

Catching a glimpse: Aortic Stenosis and Mitral Regurgitation



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Disclosures

No relevant disclosures



Agenda

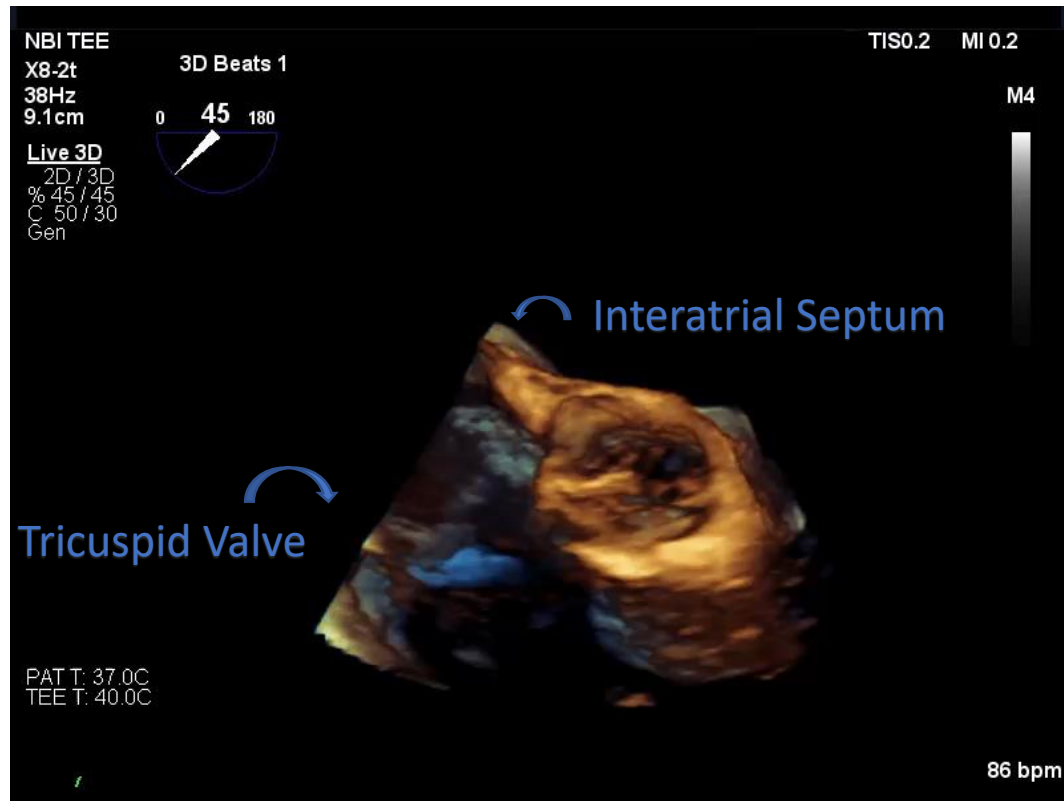
- Spectrum of disease: from normal to abnormal
- Establishing the severity of valvular pathology
- Guiding future management

Aortic Valve: normal to abnormal

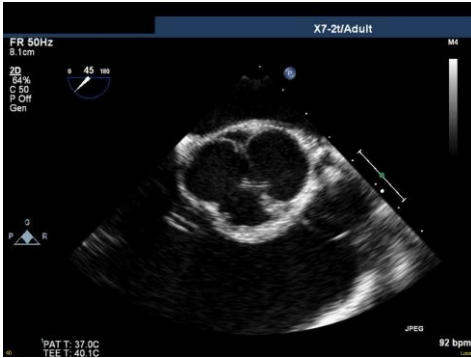


Three leaflets, mobile, thin. No evidence of restricted motion or hypermobile motion.

Normal 3D of the Aortic Valve

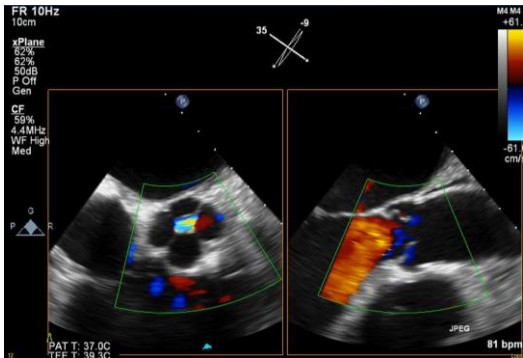


Types of aortic valve

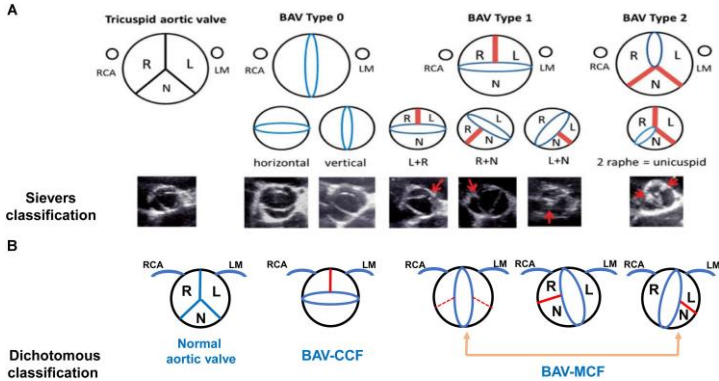


Unicuspid Aortic Valve
 Rare in adults (0.02%)
 Fatal AS in children who are <1 years old
 Sometimes 1 commissure, sometimes no commissure
 Aortic Stenosis associated with ascending aortic aneurysm

Bicuspid Aortic Valve
 Fusion of the 2 cusps
 Associated with aortic stenosis and aortic regurgitation
 Frequent 1-2% of the patients
 Associated with aortic pathology

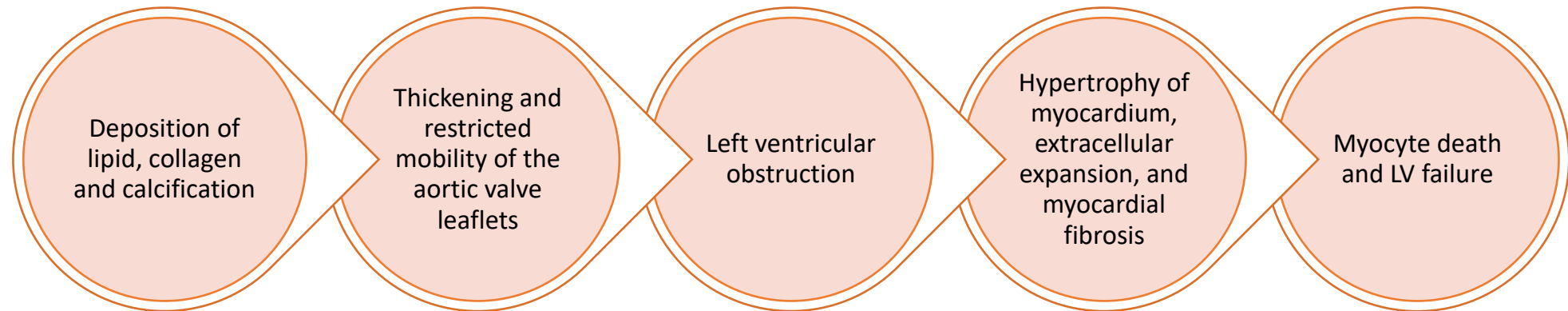


Quadracuspid Aortic Valve
 Unequal cusp size
 More aortic regurgitation than stenosis
 Associated with anomalous coronaries

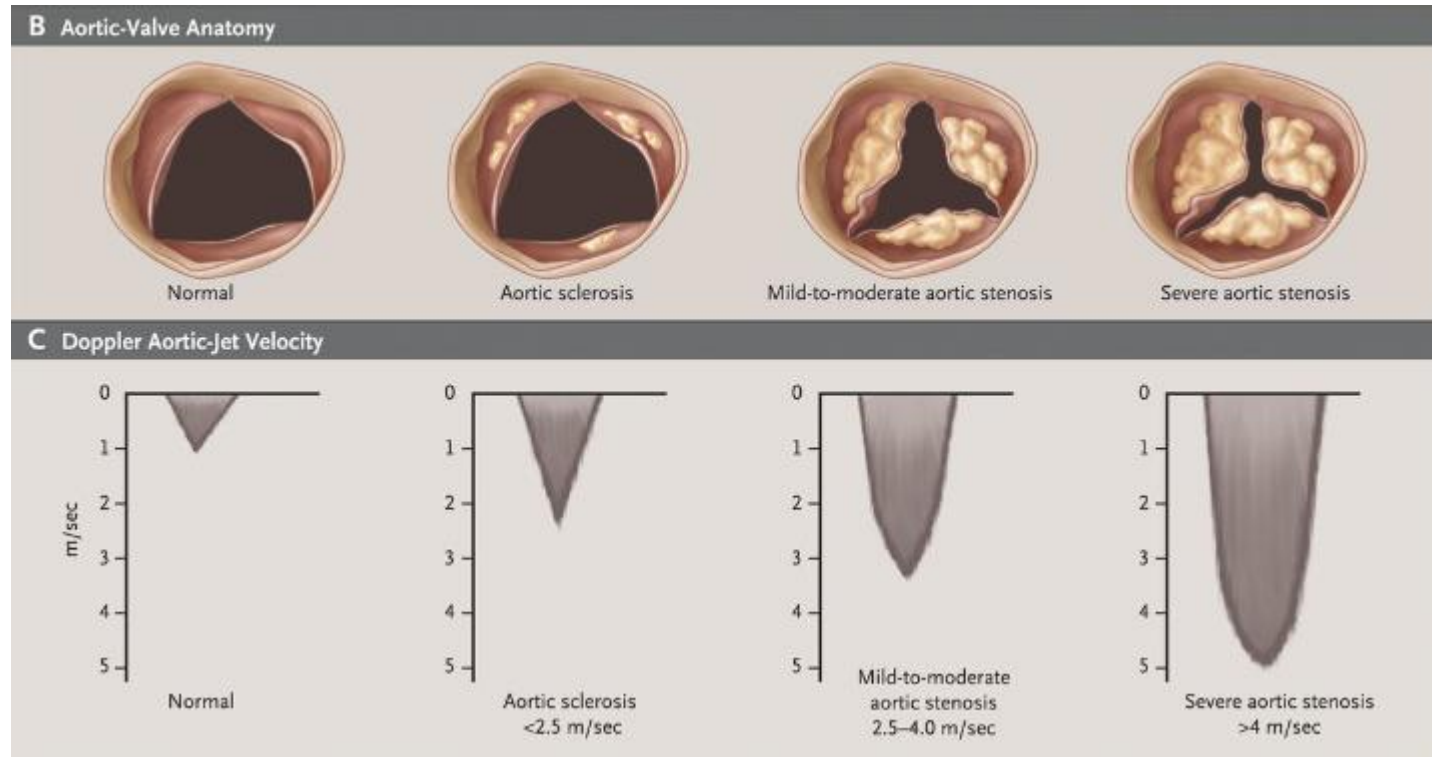


Aortic Stenosis

- Affects more than 7 million people over 75 years of age in Europe and North America
- Most common valvular pathology in developed countries



Progression of aortic stenosis



- Transthoracic Echo is the clinical standard for screening for AS and evaluating severity
- ACC/AHA guidelines: $V_{max} >4\text{m/s}$, mean gradient $>40\text{mmHg}$ and $AVA <1\text{cm}^2$
- AS severity assessment is challenging in patients with LV dysfunction

Discordant Aortic Stenosis

Is this Moderate vs Severe Aortic Stenosis?

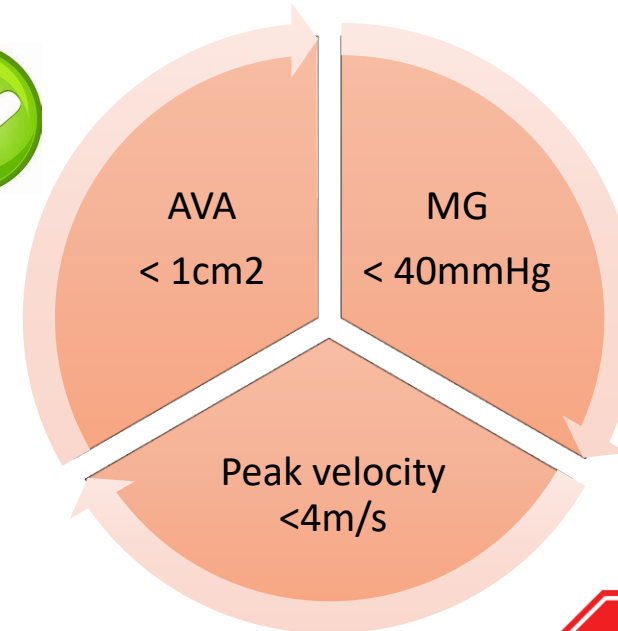
65-70% of patients have concordant findings
BUT
25-30% have discordant

Step 1: Make sure LVOT is not underestimated.
LVOT measurement is the most common error.

Use 3D or CT to confirm LVOT size

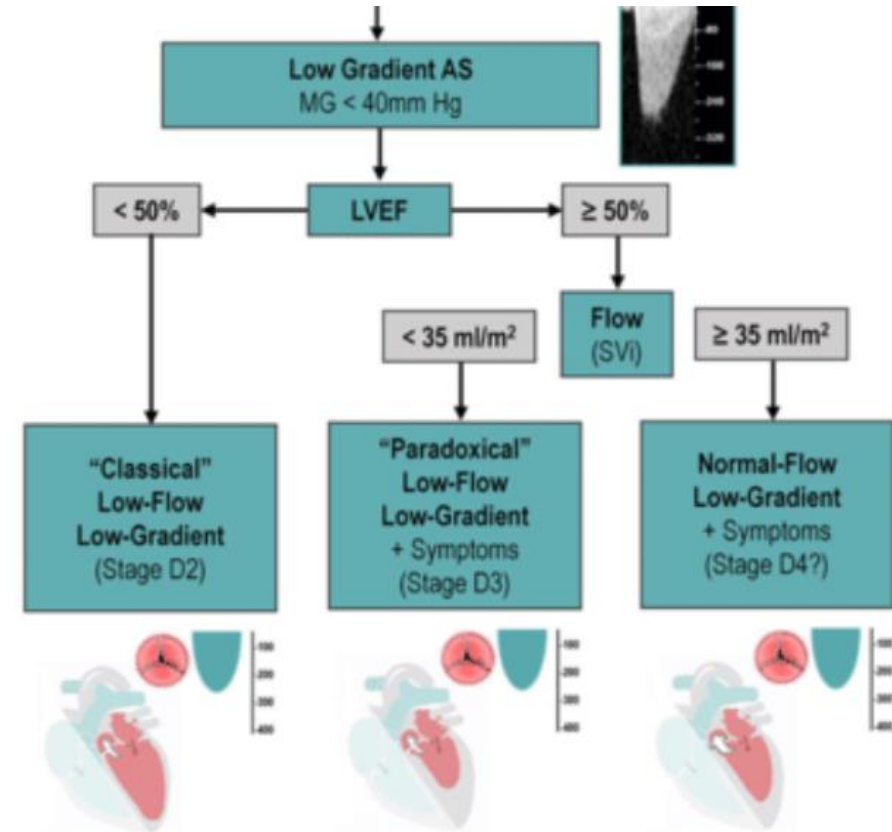
Step 2: Confirm low gradient: multi-window interrogation (suprasternal and right parasternal)

Step 3: Confirm low flow state and identify cause

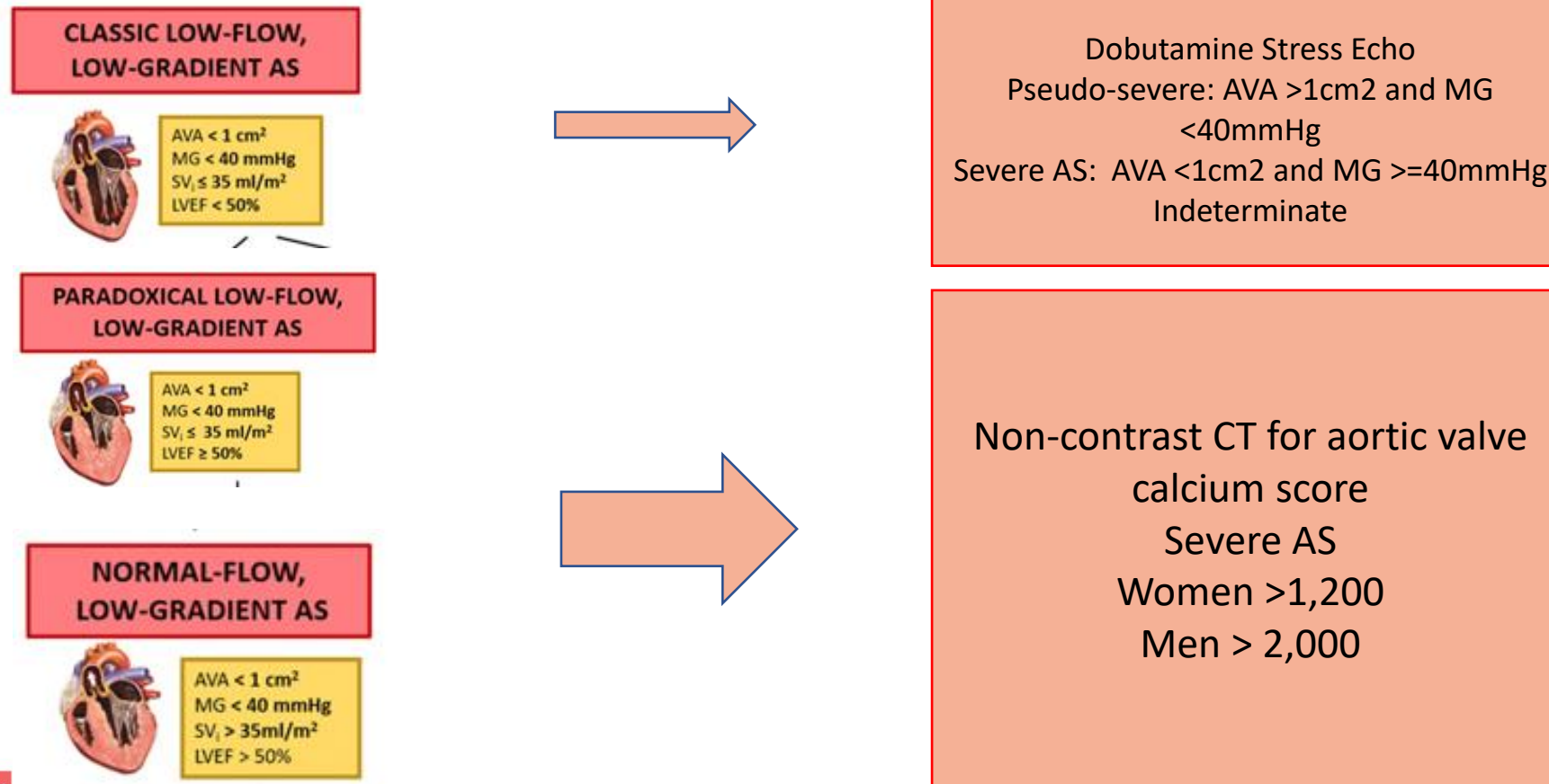


Classification of AS

- Classical Low-Flow Low-Gradient AS
 - Pseudosevere vs severe AS
 - LVEF <50%
- Paradoxical Low-Flow Low-Gradient AS
 - Normal LVEF (>50%)
 - Low stroke volume indexed (<35cc/m²)
- Normal-flow Low-Gradient
 - Normal LVEF and normal stroke volume



Diagnostic Pathways



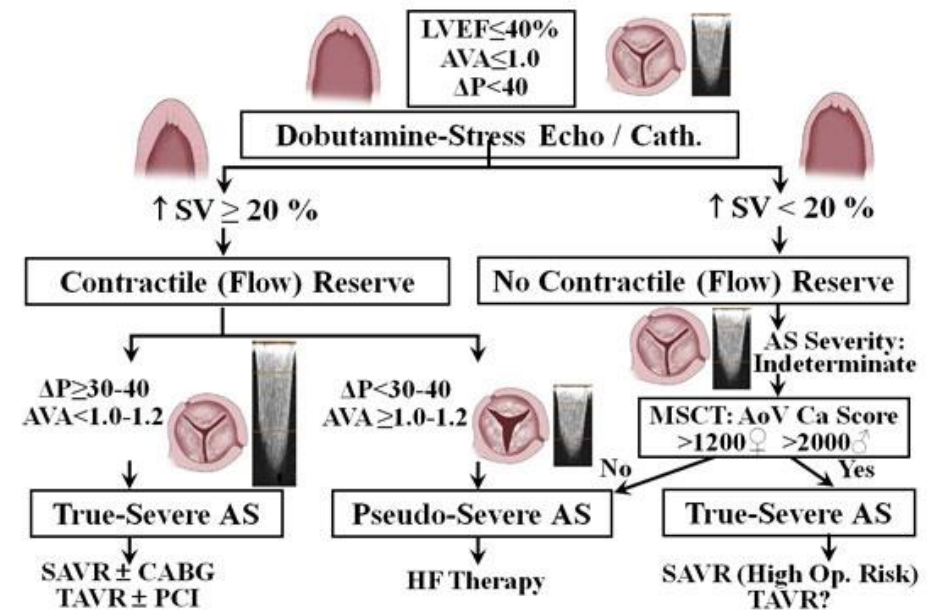
Goes back to 1995

- DeFilippe et al. evaluated 18 patients with dobutamine stress echo: $AVA \leq 0.5 \text{ cm}^2/\text{m}^2$, mean gradient $\leq 30 \text{ mm Hg}$ and $LVEF \leq 45\%$.
- Dobutamine was started at $5 \mu\text{g}/\text{kg}/\text{min}$ and increased to a peak of $20 \mu\text{g}/\text{kg}/\text{min}$ at 3-min intervals.

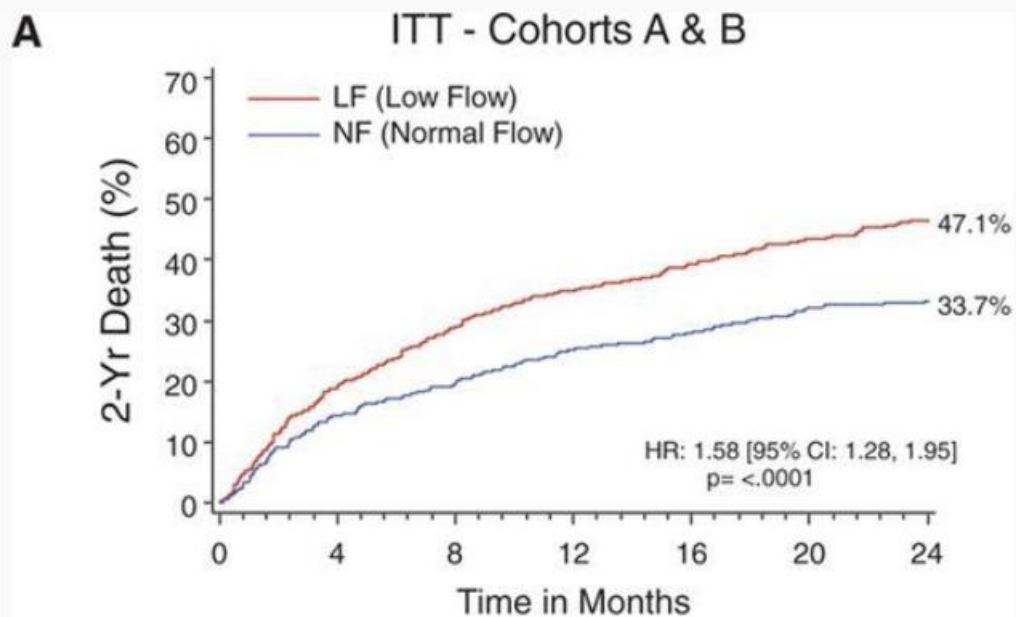
Contractile reserve - increase in peak velocity of 0.6m/s, stroke volume 20% and MPG 10mmHg with dobutamine

True SEVERE AS – small change/no change in AVA but gradients across AV increase

Pseudo-severe AS – AVA increases by 0.3 cm^2 . AV gradient increases slightly or no change

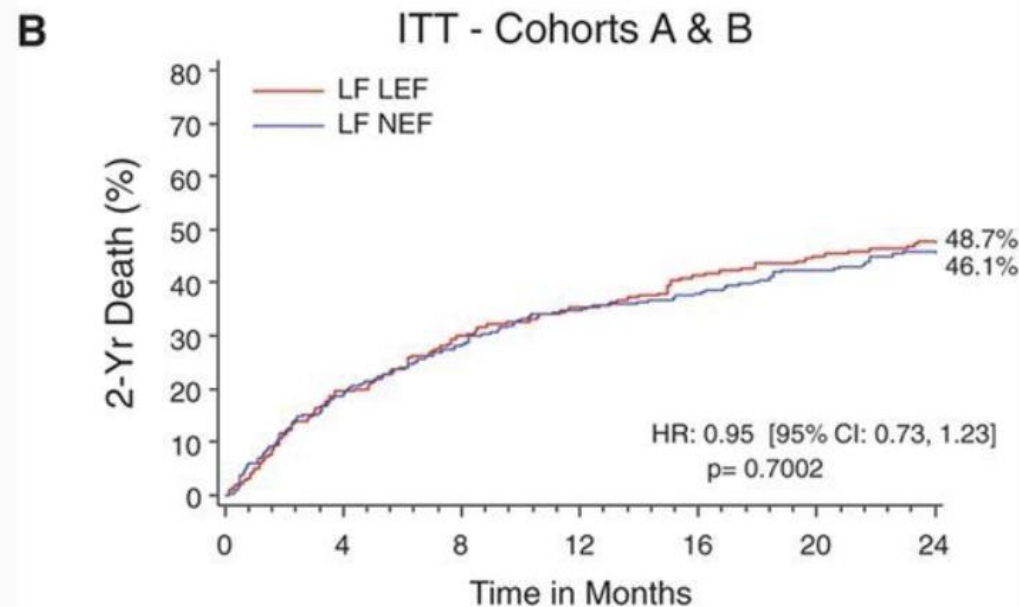


Why is low flow so important?



Number at risk

	0	4	8	12	16	20	24
LF	530	422	368	336	312	287	265
NF	441	368	342	318	302	282	270



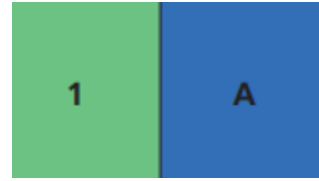
Number at risk

	0	4	8	12	16	20	24
LF LEF	225	177	154	142	128	119	109
LF NEF	304	244	213	193	183	167	155

AS management



3. In asymptomatic patients with severe AS (Stage C1) who are undergoing cardiac surgery for other indications, AVR is indicated.¹²⁻¹⁶



1. In adults with severe high-gradient AS (Stage D1) and symptoms of exertional dyspnea, HF, angina, syncope, or presyncope by history or on exercise testing, AVR is indicated.¹⁻⁷



4. In symptomatic patients with low-flow, low-gradient severe AS with reduced LVEF (Stage D2), AVR is recommended.¹⁷⁻²⁴

Stage C:

C1: Asymptomatic Severe

$V_{max} \geq 4\text{m/s}$ and $MG \geq 40\text{mmHg}$, $AVA < 1\text{cm}^2$

C2: Asymptomatic Severe with LV dysfunction

$LVEF < 50\%$

Stage D:

D1: Symptomatic severe High gradient AS

D2: Symptomatic severe low-flow, low-gradient AS with reduced LVEF

D3: Symptomatic severe low gradient AS with normal LVEF or Paradoxical low-flow severe AS



2. In asymptomatic patients with severe AS and an $LVEF < 50\%$ (Stage C2), AVR is indicated.⁸⁻¹¹



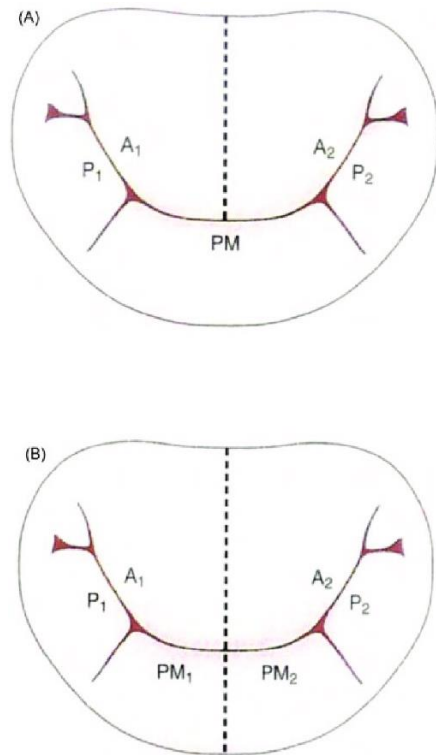
5. In symptomatic patients with low-flow, low-gradient severe AS with normal LVEF (Stage D3), AVR is recommended if AS is the most likely cause of symptoms.²⁵⁻²⁷

Moving on to Mitral Valve

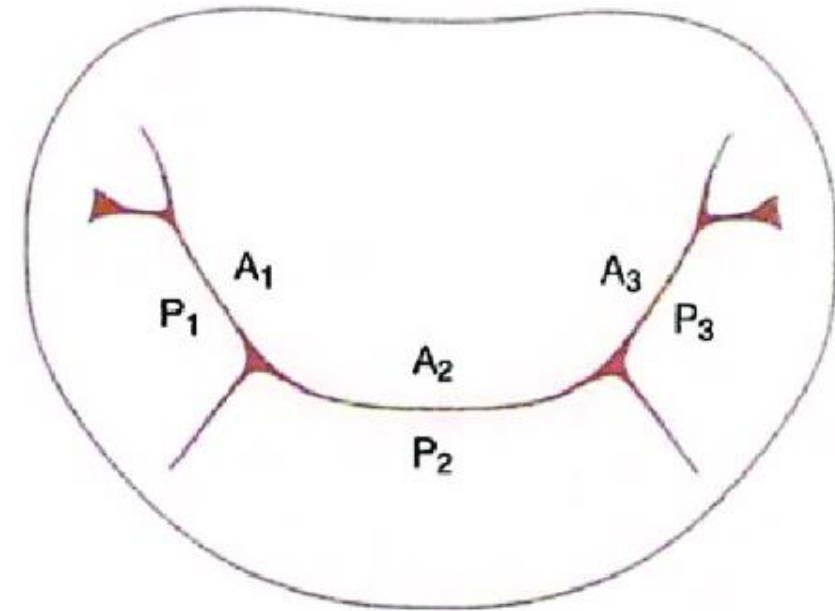


Mitral Valve Nomenclature

- Duran classification



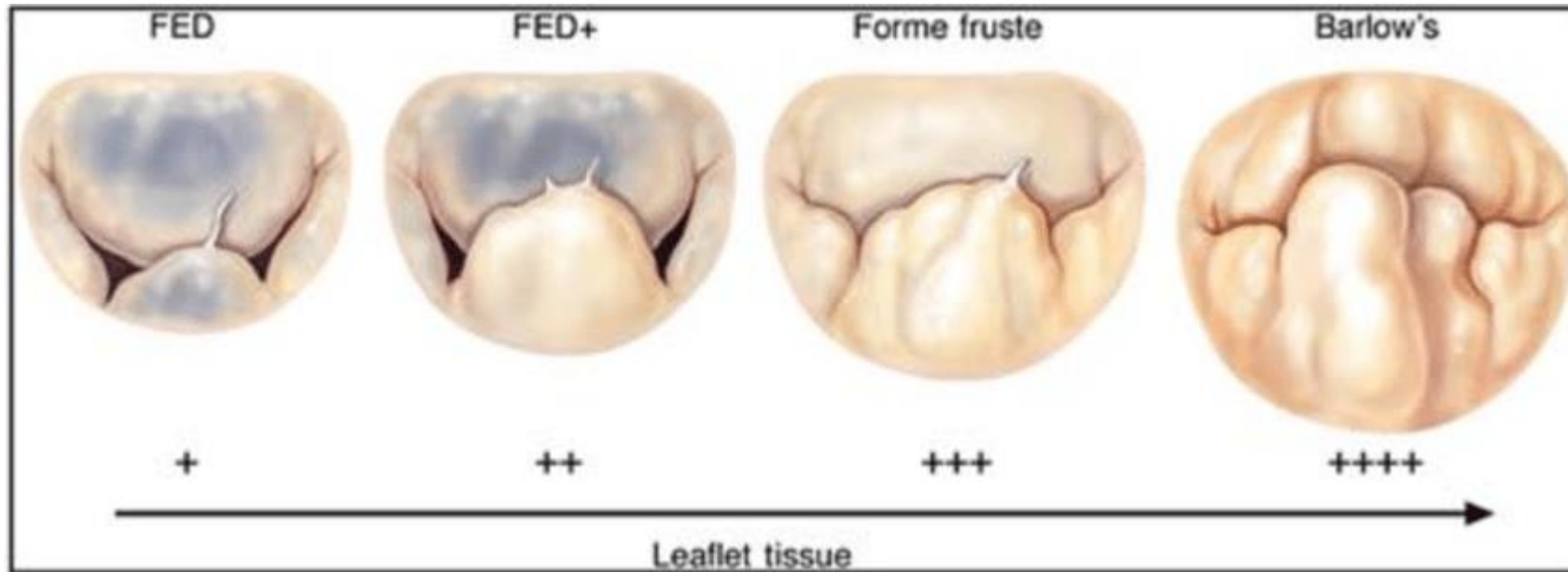
- Carpentier classification



Primary Mitral Regurgitation

1. Abnormality of the MV apparatus
2. Most common cause is myxomatous degeneration
Ranges from focal prolapse to Barlow's disease
3. Other causes: leaflet perforation, cleft mitral leaflet, rheumatic heart disease, radiation and connective tissue disease

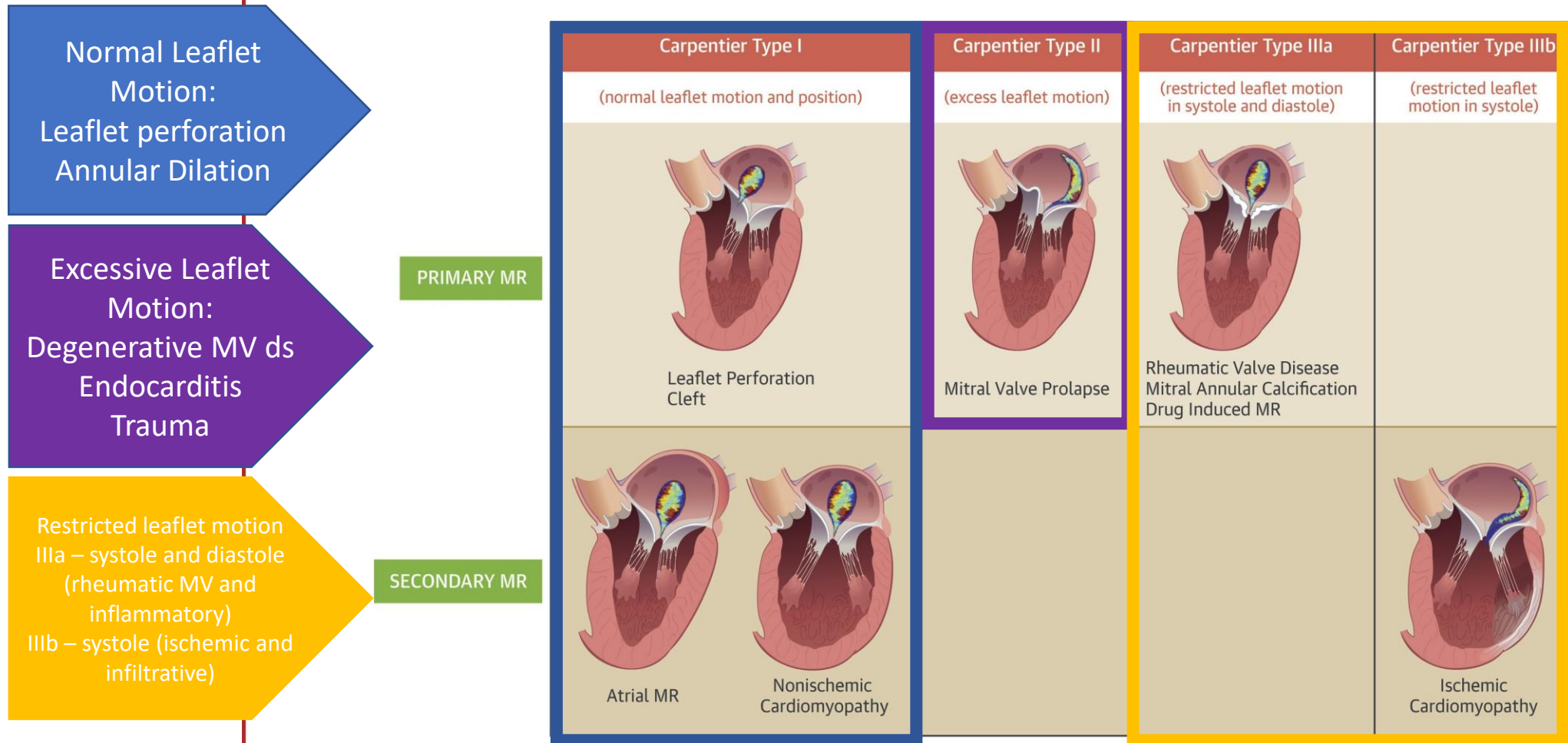
Spectrum of Mitral Valve Prolapse



Secondary Mitral Regurgitation

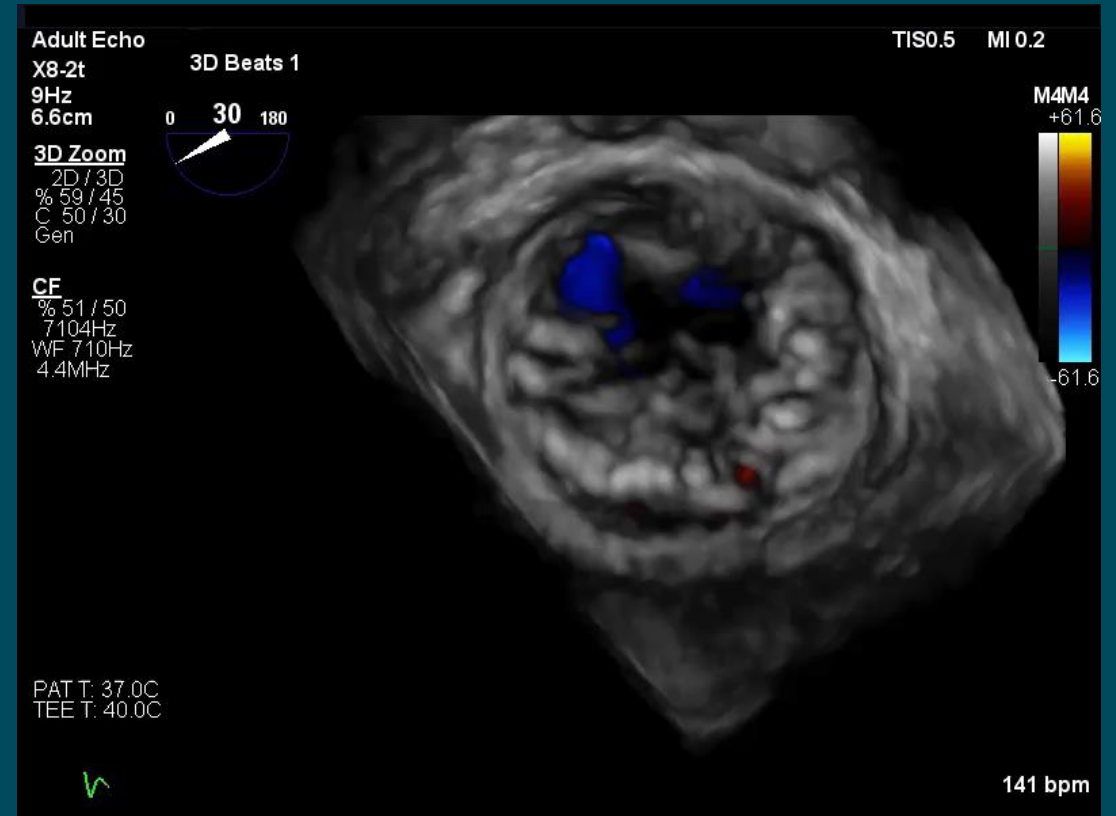
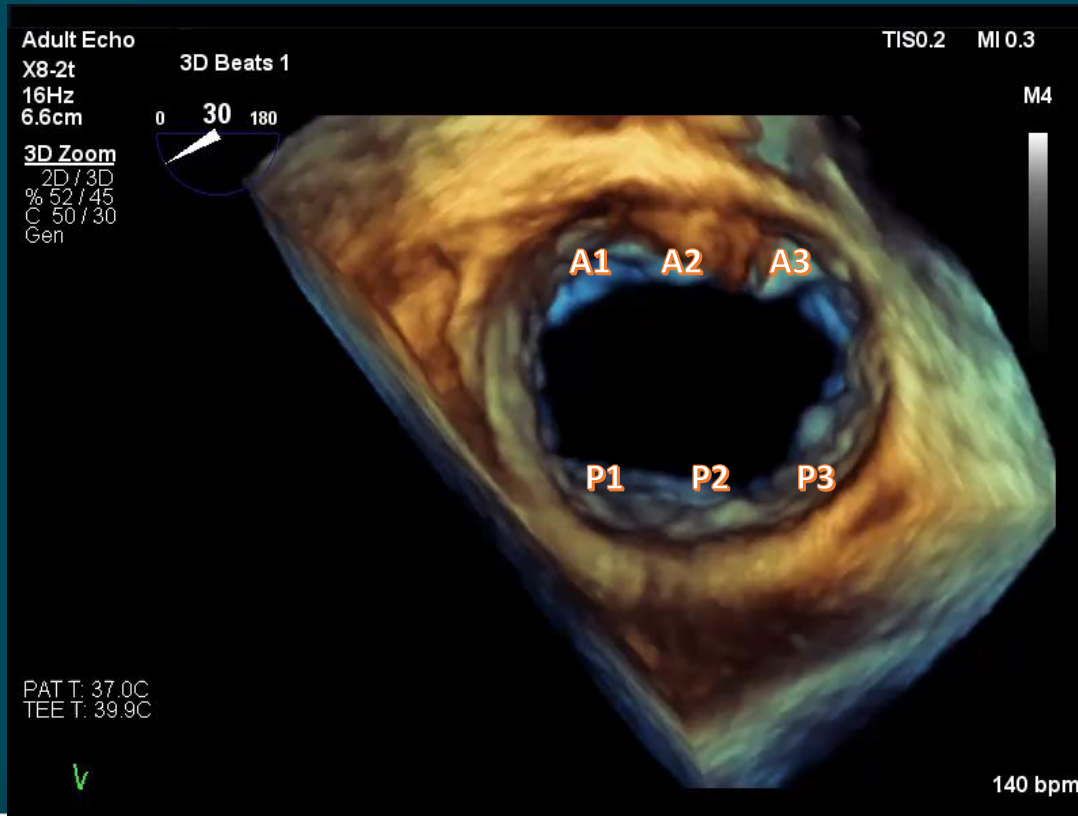
1. Disease of the atrium or ventricle
2. Non-ischemic: increased EROA from annular dilation and loss of annular contraction. Lack of coaptation
3. Atrial MR: severe left atrial enlargement (due to persistent atrial fibrillation)
4. Ischemic: regional inferior wall motion abnormality leading to posterior leaflet tethering and posterior MR. Anterior leaflet override. Can also have central MR due to global wall motion abnormalities

CENTRAL ILLUSTRATION: Classification of the Etiology of MR

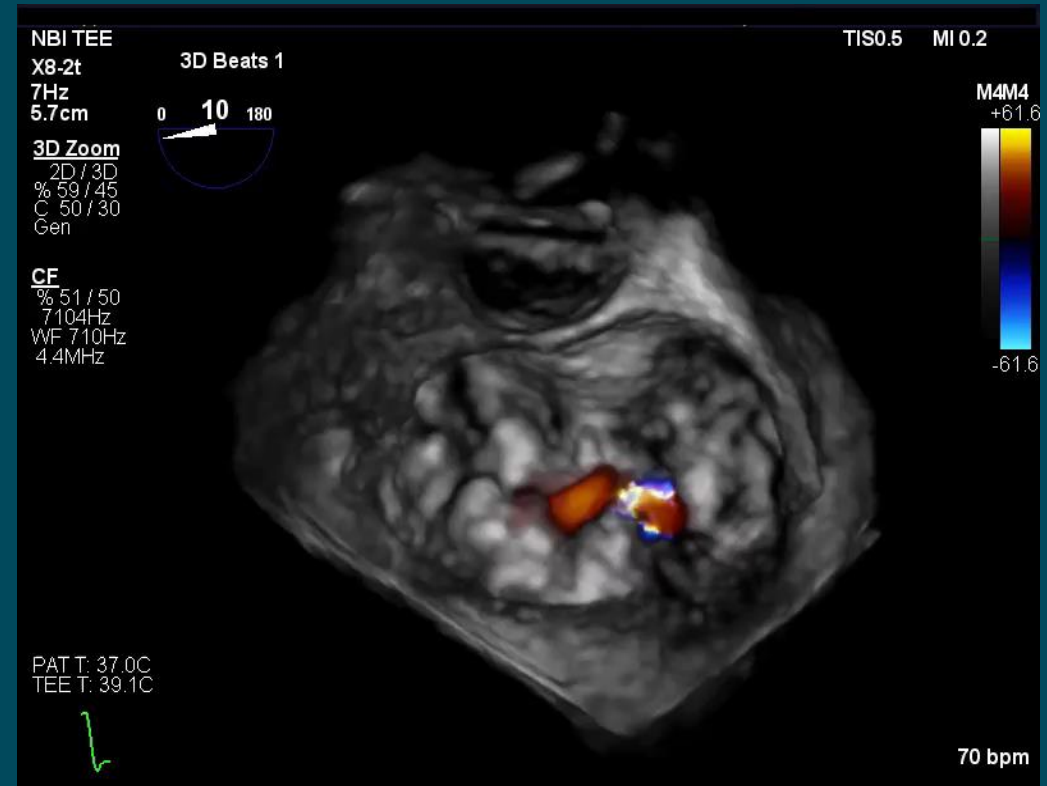
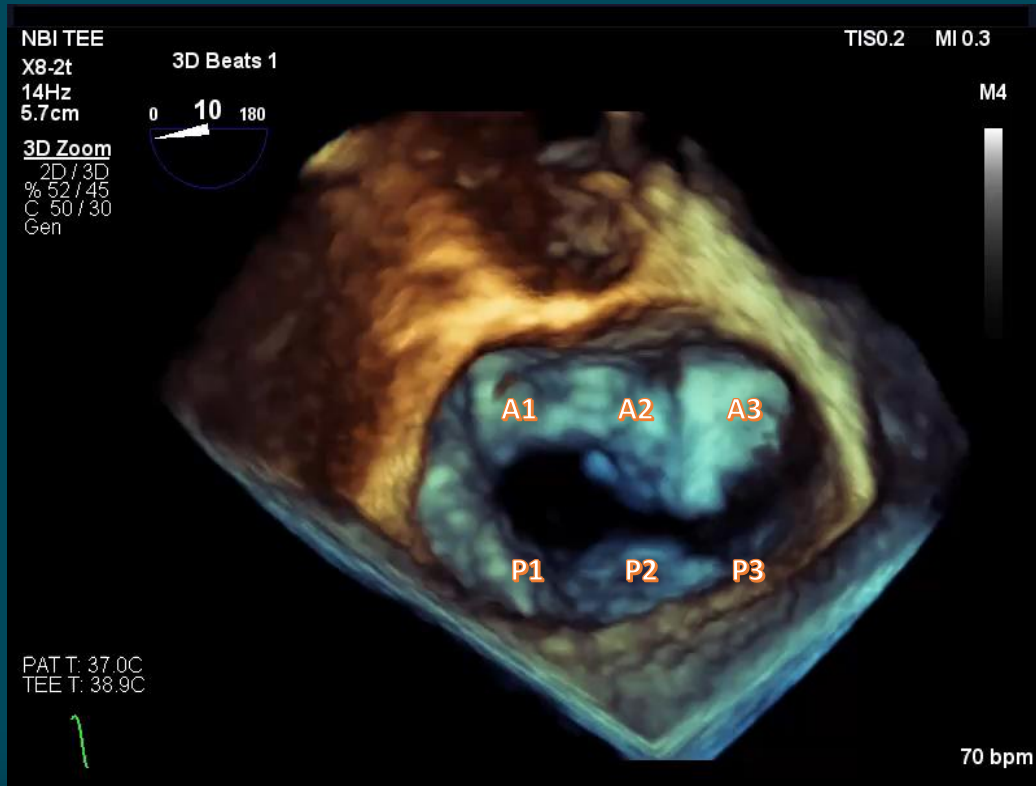


El Sabbagh, A. et al. J Am Coll Cardiol Img. 2018;11(4):628-43.

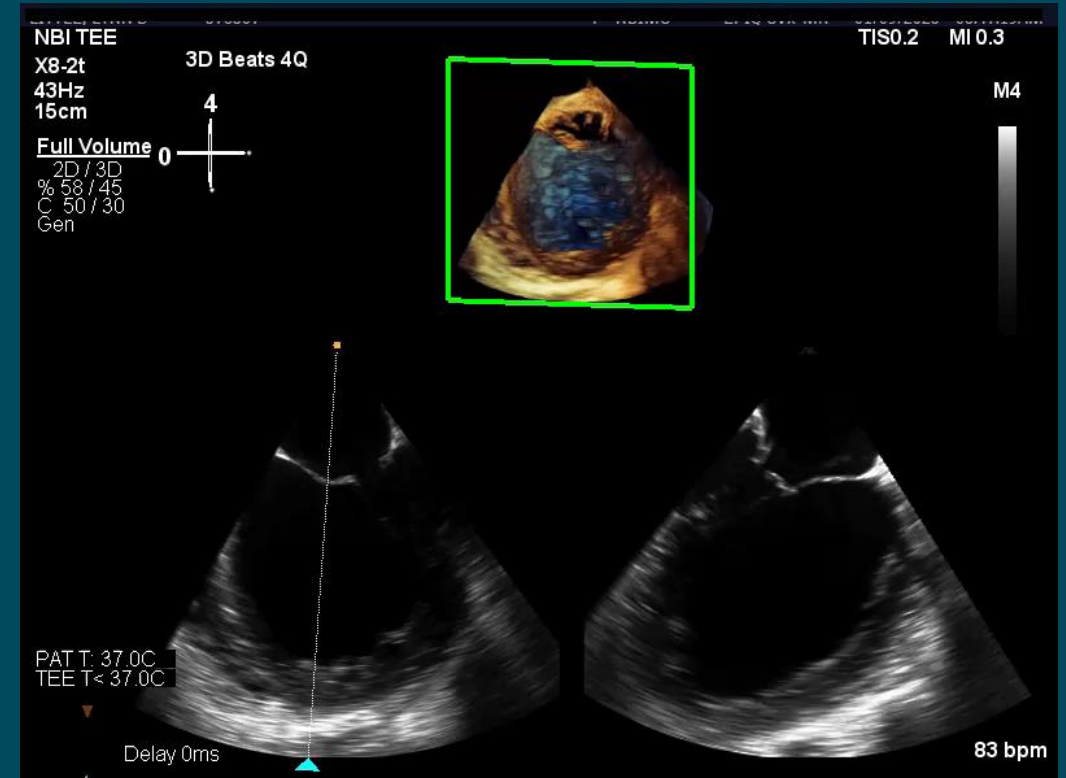
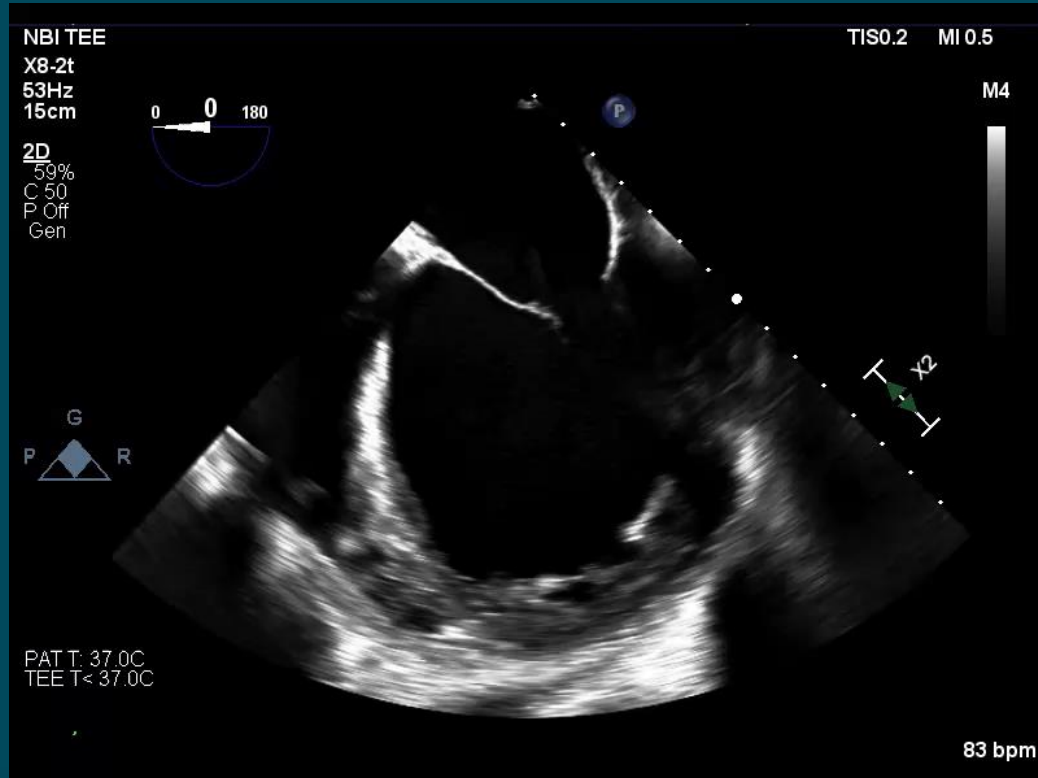
Normal Mitral Valve 3D



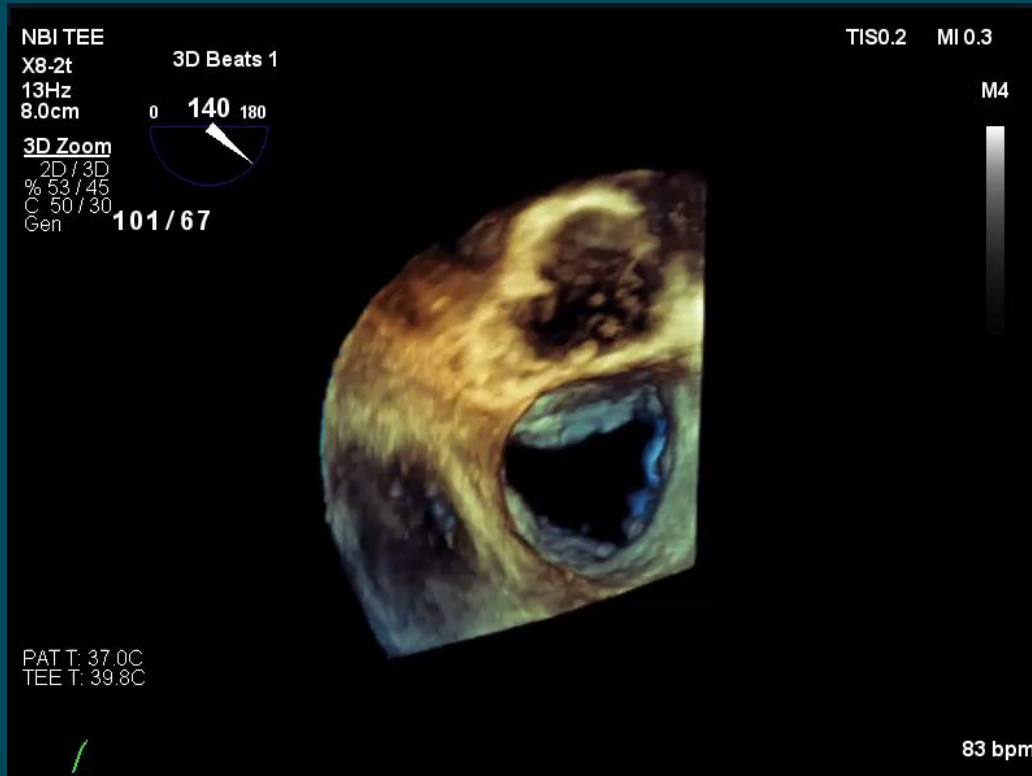
Focal Prolapse (P2)



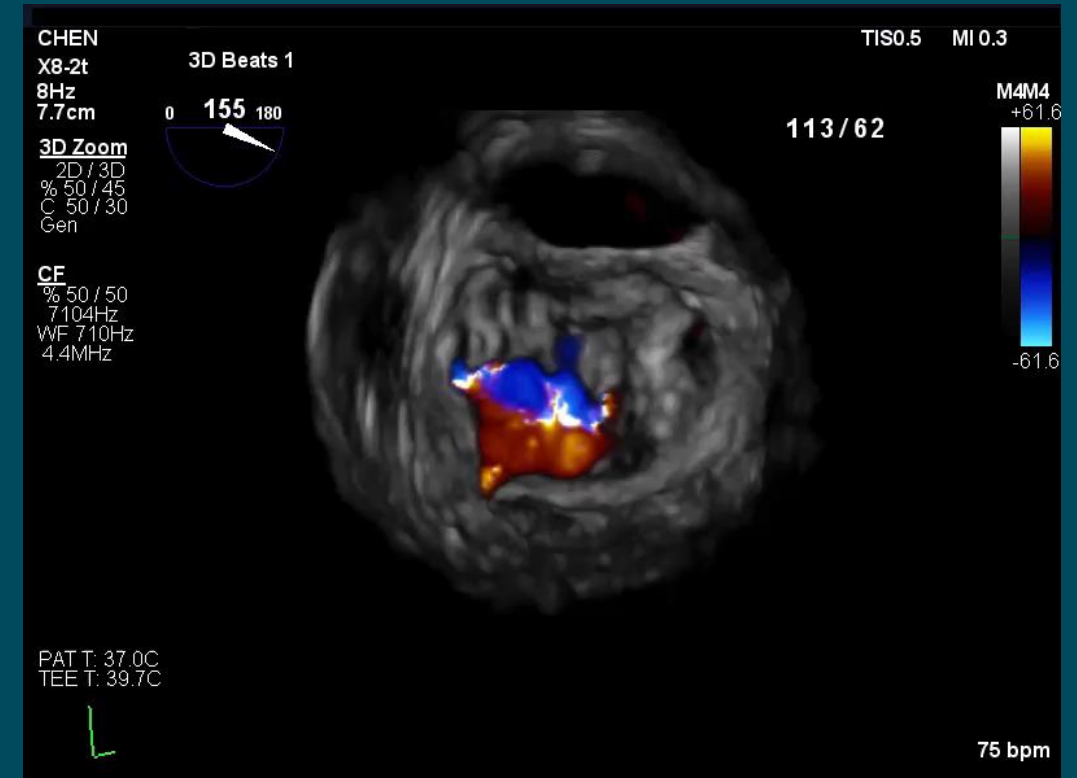
Non-ischemic CM



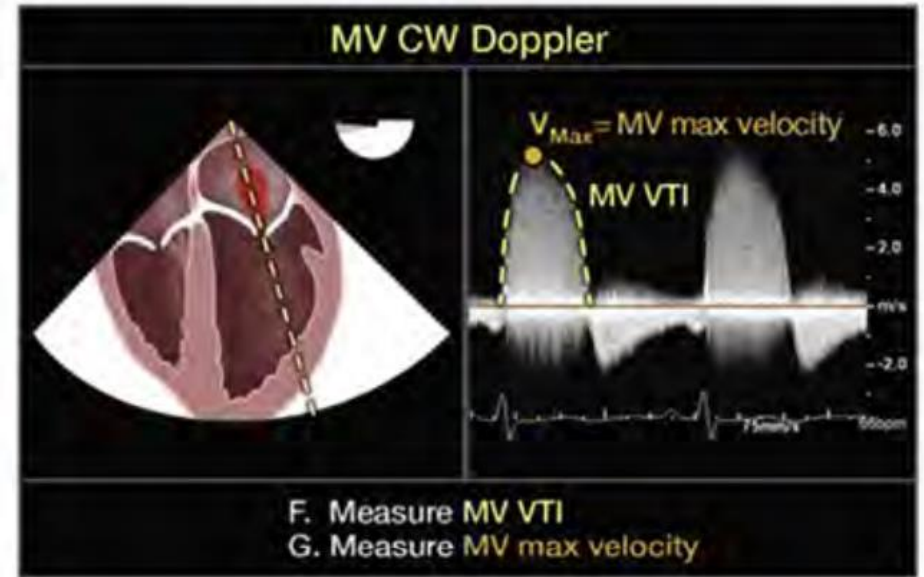
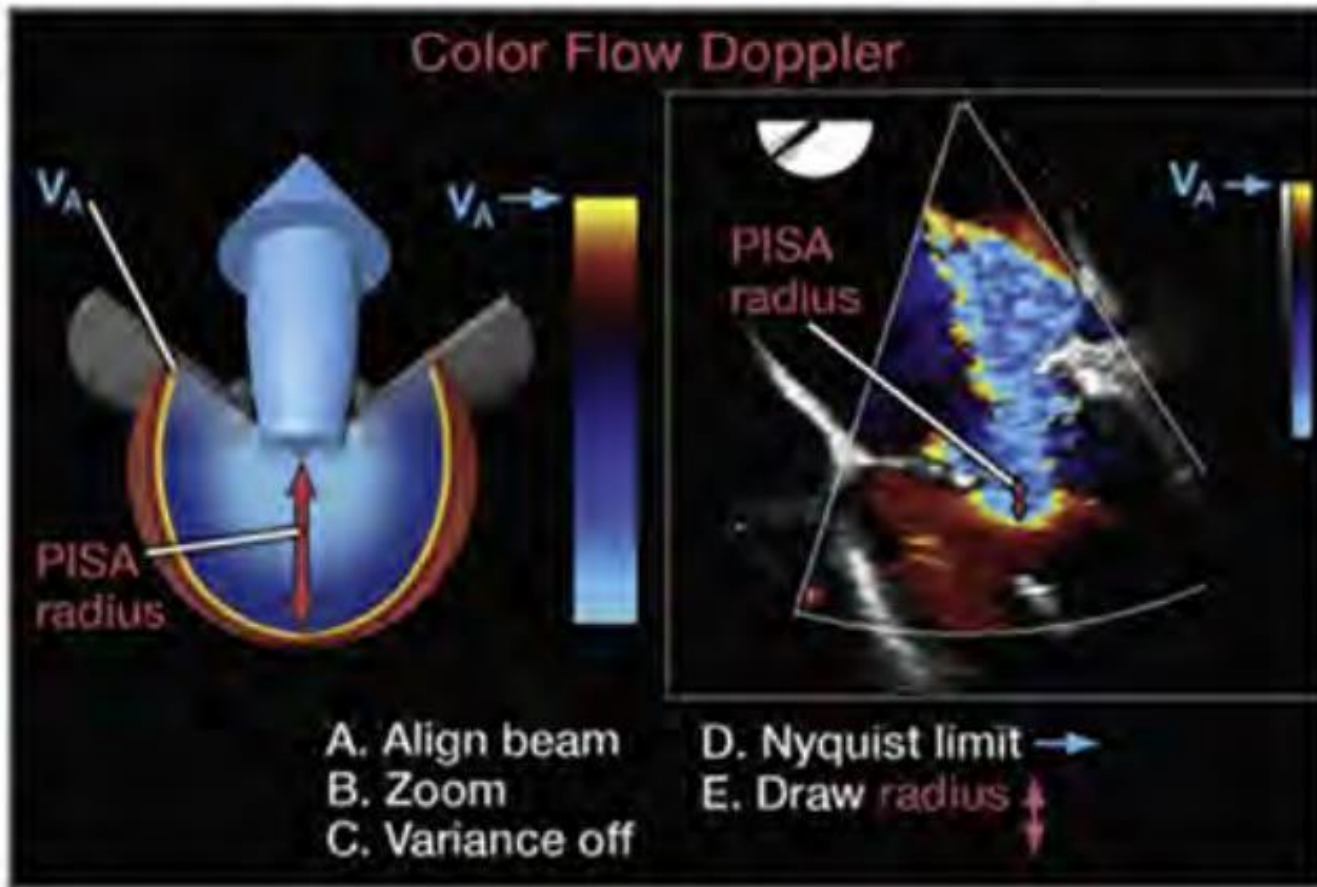
Using 3D to identify pathology



Myxomatous Mitral Valve



Establishing Severity of MR: PISA



$$PISA = 2 \times \pi \times r^2$$

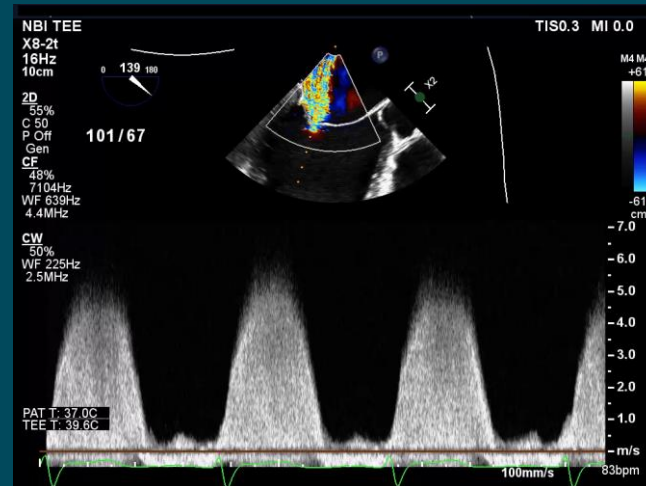
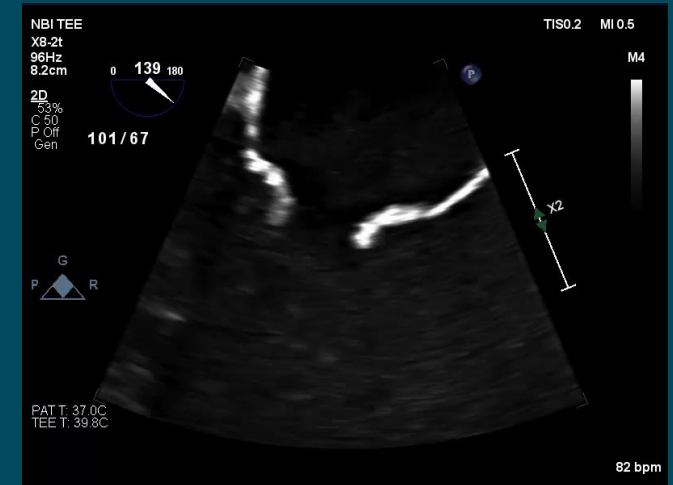
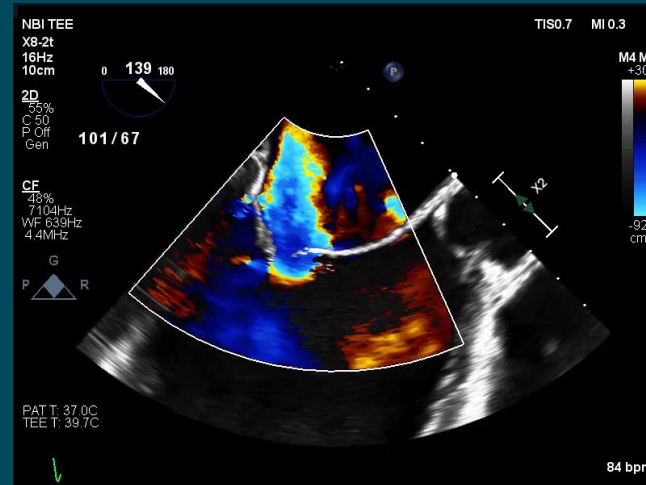
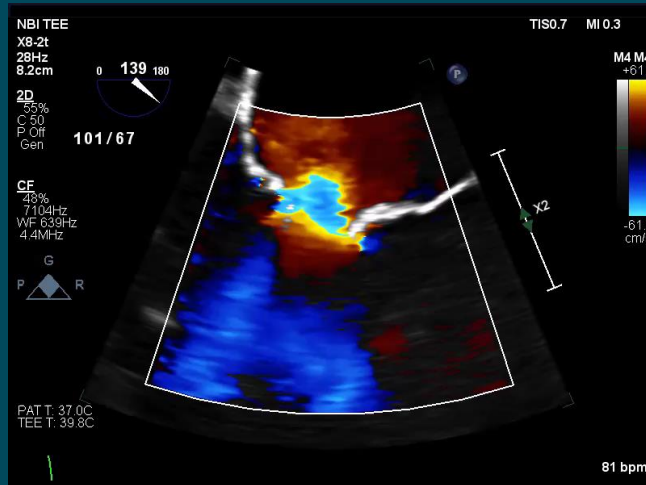
$$\text{Regurgitant flow} = 2 \times \pi \times r^2 \times V_A$$

$$ERO = \frac{\text{Reg. flow}}{V_{Max}} = \frac{2 \times \pi \times r^2 \times V_A}{V_{Max}}$$

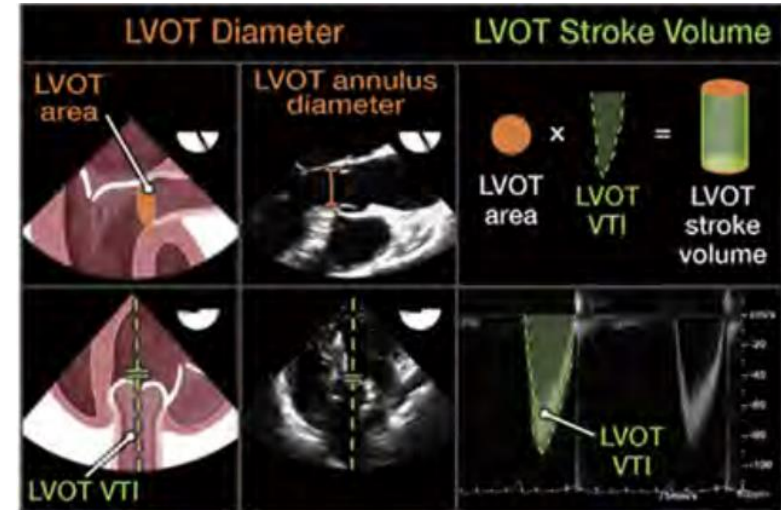
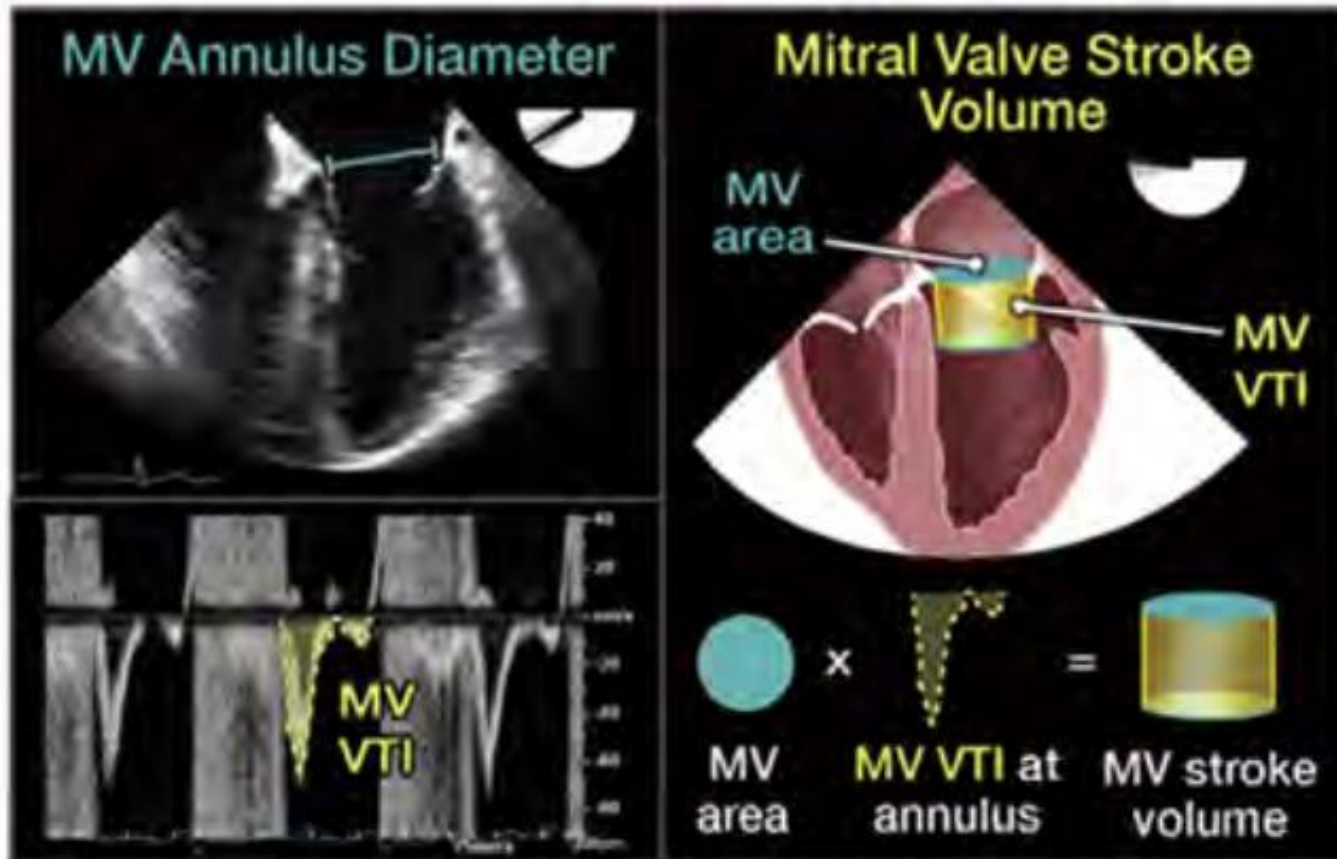
$$RVol = ERO \times VTI \quad RF = \frac{RVol}{\text{Stroke volume}^*}$$

* Calculated as forward stroke volume (either using transmitral inflow or the sum of LVOT flow and RVol)

Quantitative Evaluation - PISA



Establishing Severity of MR



$$RVol = \text{Mitral valve SV} - \text{LVOT SV}$$

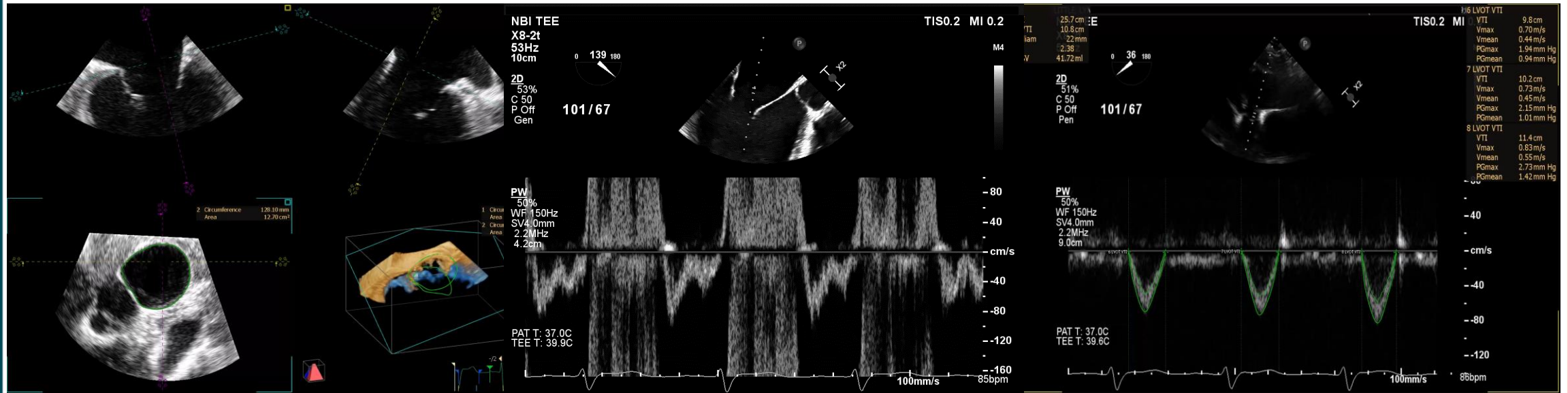
$$RVol = (MV \text{ area} \times MV \text{ VTI}) - (LVOT \text{ area} \times LVOT \text{ VTI})$$

$$RVol = \left\{ \pi \times \left(\frac{MV \text{ annulus diameter}}{2} \right)^2 \times MV \text{ VTI} \right\} - \left\{ \pi \times \left(\frac{LVOT \text{ diameter}}{2} \right)^2 \times LVOT \text{ VTI} \right\}$$

$$EROA (cm^2) = \frac{RVol}{MR \text{ VTI}} \quad RF (\%) = \frac{RVol}{MV \text{ SV}}$$

Hahn et al. Recommended Standards for the Performance of TEE for SHI. JASE 2022

Quantitative Evaluation

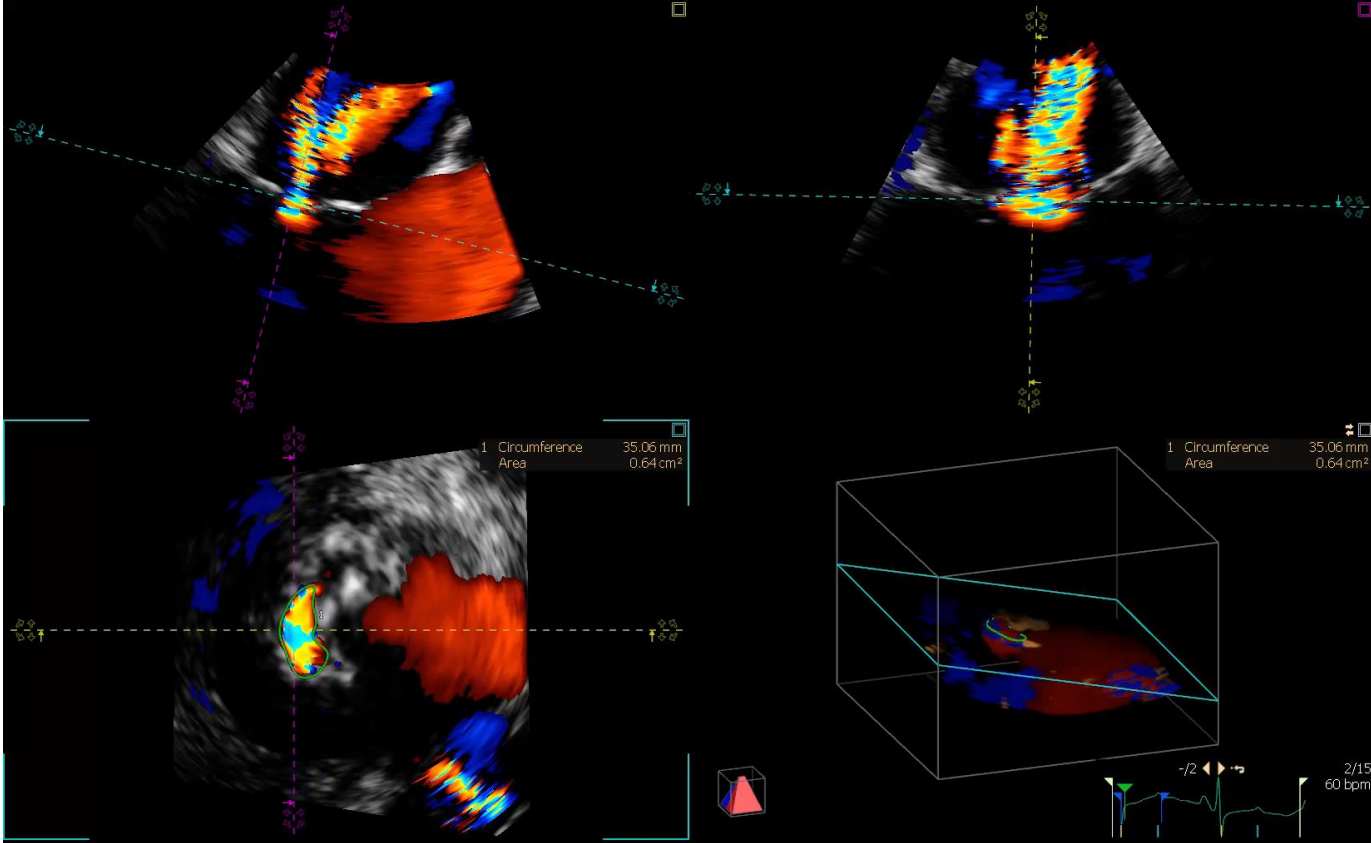


3D mitral annulus area is 12.7cm and MV annulus VTI is 14cm

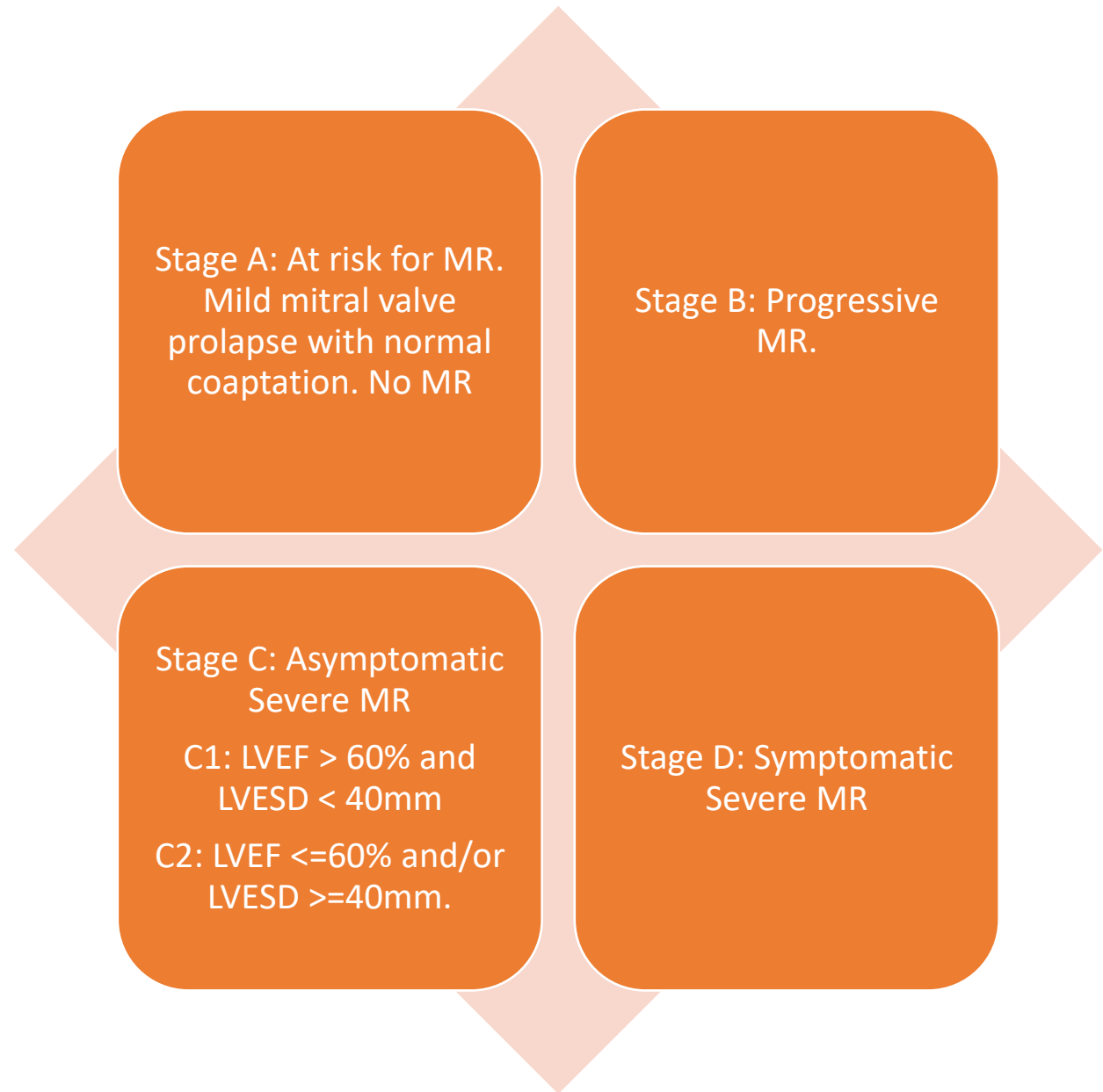
LVOT diameter is 2.2cm and LVOT VTI is 11cm

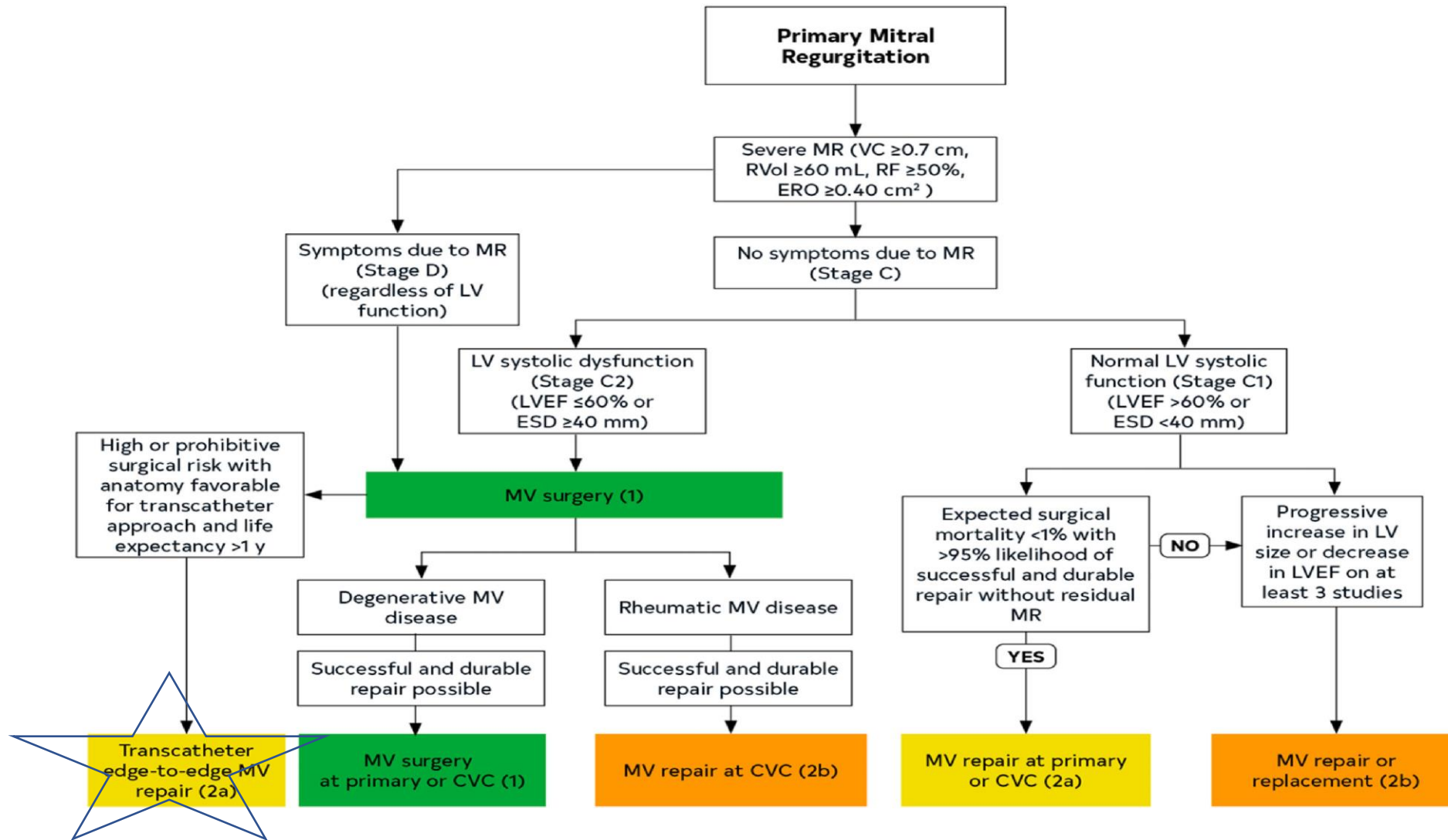
LVOT stroke volume = 42cc. EROA by CE is 0.8cm² and regurgitant volume is 136cc.

3D Vena Contracta



Stages of Primary Mitral Regurgitation





Stages of Secondary Mitral Regurgitation

Stage A: At risk for MR. Normal valve leaflets, chords and annulus in patient with CAD or CM

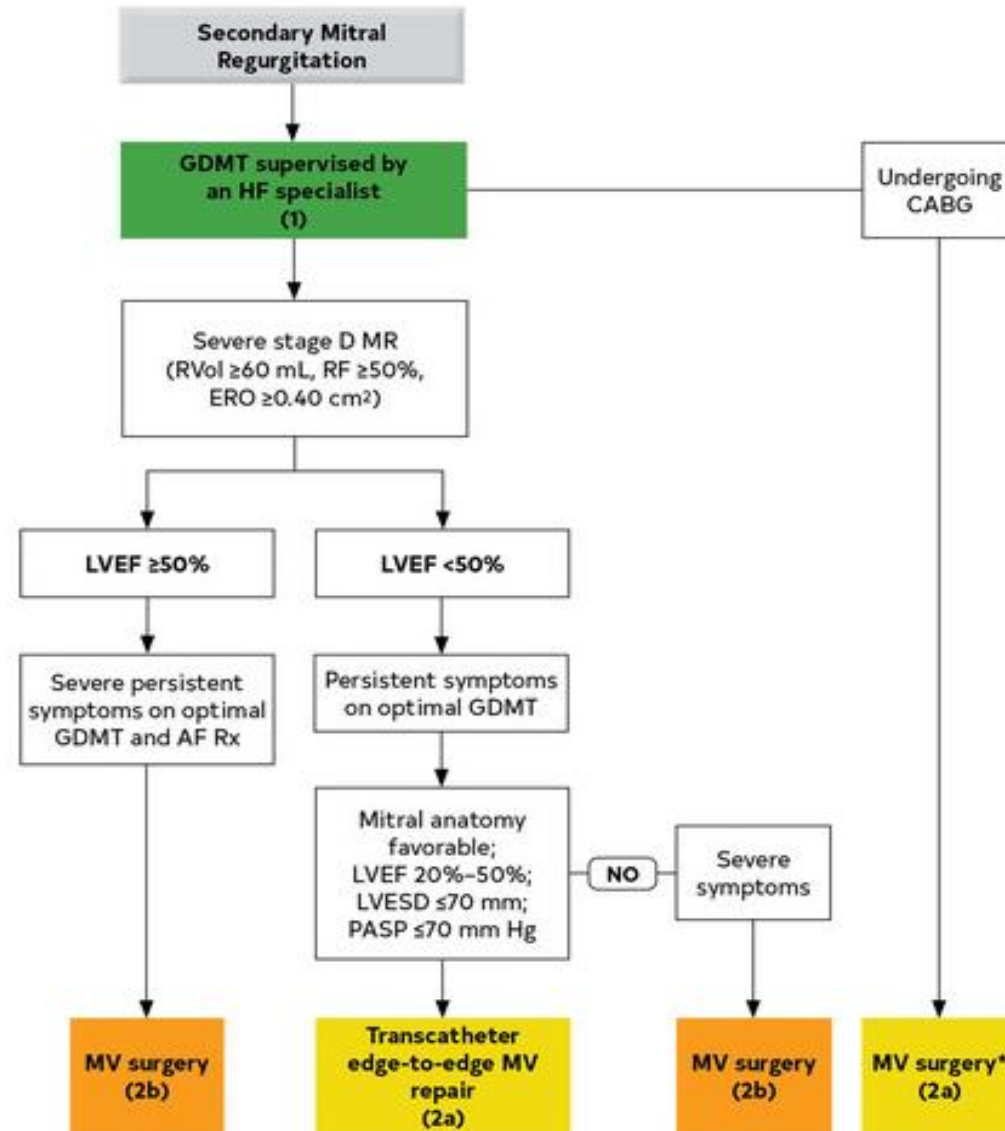
Stage B: Progressive MR. Regional wall motion abnormalities with mild tethering

Stage C: Asymptomatic Severe MR. Regional wall motion abnormalities with severe tethering. Annular dilation and loss of coaptation

Stage D: Symptomatic Severe MR. Regional wall motion abnormalities with severe tethering. Annular dilation and loss of coaptation + HF symptoms, dyspnea and decrease in exercise tolerance



Management of Secondary Mitral regurgitation



Summary

1. Aortic Stenosis is not only pathology of the valve, but disease of myocardium
2. Classification and grading of aortic stenosis is tricky and requires systematic approach
3. Important to describe mitral valve anatomy and quantify regurgitation severity
4. Team approach to patient care