


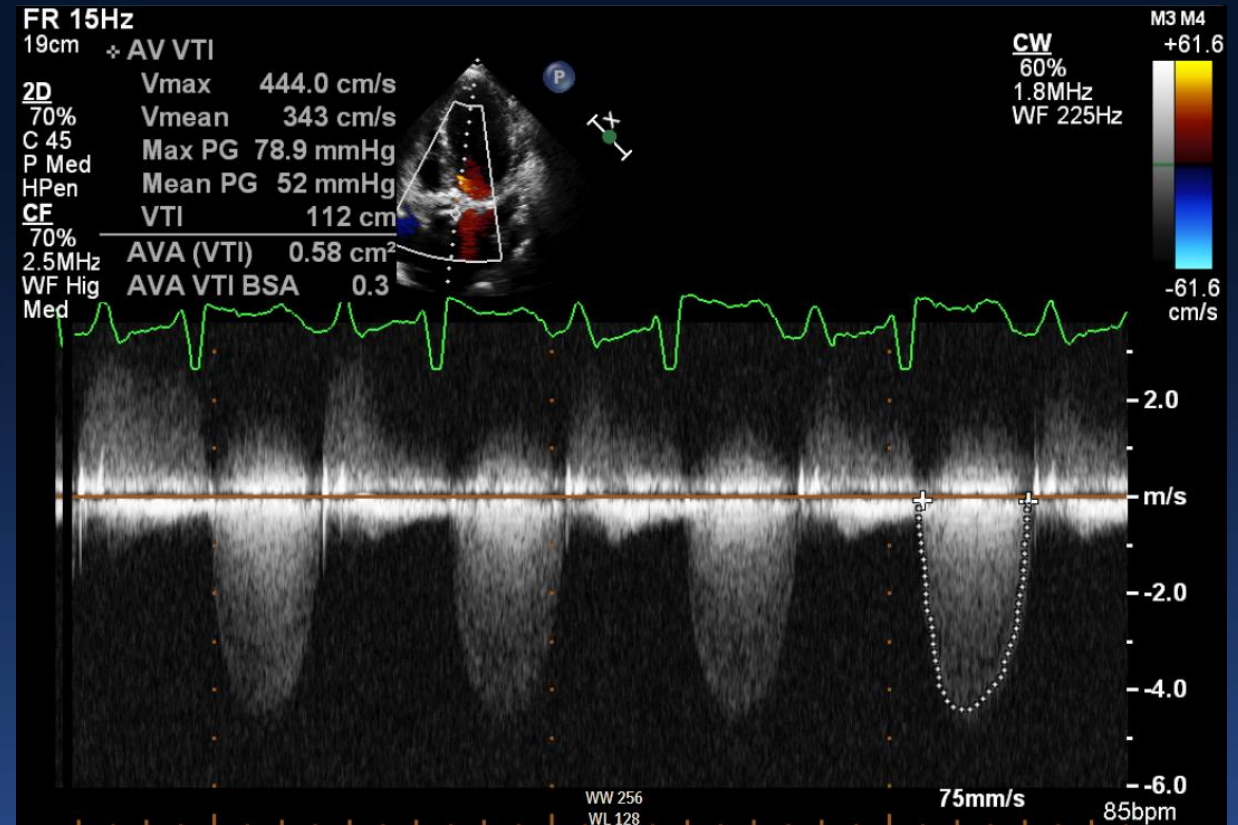
Implante valvular aortico percutaneo sobre valvula protésica (VIV). Caso Educativo Interactivo

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Director Cardiac Cath Lab
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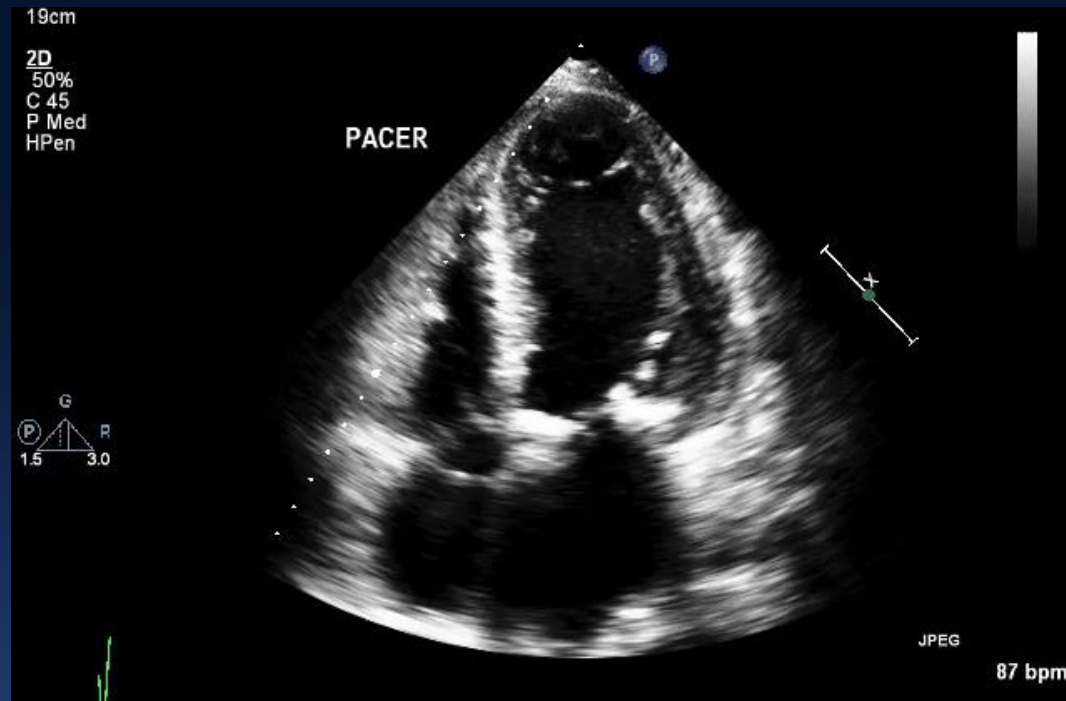


Case #1

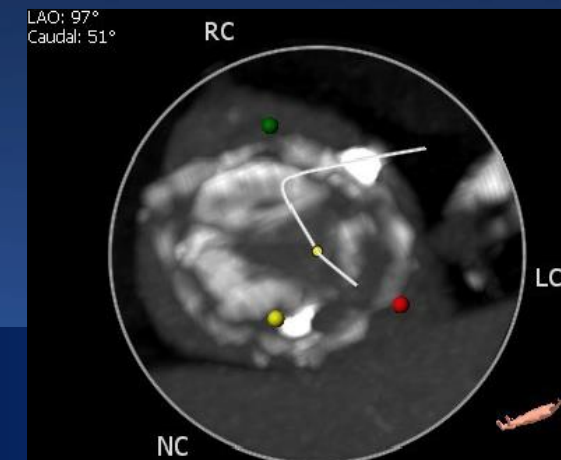
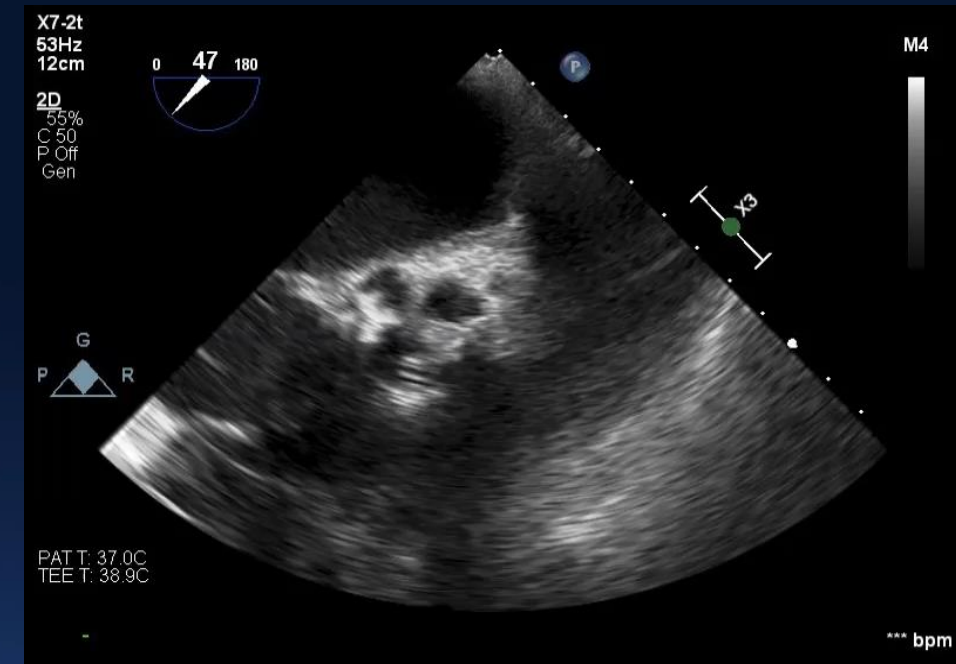
- 63 y/o male s/p SAVR (25mm Mosaic) Porcine in 2015
- Cath 7/02/19: 90% mid RCA and severe bioprosthetic AS
- ESRD s/p 2 failed kidney Tx (1987 and 2001)
 - R arm AVF for dialysis
- Paroxysmal A-fib (on apixaban), SSS s/p PPM (1/2019)
- Chronic steroid therapy; TIA, HTN, HLD, asthma, gout
- STS PROM 10.6%



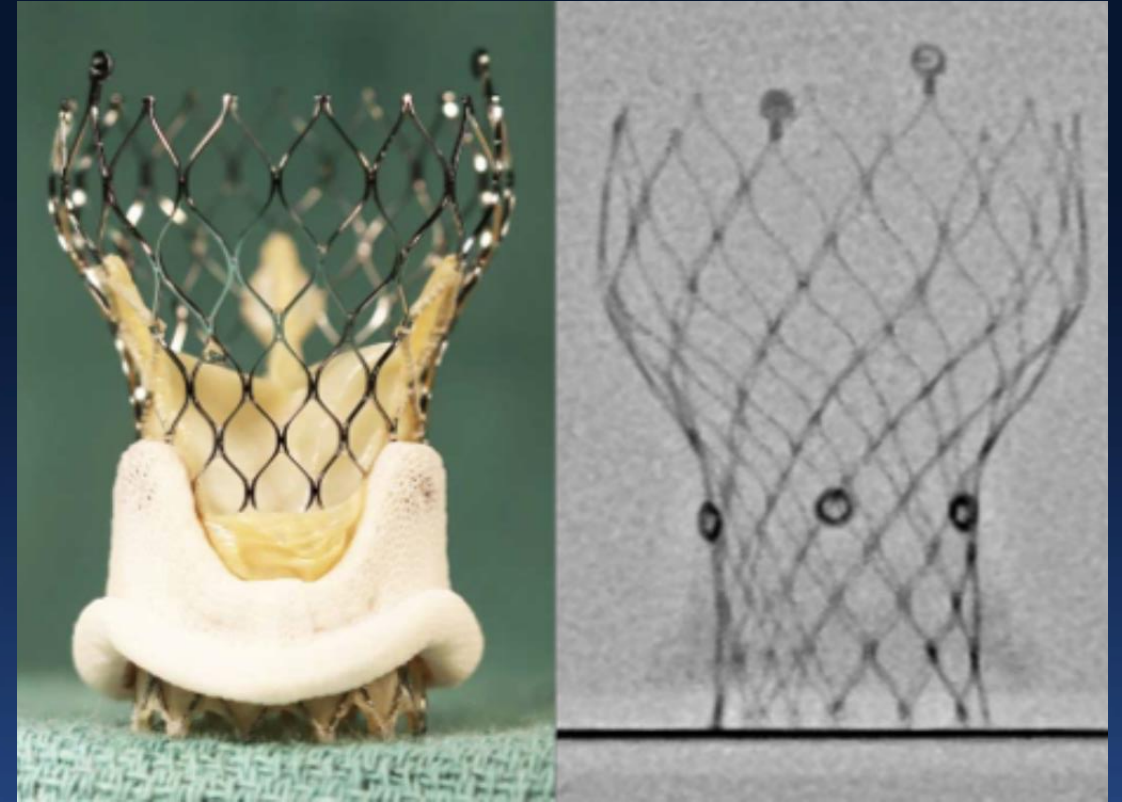
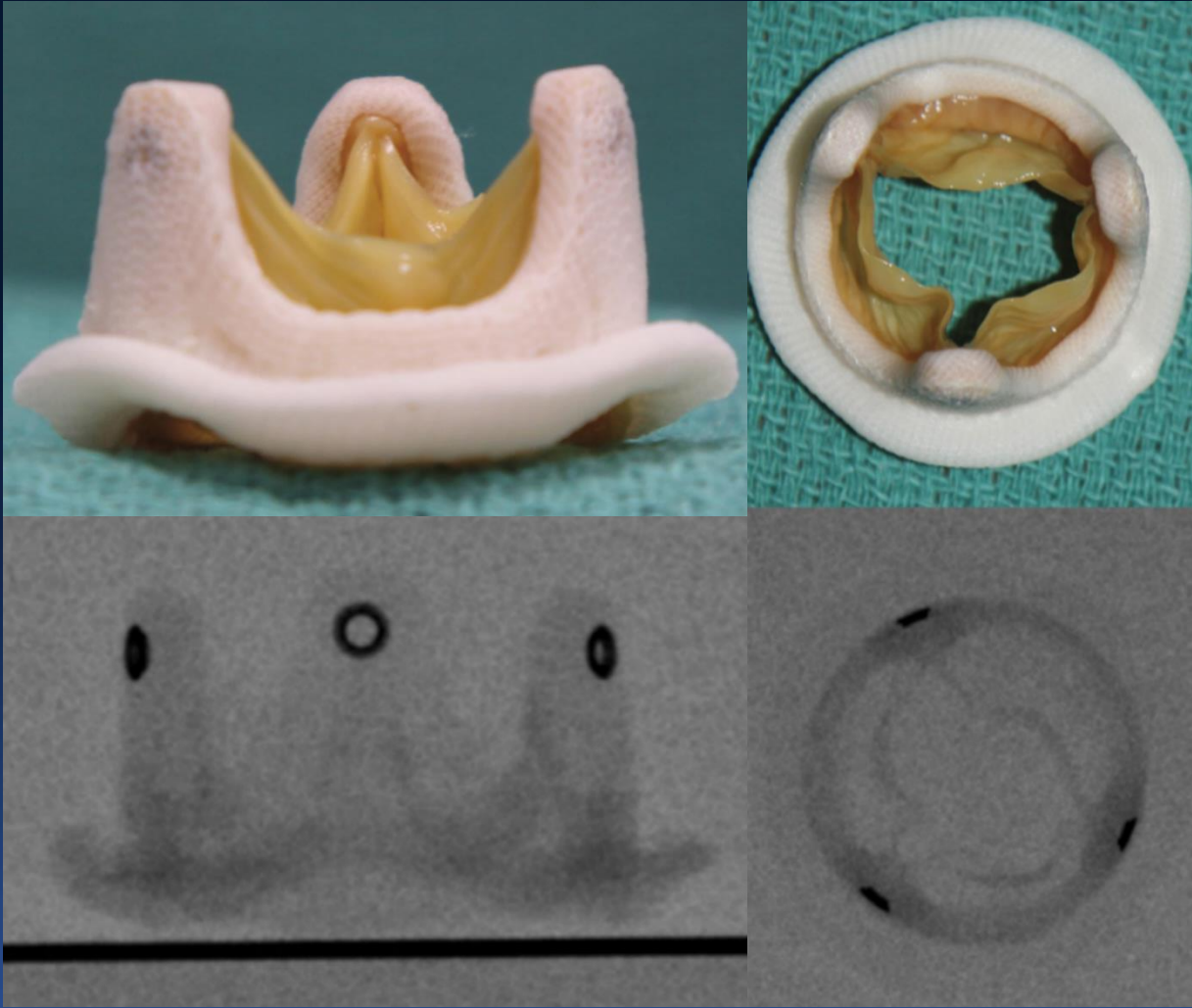
ECHO (TTE + TEE) and CT of Ao valve



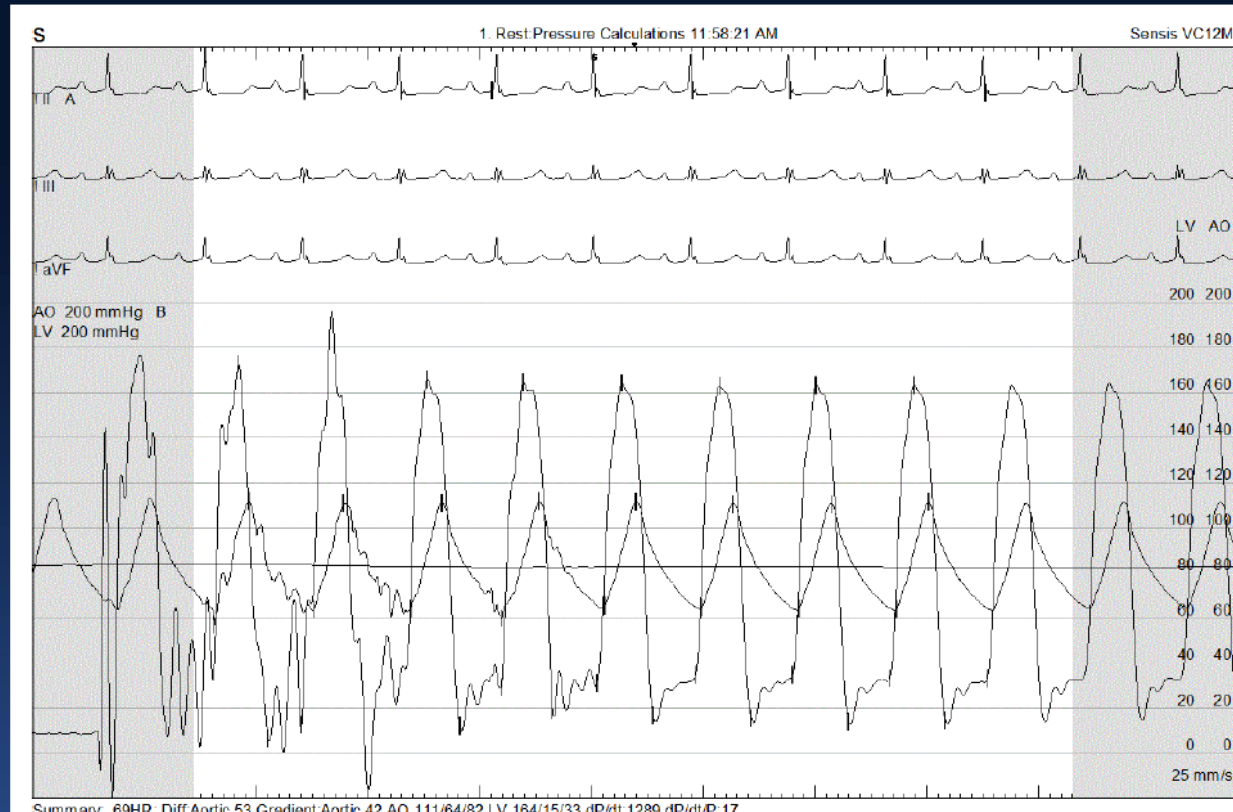
EF 35-40%, mean mitral gradient 5 mmHg



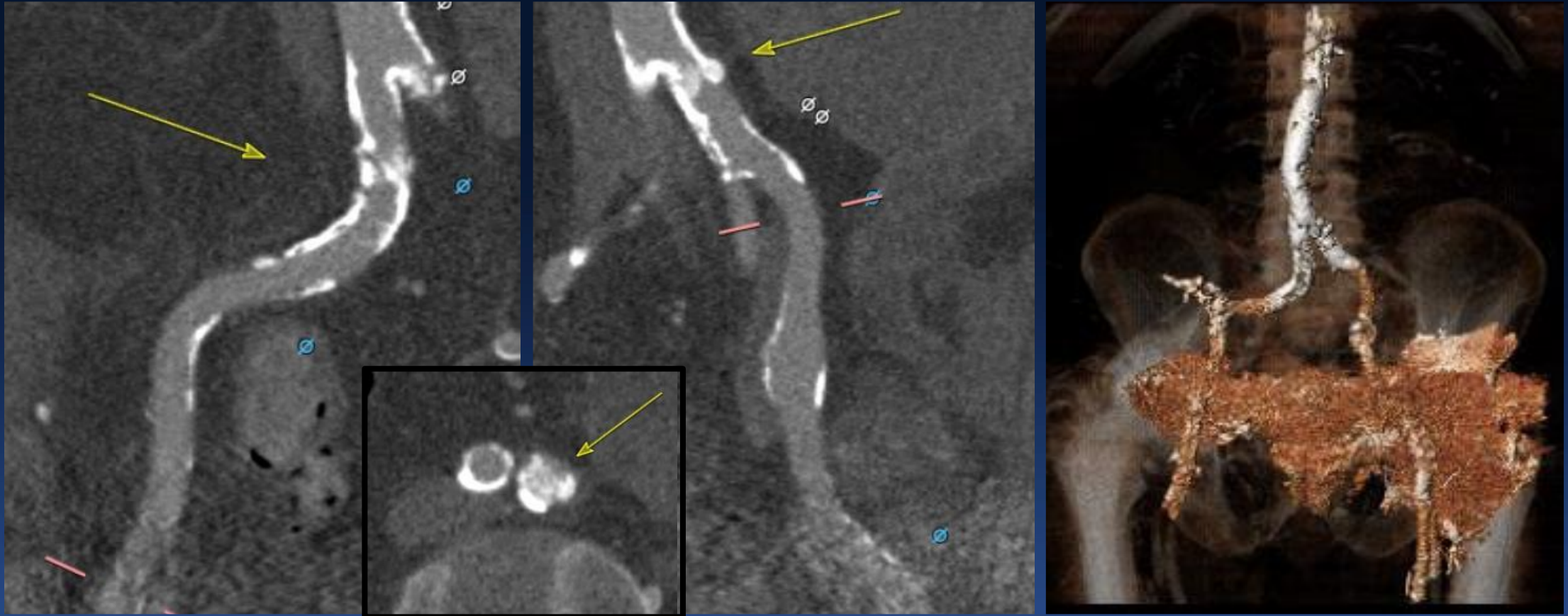
Mosaic 25 mm Bioprosthesis



Cath: Severe Bioprosthetic AS + 1-v CAD

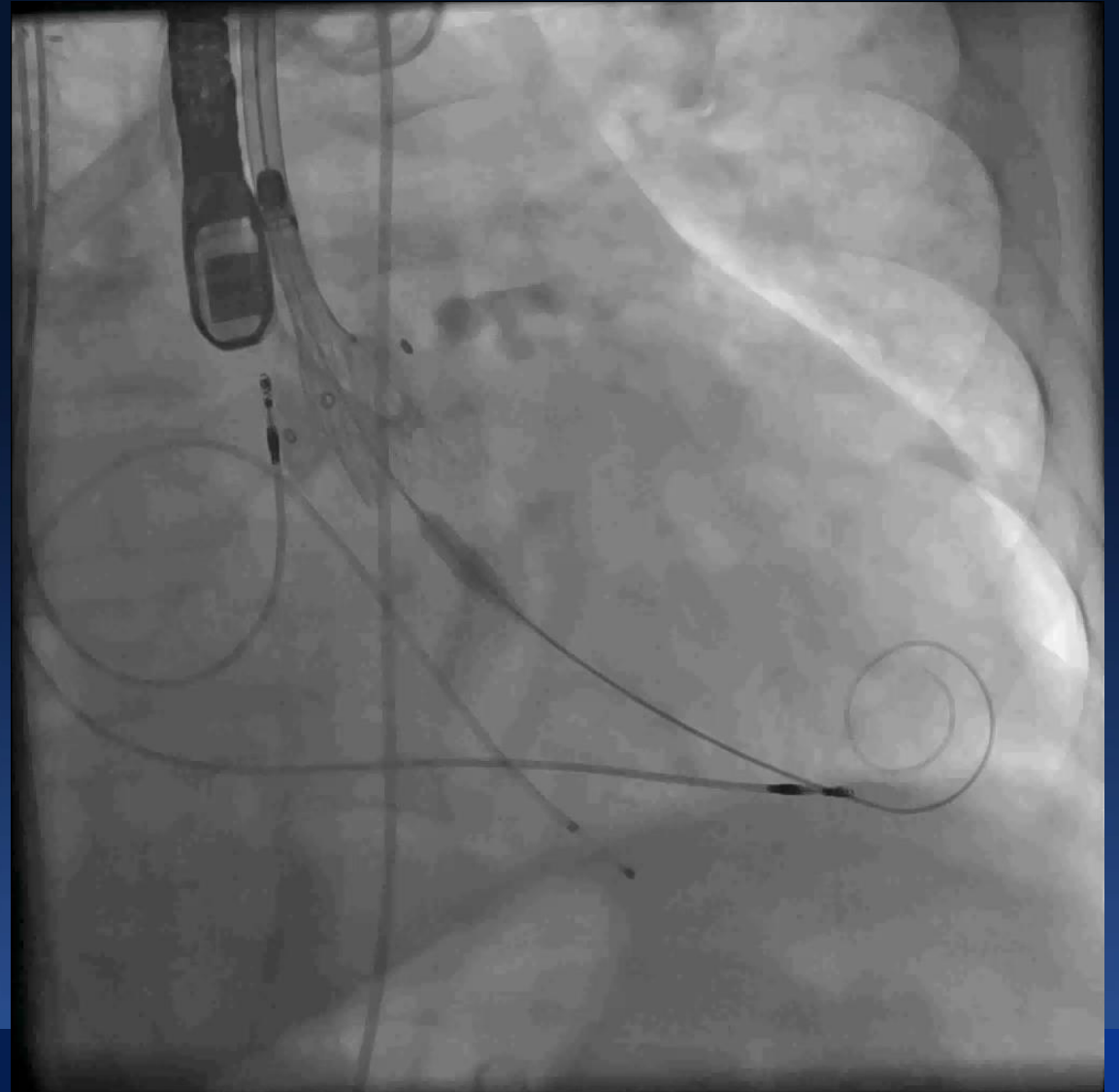
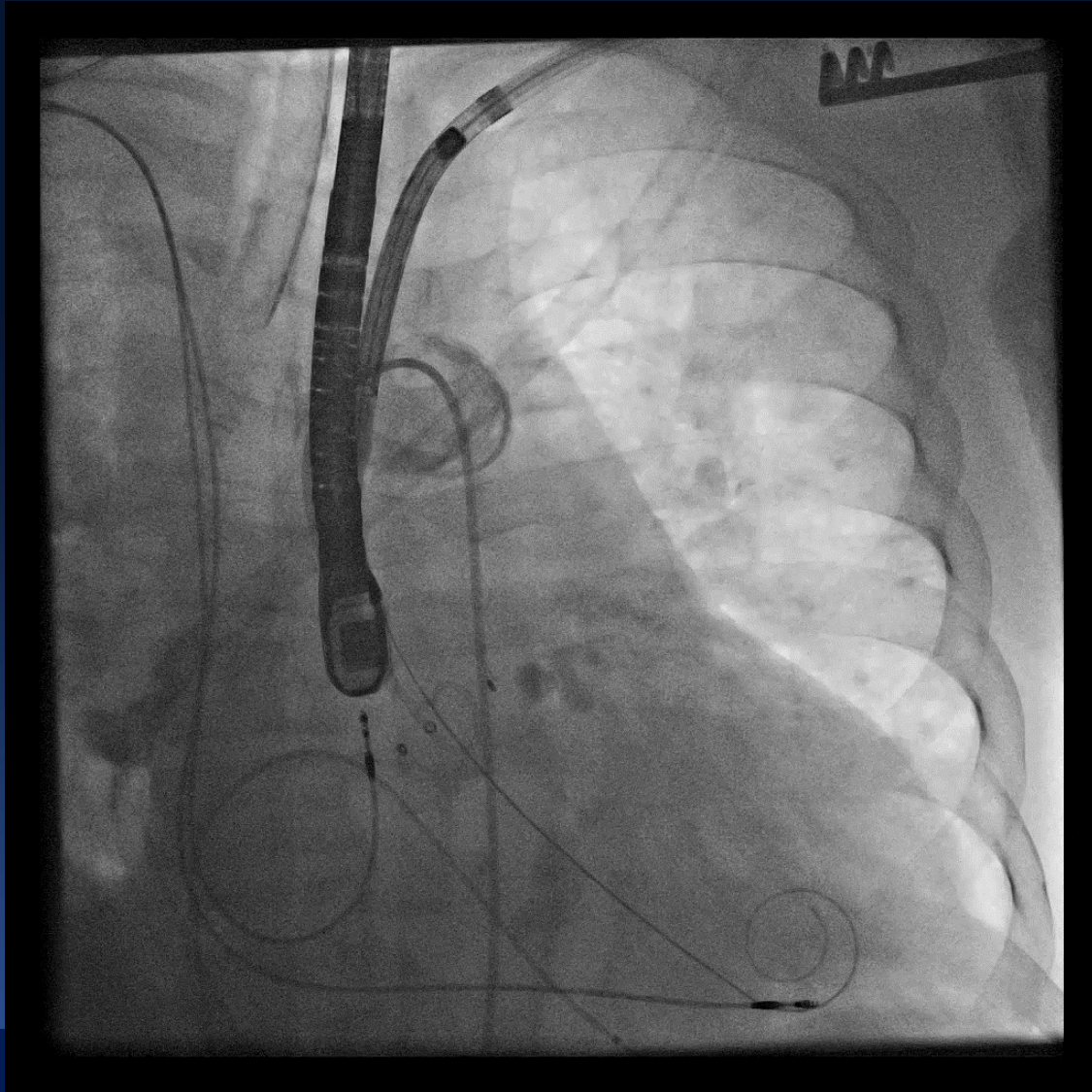


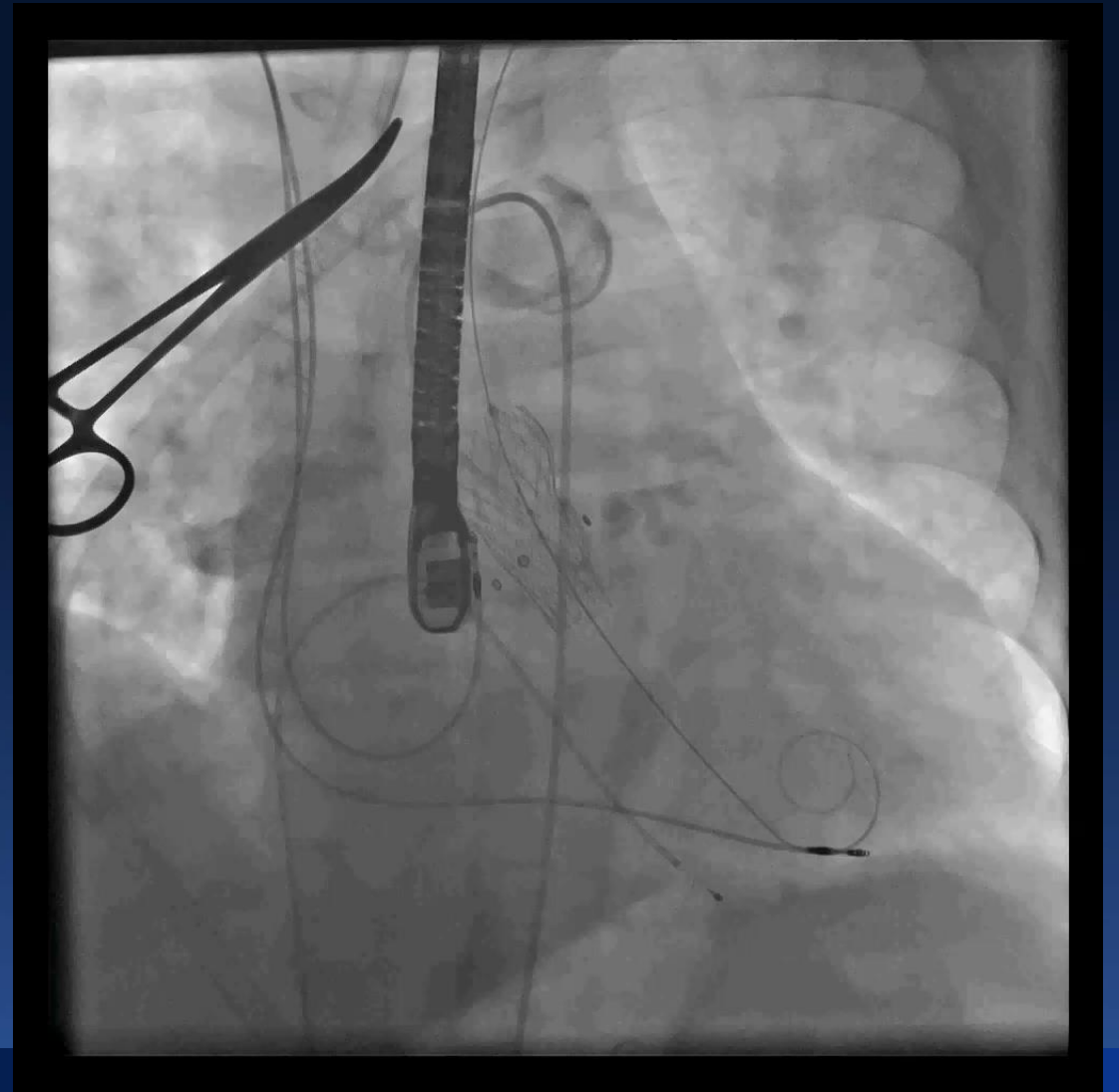
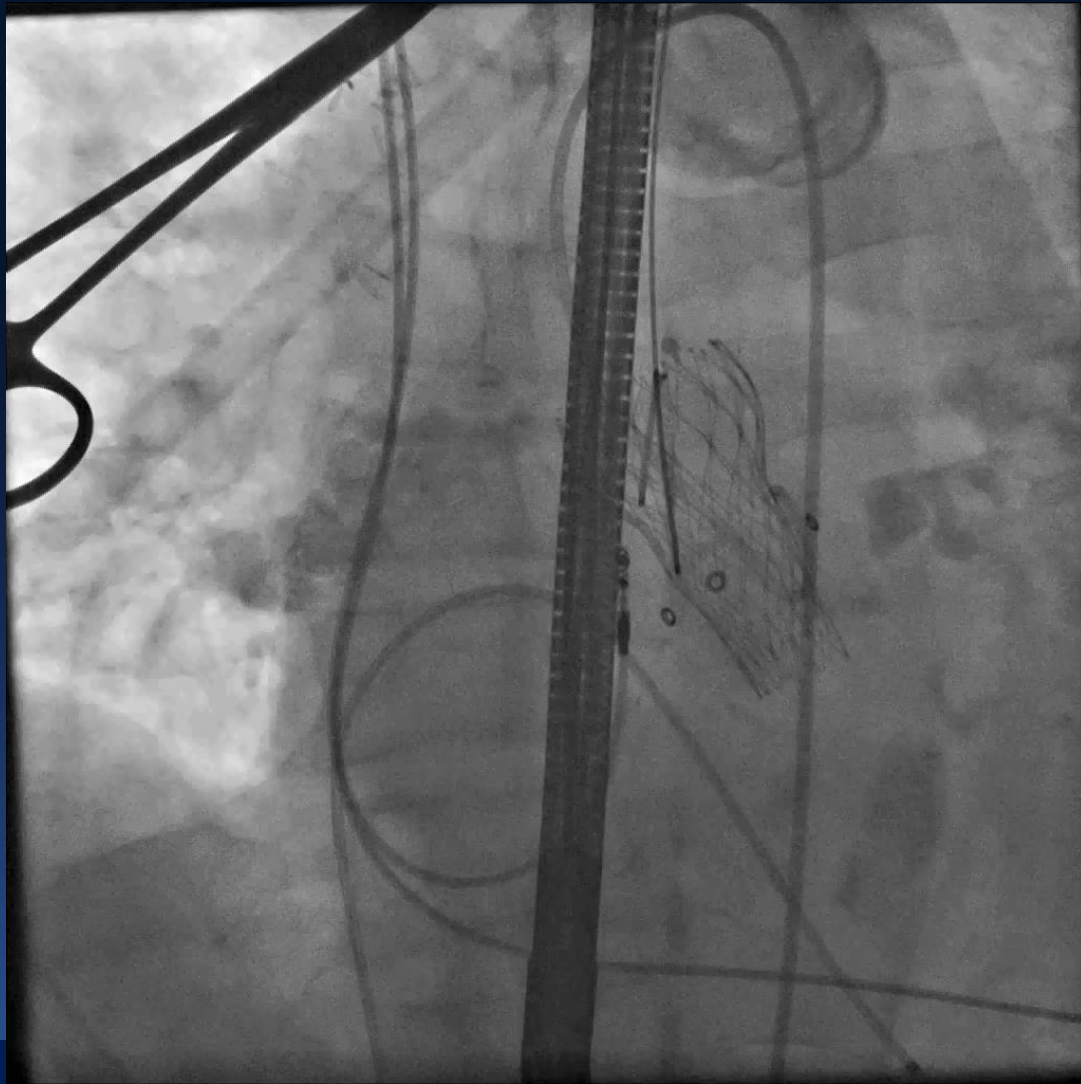
Femoral Access?



Cath Lab Setup





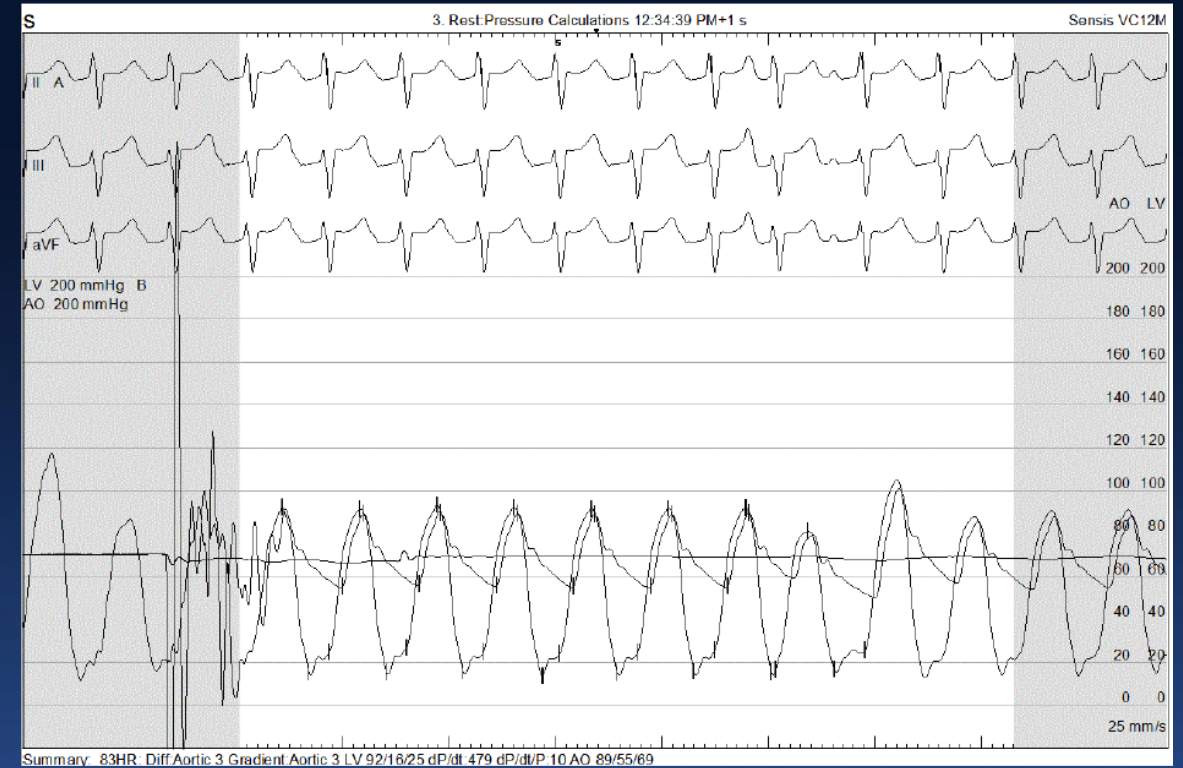


Bioprosthetic Valve Fracture During Valve-in-valve TAVR: Bench to Bedside

John T Saxon,^{1,2} Keith B Allen,^{1,2} David J Cohen^{1,2} and Adnan K Chhatriwalla^{1,2}



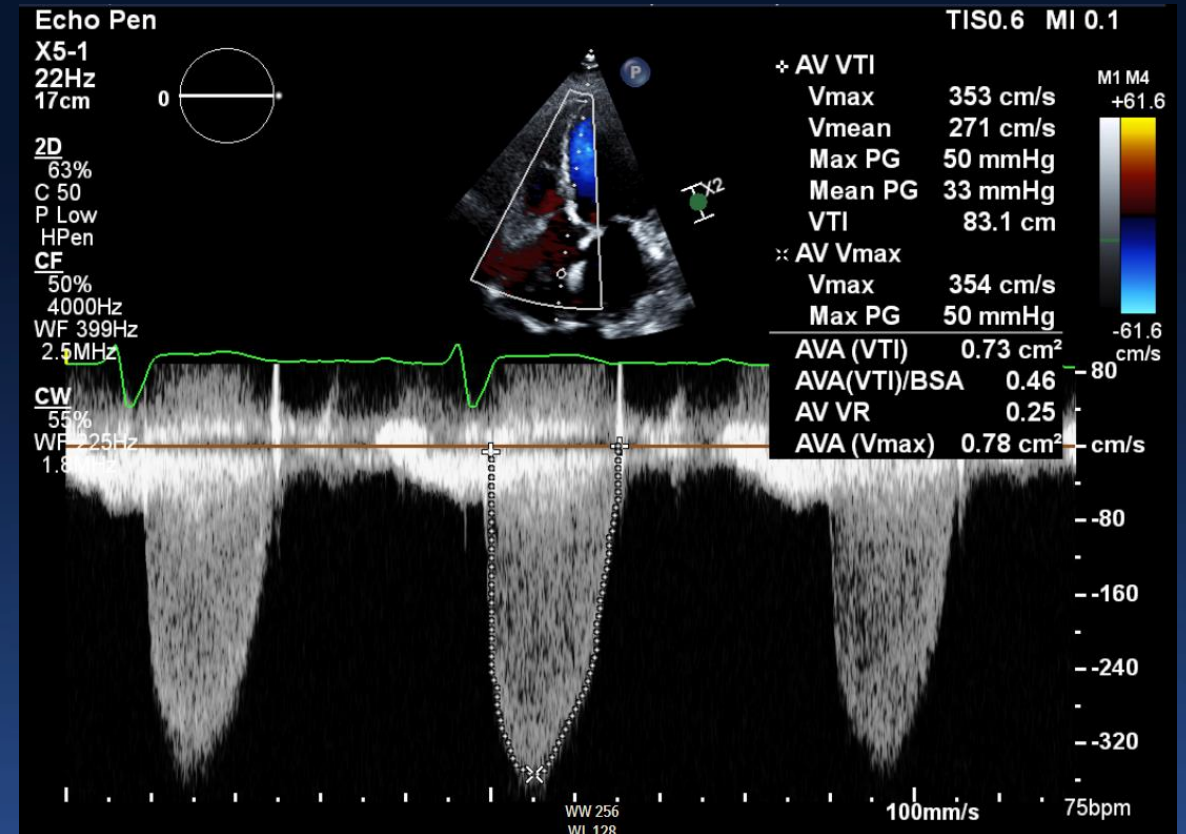
(1) A high pressure stopcock connects the valvuloplasty balloon to a syringe of dilute contrast and an indeflator. (2) The syringe is used to inflate the balloon manually. (3) The stopcock is turned so that the syringe is off and the indeflator is on. (4) The indeflator is dialed to the desired pressure, until the bioprosthetic valve fractures or the balloon ruptures.

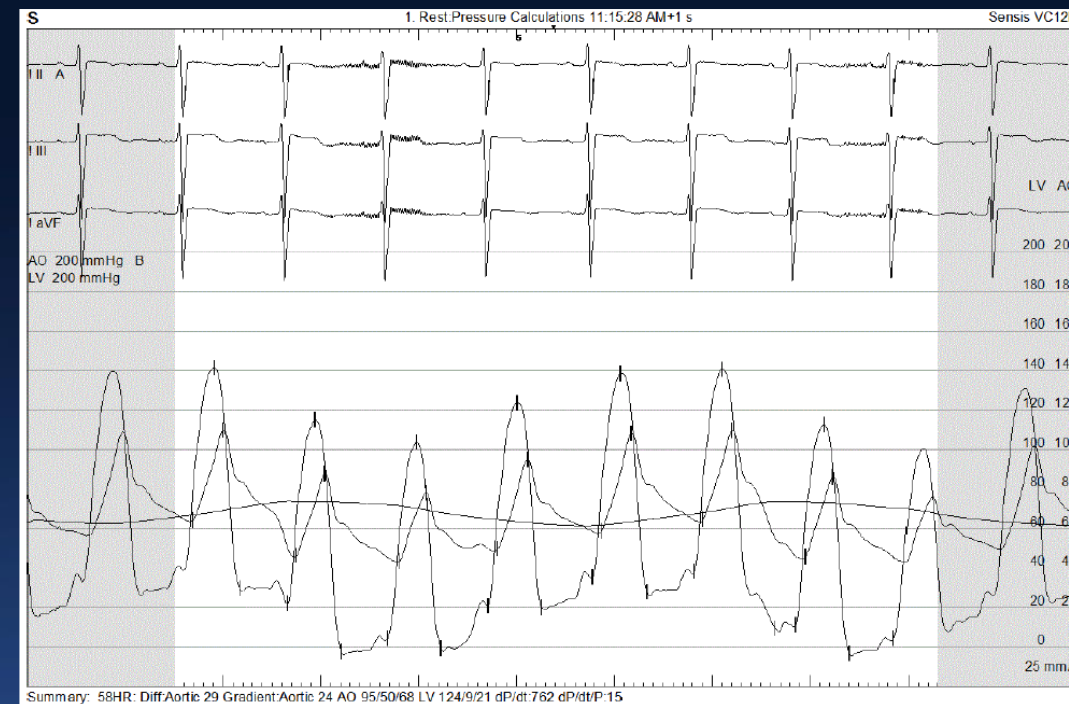
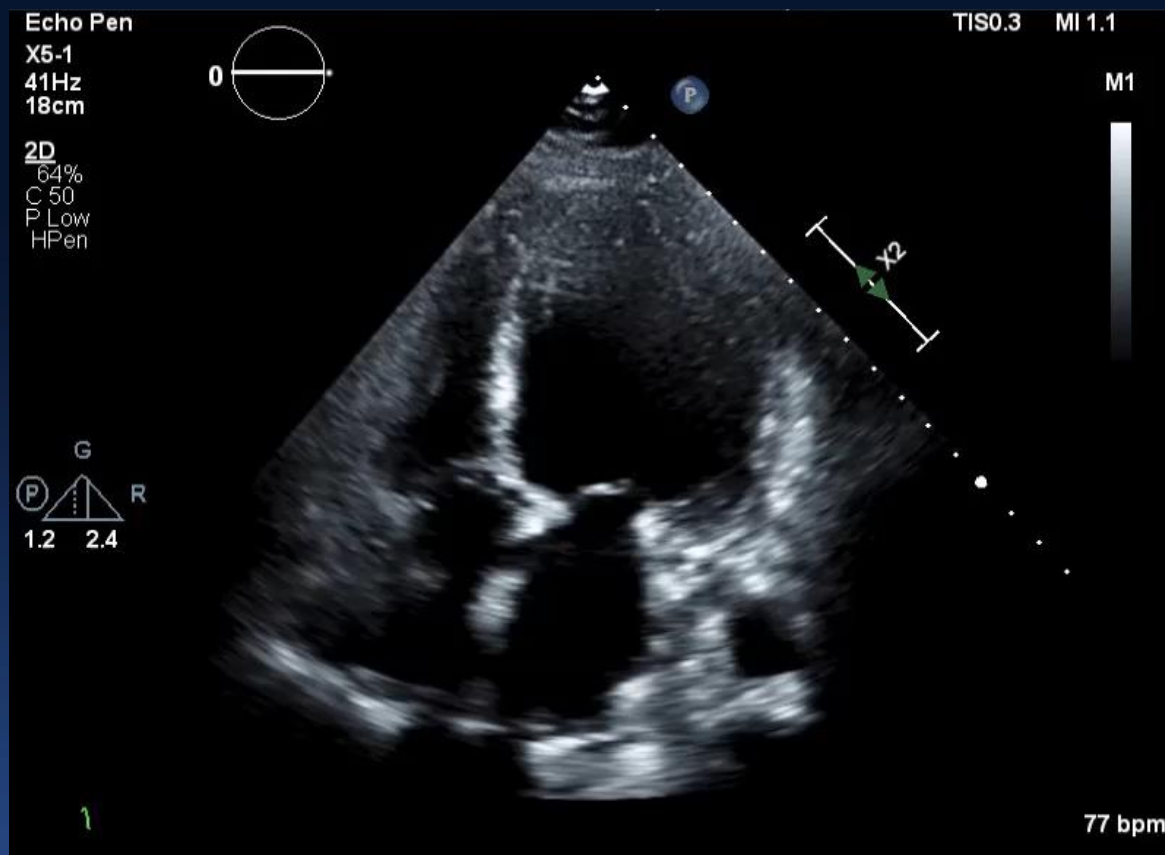




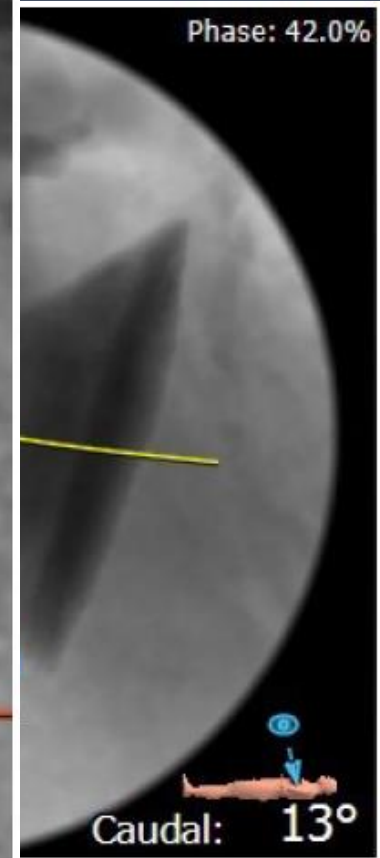
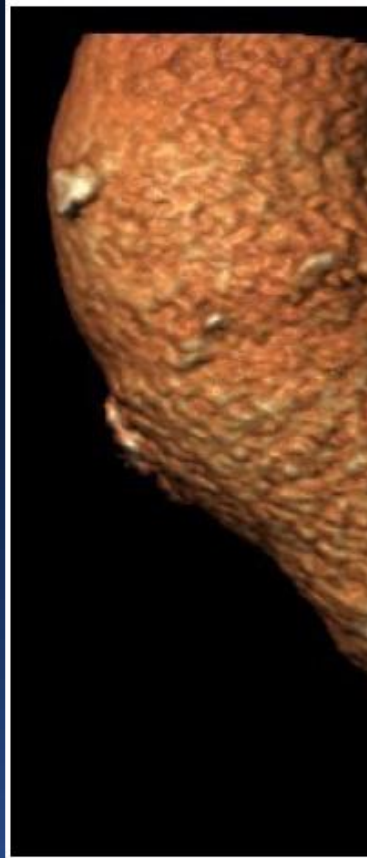
Case #2

- 78 yo female with severe bioprosthetic AS, SAVR in 2005, CHF NYHA class III
 - Medtronic Mosaic Valve 21 mm
 - Annulus perimeter measures 53.3mm, sizing 23 mm Evolut R (based on IFU)
- HTN, hyperlipidemia (intolerant to statins), NIDDM
 - 62.1 kg/149 cm
- CAD s/p MI and LAD PCI 2017
- STS PROM 6.23%

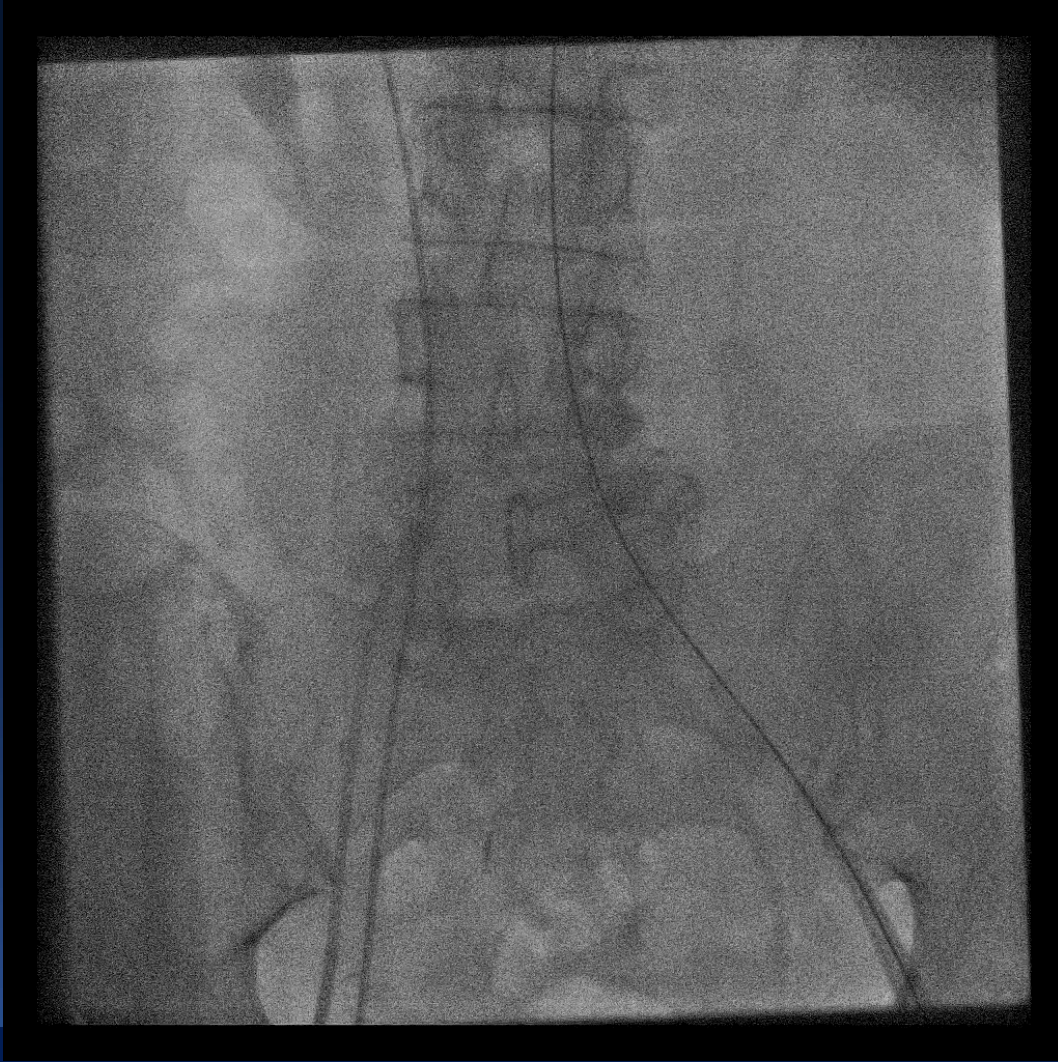


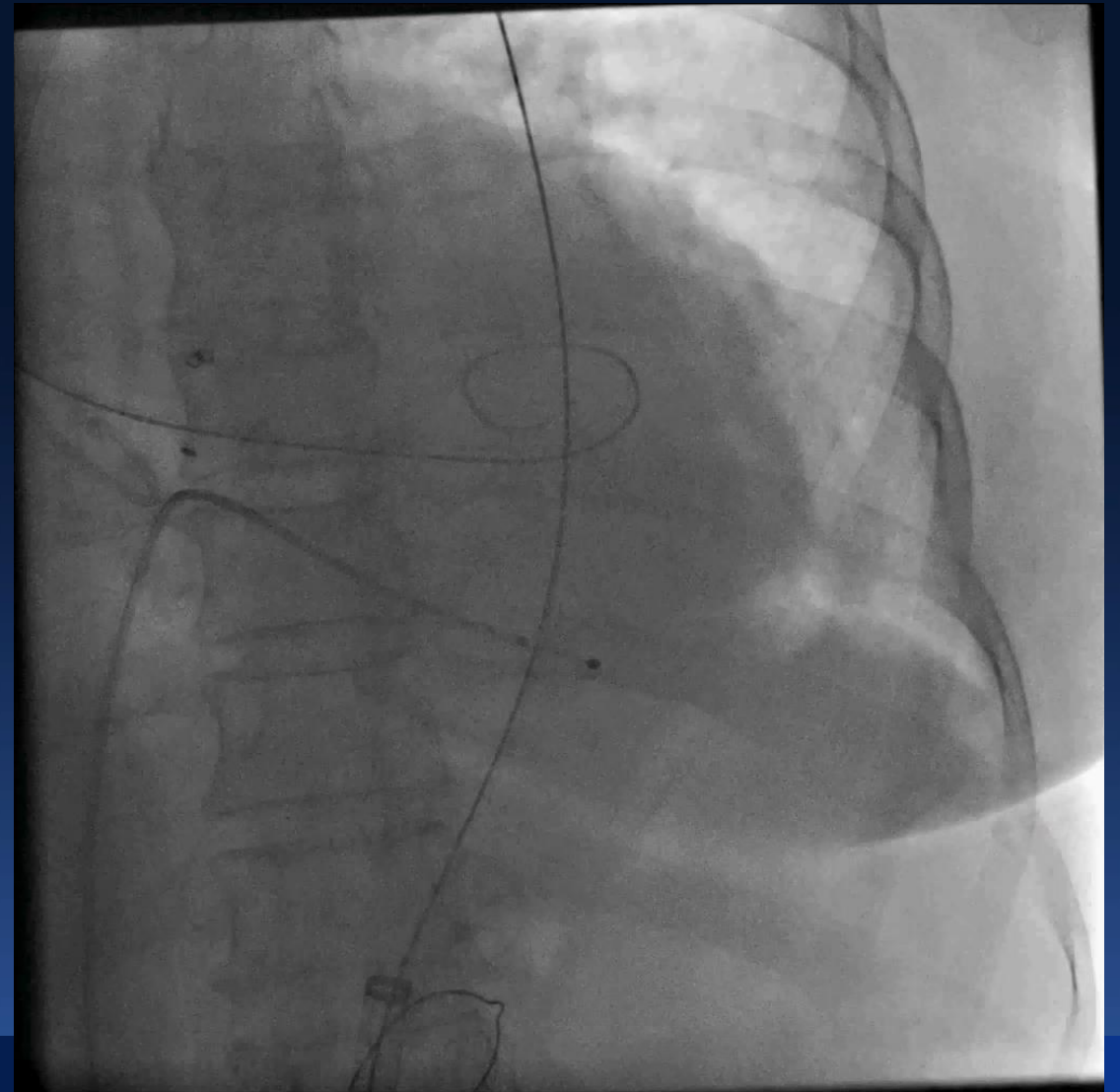
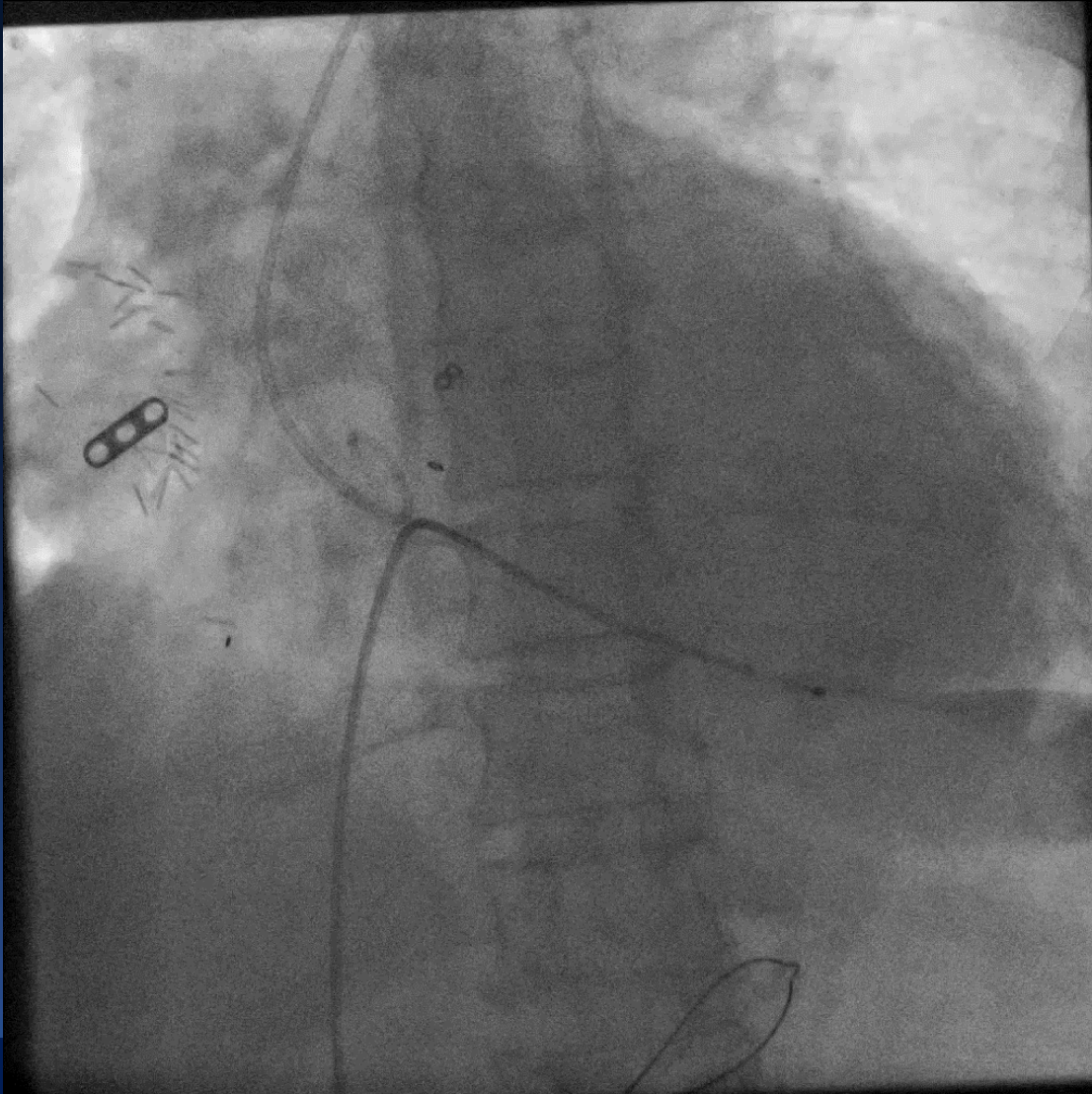


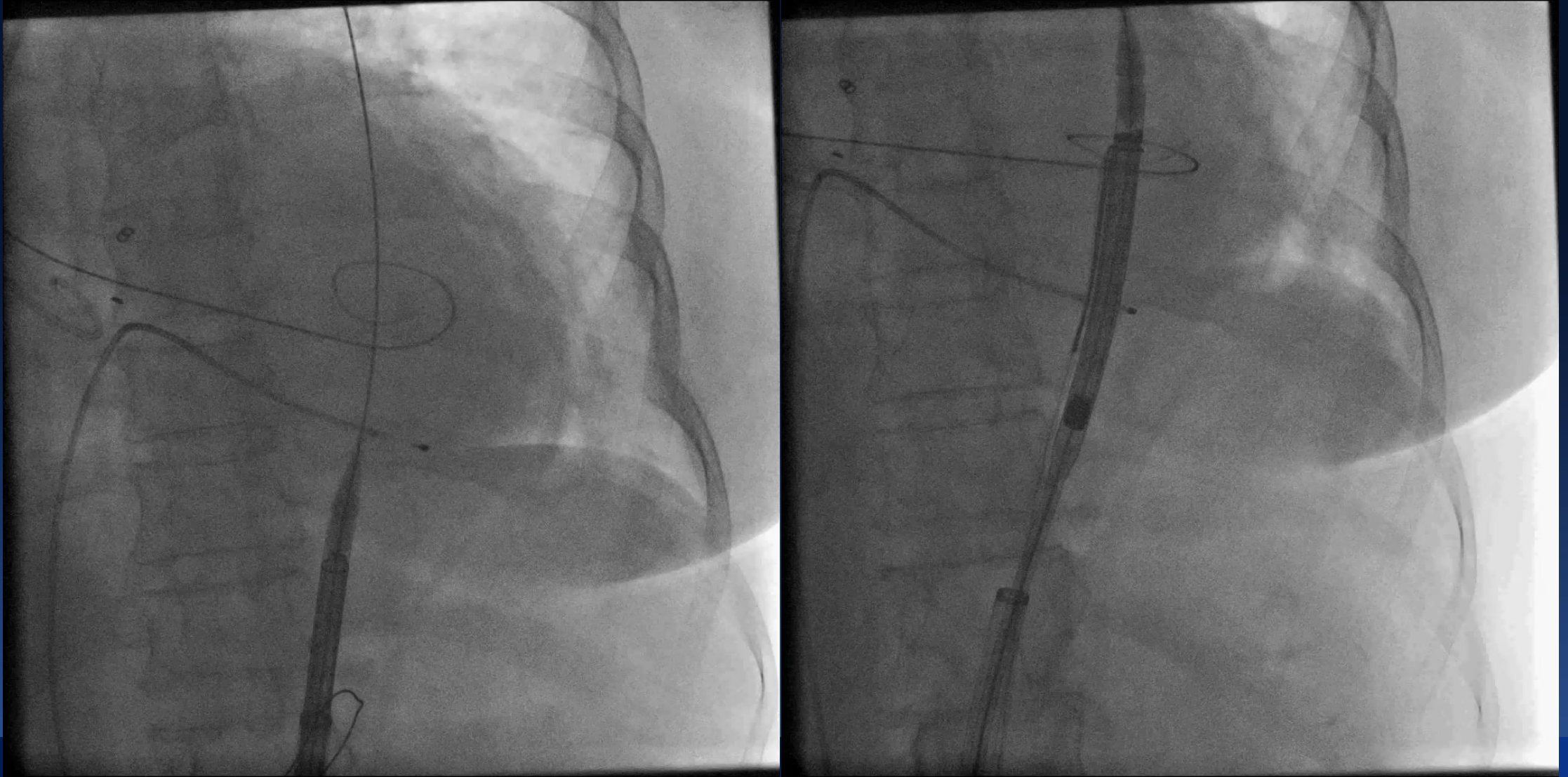
The Challenge

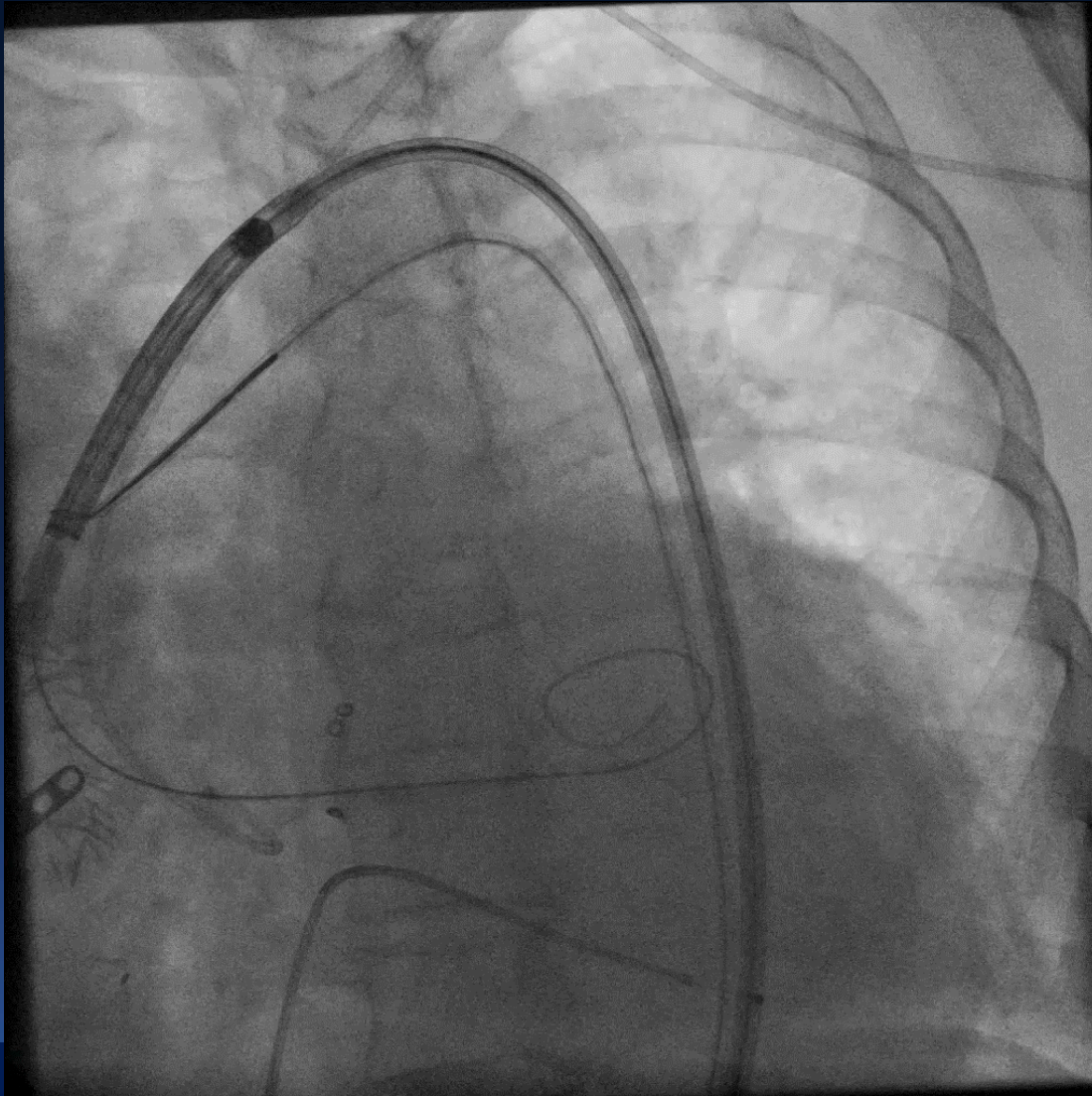


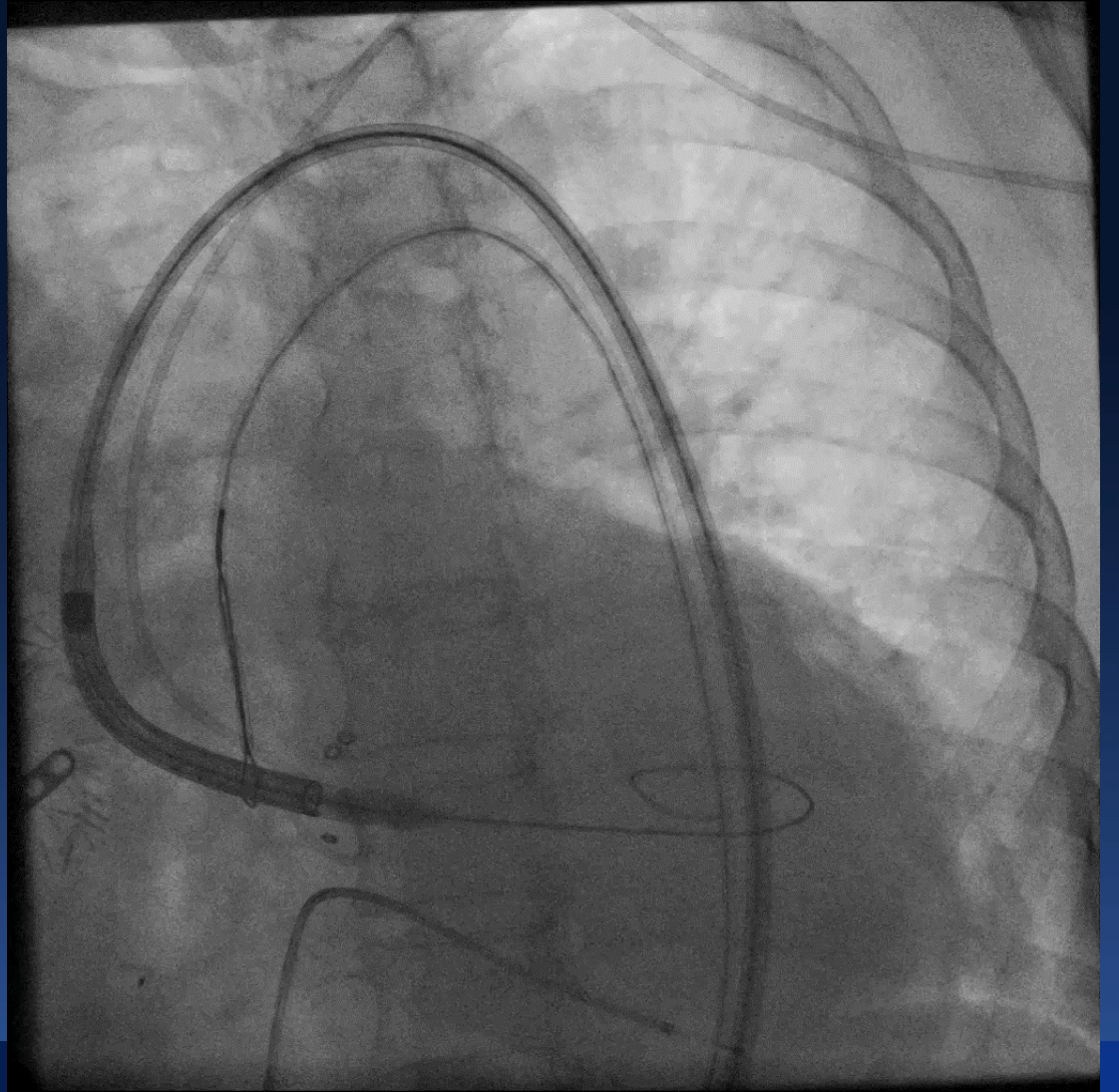
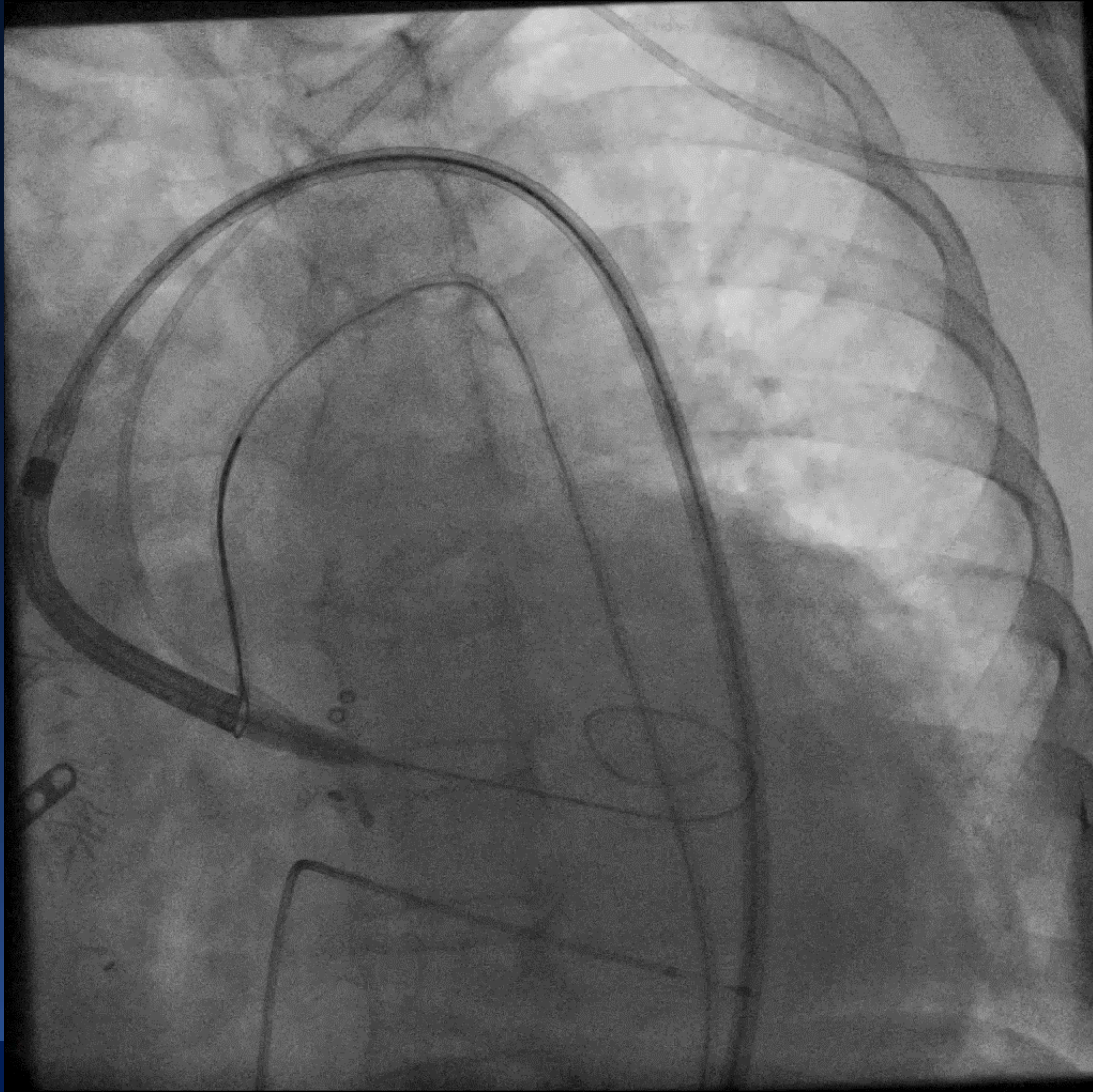
Strategy: Bifemoral Access

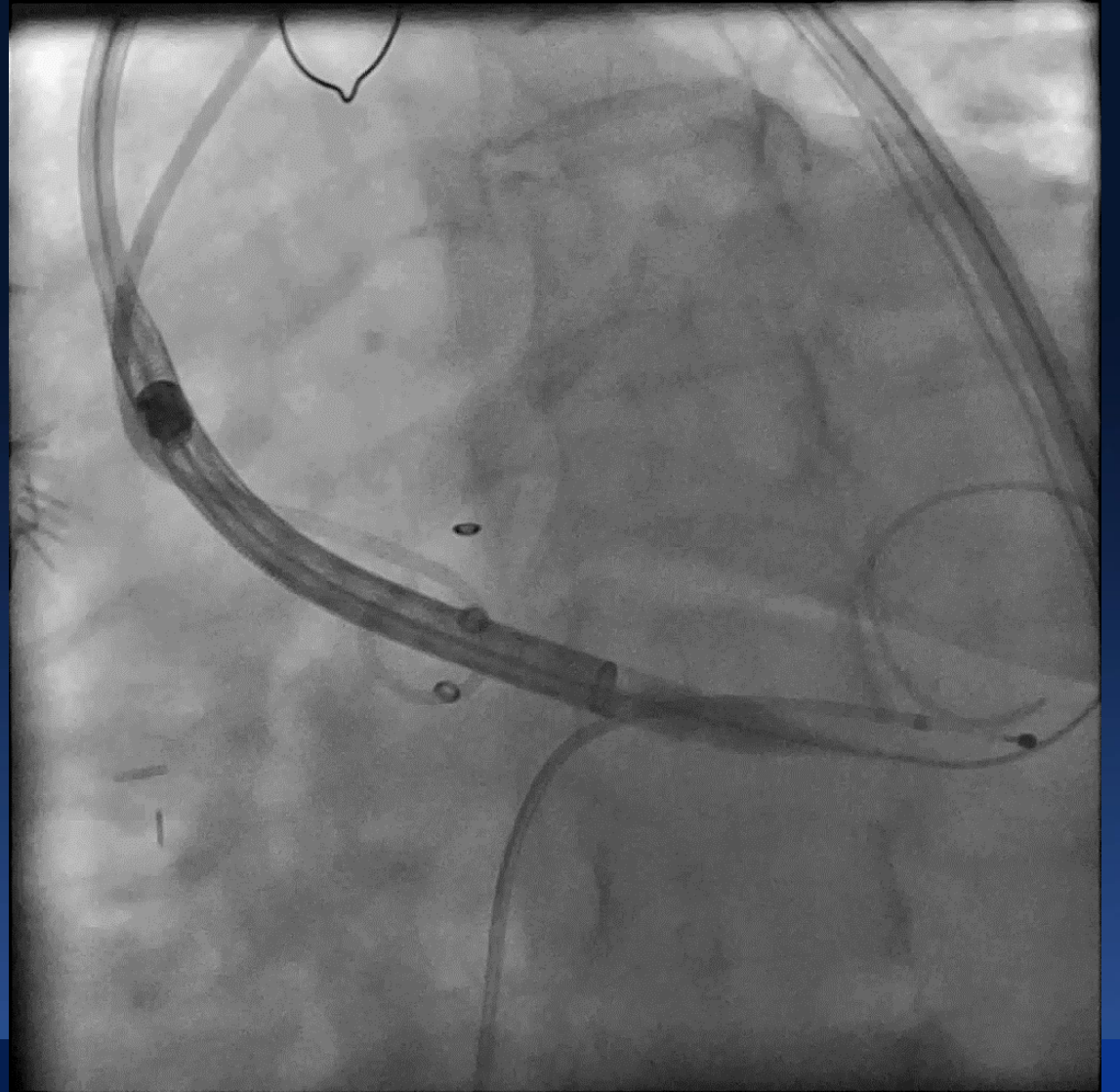
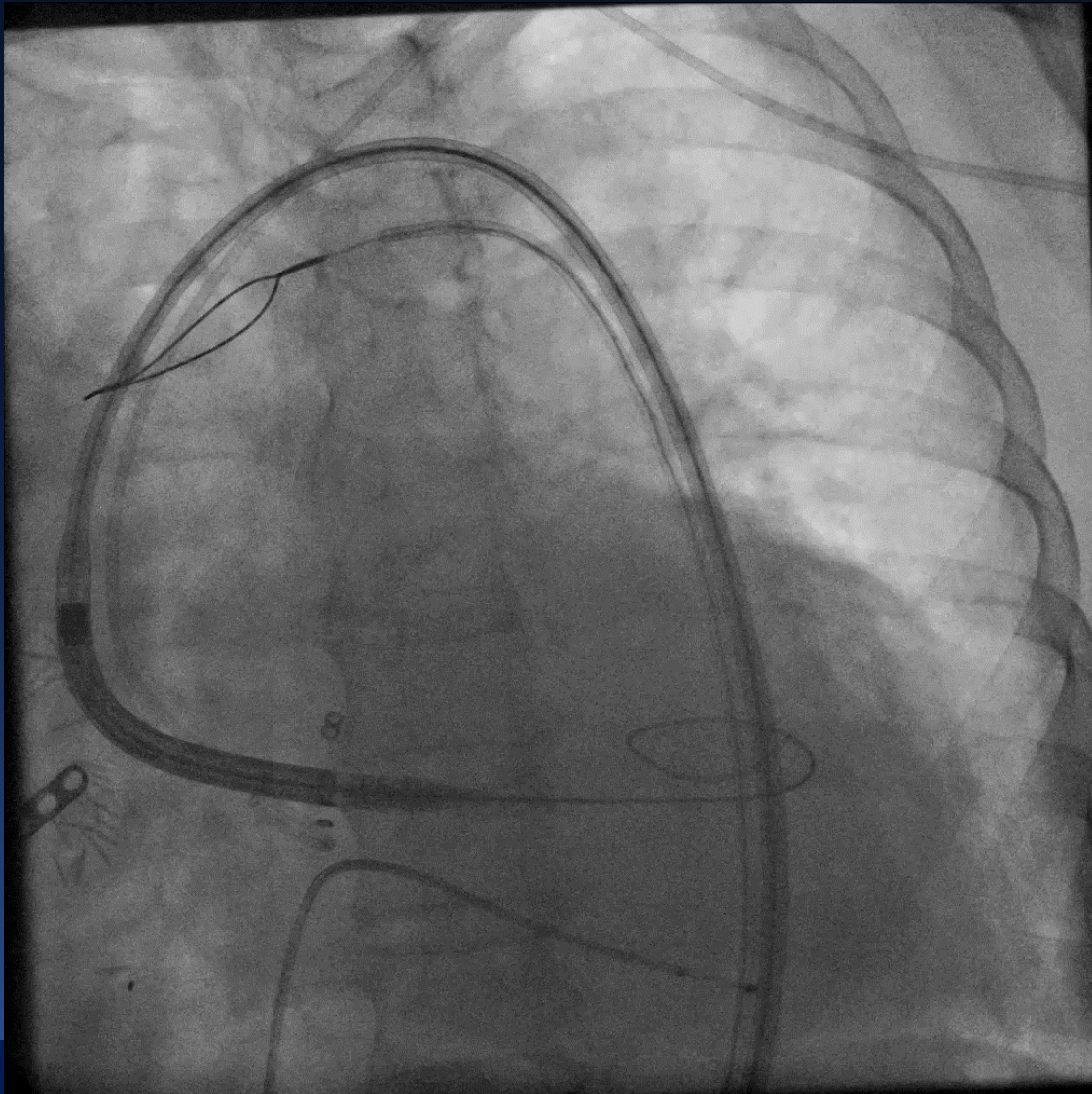


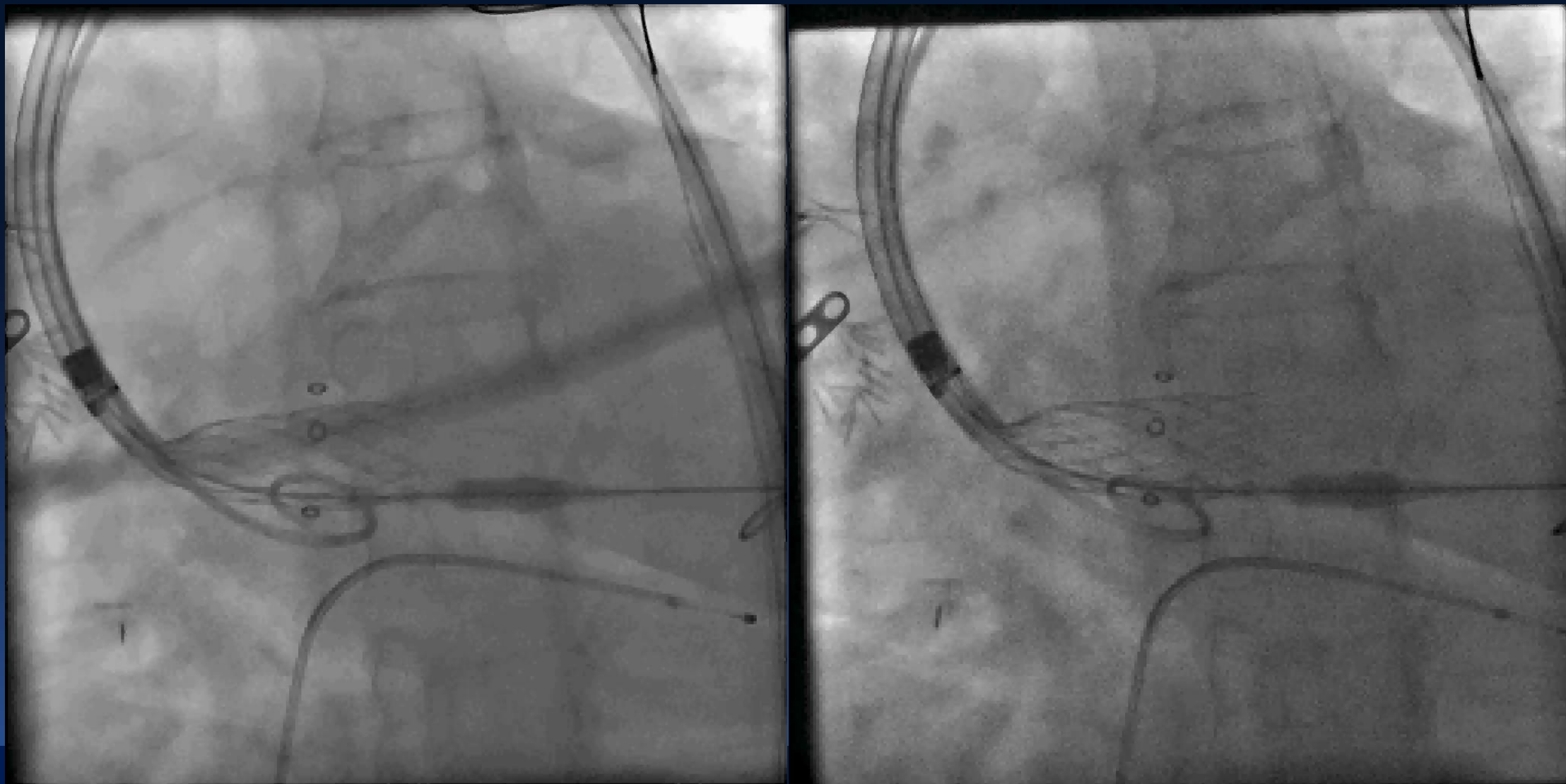


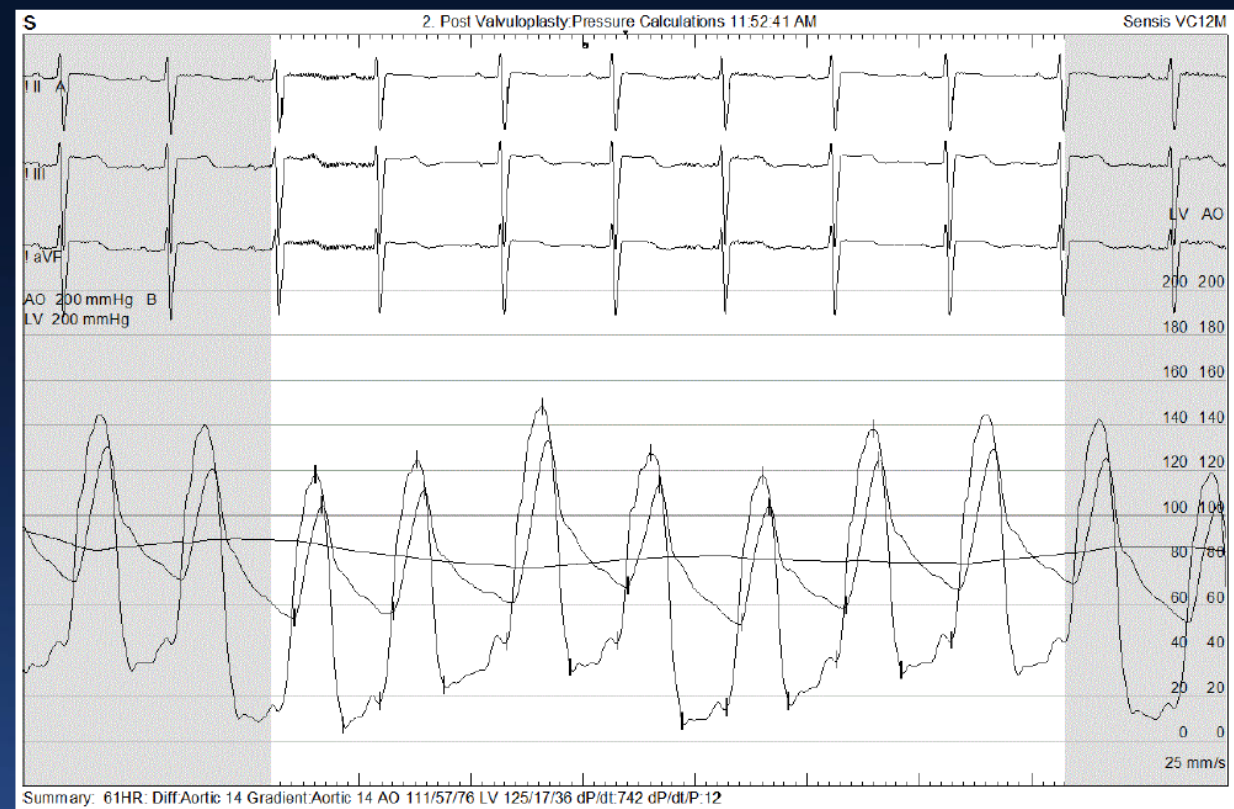


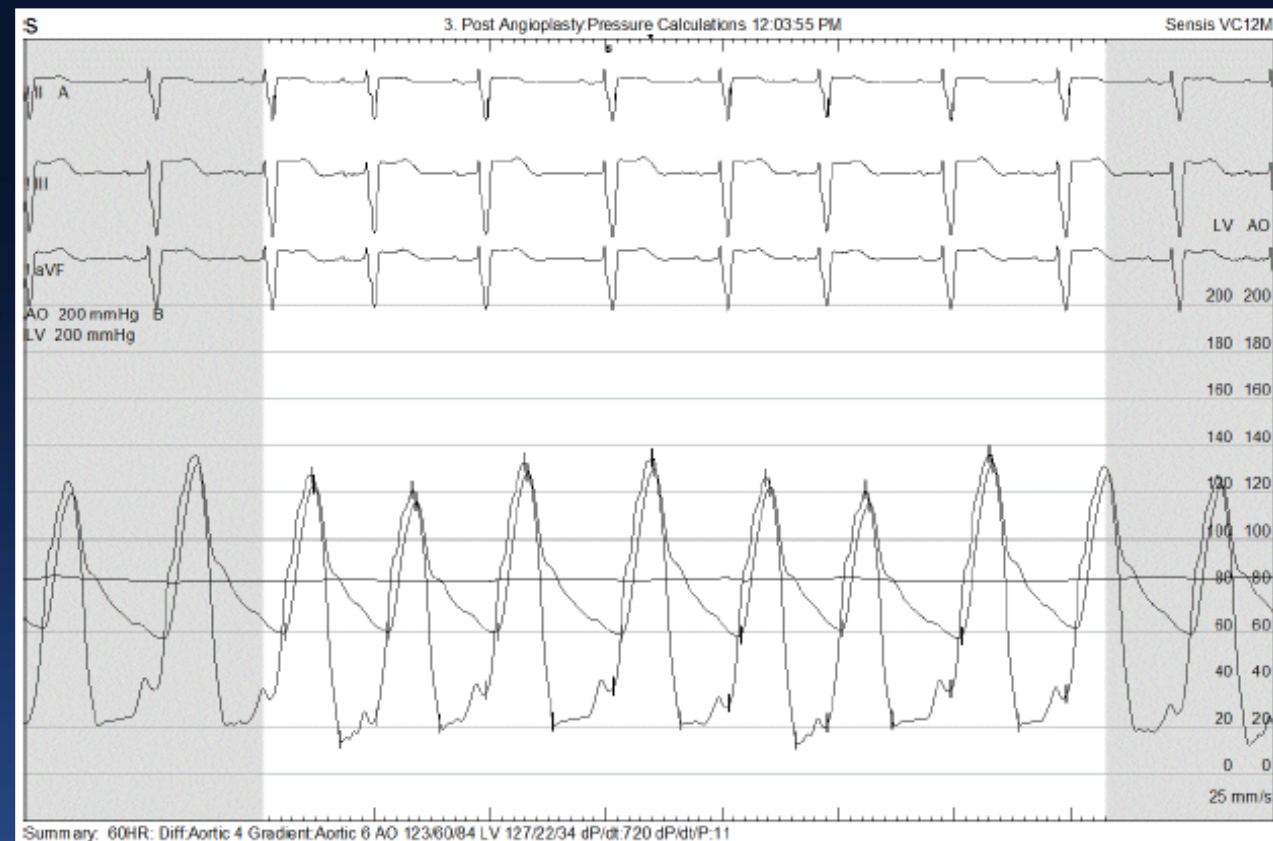
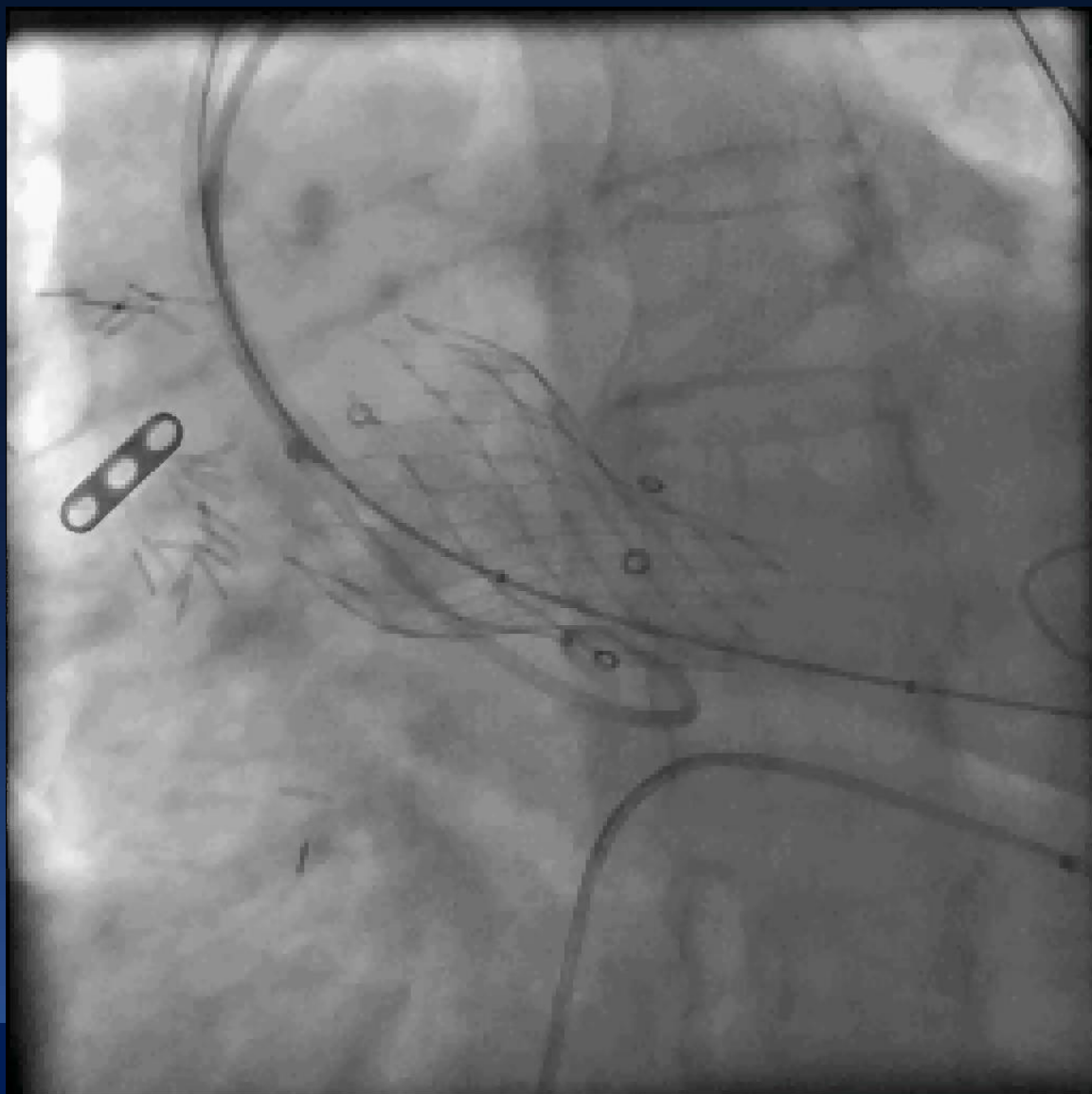




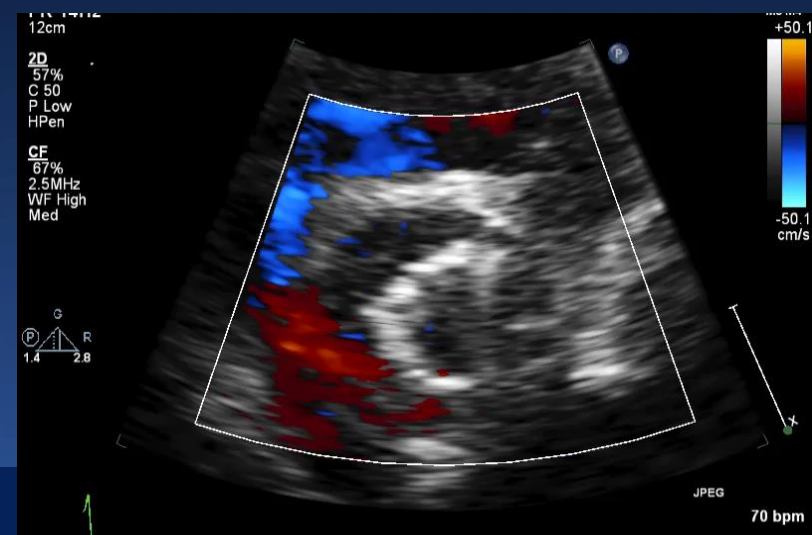
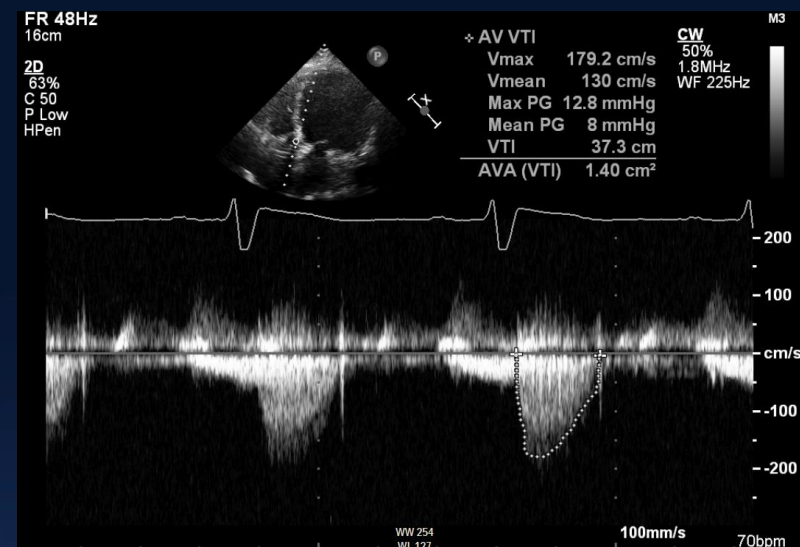
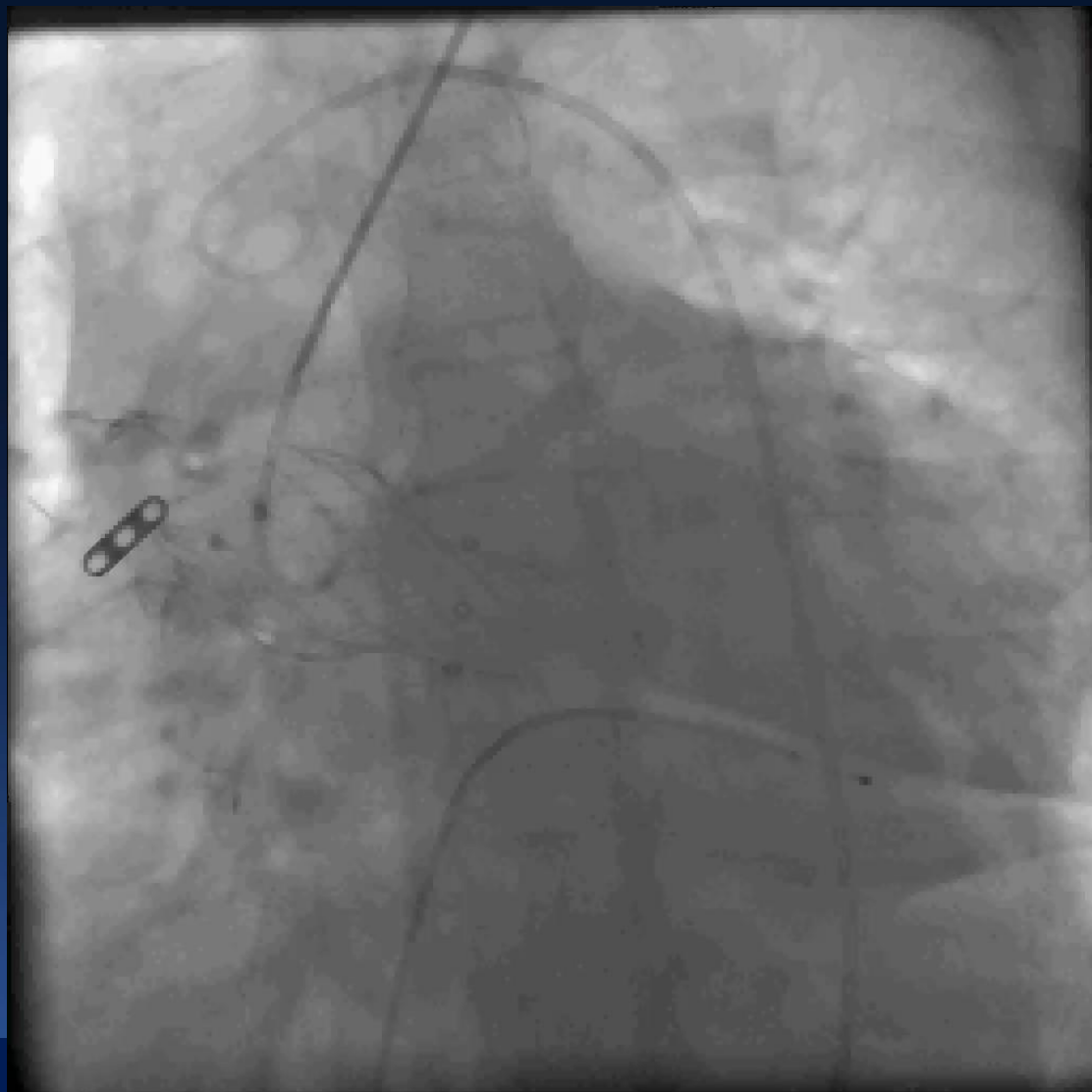




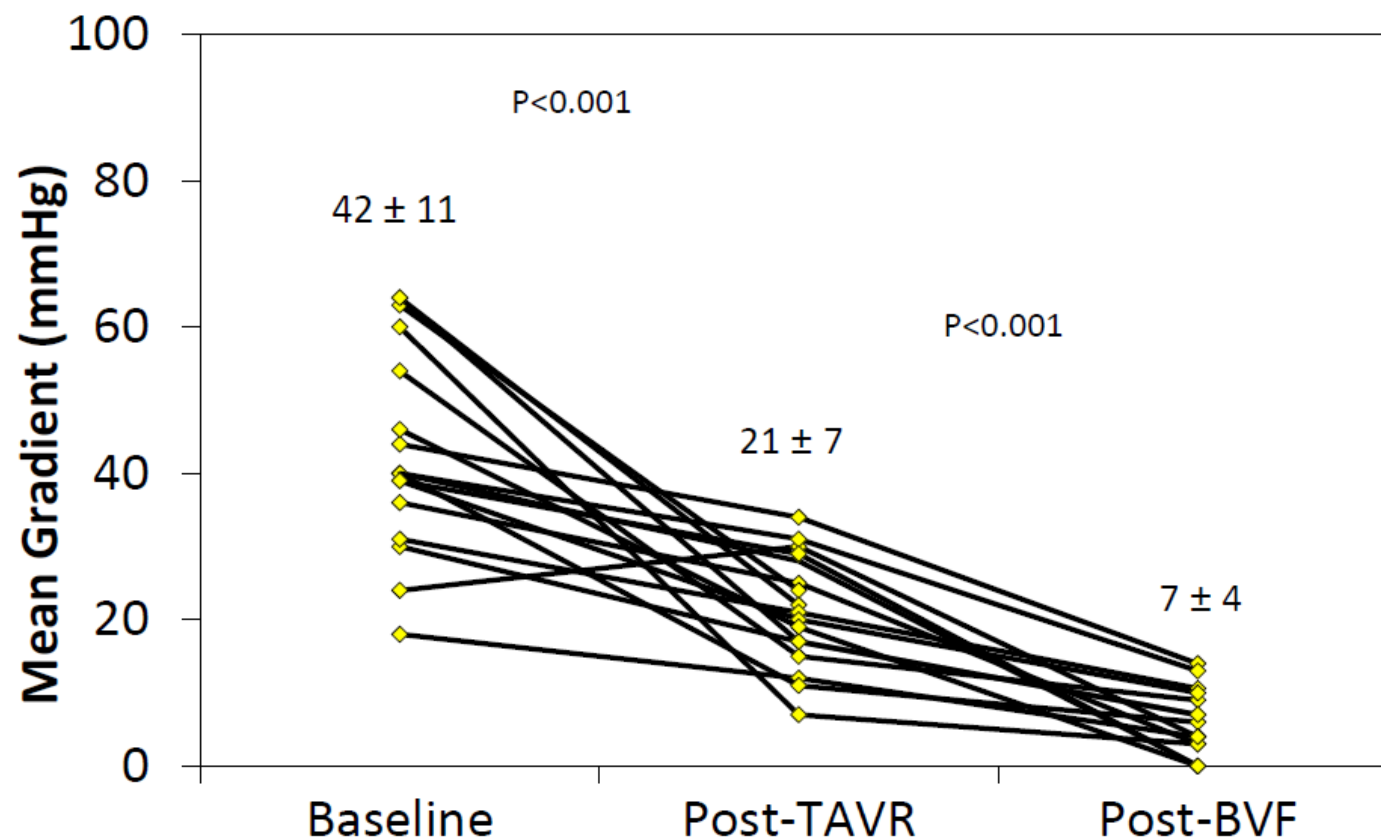




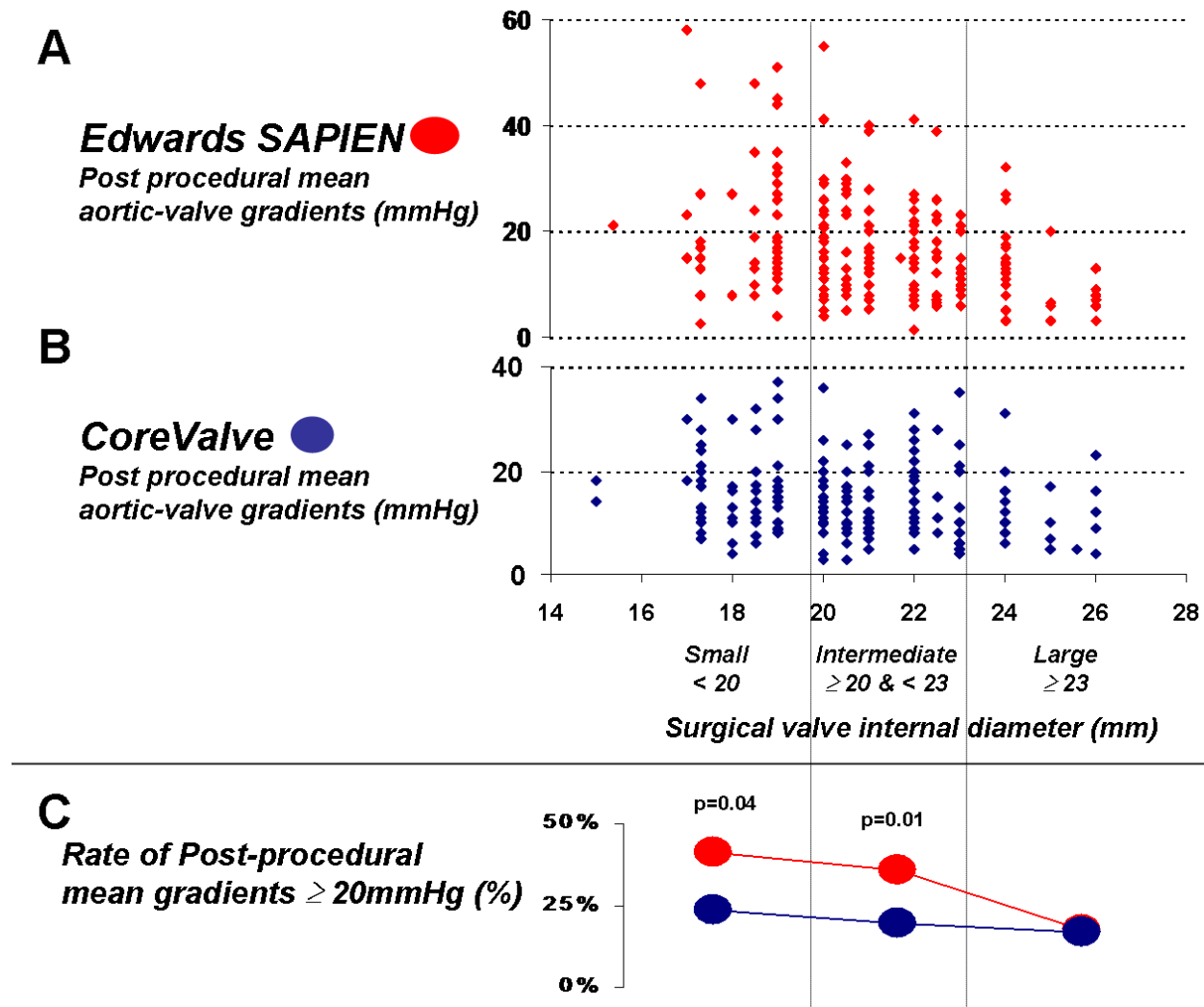
Final Aortogram



Bioprosthetic Valve Ring Fracture



Residual stenosis: Consider supra-annular design



Predictive Factors, Management, and Clinical Outcomes of Coronary Obstruction Following Transcatheter Aortic Valve Implantation

Insights From a Large Multicenter Registry

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Objectives This study sought to evaluate the main baseline and procedural characteristics, management, and clinical outcomes of patients from a large cohort of patients undergoing transcatheter aortic valve implantation (TAVI) who suffered coronary obstruction (CO).

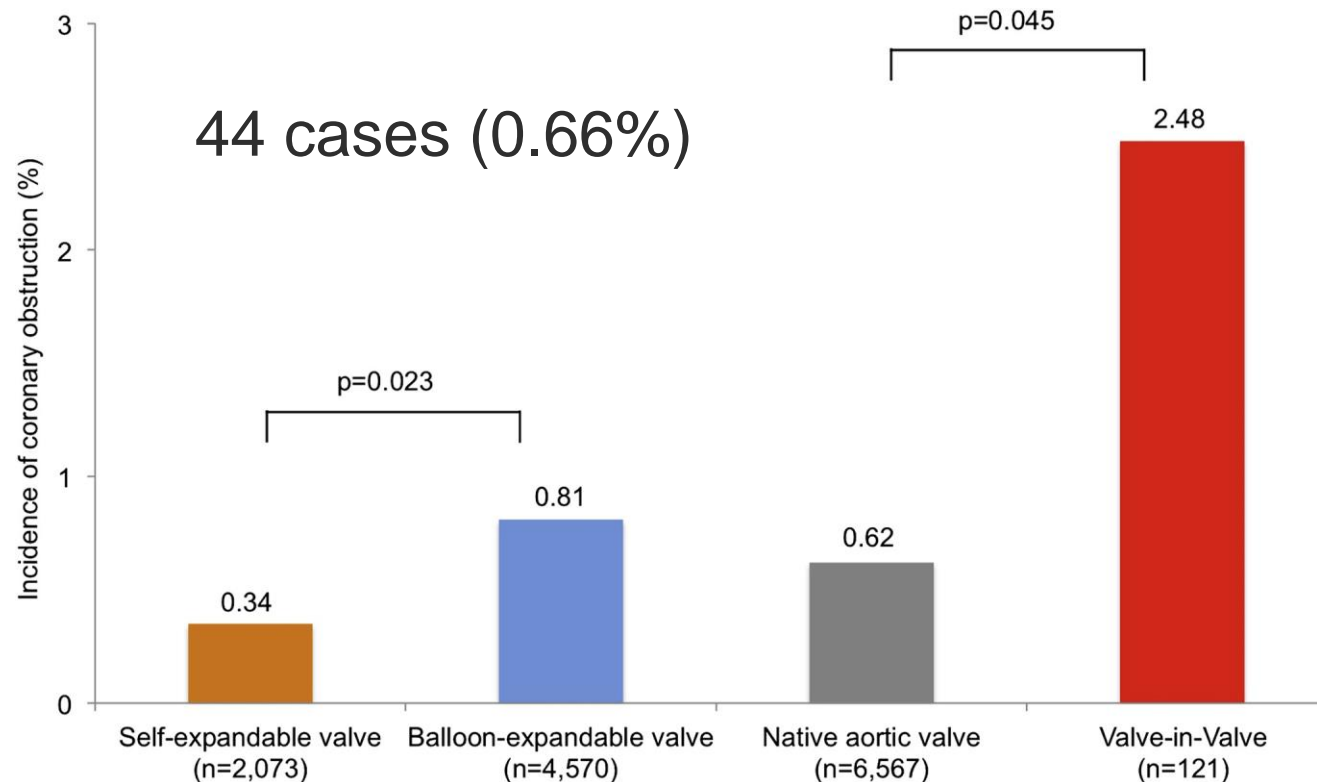
Background Very little data exist on CO following TAVI.

Methods This multicenter registry included 44 patients who suffered symptomatic CO following TAVI of 6,688 patients (0.66%). Pre-TAVI computed tomography data was available in 28 CO patients and in a control group of 345 patients (comparisons were performed including all patients and a cohort matched 1:1 by age, sex, previous coronary artery bypass graft, transcatheter valve type, and size).

Results Baseline and procedural variables associated with CO were older age ($p < 0.001$), female sex ($p < 0.001$), no previous coronary artery bypass graft ($p = 0.043$), the use of a balloon-expandable valve ($p = 0.023$), and previous surgical aortic bioprosthesis ($p = 0.045$). The left coronary artery was the most commonly involved (88.6%). The mean left coronary artery ostia height and sinus of Valsalva diameters were lower in patients with obstruction than in control subjects (10.6 ± 2.1 mm vs. 13.4 ± 2.1 mm, $p < 0.001$; 29.1 ± 3.8 mm vs. 31.9 ± 4.1 mm, $p < 0.001$). Differences between groups remained significant after the case-matched analysis ($p < 0.001$ for coronary height; $p = 0.01$ for sinus of Valsalva diameter). Most patients presented with persistent severe hypotension (68.2%) and electrocardiographic changes (56.8%). Percutaneous coronary intervention was attempted in 75% of the cases and was successful in 81.8%. Thirty-day mortality was 40.9%. After a median follow-up of 12 (2 to 18) months, the cumulative mortality rate was 45.5%, and there were no cases of stent thrombosis or reintervention.

Conclusions Symptomatic CO following TAVI was a rare but life-threatening complication that occurred more frequently in women, in patients receiving a balloon-expandable valve, and in those with a previous surgical bioprosthesis. Lower-lying coronary ostium and shallow sinus of Valsalva were associated anatomic factors, and despite successful treatment, acute and late mortality remained very high, highlighting the importance of anticipating and preventing the occurrence of this complication. (J Am Coll Cardiol 2013;62:1552–62) © 2013 by the American College of Cardiology Foundation

Coronary Obstruction Registry



81 centers in North America, Europe, South America, and Asia, 2007-13

N=6,688, 30-day mortality 40.9%

Ribeiro HB et al. J Am Coll Cardiol 2013;62:1552–62



Incidence, predictors, and clinical outcomes of coronary obstruction following transcatheter aortic valve replacement for degenerative bioprosthetic surgical valves: insights from the VIVID registry

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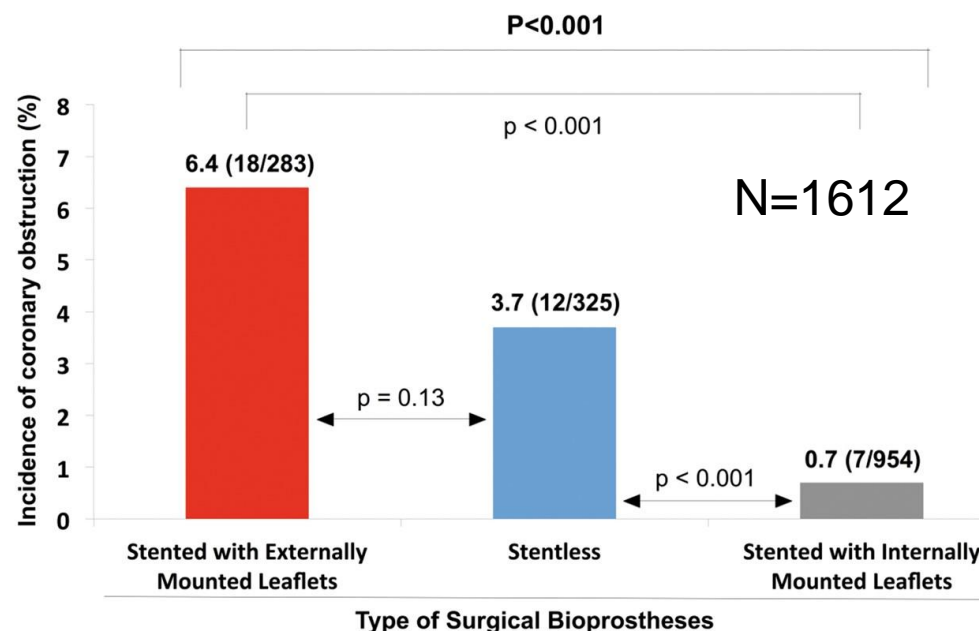
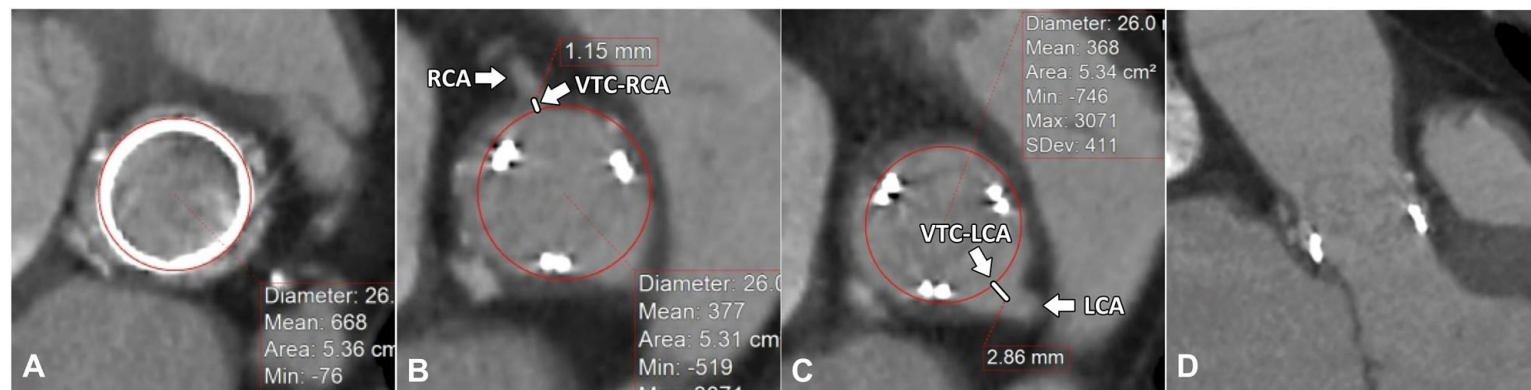
See page 696 for the editorial comment on this article (doi:10.1093/eurheartj/ehx529)

Aims

There are limited data on coronary obstruction following transcatheter valve-in-valve (ViV) implantation inside failed aortic bioprostheses. The objectives of this study were to determine the incidence, predictors, and clinical outcomes of coronary obstruction in transcatheter ViV procedures.

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VIVID Registry



Stentless or stented bioprosthesis with externally mounted leaflets (OR 7.67; 95% CI: 3.14–18.7, $P < 0.001$)

Shorter VTC distance (OR: 0.22 per 1 mm increase; 95% CI: 0.09–0.51; $P < 0.001$), with an optimal cut-off level of 4 mm (area under the curve: 0.943; $P < 0.001$).



