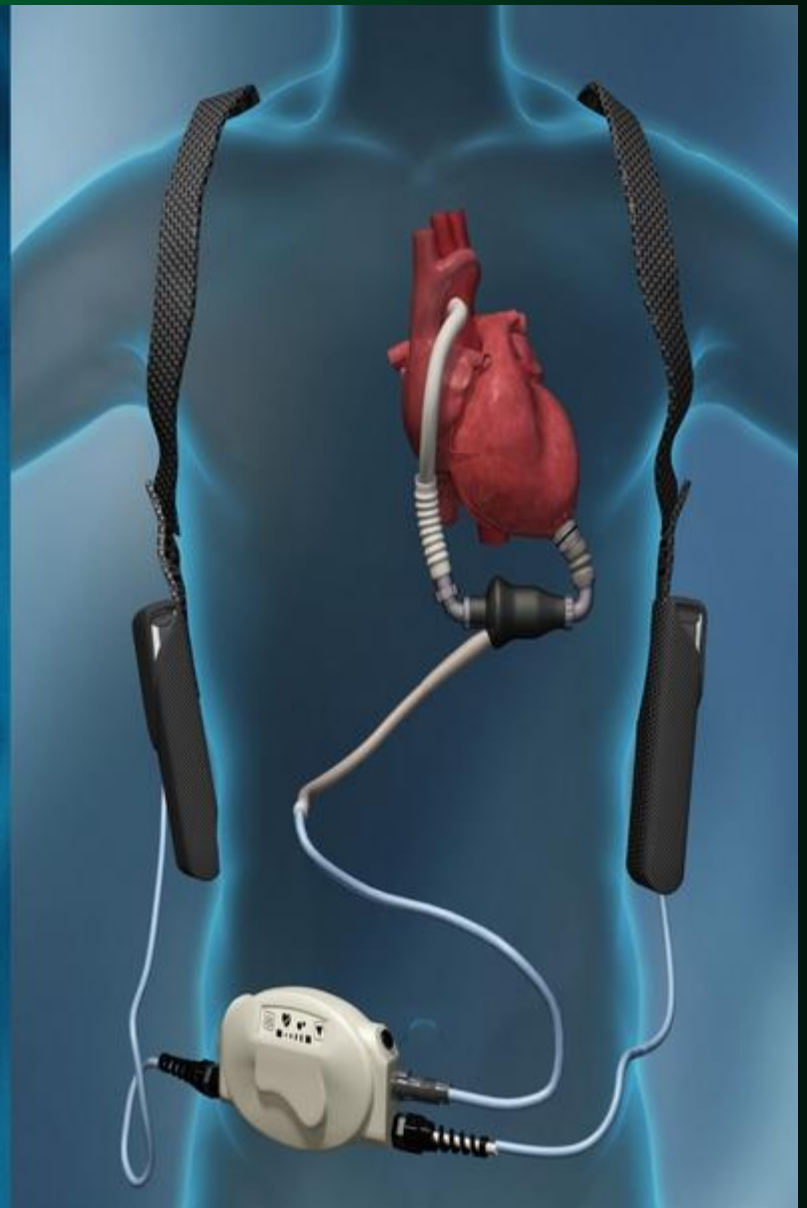
# Contraindications to TAVI

- ✓ Inadequate annulus size (<18 mm, >29 mm),
- ✓ Active endocarditis,
- ✓ Symmetric valve calcification,
- ✓ Short distance between the annulus and the coronary ostium,
- ✓ Plaques with mobile thrombi in the ascending aorta,
- ✓ Inadequate vascular access ,
- ✓ Haemodynamic instability,
- ✓ Severe LV dysfunction.



# Ventricular Assist Device (VAD)

- A mechanical pump that is surgically attached to one of the heart's ventricles to augment or replace native ventricular function
- Can be used for the left (L VAD), right (R VAD), or both ventricles (Bi VAD)
- Are powered by external power sources that connect to the implanted pump via a percutaneous lead (driveline) that exits the body on the right abdomen
- Pump output flow can be pulsatile or nonpulsatile







# ENHANCED EXTERNAL COUNTERPULSATION (EECP)



- ✓ Have chronic stable angina
- ✓ Are not receiving adequate relief from angina by taking nitrates
- ✓ Do not qualify as a candidate for invasive procedures (bypass surgery, angioplasty, or stenting)



- ✓ A set of cuffs is wrapped around the calves, thighs and buttocks. These cuffs are sequentially inflated followed by the rapid release of pressure on deflation.
- ✓ Inflation and deflation are electronically synchronized with the heartbeat and blood pressure using the ECG and blood pressure monitors.



- ✓ EECP may encourage blood vessels to open small channels or collaterals that may eventually create "natural bypass" vessels to provide blood flow to heart muscle. This contributes to the relief of angina symptoms.





# Novel Oral Anticoagulants

## New Oral Anticoagulants

	<b>Dabigatran Pradaxa®</b>	<b>Rivaroxaban Xarelto®</b>	<b>Apixaban Eliquis®</b>	<b>Edoxaban Savaysa®</b>
<b>Atrial fibrillation</b>	FDA approved	FDA approved	FDA approved	Applied for FDA approval 1/8/2014
<b>VTE Treatment</b>	FDA approved	FDA approved	FDA approved	Applied for FDA approval 1/8/2014
<b>VTE Prevention, Ortho surgery</b>	No FDA activity	FDA approved	FDA approved	No FDA activity



# NOACs - Advantages

- ✓ Fixed dose regimen
- ✓ Less drug interaction
- ✓ No food interaction
- ✓ No need for regular blood test



And miles to go...









..... *Thank You*

Filter: Filter 3



Filter: Filter 3



