

Echo to Guide Percutaneous Interventions

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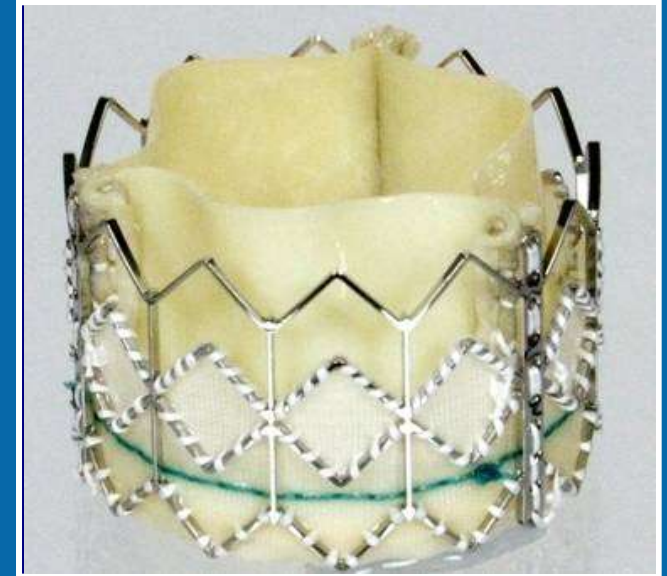
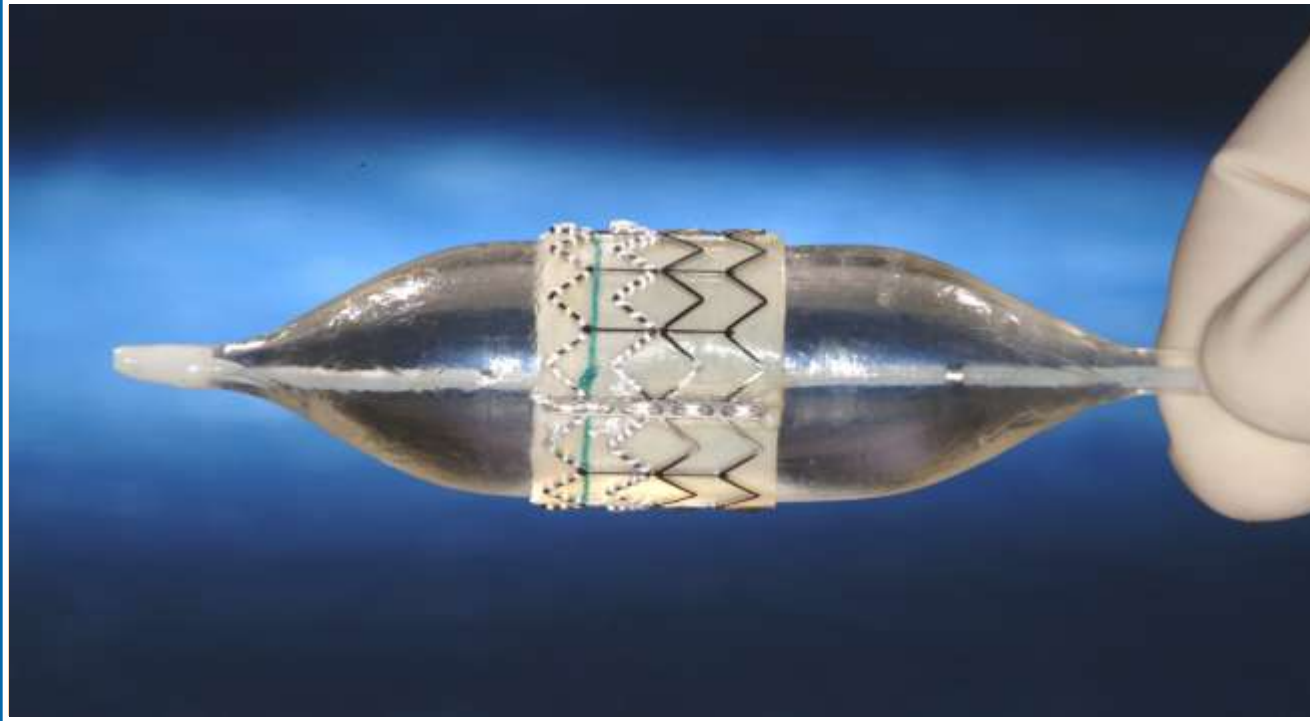
Echocardiography in Guiding Percutaneous Interventions

- Myxomatous mitral valve disease
- Aortic valve disease
- Per-valvular regurgitation

Cribier-Edwards Percutaneous Heart Valve *SYSTEM*

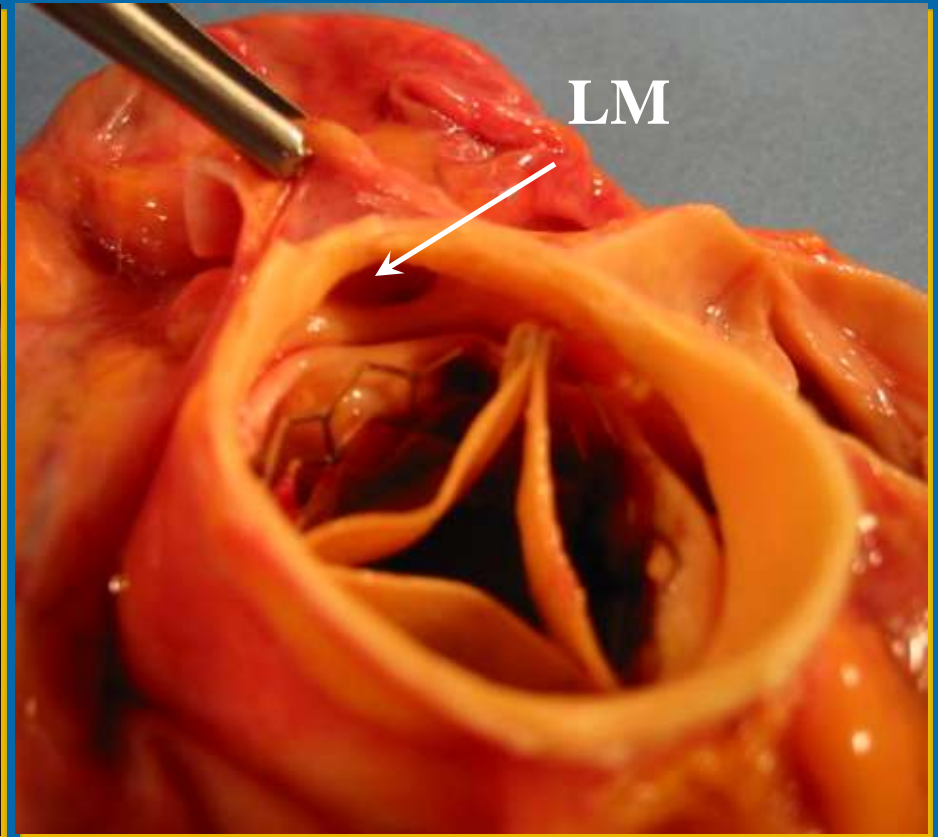
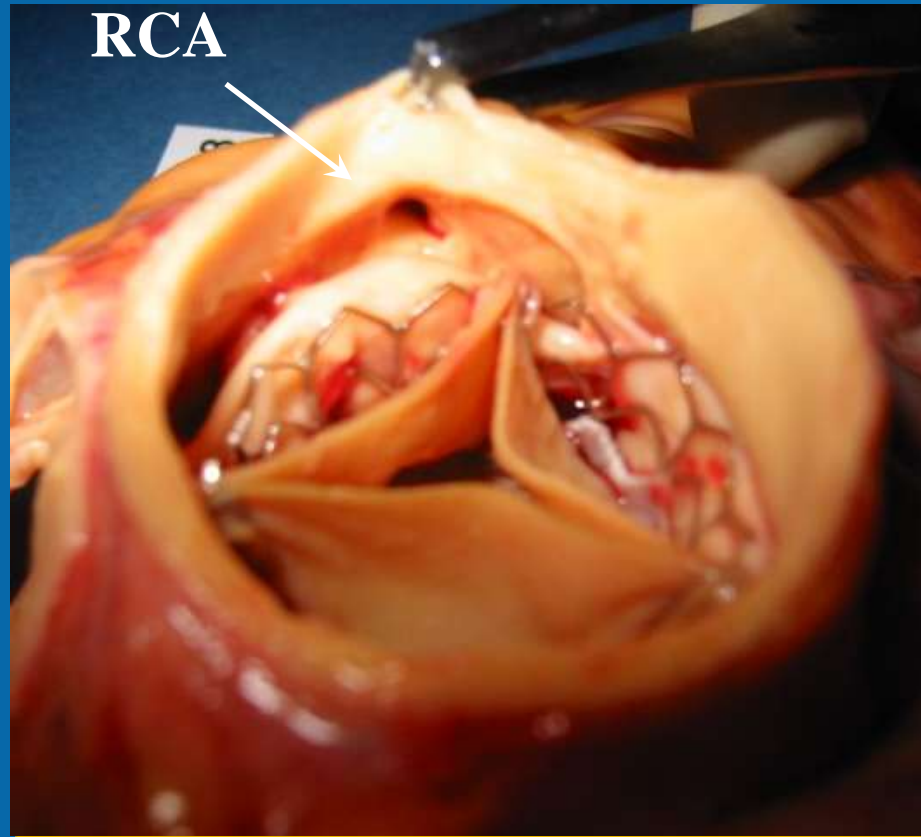


Transcatheter Heart Valve

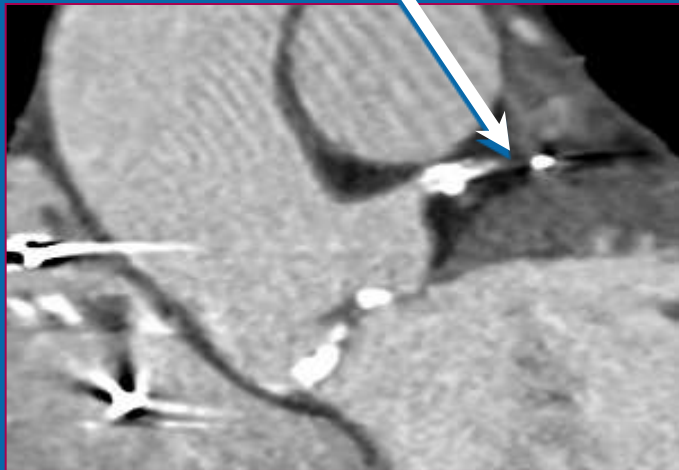


- Balloon expandable
- Stainless steel stent
- Bovine (equine) pericardium
- Optimal hemodynamics

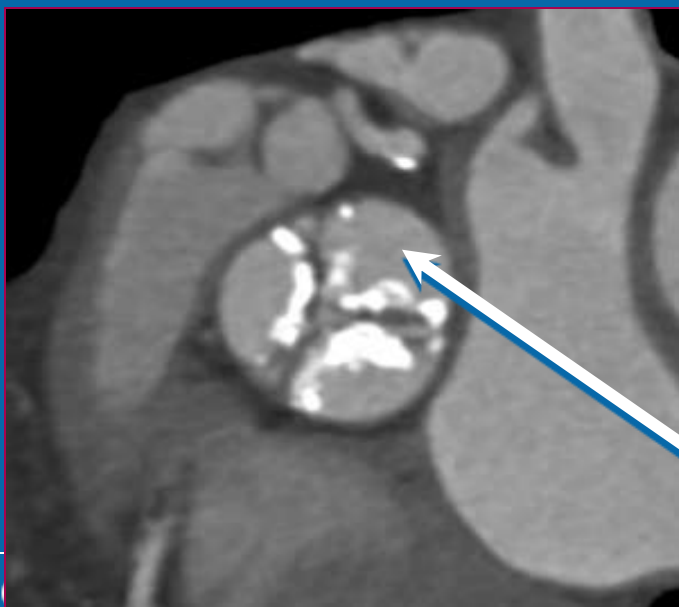
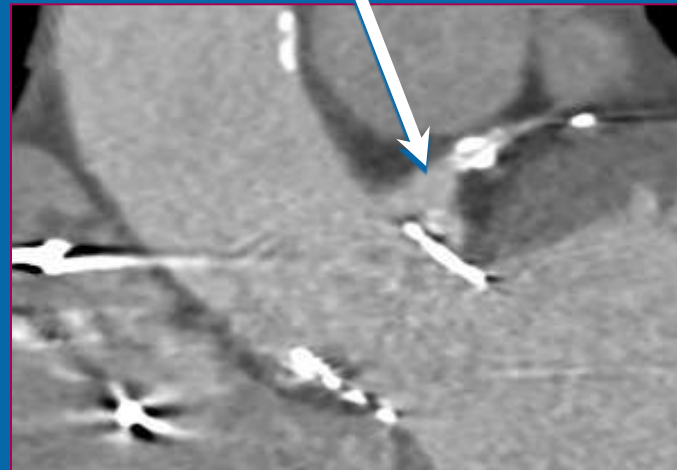
Patent Coronary Arteries



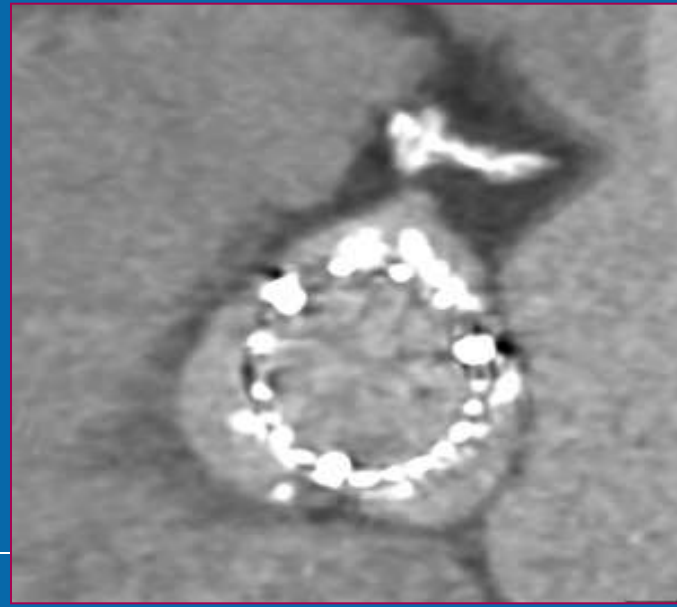
Left main ostium

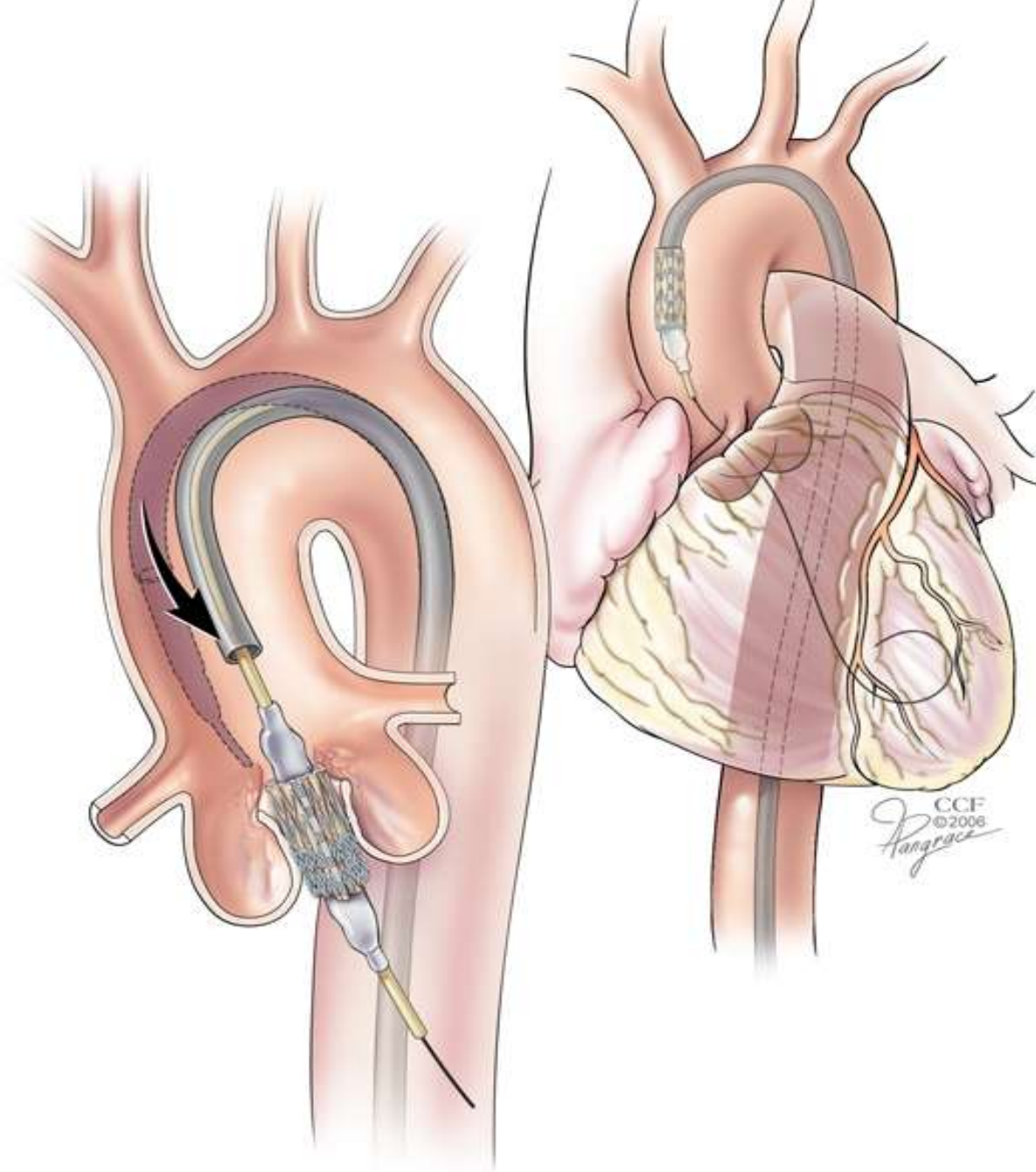


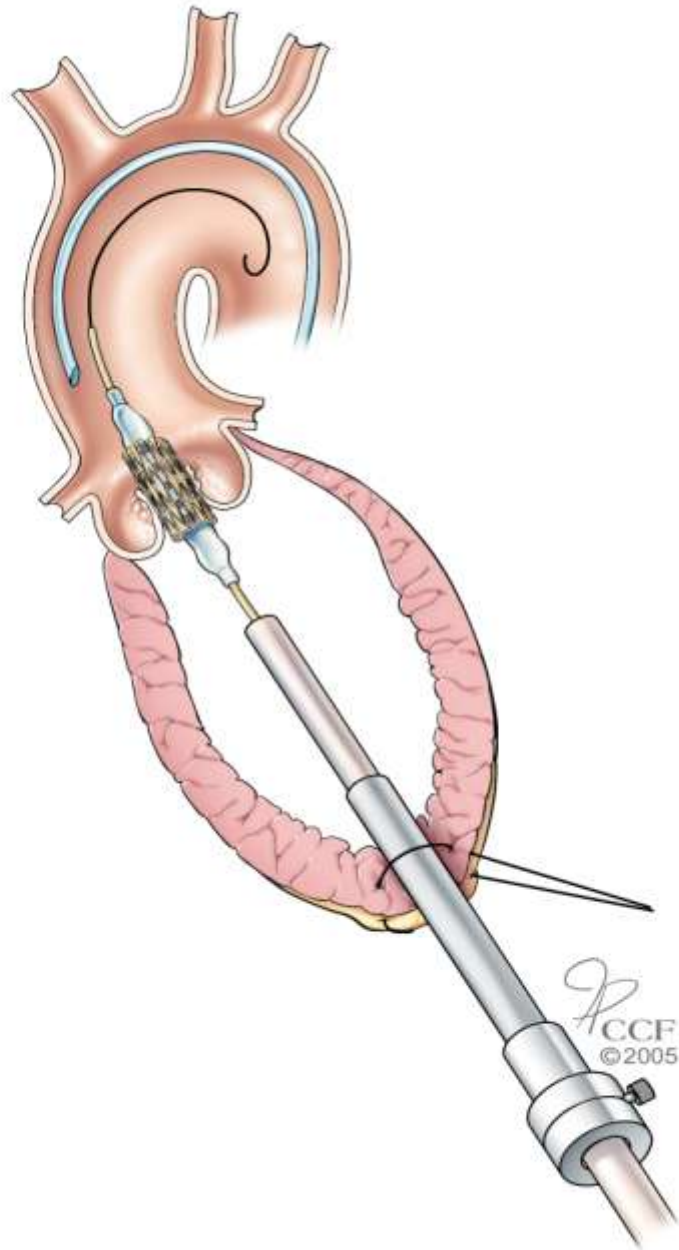
Left main ostium

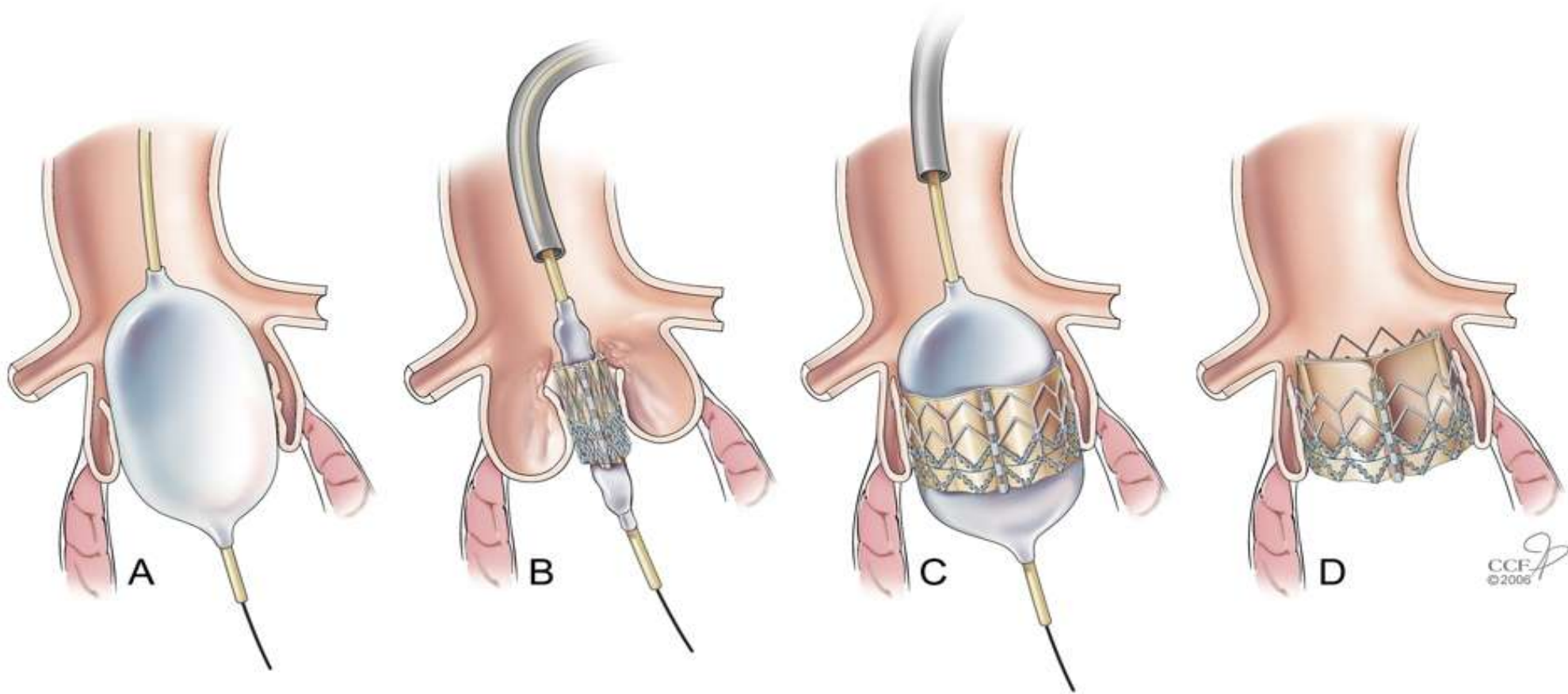


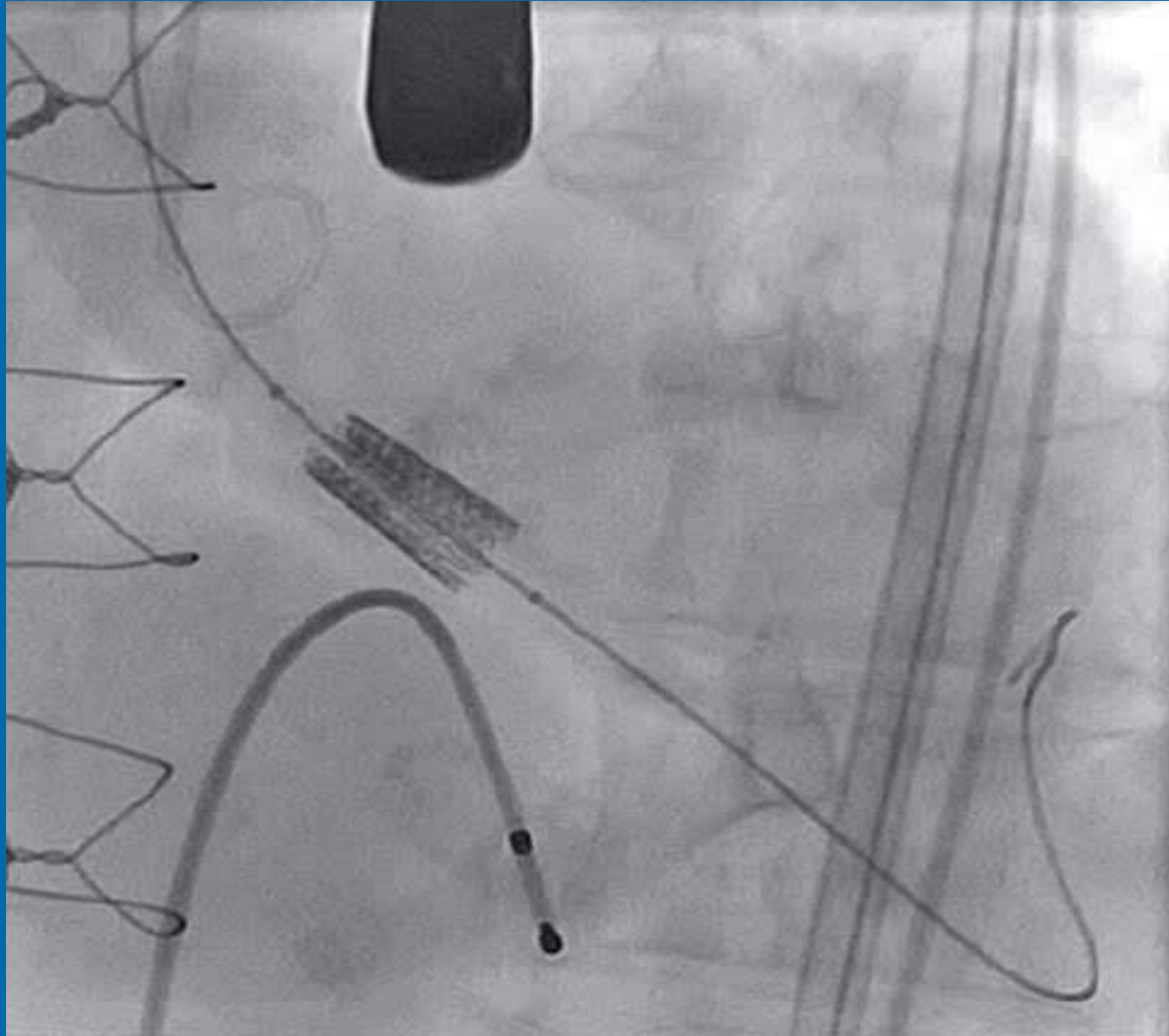
Left cusp









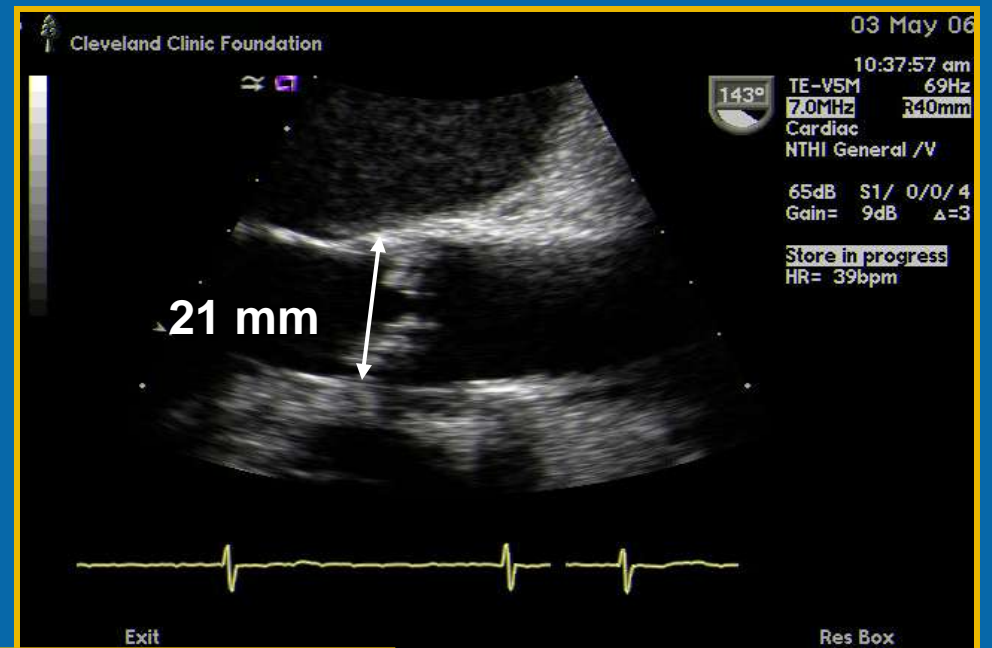
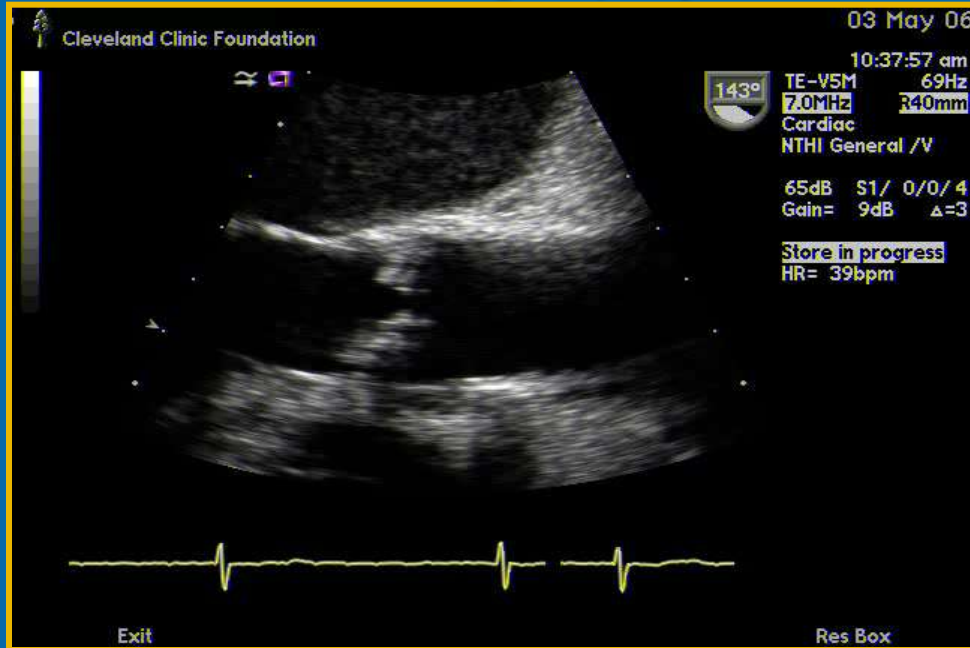


Patient #1 R.M.

90 yo Man

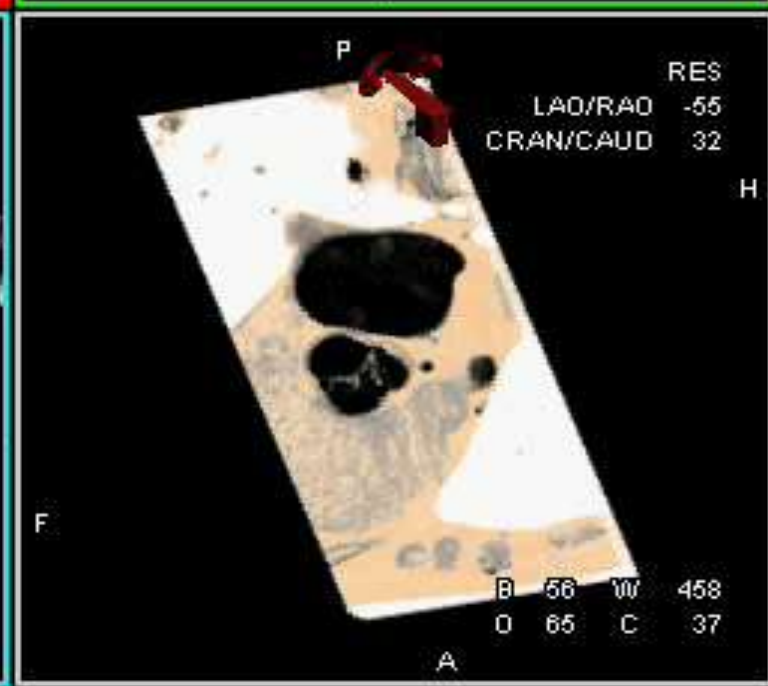
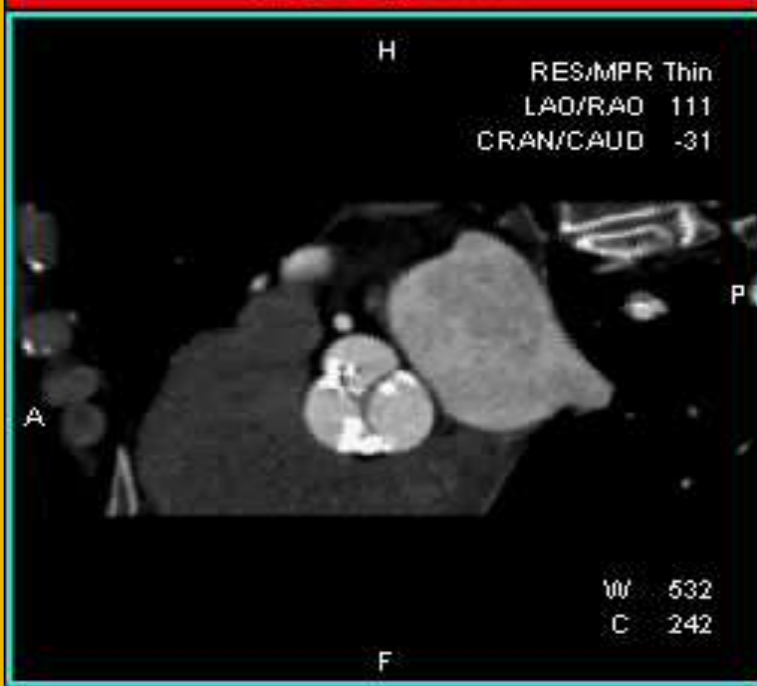
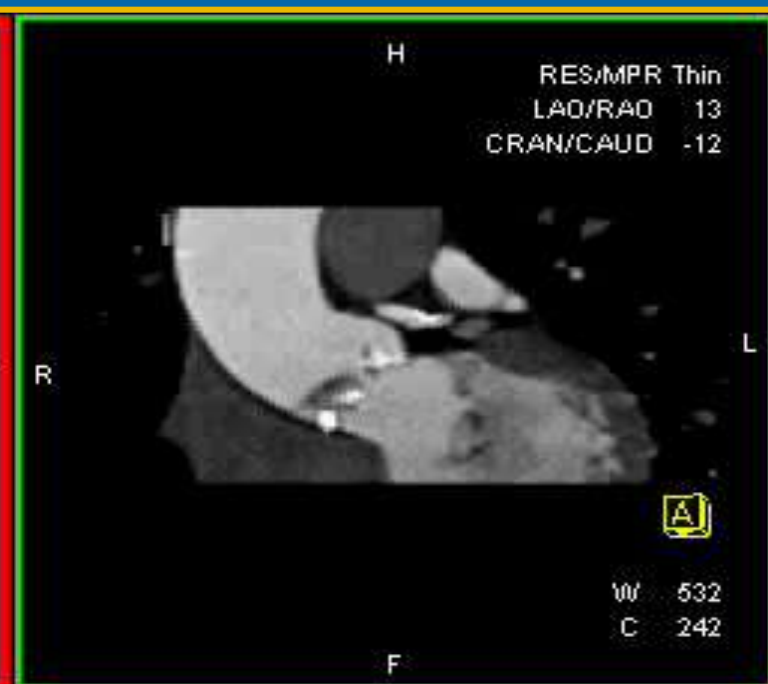
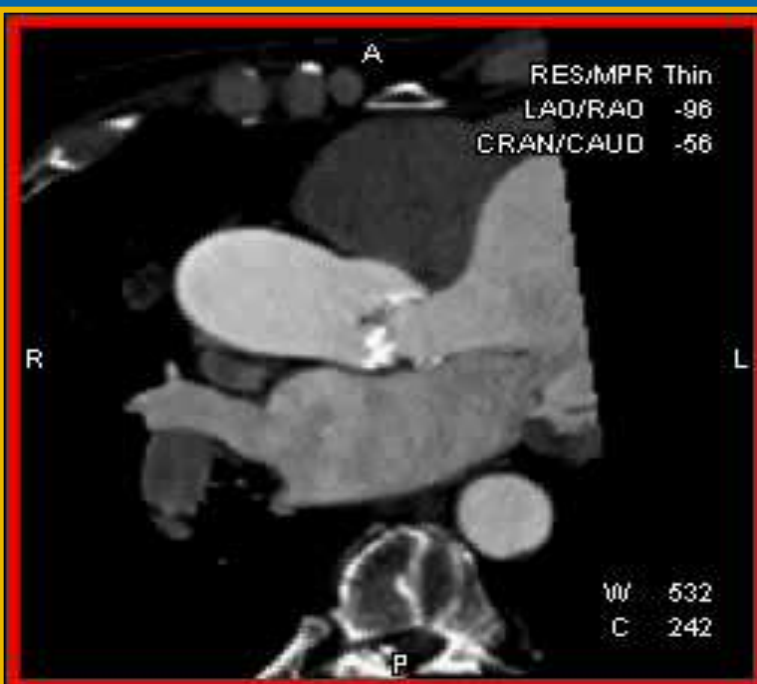
- 90 year-old male
- Severe aortic stenosis with NYHA Class III symptoms
- **Past Medical History:**
 - Coronary Artery Disease with CABGx4 in 1998
 - Right Carotid Endarterectomy
 - Atrial fibrillation
 - Hypertension
 - Marked Thoracic Kyphosis

Echocardiography



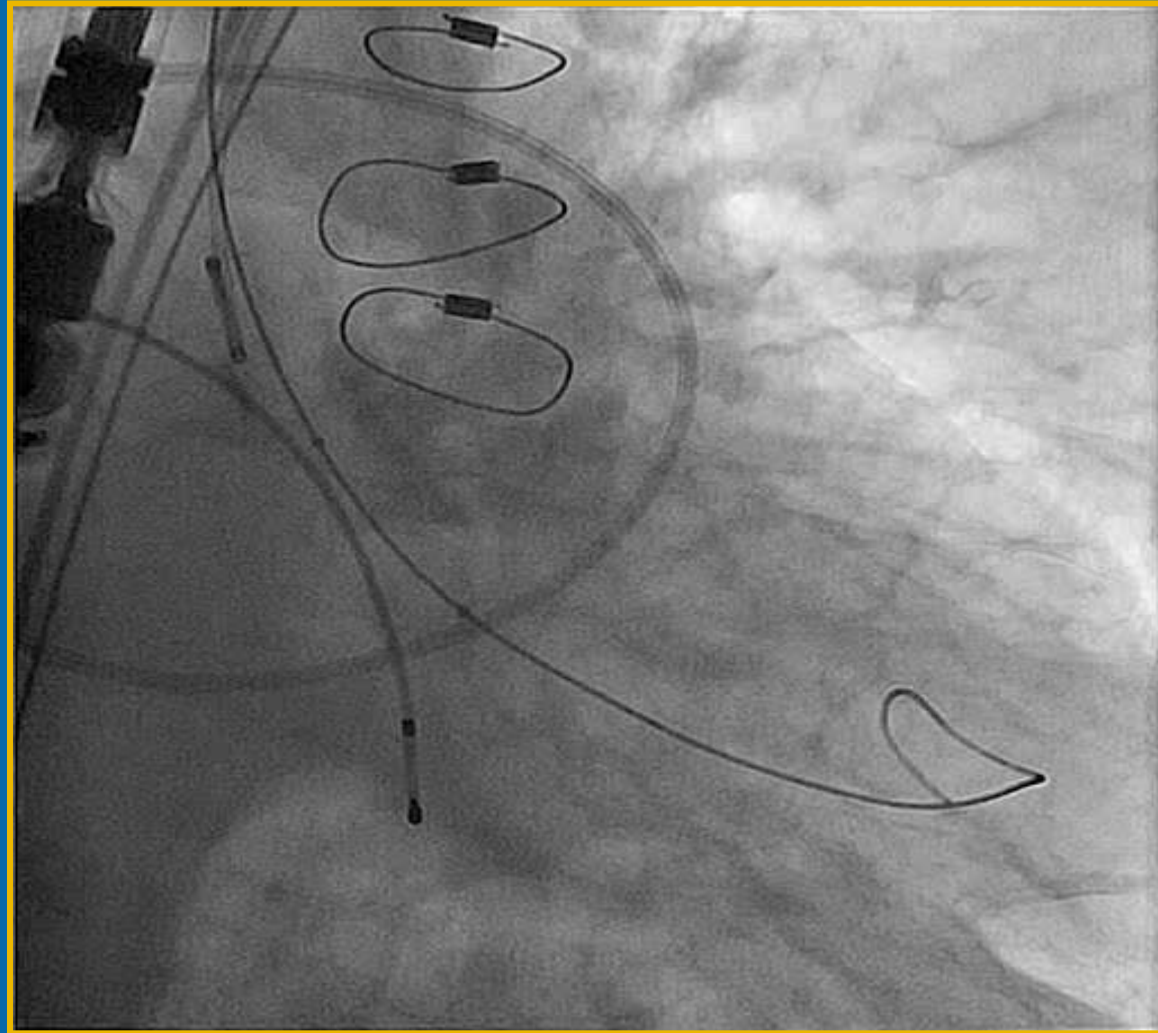
Echocardiogram

- LVEF 50%
- Stage II Diastolic Dysfunction
- Mild left ventricular hypertrophy
- Severe Aortic Stenosis (57/33 mmHg) valve area 0.7cm². 1-2+ AI.
- 1-2+ Mitral Regurgitation
- RVSP 69mm Hg

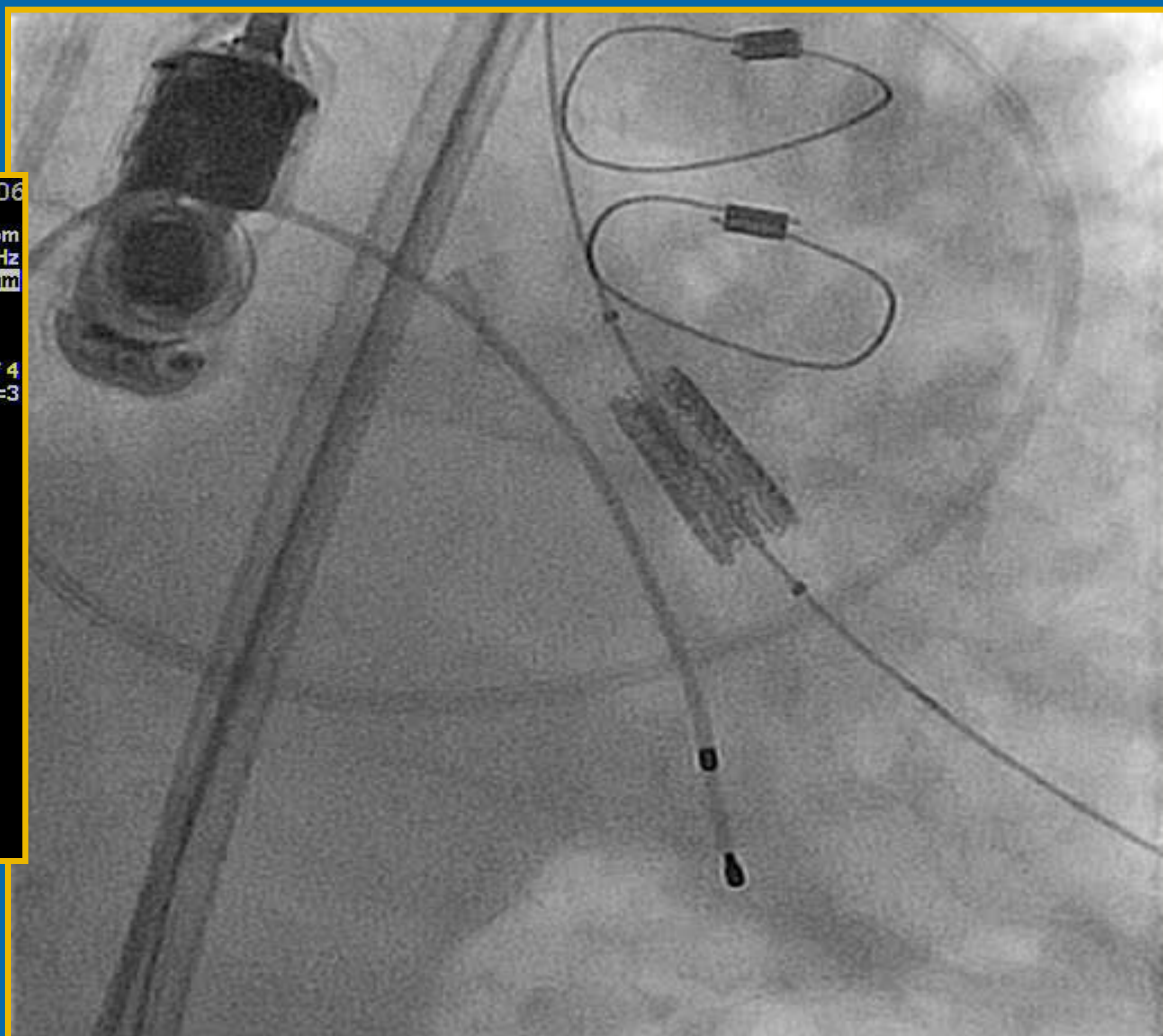
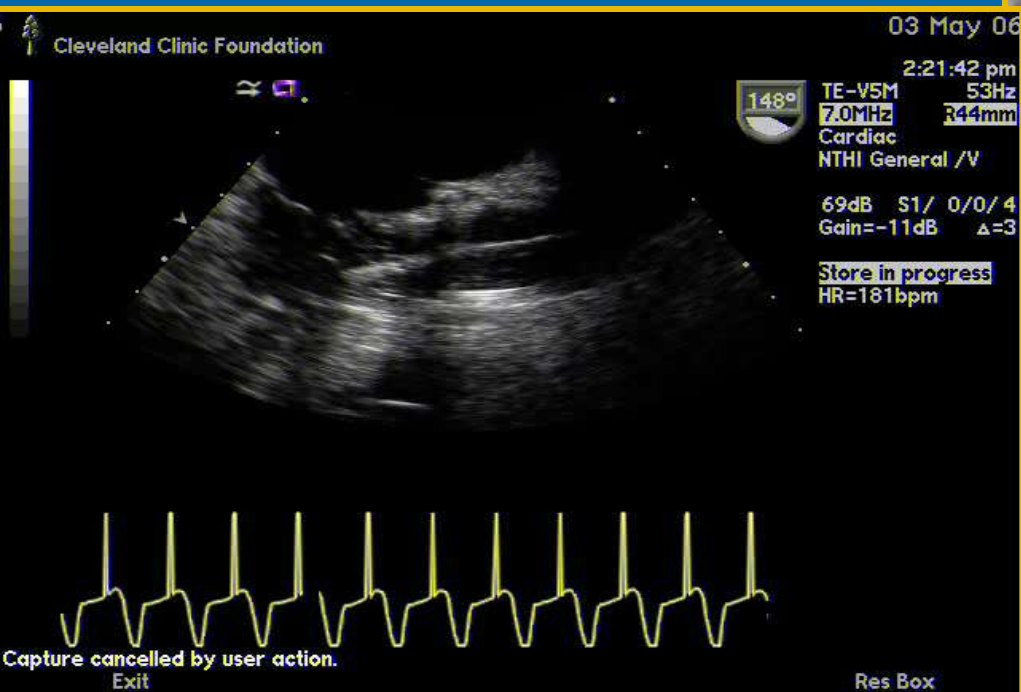


Balloon Valvuloplasty

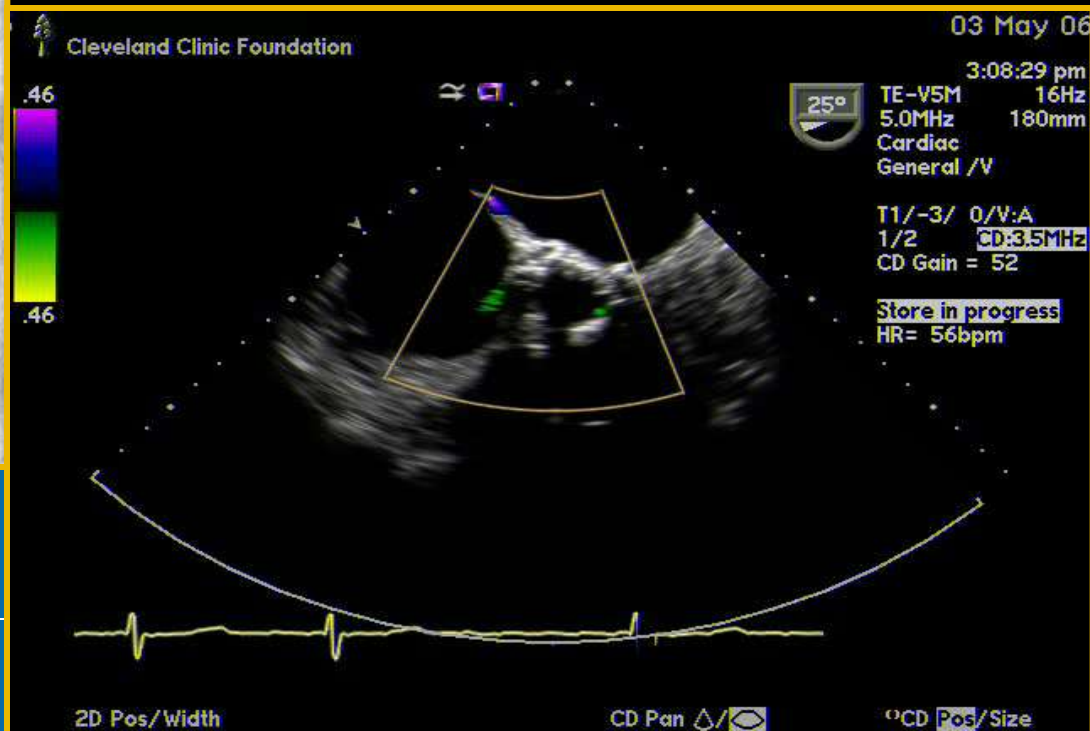
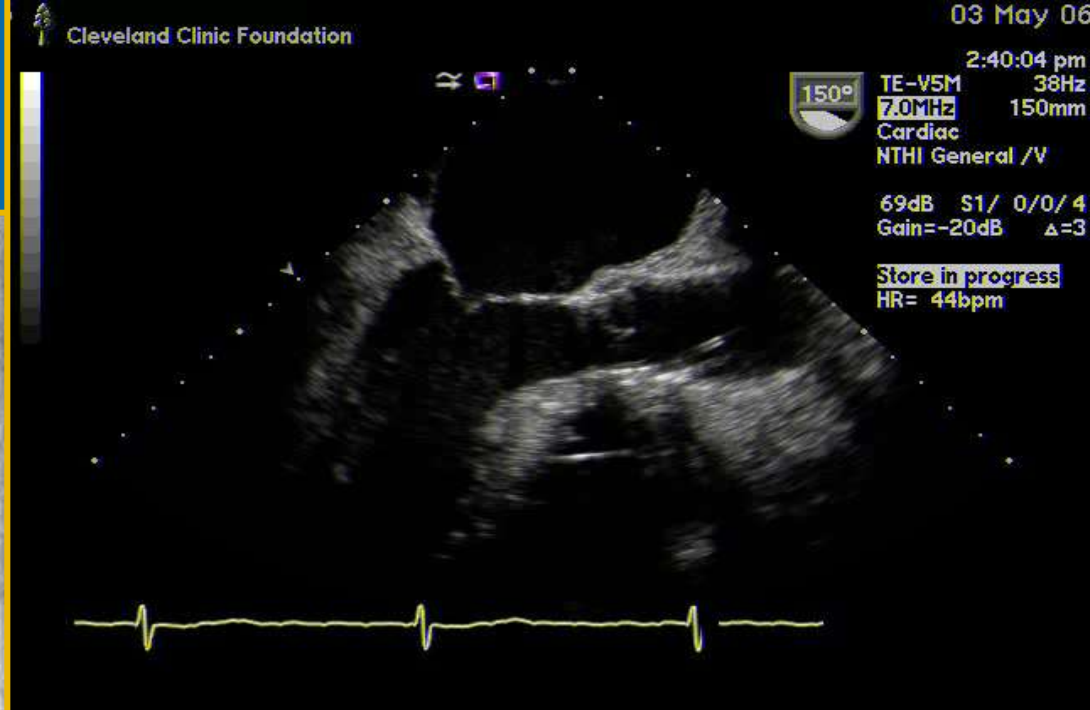
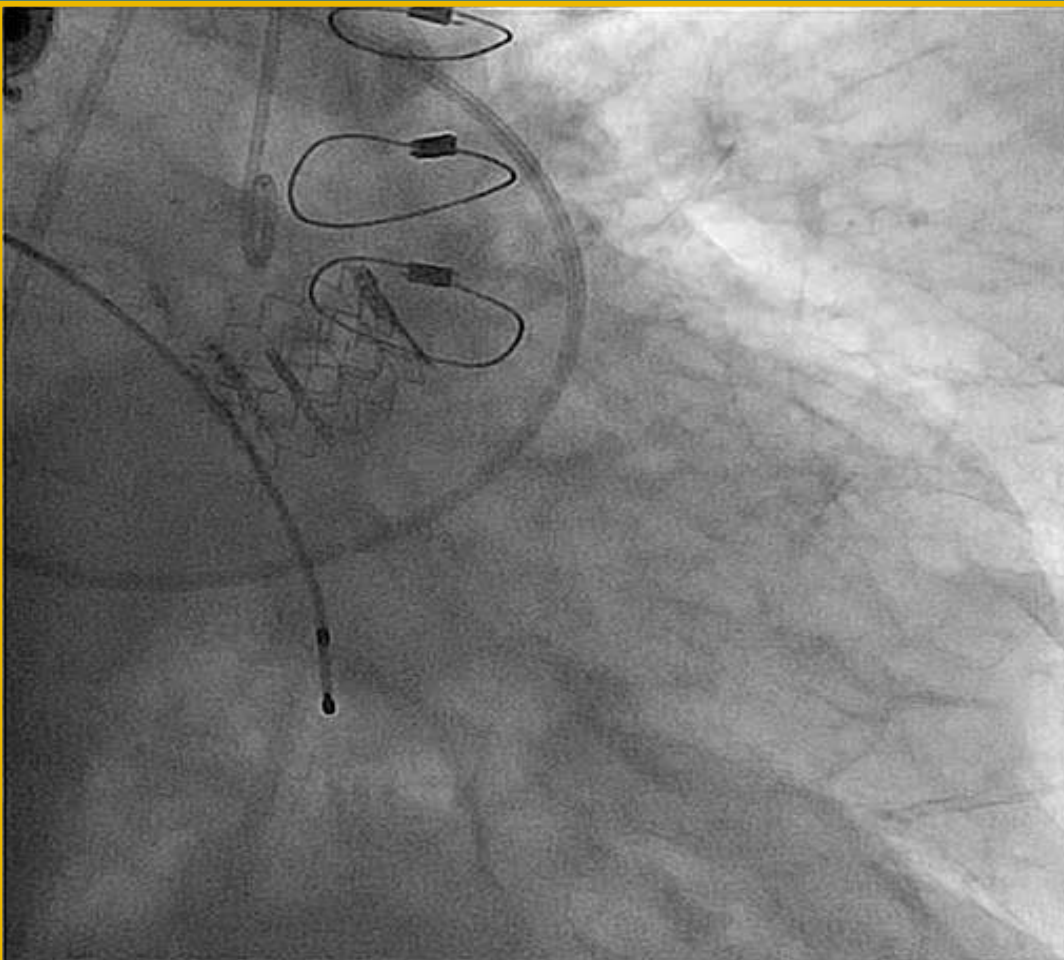
Used to Predilate the Valve



Valve Deployment



Result



Patient is playing golf 1 year later!

3D TEE in guiding PAVR

Live 3D Guidance of PAVR



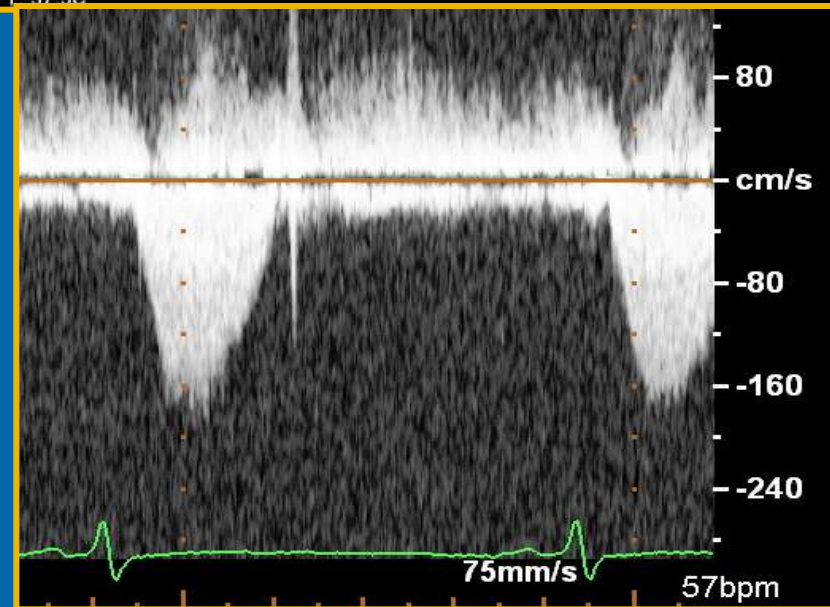
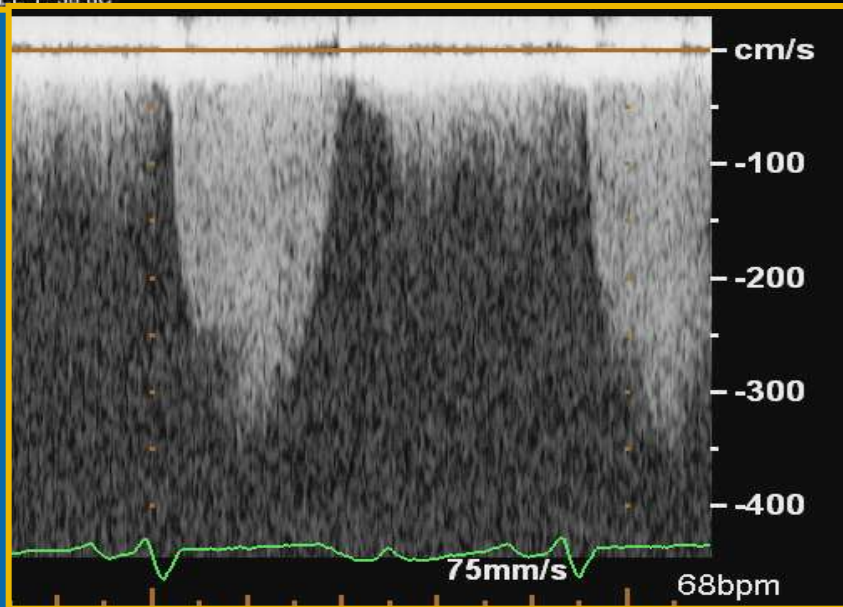
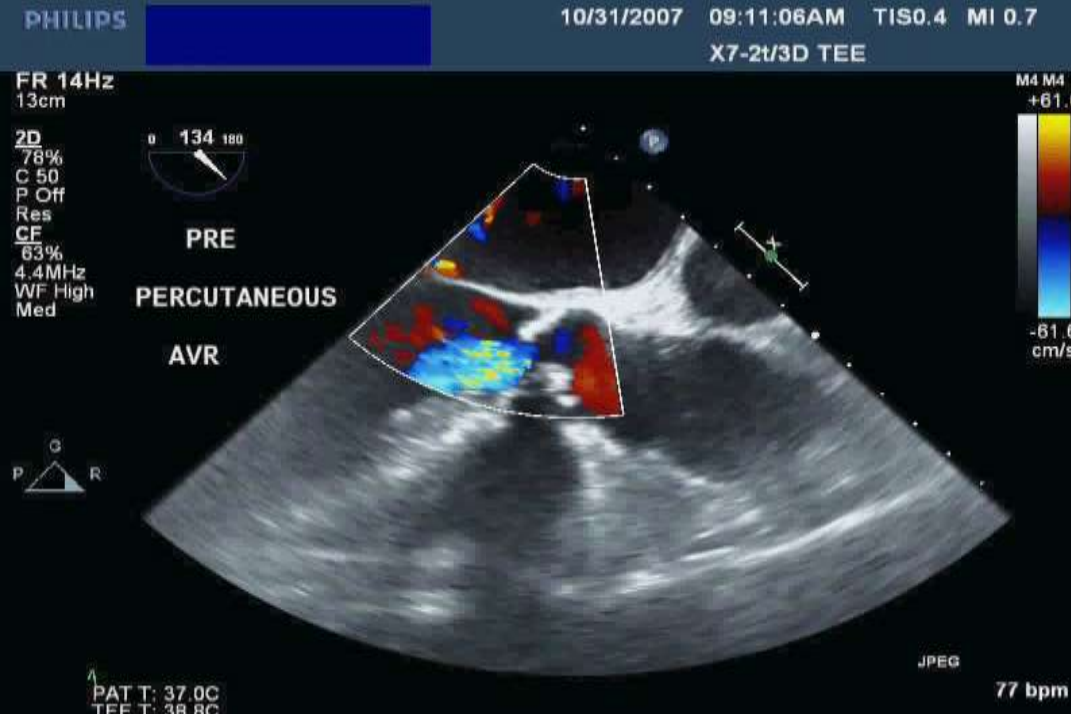
Live 3D Guidance of PAVR



Rapid pacing \Rightarrow balloon inflation of bioprosthesis

Post-Deployment Assessment





 Cleveland Clinic
2-3+ AR, $\Delta p = 54/38$ mmHg

1+ AR, $\Delta p = 12/8$ mmHg

PARTNER Study Design

Symptomatic Severe Aortic Stenosis

ASSESSMENT: High Risk AVR Candidate
3105 Total Patients Screened

Total = 1058 patients
**2 Parallel Trials:
Individually Powered**

n= 700

High Risk

ASSESSMENT:
Transfemoral Access

High Risk TF

High Risk TA

1:1 Randomization

1:1 Randomization

TAVI
Trans
femoral

vs

Surgical
AVR

TAVI
Trans
femoral

vs

Surgical
AVR

Primary Endpoint: All Cause Mortality (1 yr)
(Non-inferiority)

Inoperable

n=358

ASSESSMENT:
Transfemoral Access

1:1 Randomization

Not In Study

TAVI
Trans
femoral

vs

Standard
Therapy
(usually BAV)

Primary Endpoint: All Cause Mortality over length of
trial (Superiority)

Primary Endpoint

- *All-cause mortality over the duration of the study*

Co-Primary Endpoint

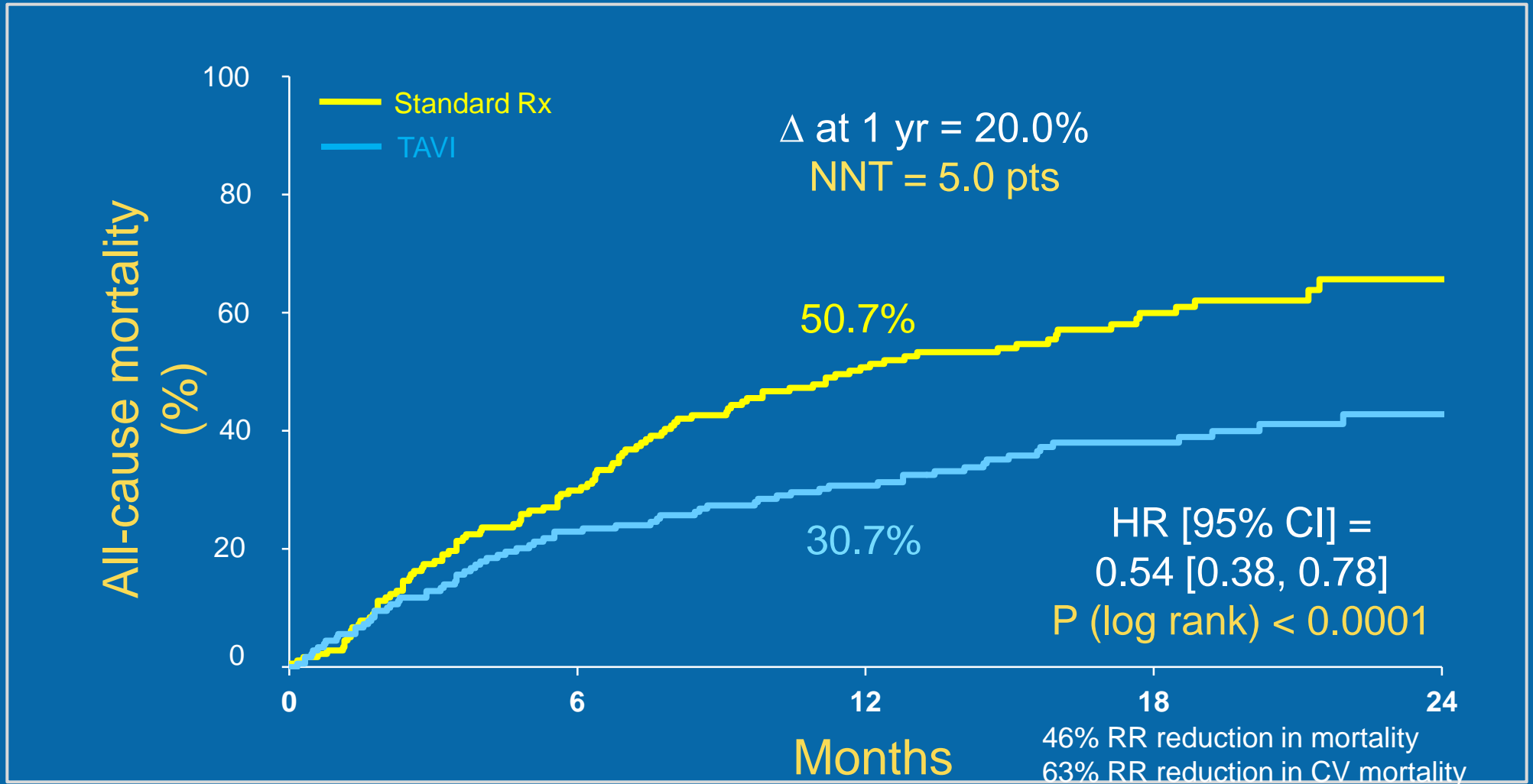
- *Hierarchical composite of all-cause mortality and repeat hospitalization over the duration of the study*

Procedural Outcomes

TAVI (179 patients)

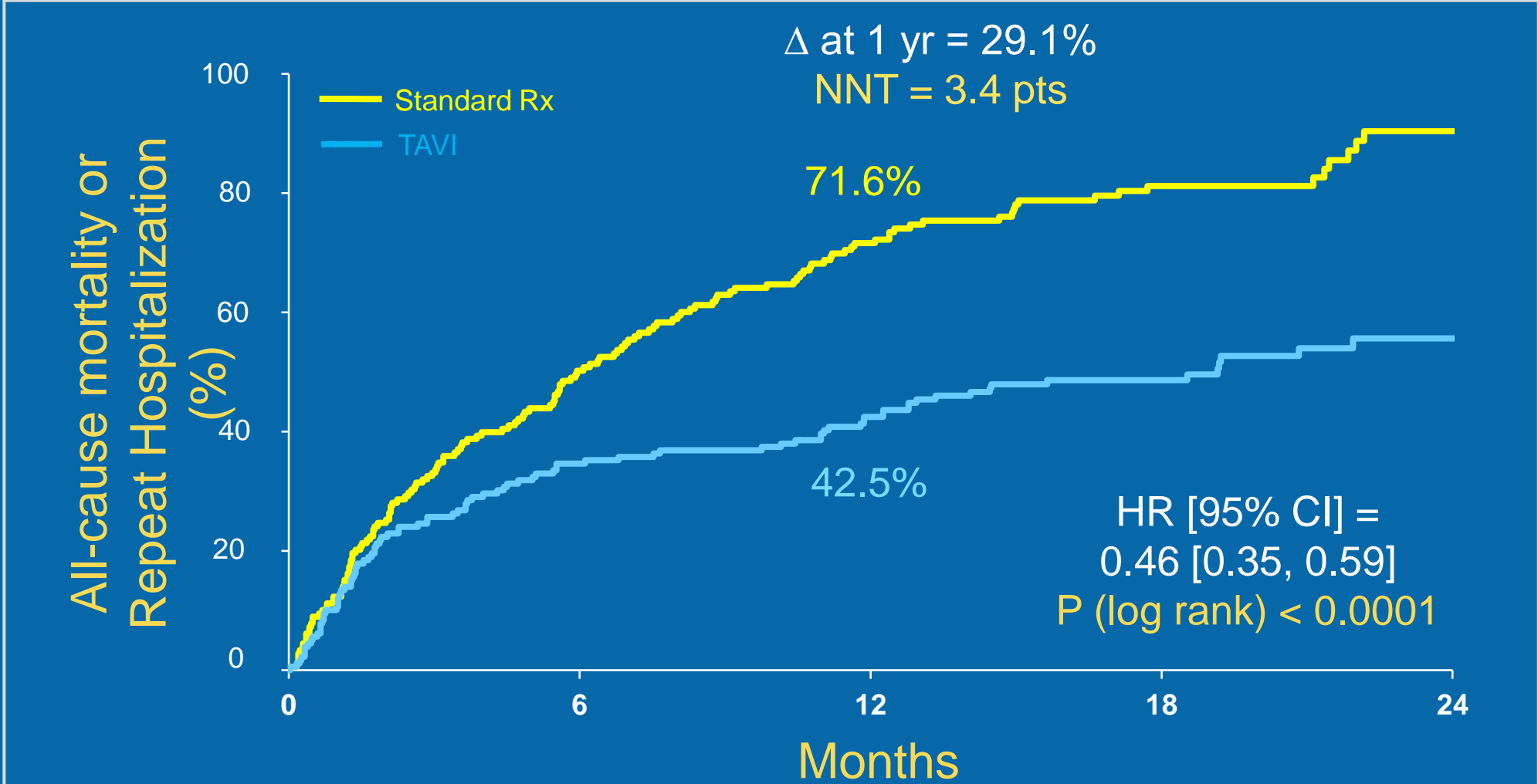
- **6 (3.4%) pts did not receive TAVI**
 - 2 died before scheduled implant
 - 2 unsuccessful transfemoral access
 - 2 intra-procedural annulus measurement too large and procedure aborted
- **After randomization, median time to TAVI was 6 days** (inter-quartile range 3 - 11 days)
- **During TAVI (first 24 hours)**
 - 2 (1.1%) deaths
 - 3 (1.7%) major strokes
 - 1 (0.6%) valve embolization
 - 2 (1.1%) pts with multiple (≥ 2) valve implants
- **In the first 30 days, 11 (6.4%) pts receiving TAVI died**

All Cause Mortality



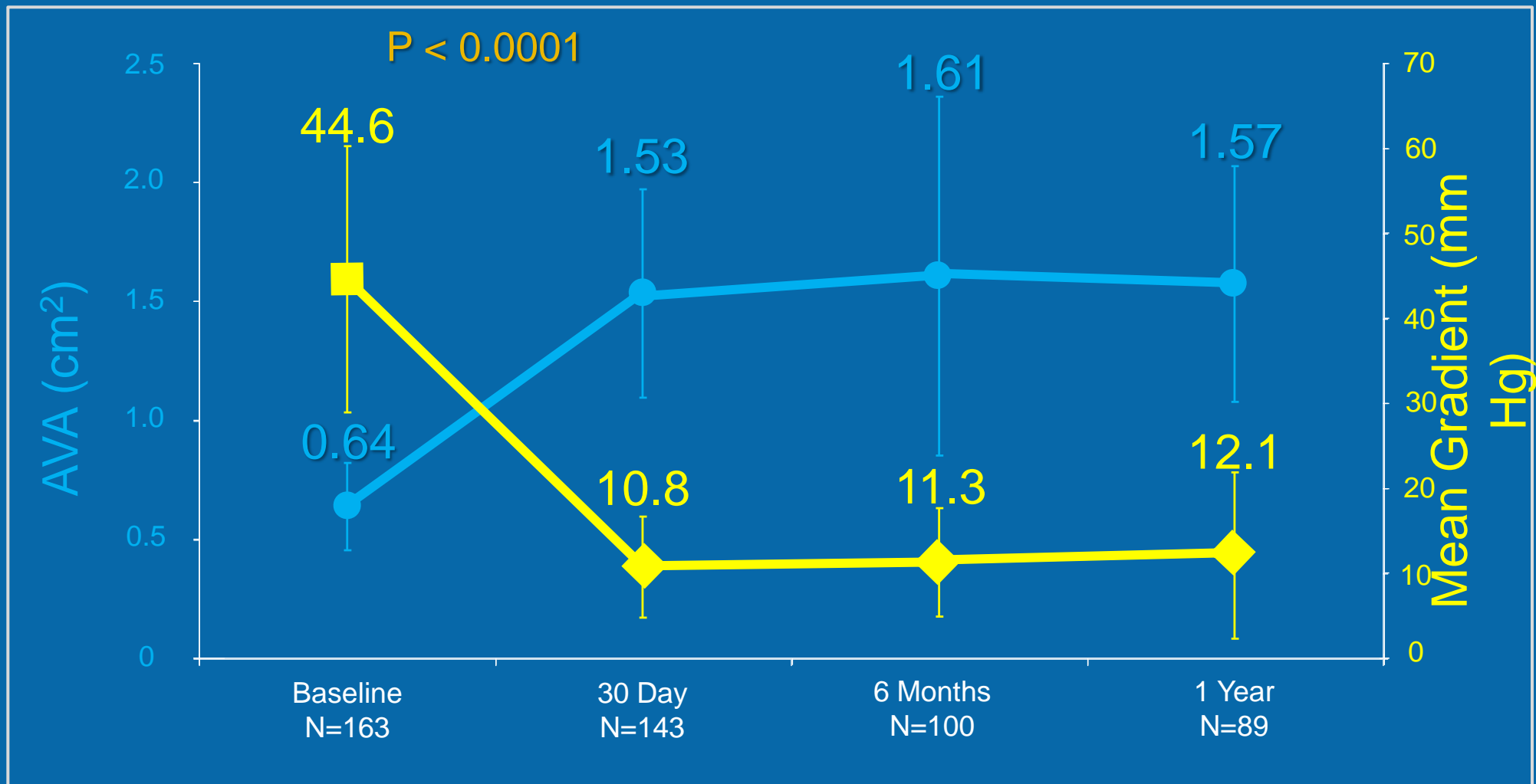
Numbers at Risk					
TAVI	179	138	122	67	26
Standard Rx	179	121	83	41	12

Mortality or Repeat Hosp

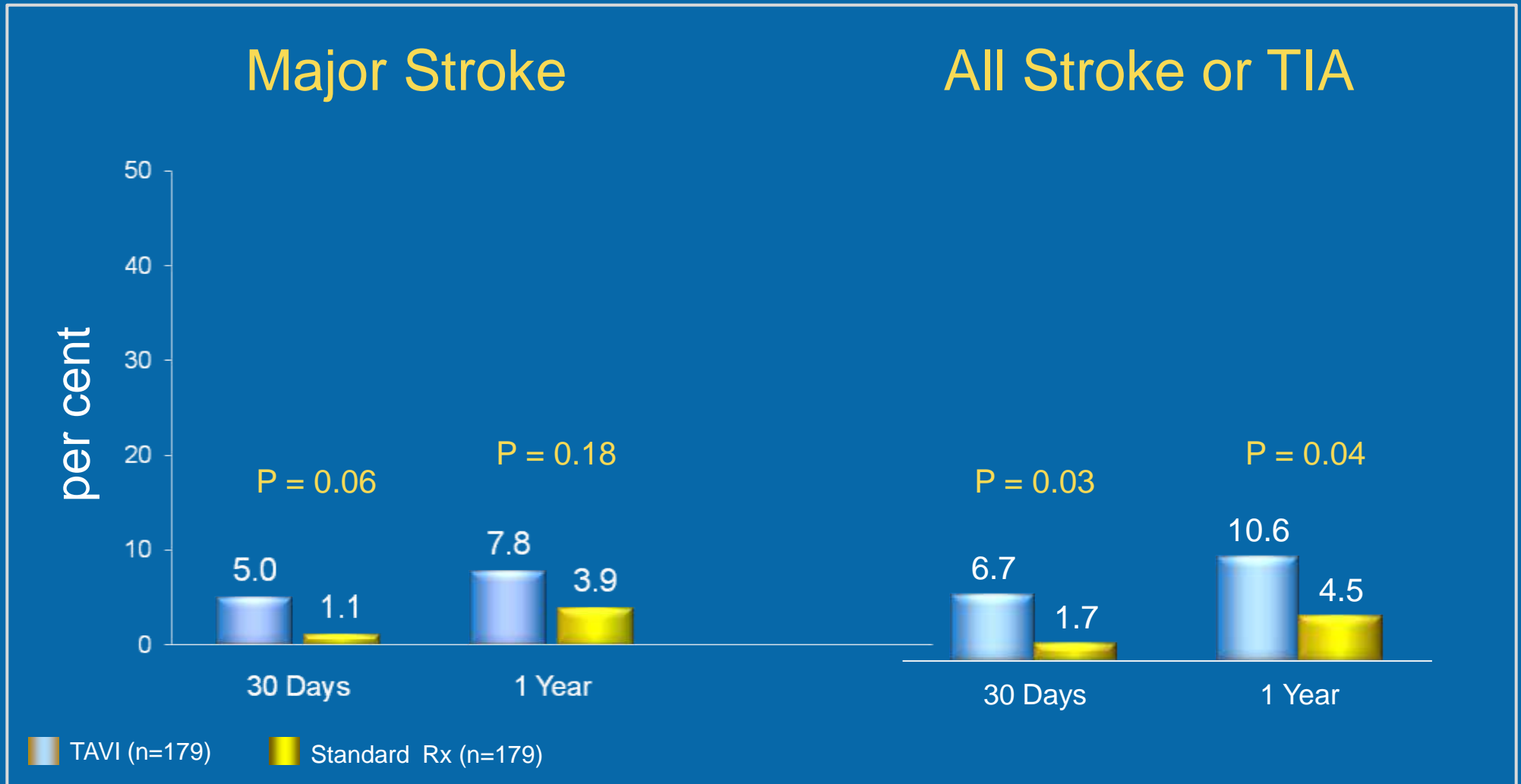


Numbers at Risk					
TAVI	179	117	102	56	22
Standard Rx	179	121	49	23	4

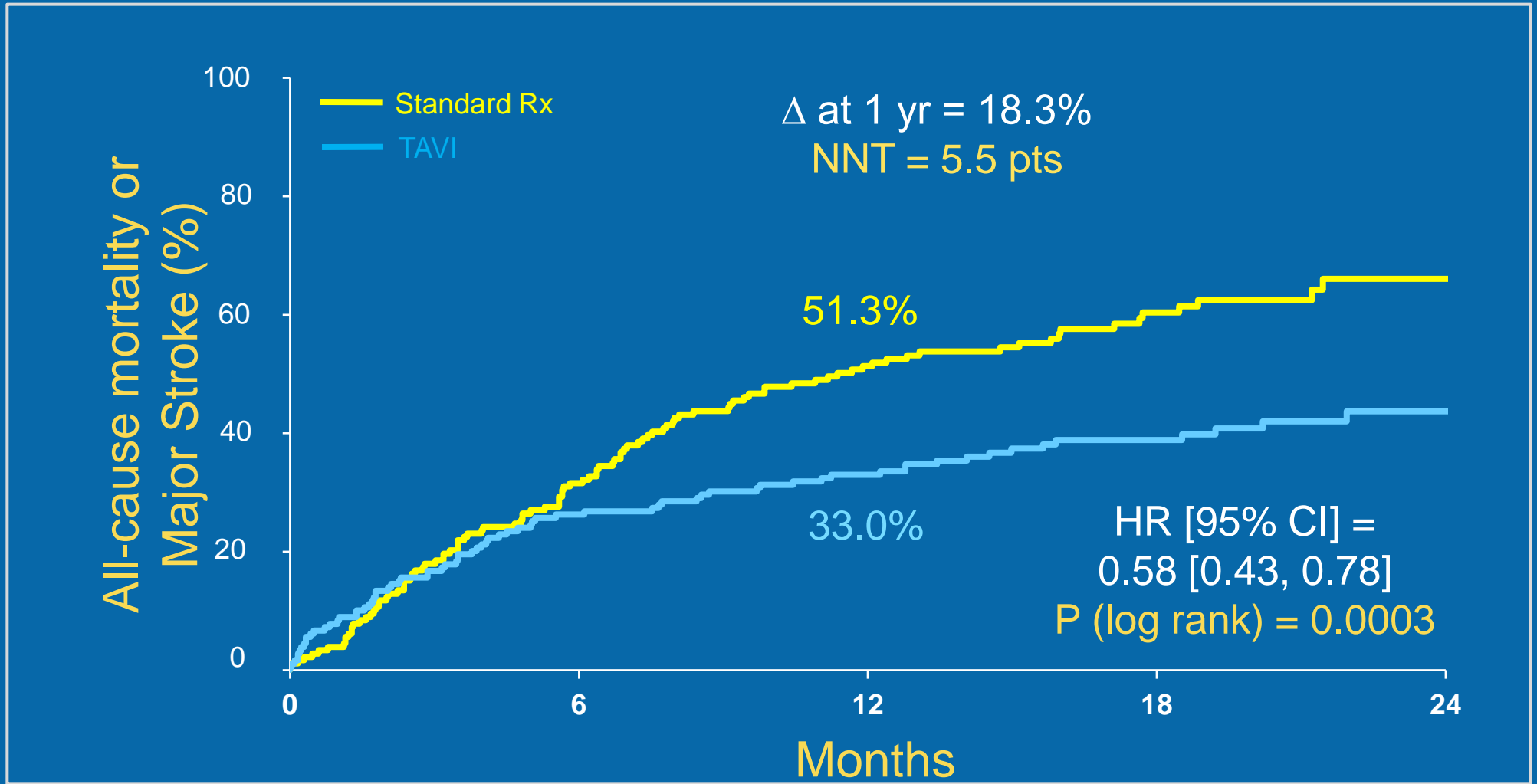
AVA and Mean Gradients All TAVI patients



Clinical Outcomes at 30 Days and 1 Year



Mortality or Major Stroke



Numbers at Risk					
TAVI	179	132	118	56	25
Standard Rx	179	118	83	41	12

Conclusions

In patients with severe AS and symptoms, who are not suitable candidates for surgery...

- TAVI was accomplished with acceptable safety
- TAVI markedly reduced the rate of...
 - all-cause mortality by 46%
 - all-cause mortality & hospitalization by 54%,
- TAVI improved cardiac symptoms
- TAVI resulted in more frequent complications
 - major vascular complications
 - major bleeding episodes
 - major strokes

Clinical Implications

- Balloon-expandable TAVI should be the new standard of care for patients with aortic stenosis who are not suitable candidates for surgery!

Echo to Guide Clinical Management in Patients with Mitral Regurgitation

Morphology

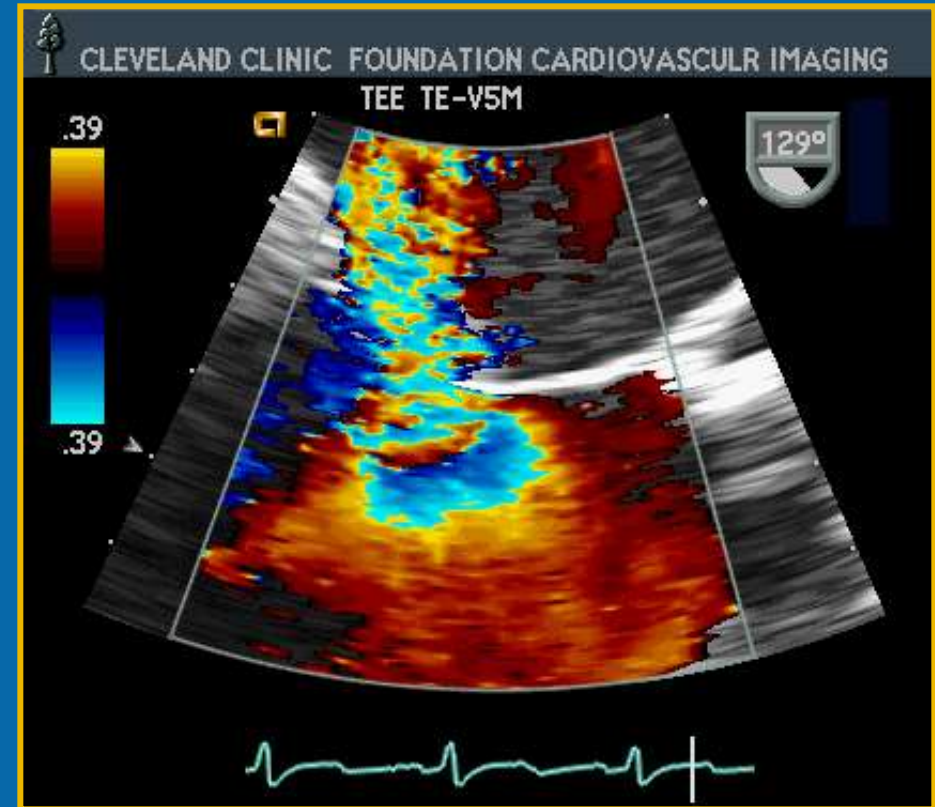
	Rheumatic	Myxomatous
LV	Normal/Small	Dilated
MV	Thick/restricted	Thick/hypermobile
Chordae	Short	Long
AV involved	Often	Rare
Ca++	+++	+

Myxomatous Mitral Valve Disease



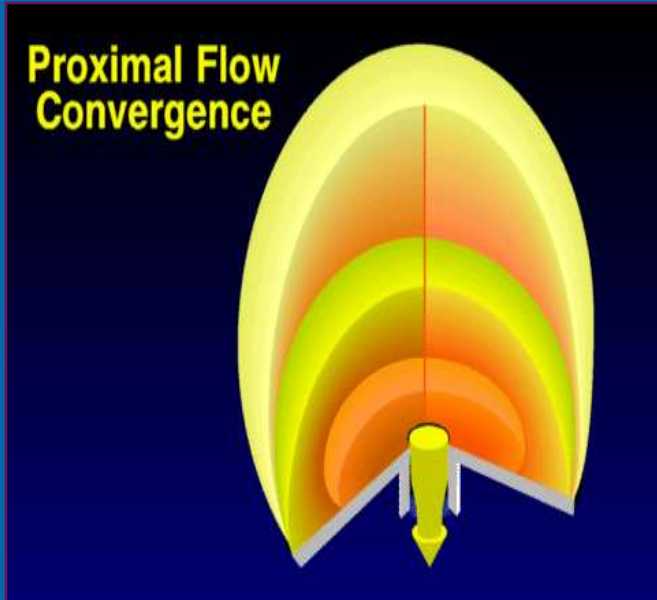
Quantitation of Mitral Regurgitation by Echo

- Color jet area
- PISA
- Vena contracta
- Volumetric Assessment
- Jet density by CW Doppler
- Chamber Size
- Pulmonary veins

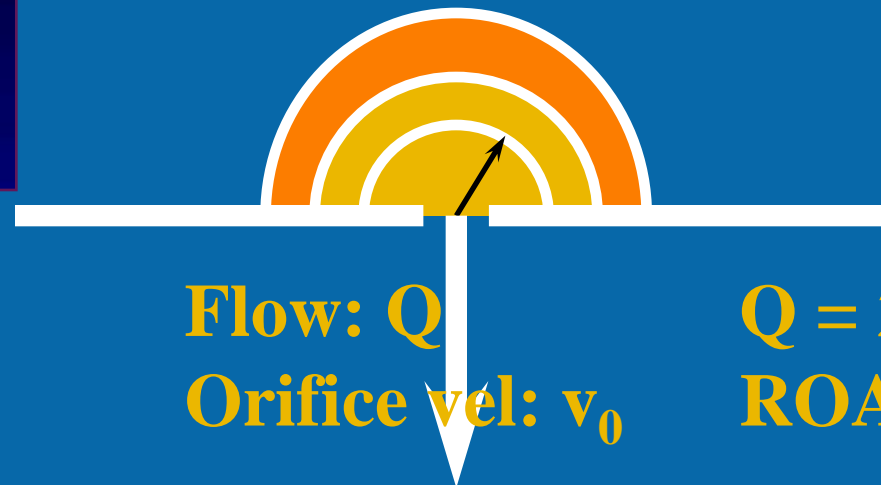


Regurgitant Orifice Area $> 0.4 \text{ cm}^2$ = Severe

Proximal Flow Convergence Method



Aliasing velocity: v ($= 42$ cm/s)
Aliasing radius: r



Flow: Q

Orifice vel: v_0

$$Q = 2\pi r^2 v$$

$$ROA = Q/v_0$$

Flow thru any isovelocity shell is equal to instantaneous orifice flow

Indications for Transthoracic Echocardiography

CLASS I

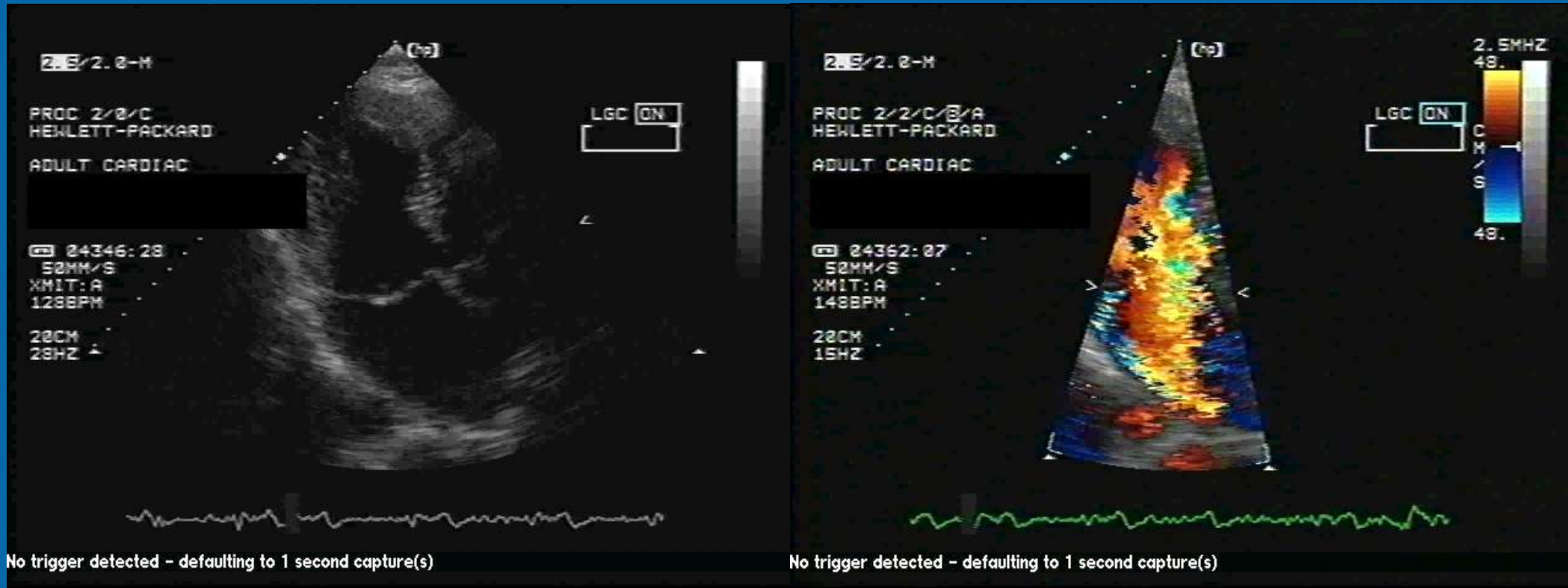
- 1. TTE is indicated for baseline evaluation of LV size and function. RV and left atrial size, pulmonary artery pressure and severity of MR (Table 4) in any patient suspected of having MR, (Level of Evidence: C)
- 2. TTE is indicated for delineation of the mechanism of MR. (Level of Evidence: B).
- 3. TTE is indicated for annual or semiannual surveillance of LV function (estimated by ejection fraction and end-systolic dimension) in asymptomatic patients with moderate to severe MR. (Level of Evidence: C)
- 4. TTE is indicated in patients with MR to evaluate the MV apparatus and LV function after a change in signs or symptoms. (Level of Evidence: C)
- 5. TTE is indicated to evaluate LV size and function and MV hemodynamics in the initial evaluation after MVR or MV repair. (Level of Evidence: C)

INDICATIONS FOR TEE

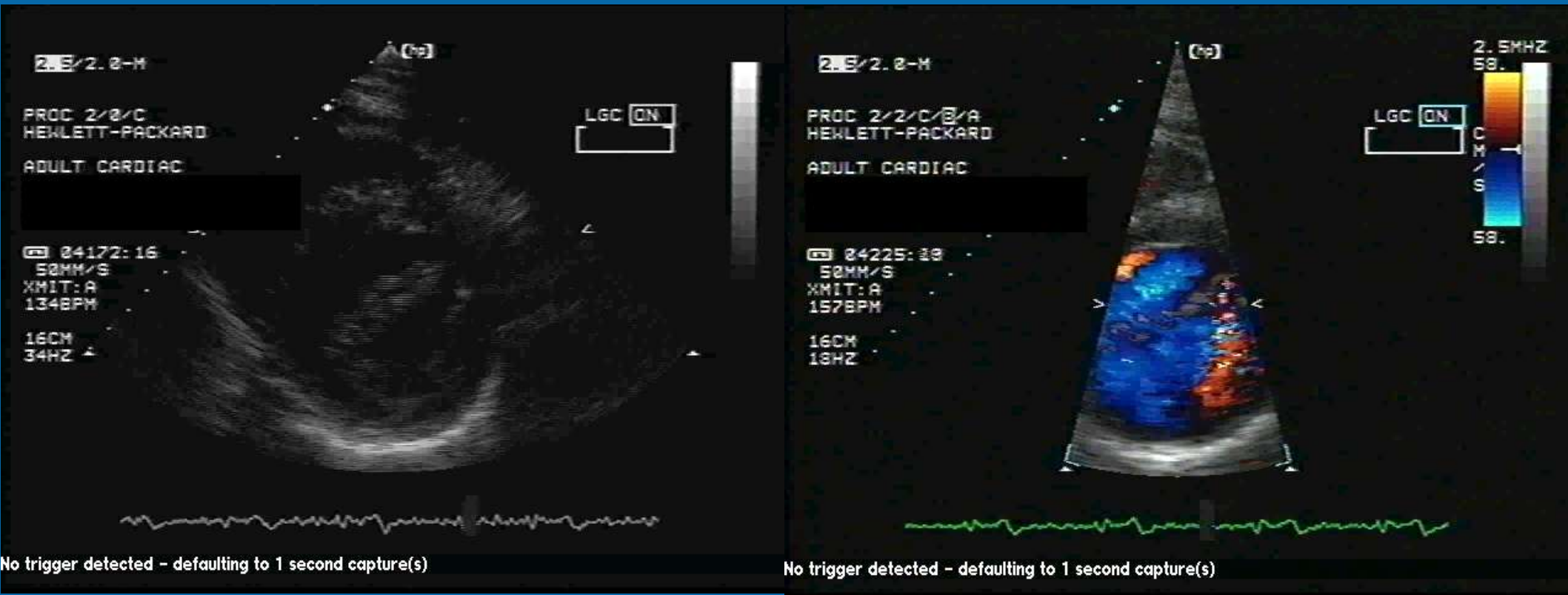
CLASS I

- 1. Preoperative or intraoperative TEE is indicated to establish the anatomic basis for severe MR in patients in whom surgery is recommended to assess feasibility of repair and to guide repair (Level of Evidence: B)
- 2. TEE is indicated for evaluation of MR patients in whom TTE provides non-diagnostic information regarding severity of MR, mechanism of MR and/or status of LV function. (Level of Evidence B)

TTE apical LAX



TTE

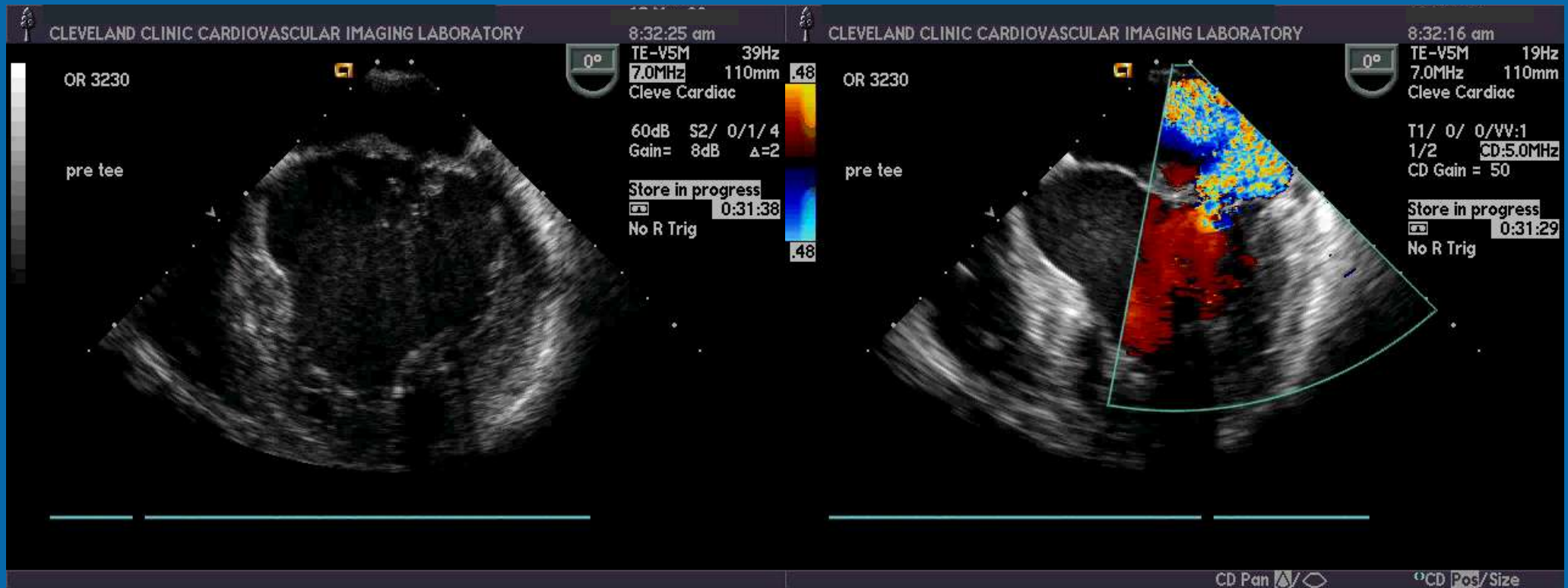


Mechanisms of MR

This echo evaluation shows ?

- a. Anterior leaflet prolapse due to myxomatous valve disease**
- b. Restriction of posterior leaflet due to RHD**
- c. Restriction of posterior leaflet due to RCA scar**
- d. Bileaflet prolapse with anterior prolapse > posterior prolapse?**
- e. Restriction of posterior leaflet due to anorexigen induced valvular disease**

Pre Intraoperative TEE

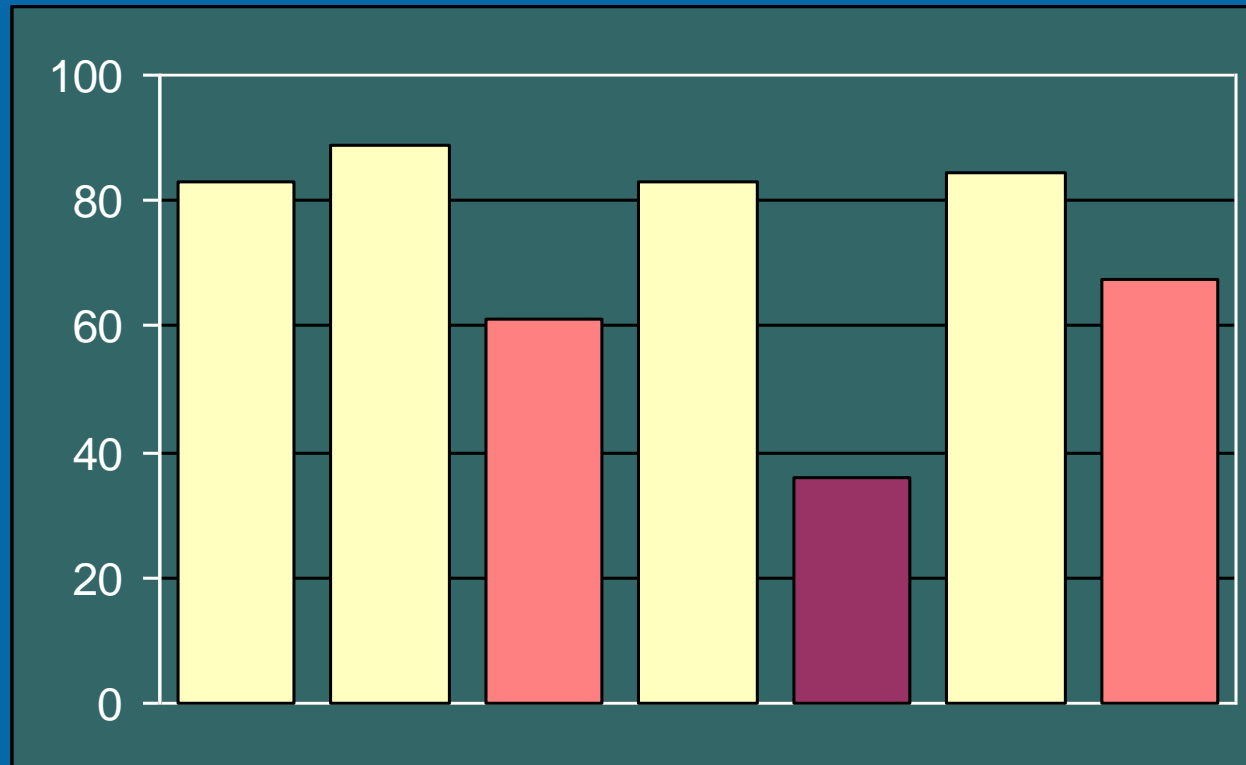


Mechanisms of Mitral Regurgitation

- Excessive Leaflet Motion
 - Prolapse
 - Leaflet, chordal, or papillary muscle disruption
- Normal Leaflet Motion
 - Perforation
- Restricted Leaflet Motion
 - Rheumatic
 - Ischemic
 - Anorexigen Induced
 - Mitral annular calcification

Feasibility of Repair - by MR mechanism

Stewart WJ: Intraoperative Echocardiography, Chap 54 in Topol's Cardiology Textbook



EXC
ALL

EXC
post

EXC
ant

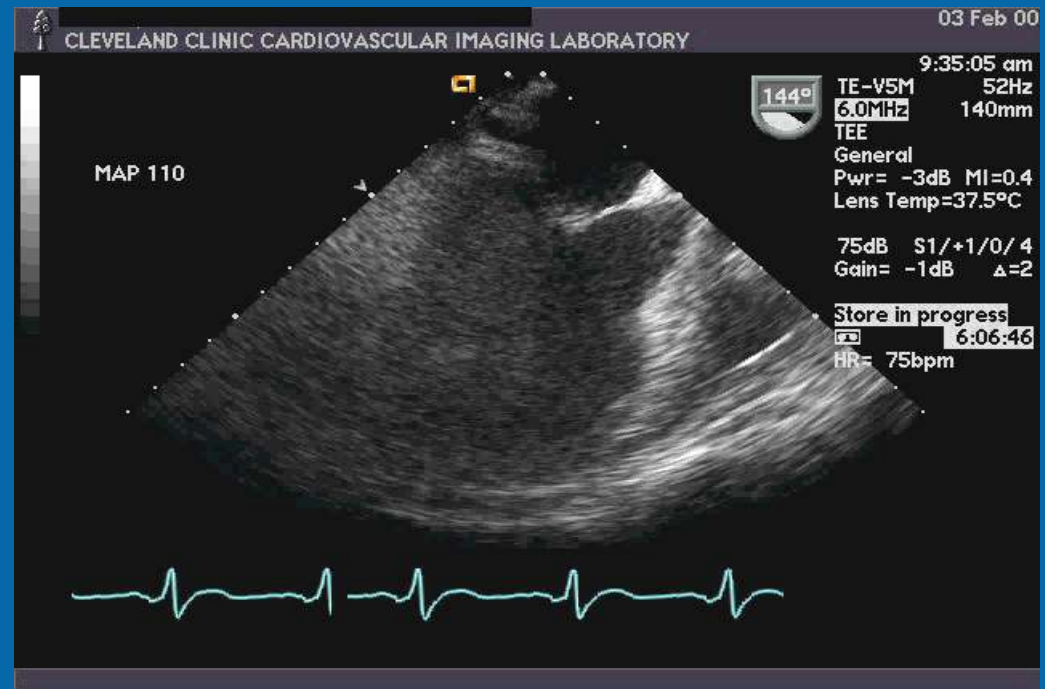
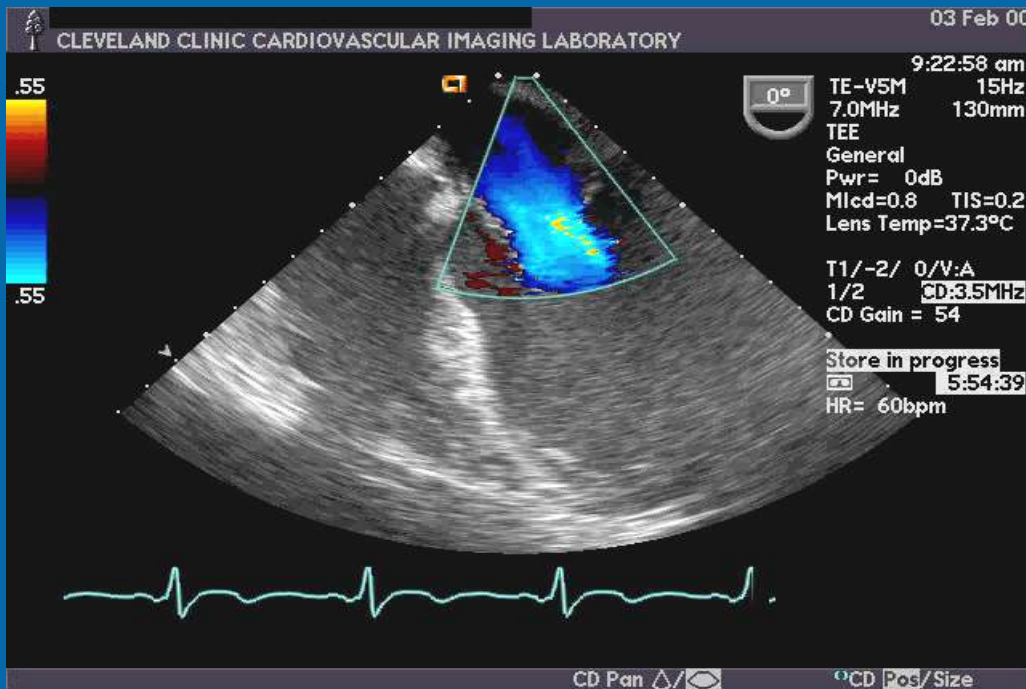
EXC
both

RES

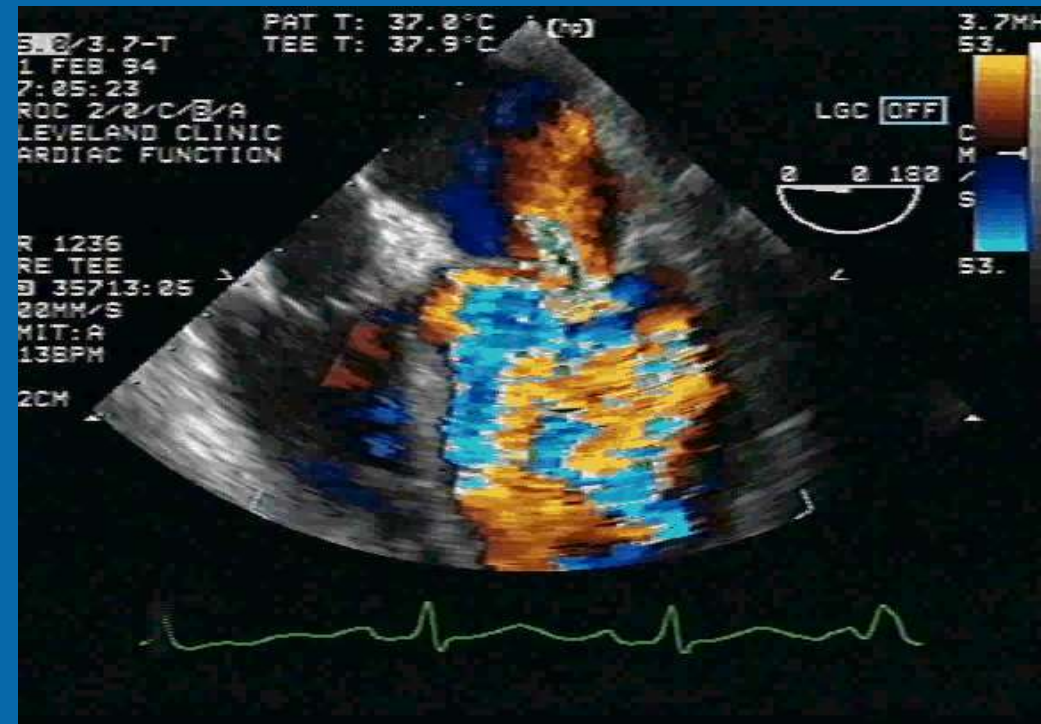
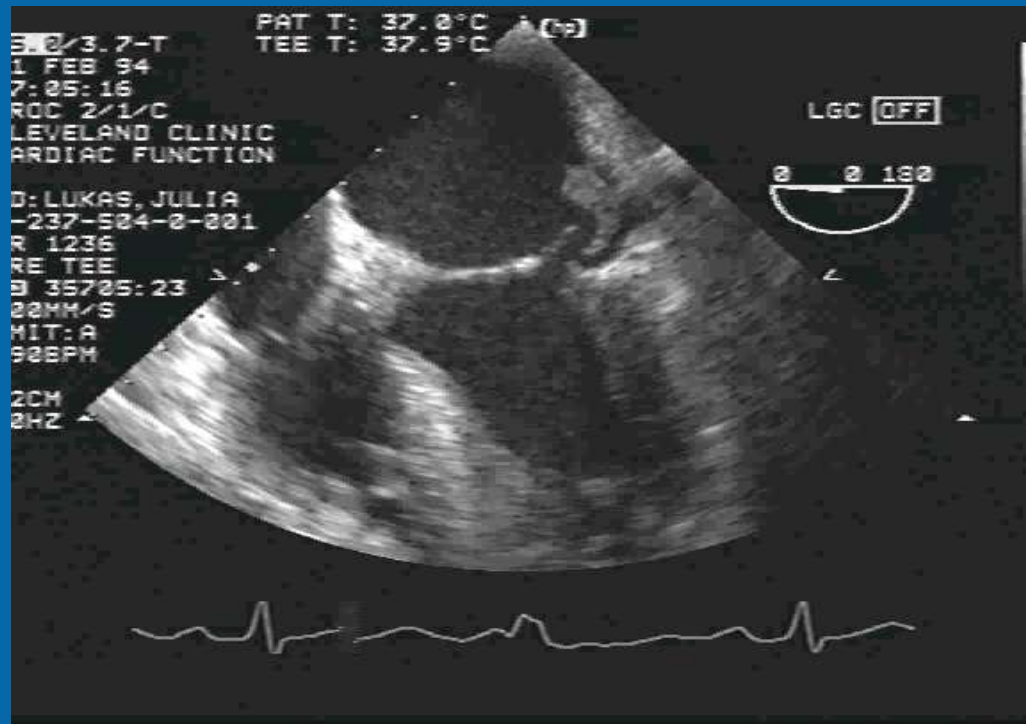
NORM
DCM

TOTAL

Papillary Muscle Dysfunction



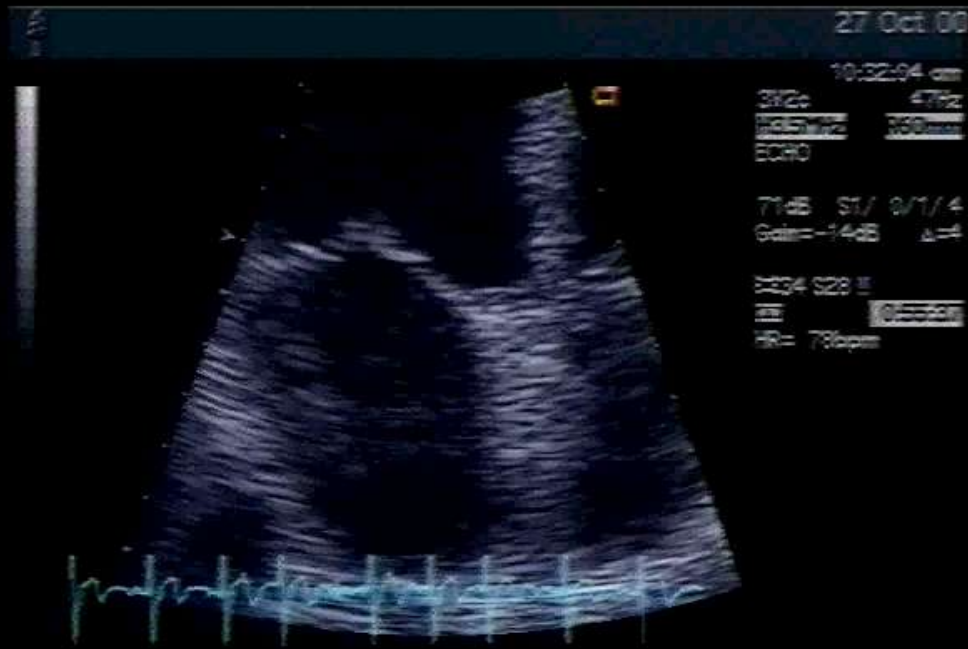
Repair or Replace?



Repair or Replace?

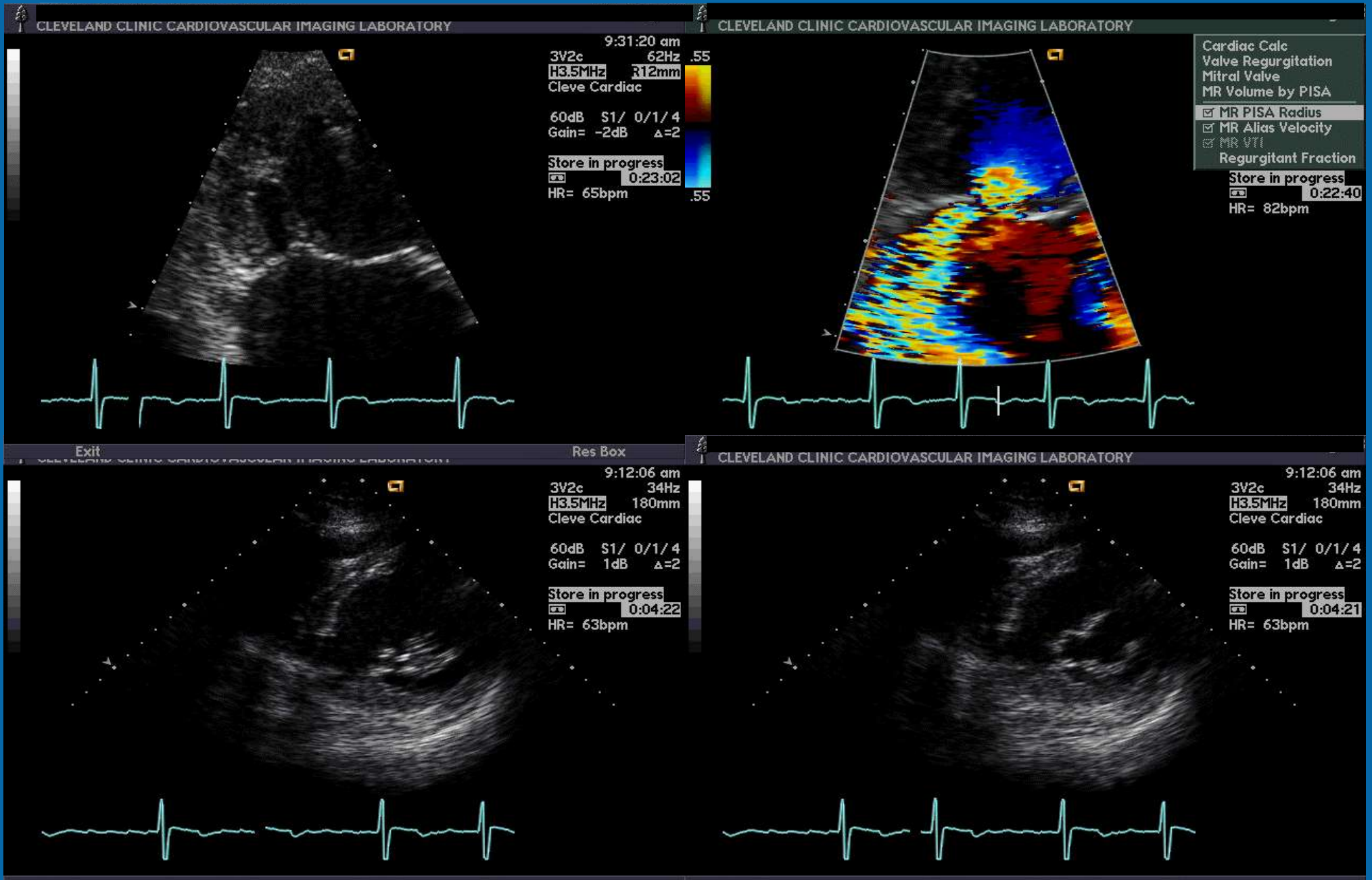


Repair or Replace

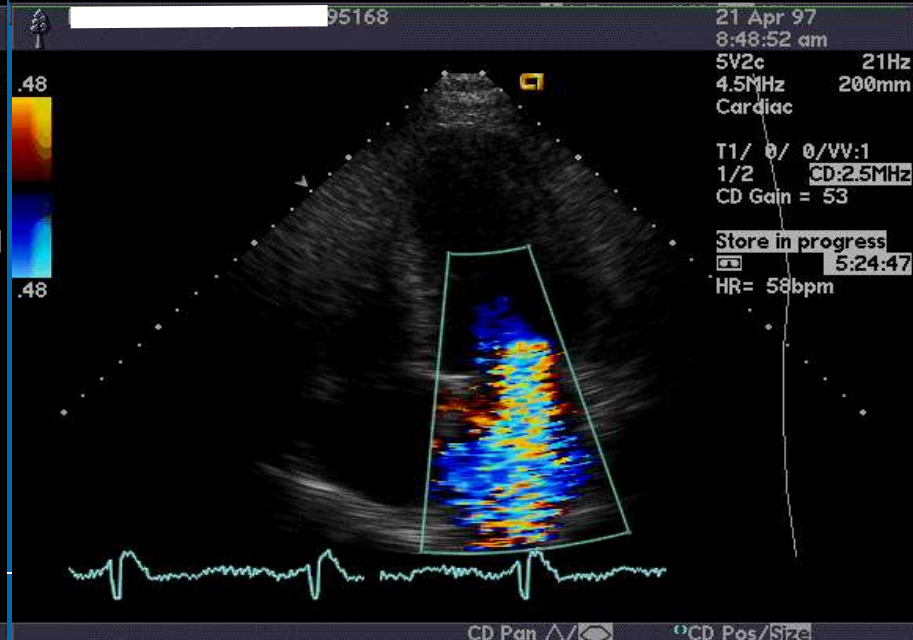
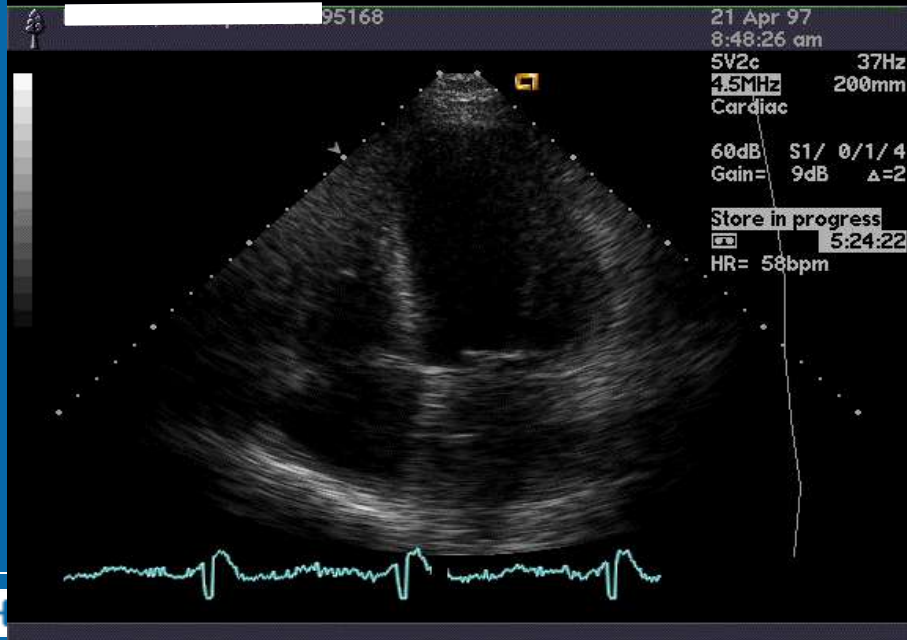
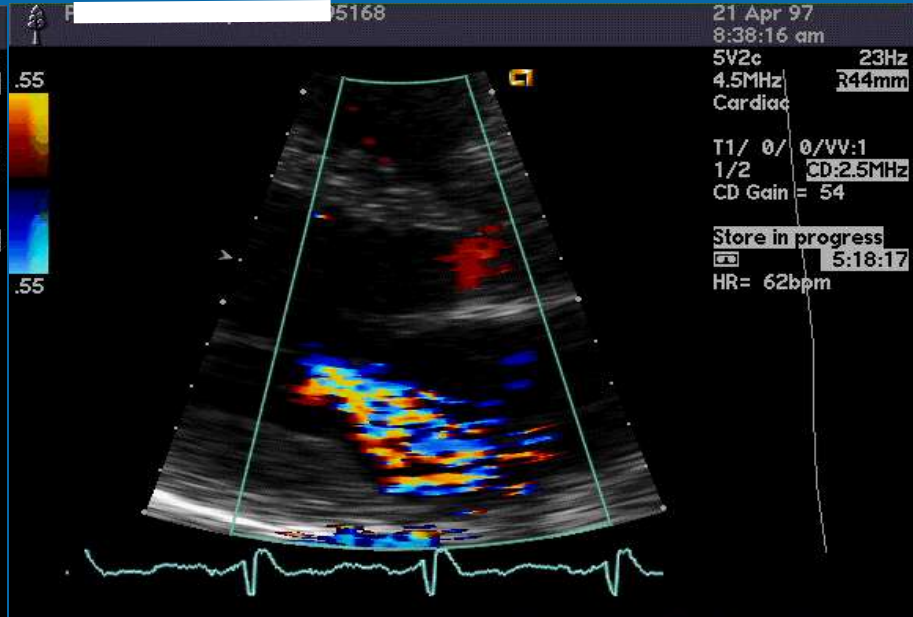
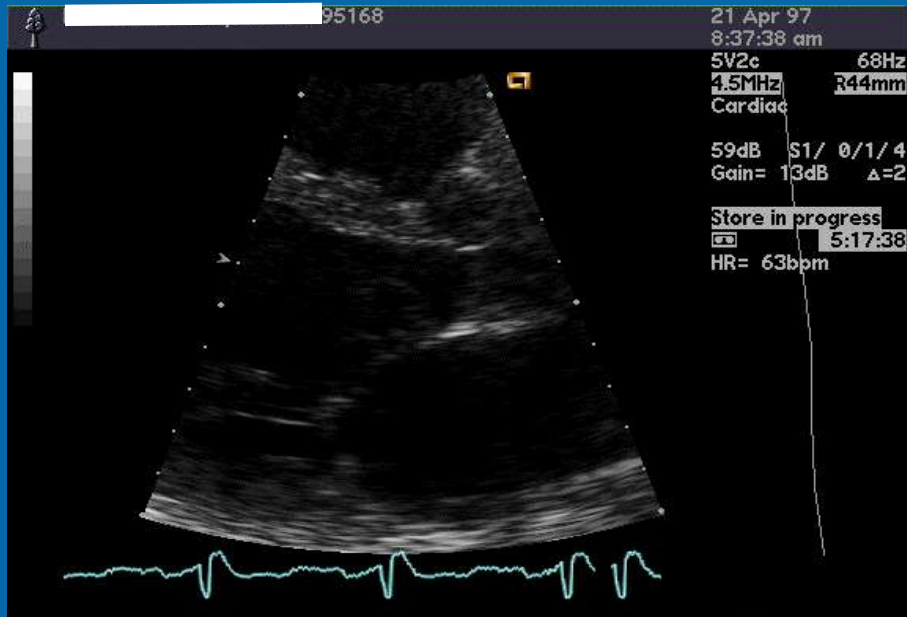


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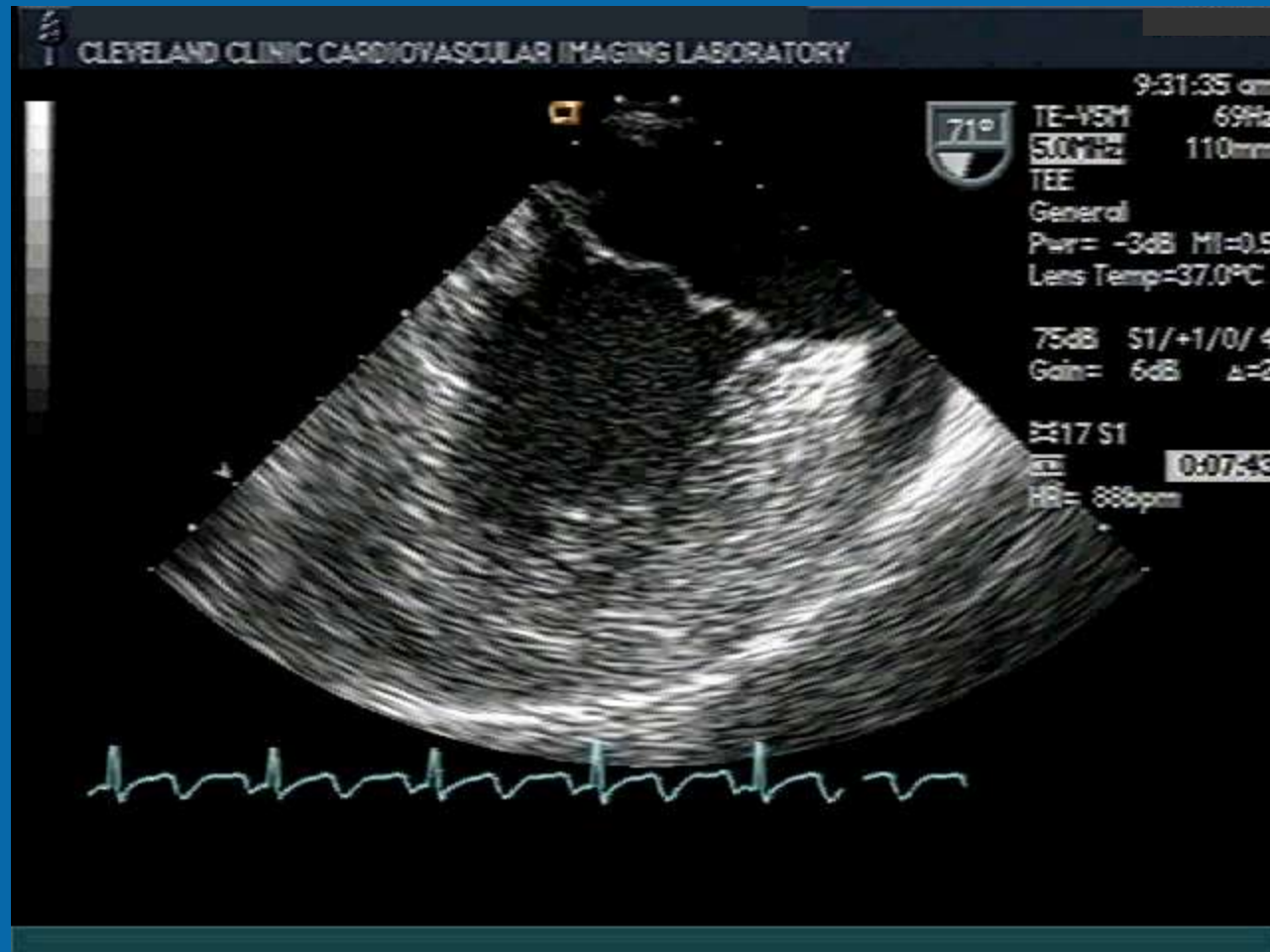
Repair or Replace?



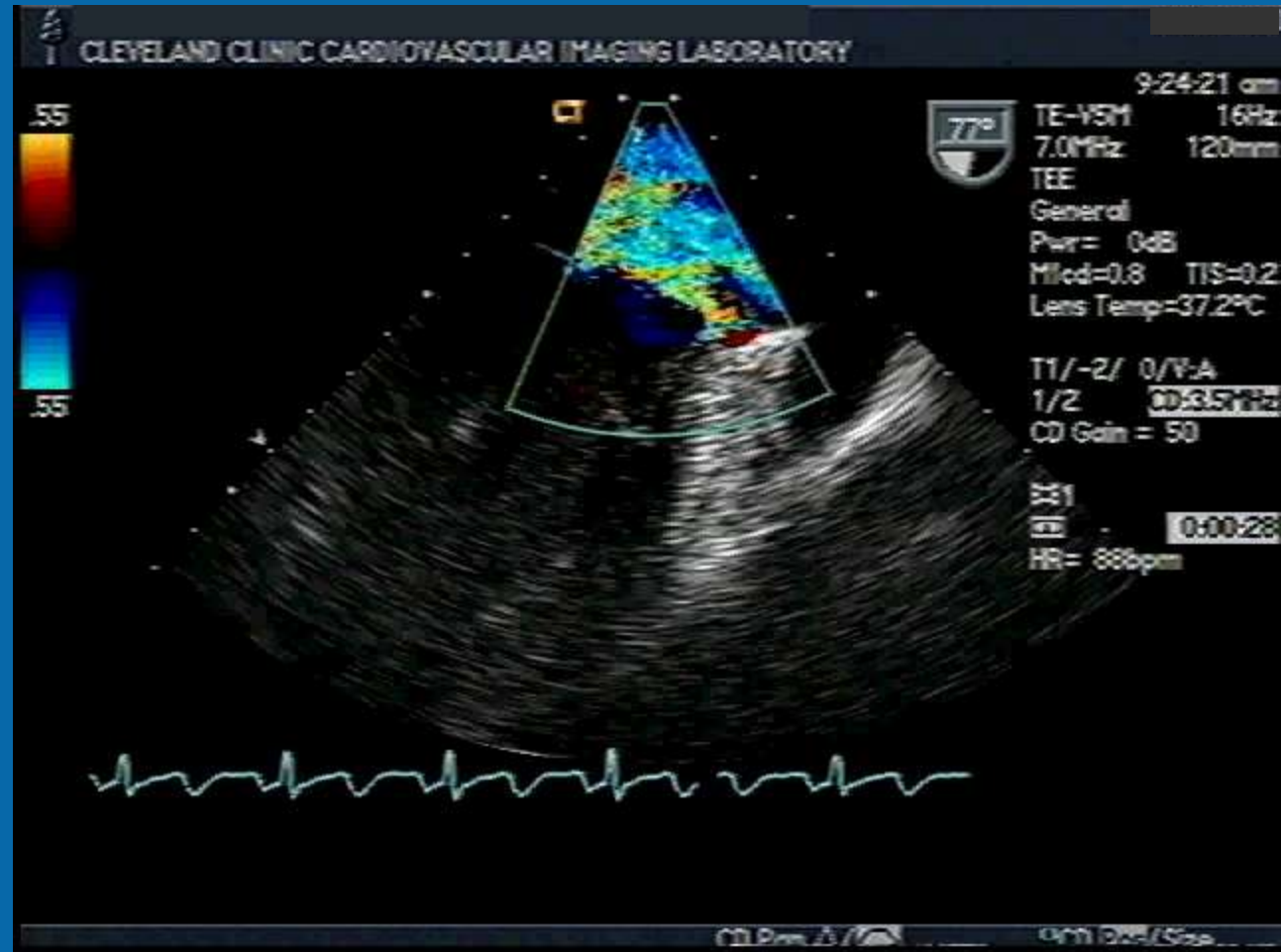
Repair or Replace?



What is the Pathology?



Mechanism?

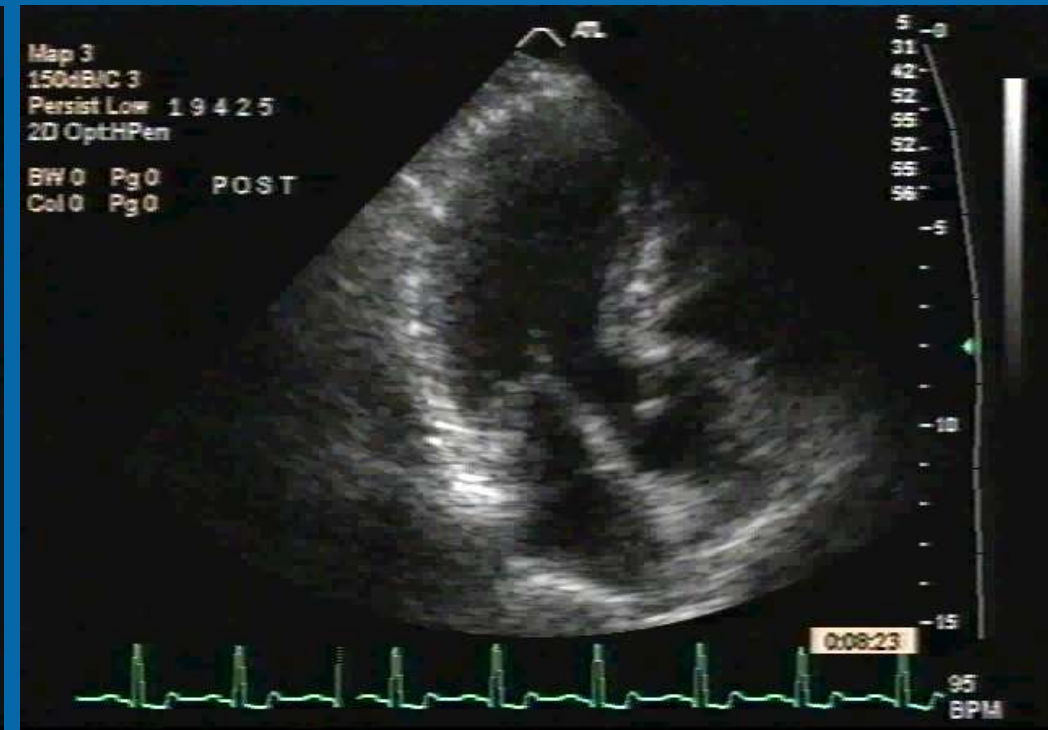


Stress Echo

Rest

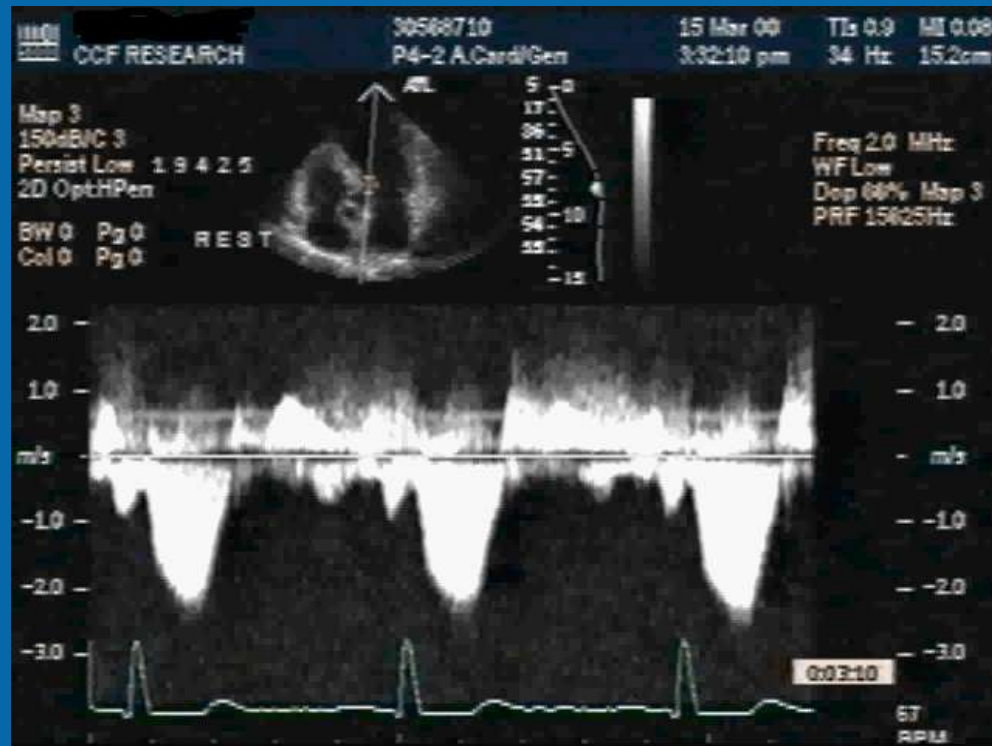


Stress



LVOT Gradients

Rest

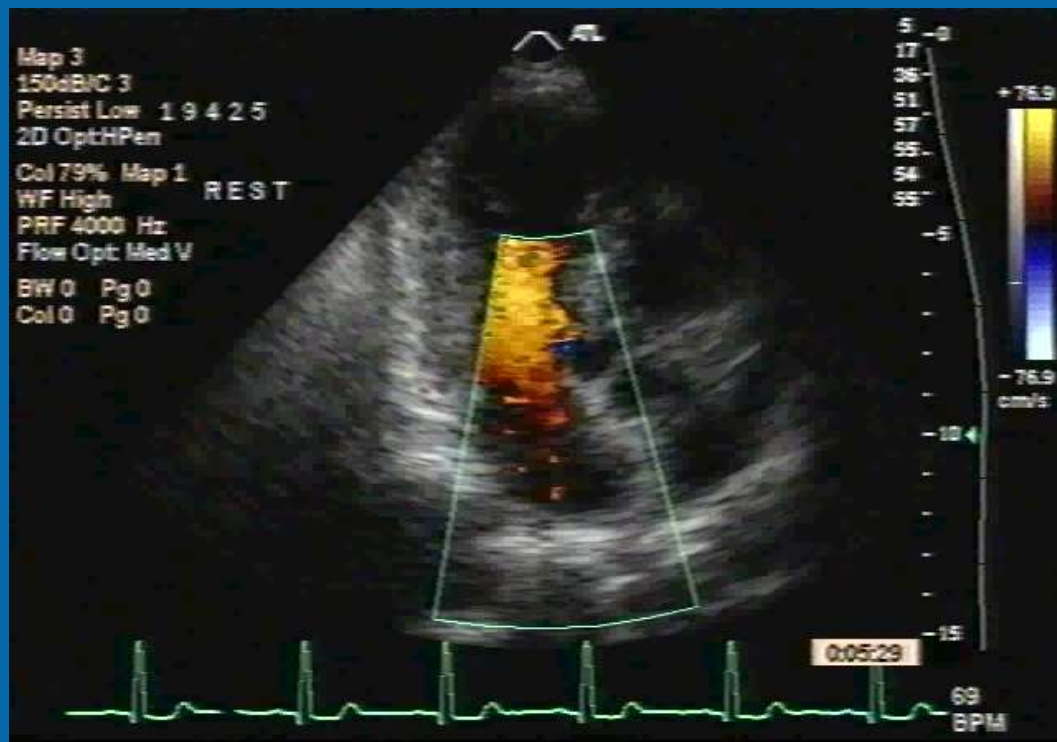


Stress

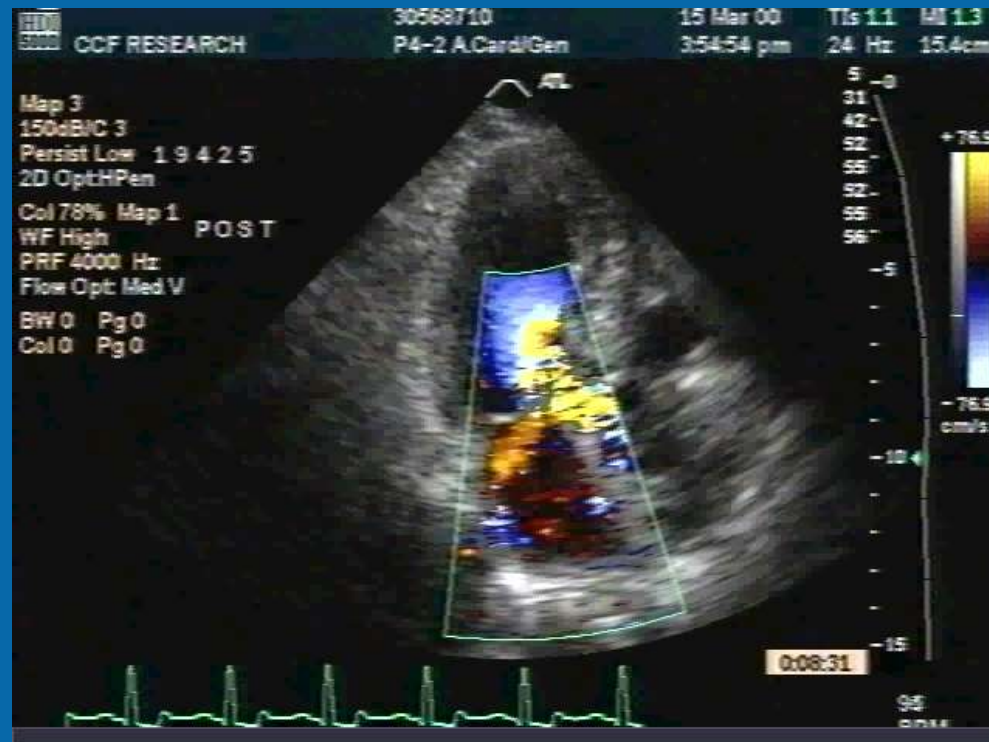


Repair or Replace?

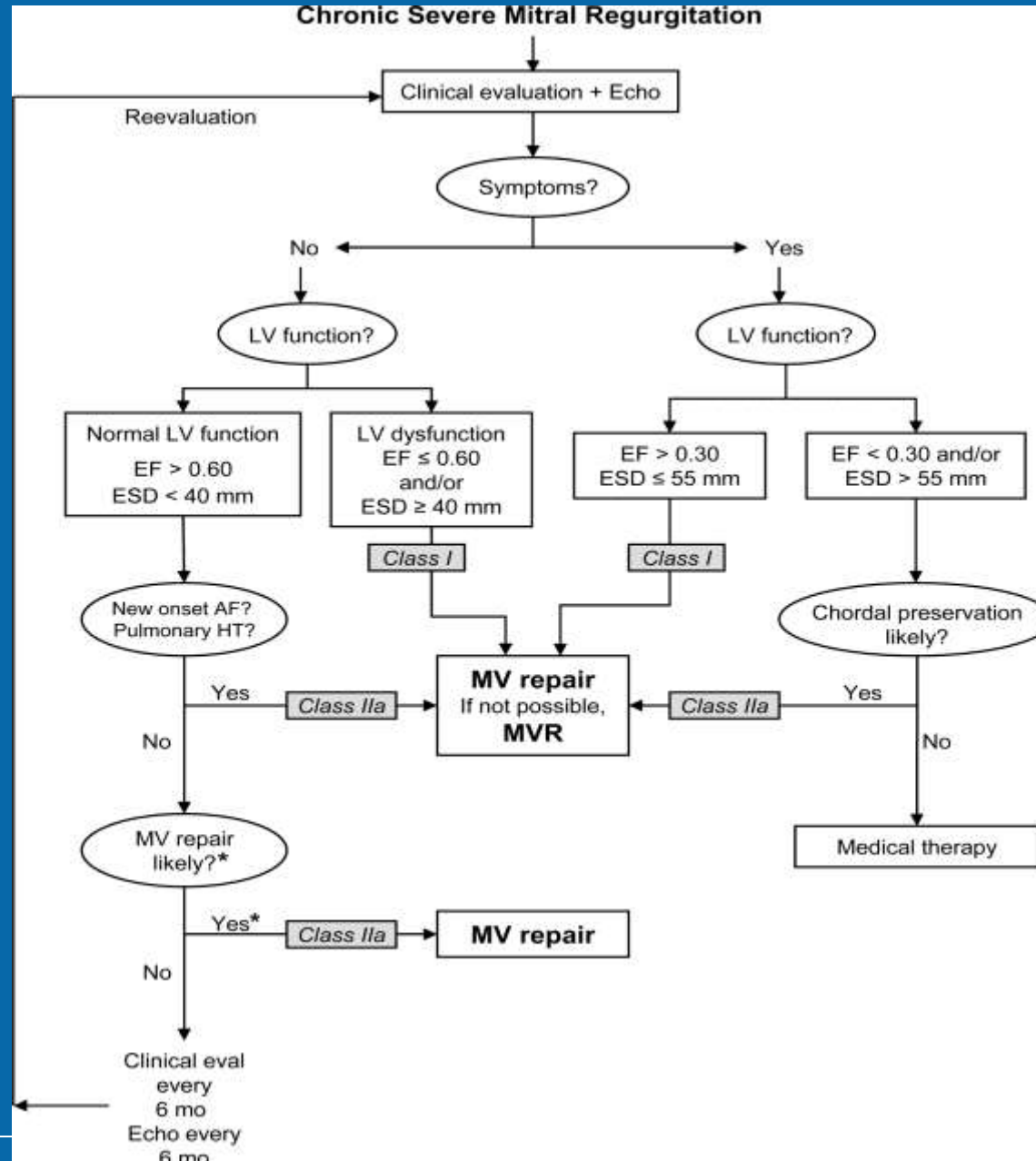
Rest



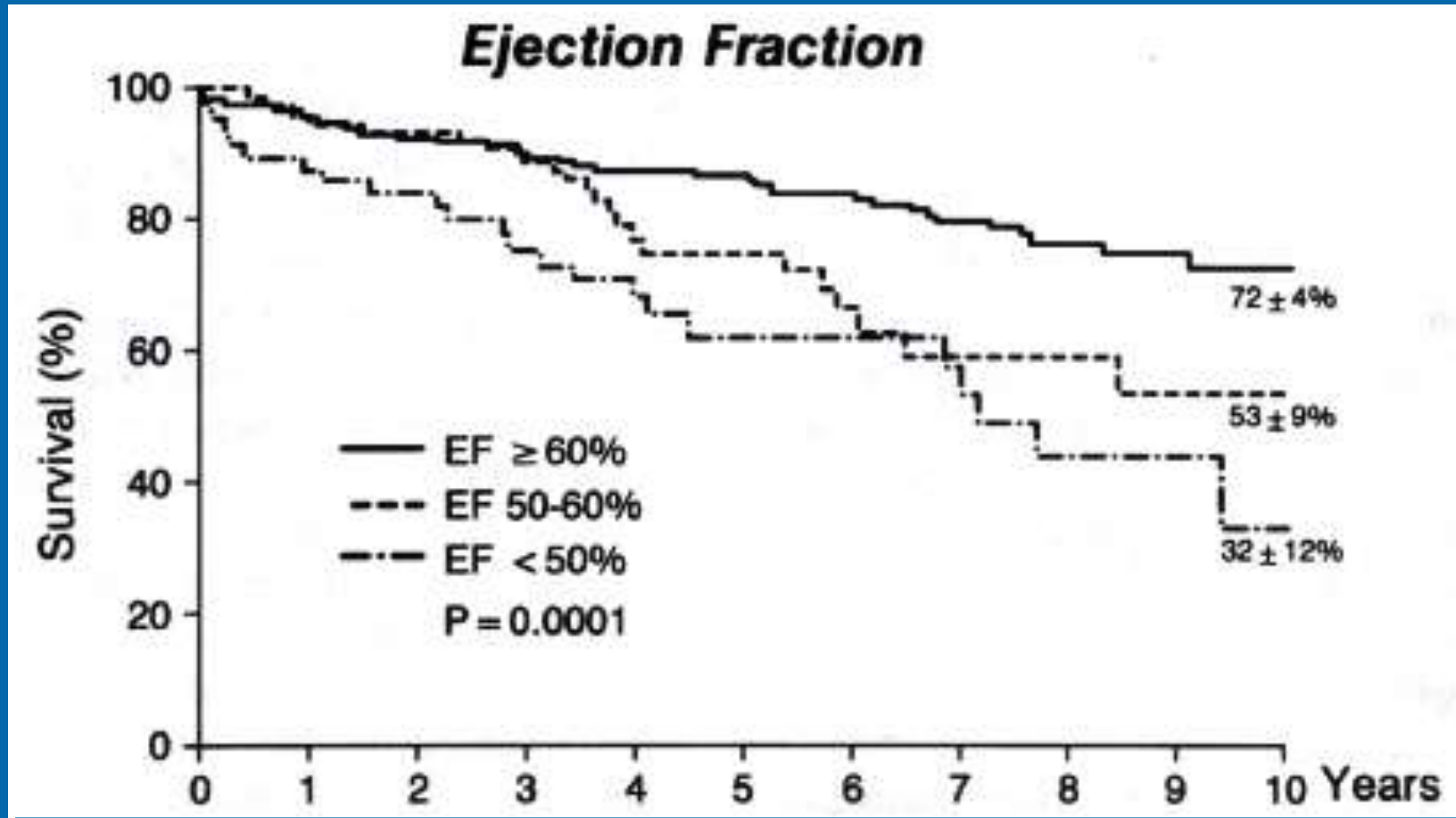
Stress



Management of Patients With Chronic Severe MR



Timing of Surgery: LV Function



Frequency of Echo Follow-up for Mitral Regurgitation

MR Severity	LV Function	Frequency
Mild	Normal ESD & EF	Q 5 yr
Moderate	Normal ESD & EF	Q 1-2 yr
Moderate	ESD >40 mm or EF < 65%	Q1 yr
Severe	Normal ESD & EF	Q 1yr
Severe	ESD > 40 mm or EF < 65%	Q 6 mo