

Selección de Pacientes para TAVI




SOLACI
SOCIEDAD
LATINOAMERICANA
DE CARDIOLOGIA
INTERVENCIONISTA

XXVII Jornadas SOLACI
9° Región Andina

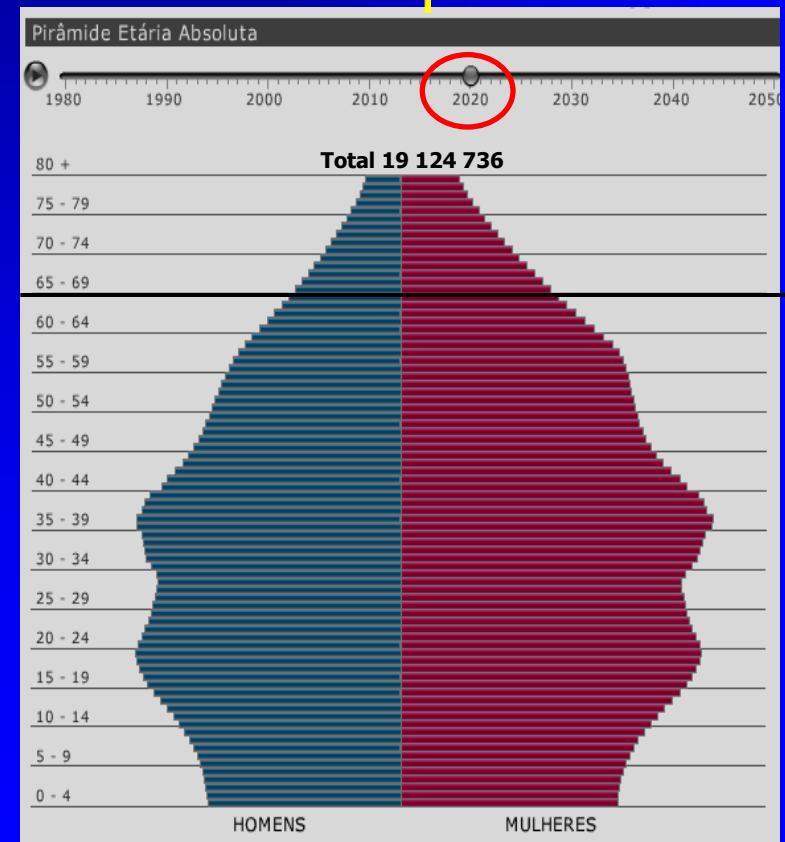
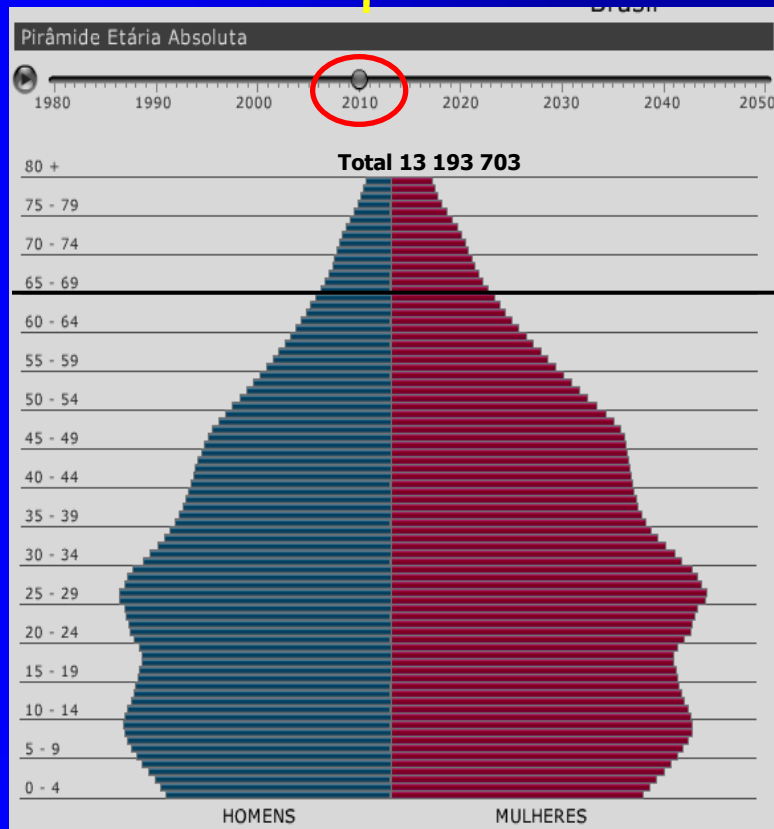
8 / 9 de Octubre 2015

Lima – Perú

População Anciana > 65 anos

Demografia - Brasil

+ 44%



Implante Transcateter Valvar Aórtico

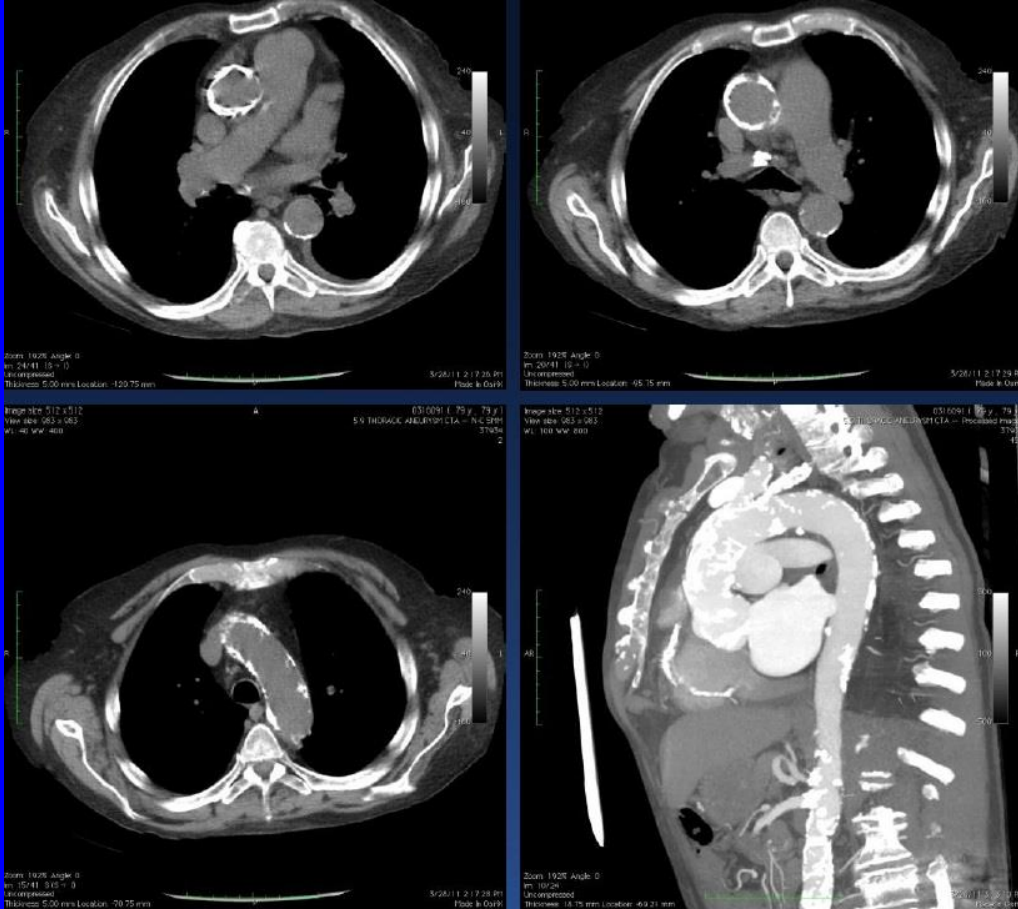
Indicaciones

- **EAO nativa calcificada grave** - AVA $< 1 \text{ cm}^2$ o $\leq 0,6 \text{ cm}^2/\text{m}^2$, Gradiente Medio $> 40 \text{ mmHg}$, Velocidad del Jato $> 4\text{m/s}$
 - **Clase Funcional (NYHA) - $\geq \text{II}$**
 - **Euroscore Logístico - $\geq 15\%$ o STS Score - $\geq 10\%$**
 - **Otros datos clínicos que se deben considerar:**

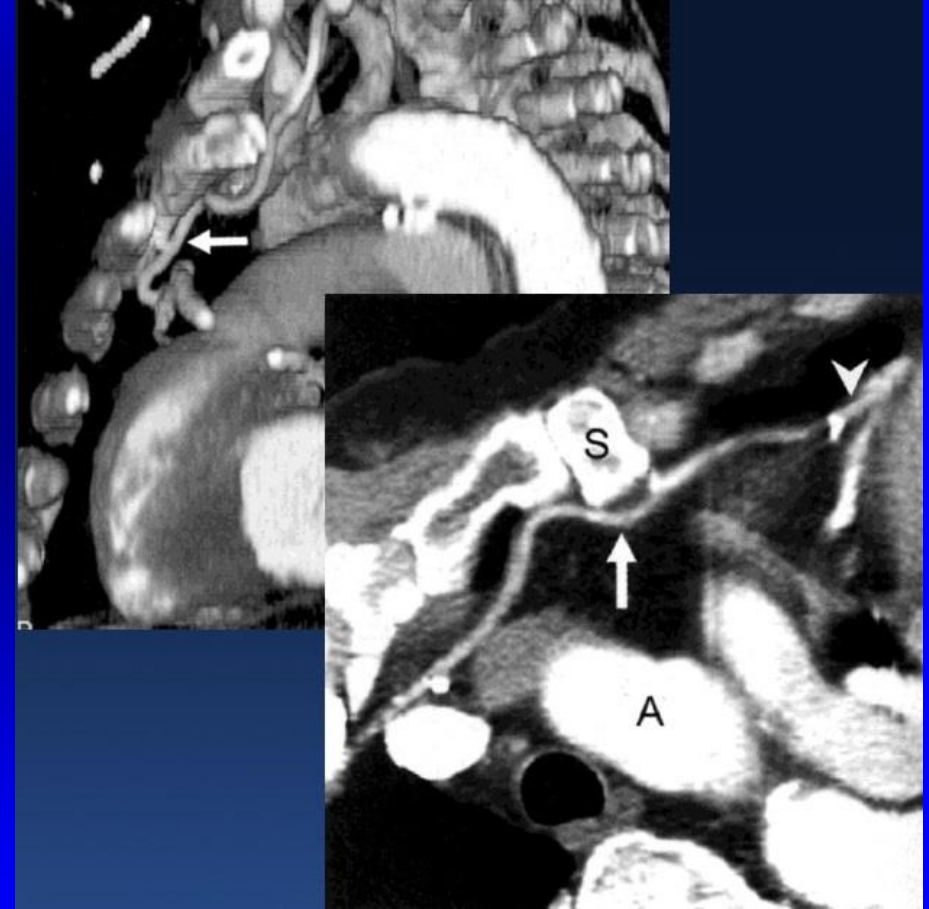
enfermedad hepática grave, aorta en porcelana, anastomosis LIMA-DA adyacente al esternón, HP $> 60 \text{ mmHg}$, insuficiencia VD, enfermedad grave del tejido conectivo, caquexia, tórax hostil (radiación, quemaduras), embolia pulmonar de repetición

Cirugía de Reemplazo Valvar

Condiciones Desfavorables



Aorta en Porcelana



LIMA adyacente al esternón

Estenosis Aórtica

Aspectos Evolutivos

Survival

Percent



Onset

severe

symptoms

Sobrevida Después del Inicio de los Síntomas

50% - 2 años

20% - 5 años



AHA/ACC Guidelines

Momento de la Intervención – TAVI o Cirugía

Recommendations	COR	LOE
AVR is recommended for symptomatic patients with severe high-gradient AS who have symptoms by history or on exercise testing (stage D1)	I	B
AVR is recommended for asymptomatic patients with severe AS (stage C2) and LVEF <50%	I	B
AVR is indicated for patients with severe AS (stage C or D) when undergoing other cardiac surgery	I	B
AVR is reasonable for asymptomatic patients with very severe AS (stage C1, aortic velocity ≥ 5.0 m/s) and low surgical risk	IIa	B
AVR is reasonable in asymptomatic patients (stage C1) with severe AS and decreased exercise tolerance or an exercise fall in BP	IIa	B

Selección de Pacientes

Evaluación de la Fragilidad

Patient A



vs.

Patient B



Same age and predicted risk
One passes the “eyeball test” – one does not

Components of Frailty Score

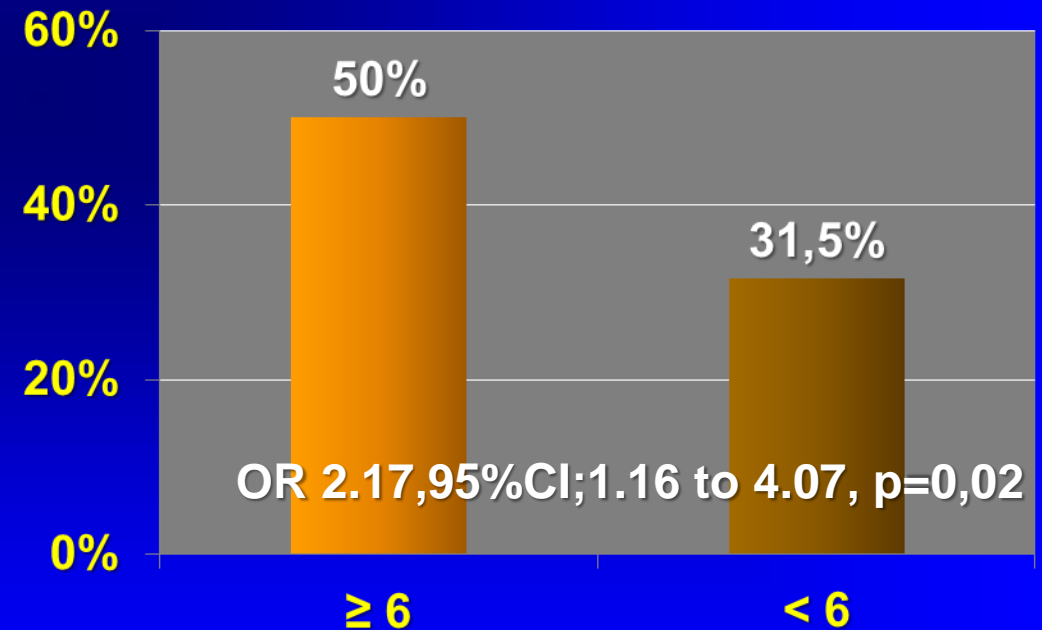
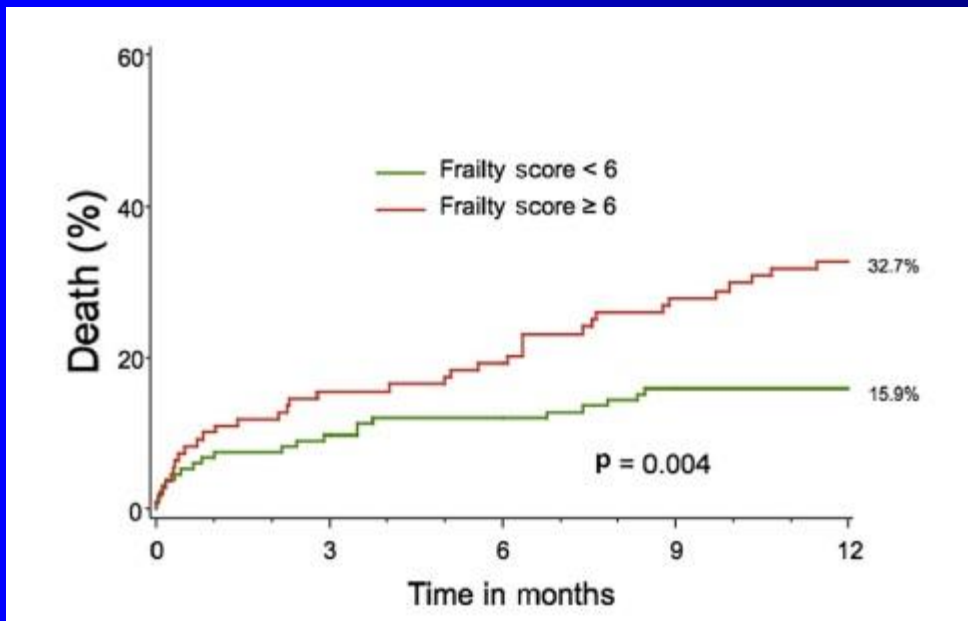
Frailty Domain	Measure	Frailty Score
Slowness	15-ft walk gait speed (m/s)	Quartiles (0-3)
Weakness	Grip strength (kg)	Sex-based quartiles (0-3)
Wasting and malnutrition	Serum albumin (g/dl)	Quartiles (0-3)
Inactivity	Katz activities of daily living	Any dependence = 3, Independent = 0

PARTNER SUBSTUDY

Evaluación de la Fragilidad

n = 244

Mortalidad y Calidad de Vida



AHA/ACC Guidelines

Recomendaciones para TAVI

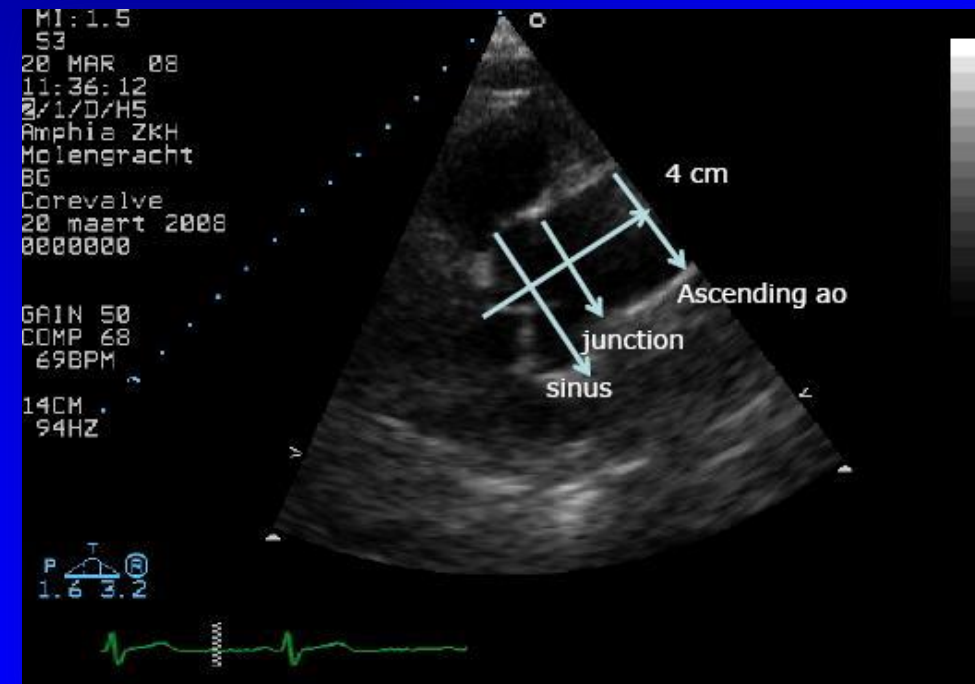
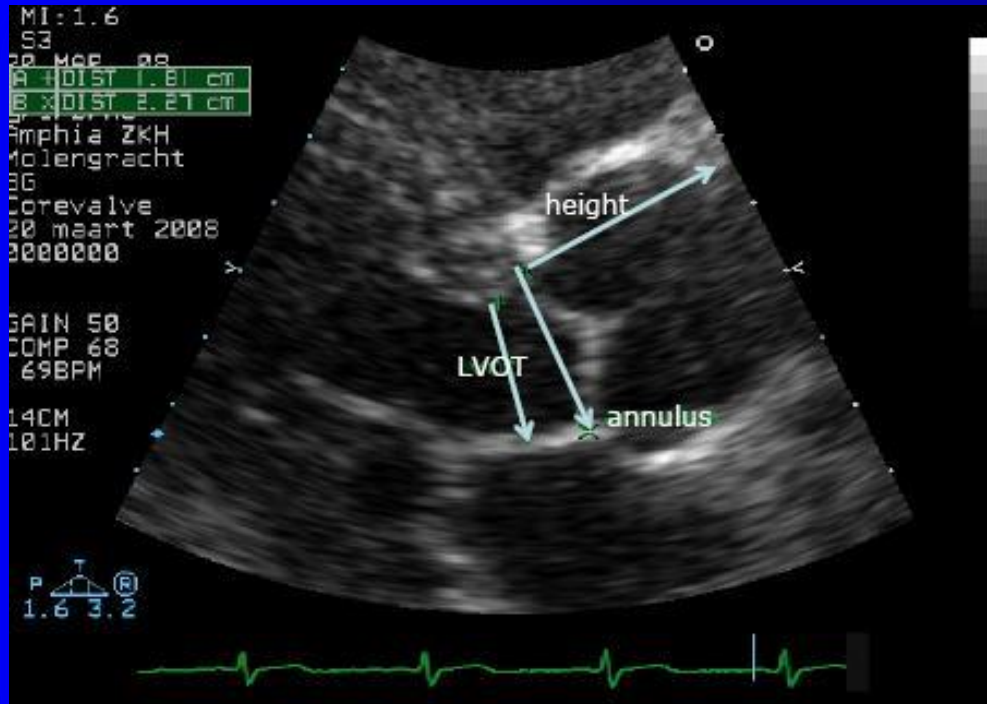
Table 8. Summary of Recommendations for AS: Choice of Surgical or Transcatheter Intervention

Recommendations	COR	LOE
Surgical AVR is recommended in patients who meet an indication for AVR (Section 3.4) with low or intermediate surgical risk (Section 2.5 in the full-text guideline)	I	A
For patients in whom TAVR or high-risk surgical AVR is being considered, members of a Heart Valve Team should collaborate to provide optimal patient care	I	C
TAVR is recommended in patients who meet an indication for AVR for AS who have a prohibitive surgical risk and a predicted post-TAVR survival >12 mo	I	B
TAVR is a reasonable alternative to surgical AVR in patients who meet an indication for AVR (Section 3.4) and who have high surgical risk (Section 2.5 in the full-text guideline)	IIa	B
Percutaneous aortic balloon dilation may be considered as a bridge to surgical or transcatheter AVR in severely symptomatic patients with severe AS	IIb	C
TAVR is not recommended in patients in whom existing comorbidities would preclude the expected benefit from correction of AS	III: No Benefit	B

Selección de Pacientes

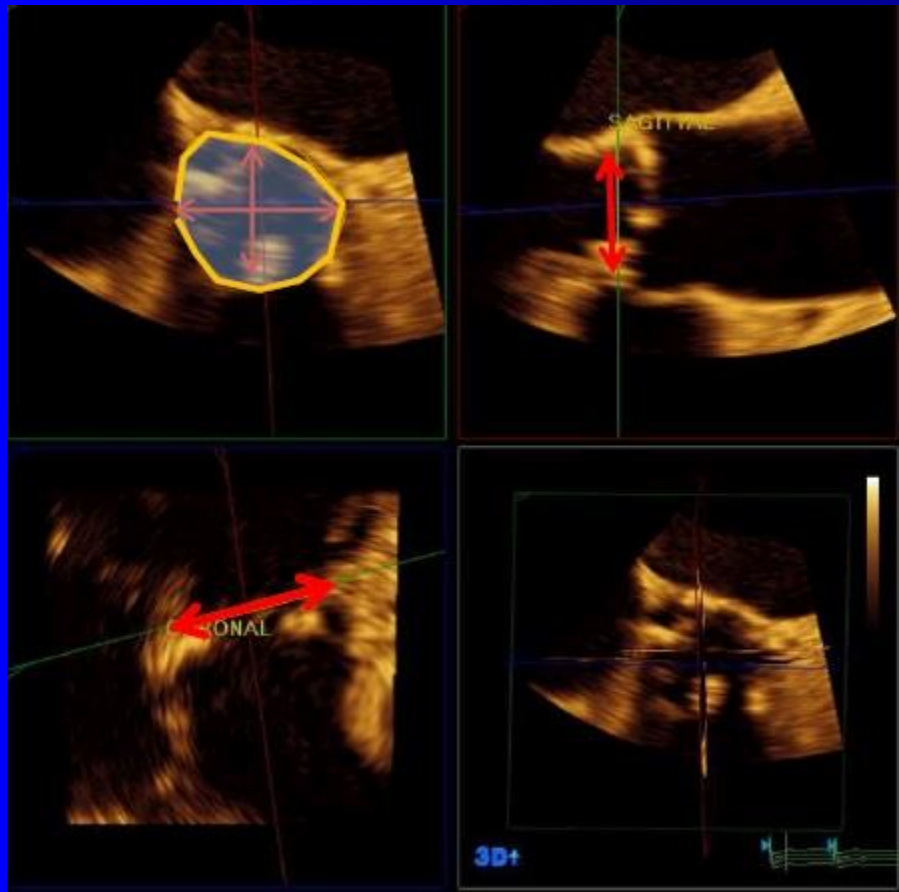
Evaluación Ecocardiografía

Proyección Paraesternal Eje Mayor



Ecocardiograma 3D

Ventajas



Annular Measurements

Advantages of 3D:

1. Ensures on-axis measurements of orthogonal views
2. Allows 3D assessment of annulus

CLINICAL RESEARCH

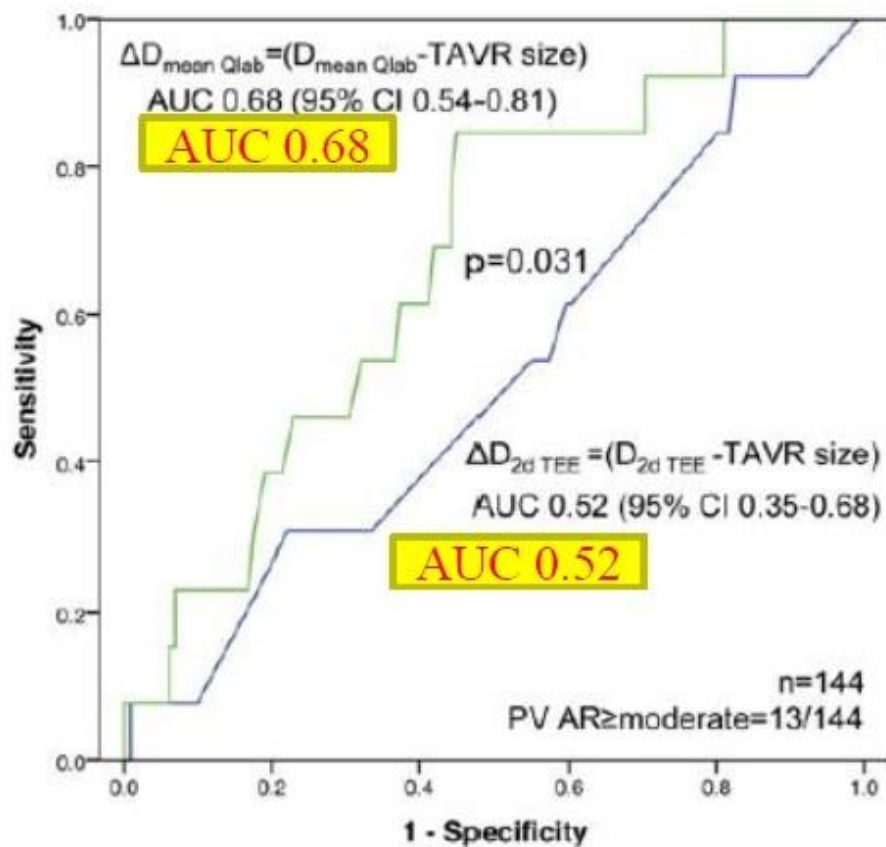
Interventional Cardiology

Aortic Annular Sizing for Transcatheter Aortic Valve Replacement Using Cross-Sectional 3-Dimensional Transesophageal Echocardiography

Cedars-Sinai Experience

256 patients undergoing TAVR with
balloon-expandable valve

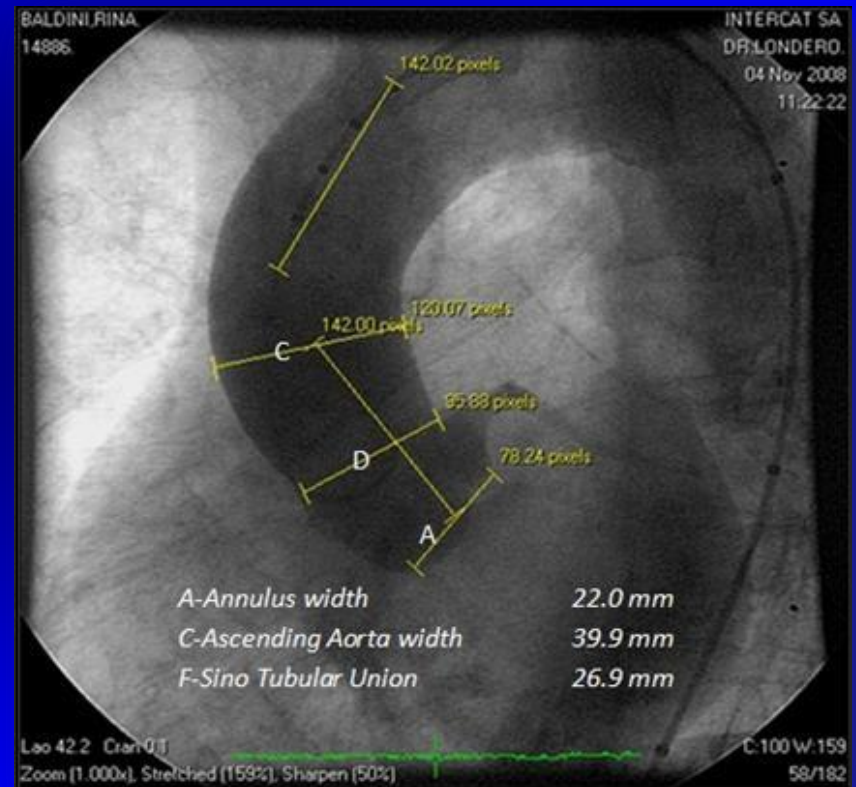
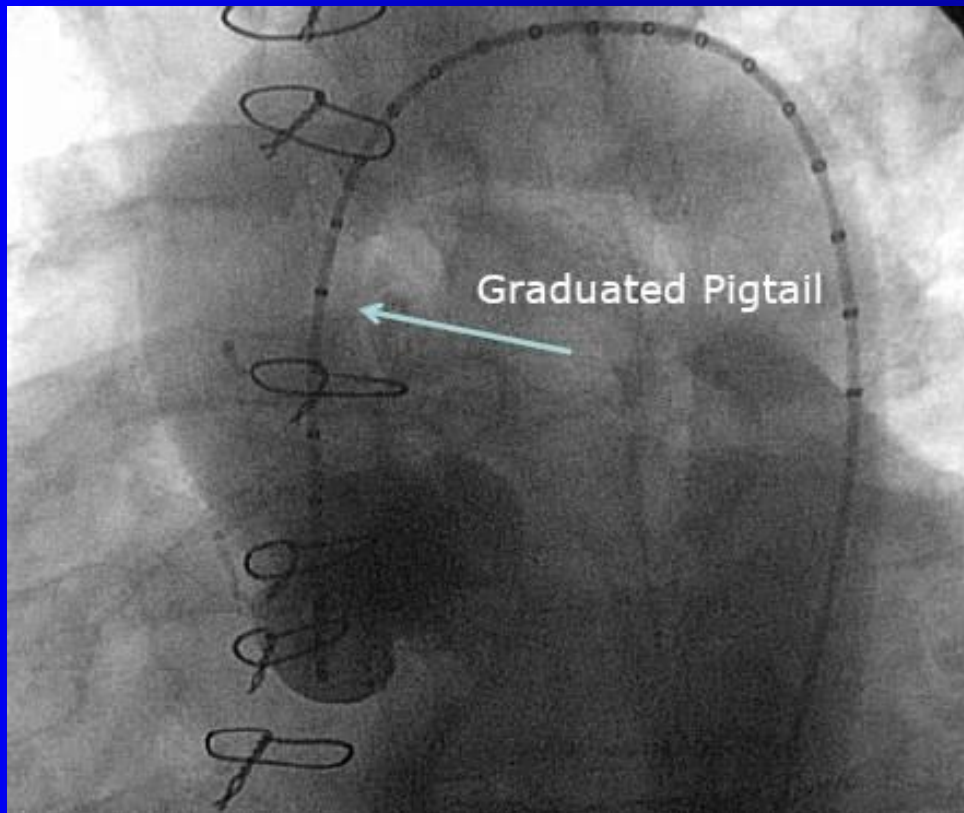
3D TEE (Qlab) vs 2D TEE



**3D TEE has greater
discriminatory value
at predicting PAVR,
compared to 2D TEE.**

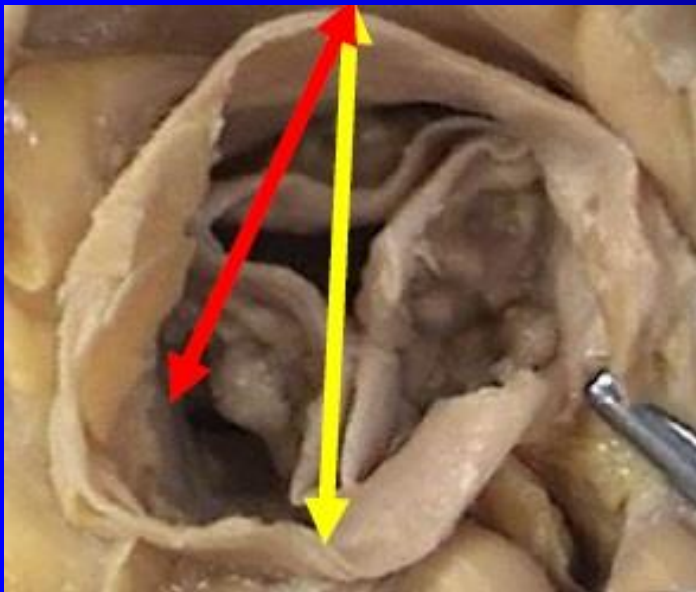
Implante Transcateter Valvar Aórtico

Evaluación Angiográfica

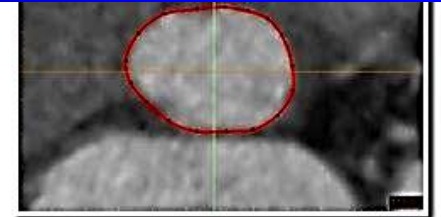


Evaluación del Anillo Valvar

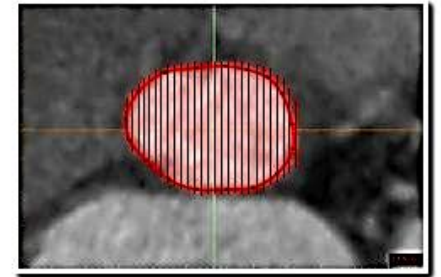
MSCT



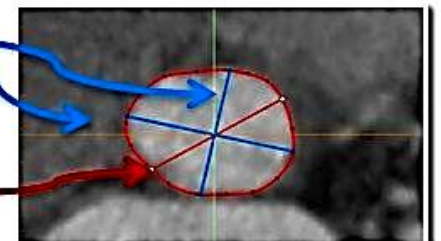
Perimeter: linear distance of tracing around the aortic annulus



Area: area contained within tracing around the aortic annulus



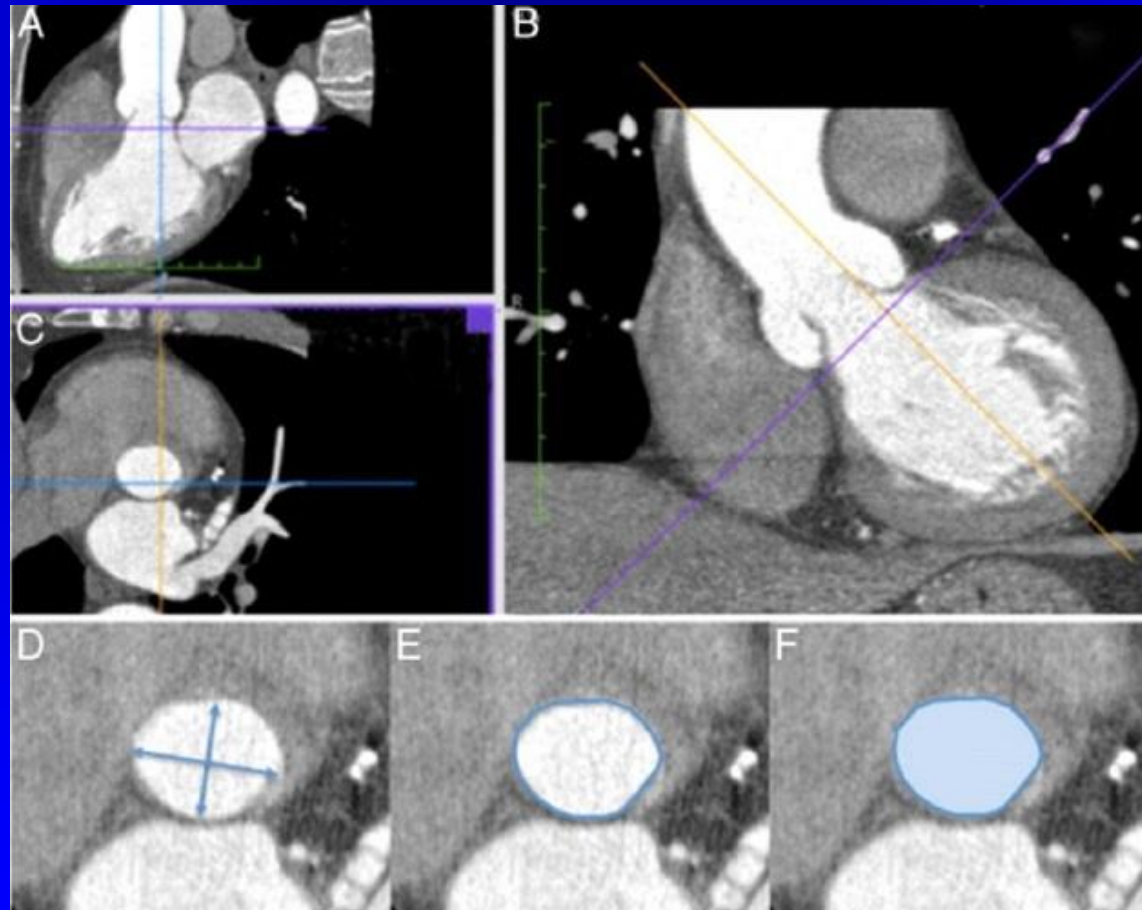
Major & Orthogonal Minor Diameters: linear distances through the center of the aortic annulus



Mean Diameter: Calculated mean of major and minor diameters

Evaluación del Anillo Valvar

MSCT



CLINICAL RESEARCH

Interventional Cardiology

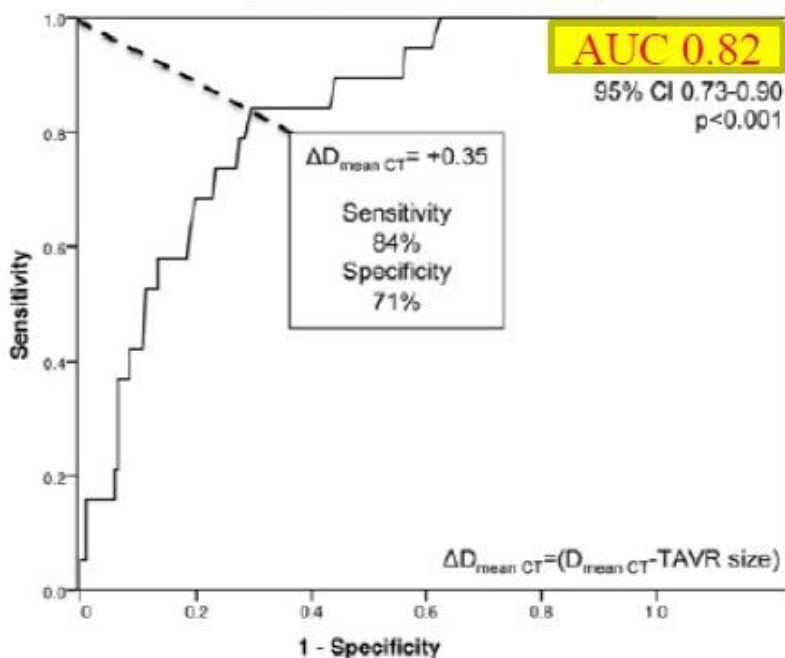
**Aortic Annular Sizing for Transcatheter
Aortic Valve Replacement Using Cross-Sectional
3-Dimensional Transesophageal Echocardiography**

Cedars-Sinai Experience

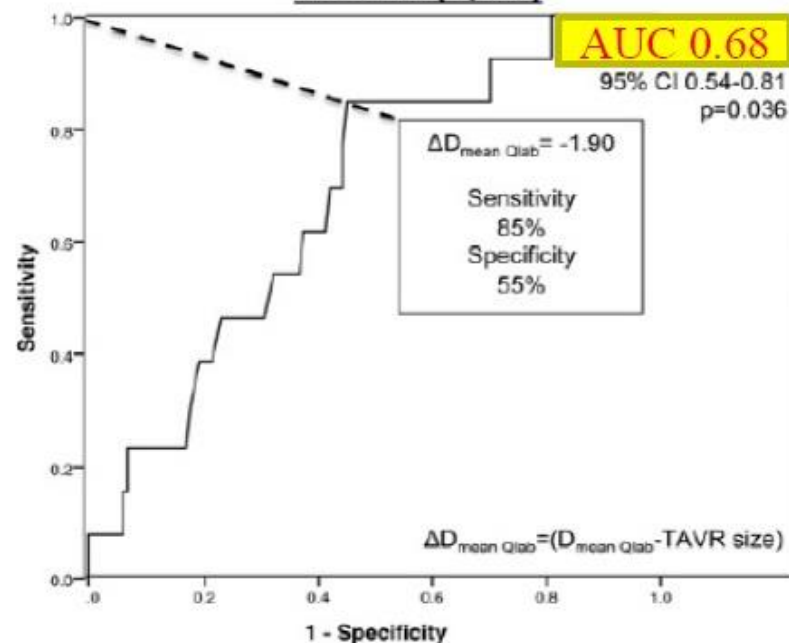
256 patients undergoing TAVR with
balloon-expandable valve

**CT-based parameters have numerically greater discriminatory
value for PVAR, compared with 3D TEE-derived parameters.**

Cross-sectional CT







3D TEE (Qlab)



Elección de la Prótesis

CoreValve

Producto	CoreValve® Evolut™	CoreValve®		
				
Tamaño	23 mm	26 mm	29 mm	31 mm
Diámetro do anel	18-20 mm	20-23 mm	23-26 mm	26-29 mm
Perímetro do anel	56.5-62.8 mm	62.8-72.3 mm	72.3-81.7 mm	81.7-91.1 mm
Diámetro da aorta ascendente	≤34 mm @ 30 mm do anel	≤40 mm @ 40 mm do anel	≤43 mm @ 40 mm do anel	≤43 mm @ 40 mm do anel
Diámetros do seio de Valsalva	≥25 mm	≥27 mm	≥29 mm	≥29 mm
Altura do seio de Valsalva	≥15 mm	≥15 mm	≥15 mm	≥15 mm

Elección de la Prótesis

Edwards Sapien

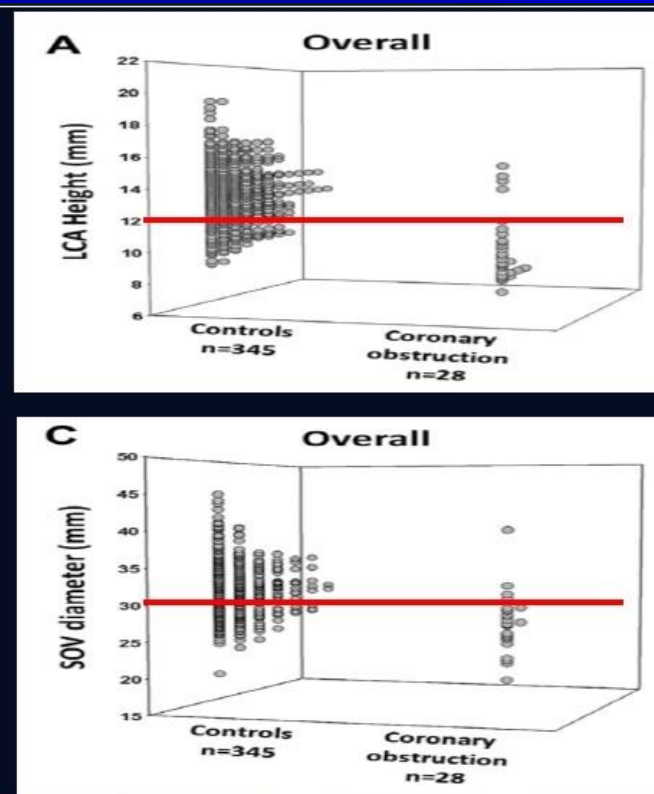
Annulus area mm ²	Perimeter (mm)	Diameter (mm)	THV size (mm)	% oversize by area
255	57	18	23	63%
285	60	19	23	46%
315	63	20	23	30%
345	66	21	23	20%
380	69	22	23-26	9%-39%
415	72	23	23-26	0%-28%
450	75	24	26	17%
490	79	25	26-29	8%-35%
530	82	26	26-29	0%-25%
575	85	27	29	15%
615	88	28	29	7%
660	91	29	29	0

Coronary Obstruction

CT Data

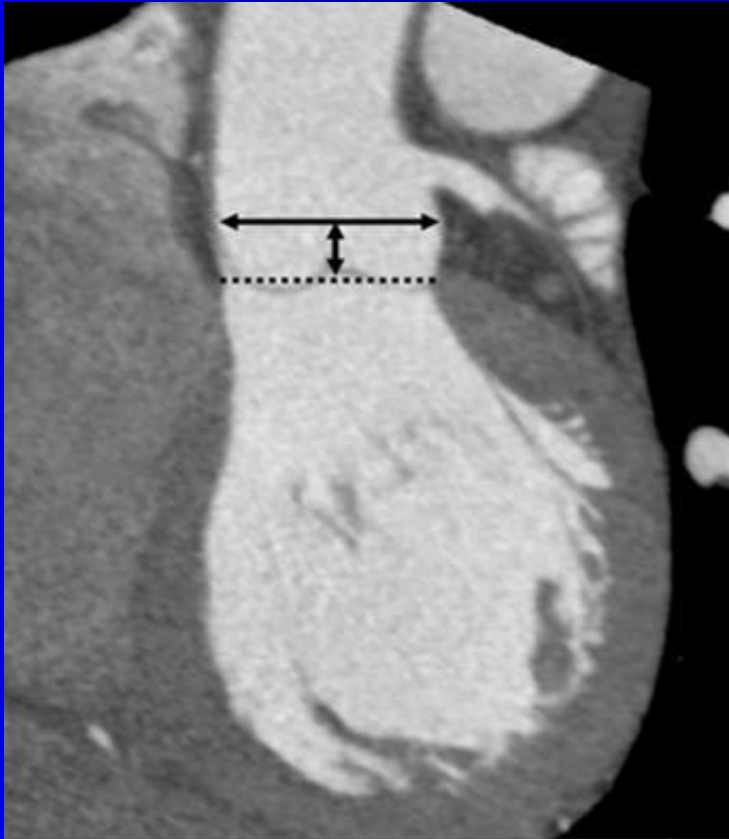
n = 6.688 81 centros n = 44 (0,66%)

- Up to 86% of the pts who had a coronary obstruction had a LCA height of < 12 mm
- The SOV diameter was <30 mm in 71.4% of the pts who had a coronary obstruction



Implante Valvar Aórtico

Altura Coronaria



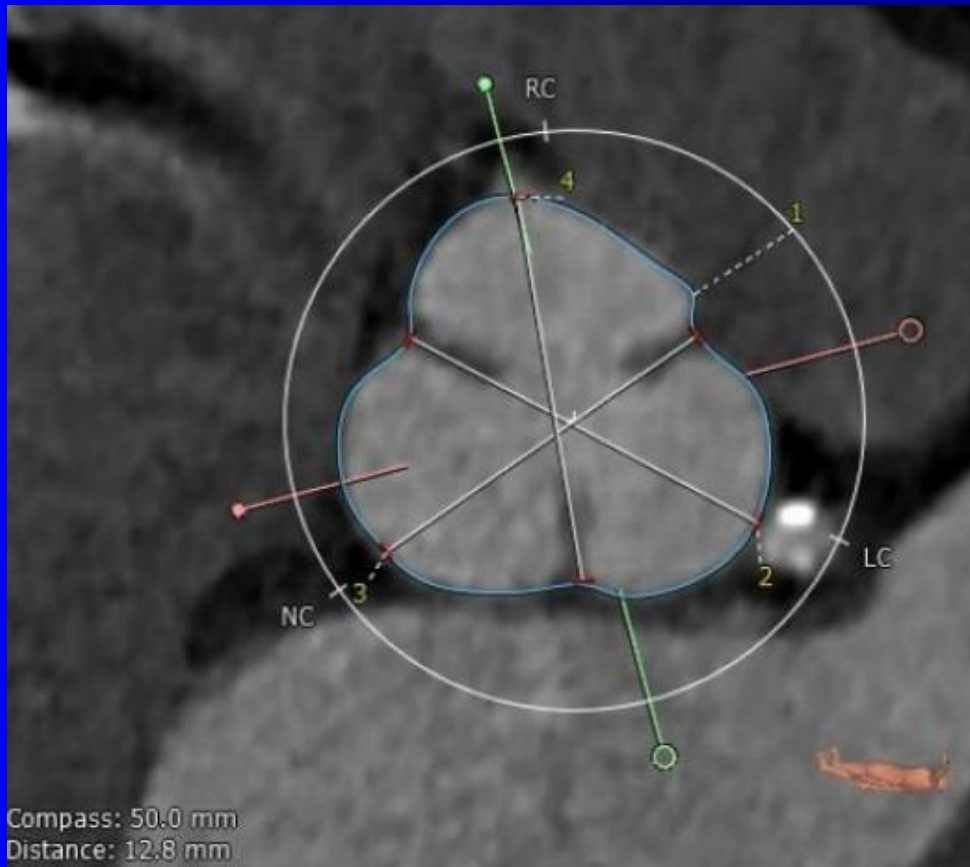
Annulus 26.3+2.8 mm
Sinus of Valsalva 32.4+4.8 mm



Annulus-L Cor. Ostium 14.4+2.9 mm
L Cor. Leaflet Length 14.2+1.8 mm

Coronary Obstruction

CT Data

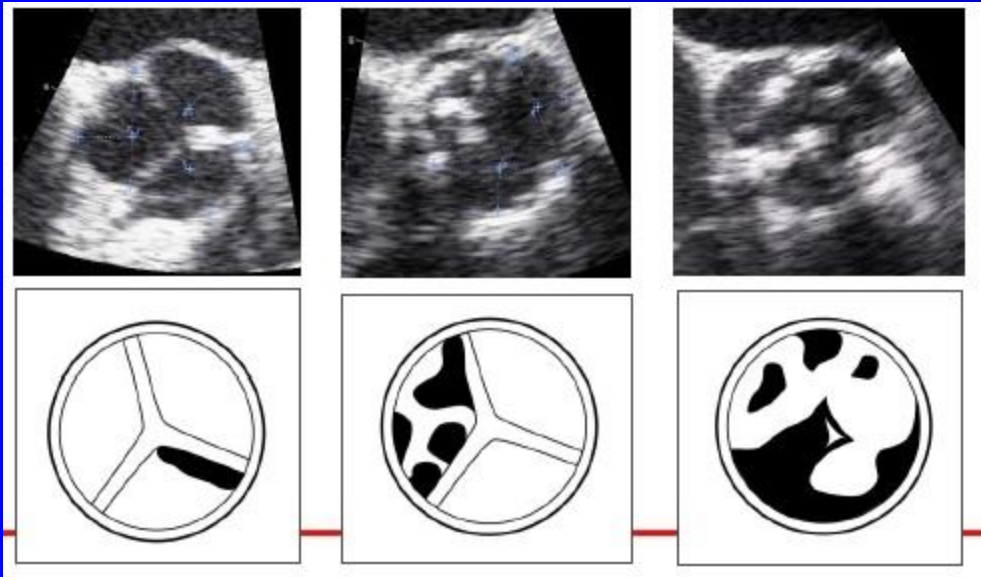


ID	Type	Label	Value
1	MasterSplineAtCurve	Area derived \emptyset	34.7 mm
		Perimeter derived \emptyset	36.6 mm
		Area	945.2 mm ²
		Perimeter	114.9 mm
2	Diameter	Sinus Of Valsava Width - LC	33.9 mm
3	Diameter	Sinus Of Valsava Width - NC	32.6 mm
4	Diameter	Sinus Of Valsava Width - RC	33.5 mm

Implante Transcateter Valvar Aórtico

Grado de Calcificación

- Calcificación Homogénea es Ideal
- Distribución heterogénea aumenta la posibilidad de “leak paravalvar”
- Calcificación voluminosa → Riesgo de obstrucción coronaria

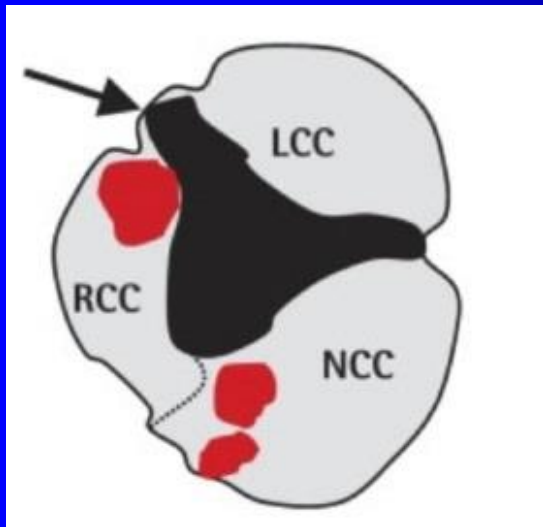


Proyección Eje Menor

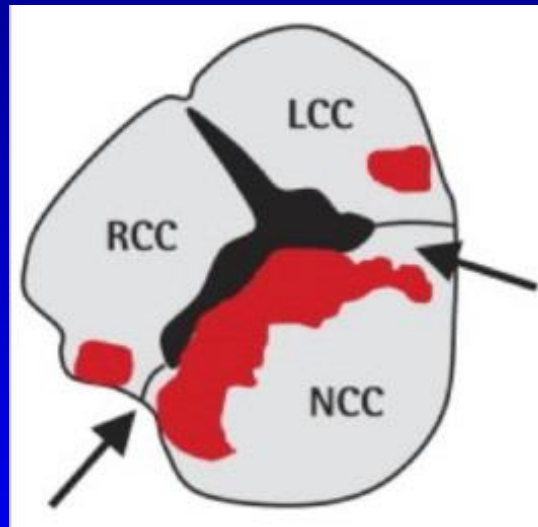


Regurgitación Aórtica Pós-Procedimiento

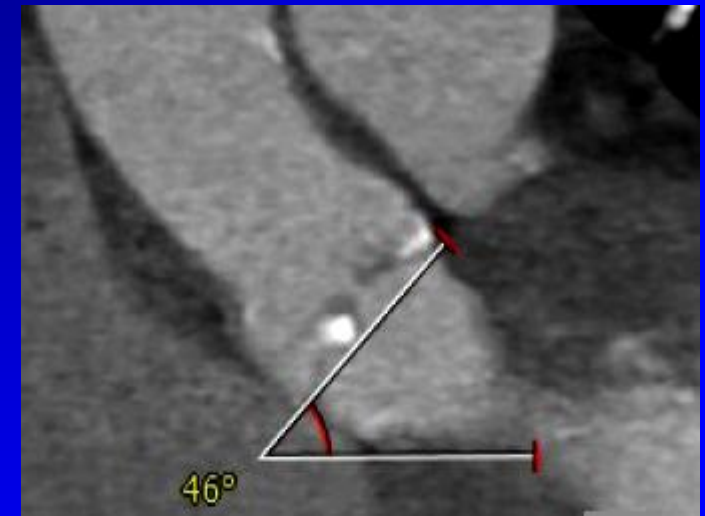
Factores Morfológicos de Riesgo



**Comisuras sin fusión
vecino a masas
voluminosas de calcio**



**Distribución asimétrica de
las masas de calcio en las
cúspides (rojo)**



**Angulación extrema entre
la vía de salida VI e la
aorta ascendente***

* CoreValve - > 45°

Rotura del Anillo Aórtico

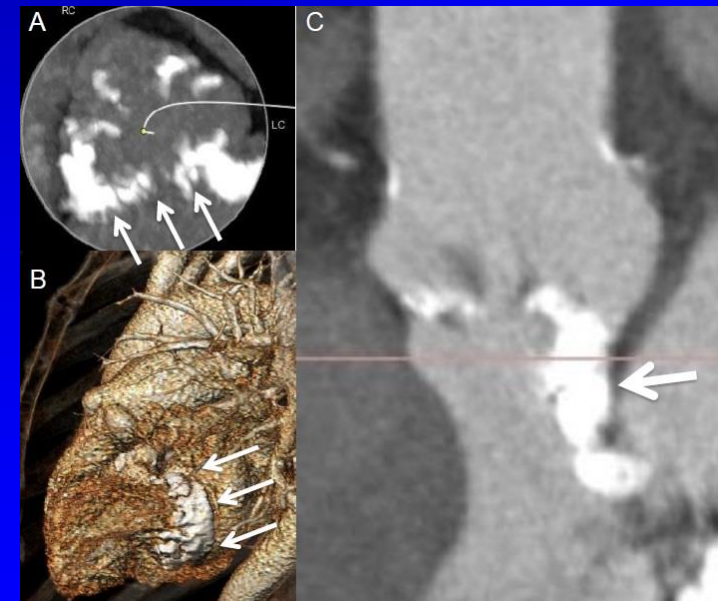
Predictores

n = 16 centers n = 31 (annular or LVOT rupture)

n = 31 (matched sample no rupture)

Conditional Logistic Regression Analysis

Factors		p value
Moderate/Severe LVOT/ Subannular Calcification	OR, 10.92; 95% CI, 3.23–36.91	< 0.001
Prosthesis Oversizing ≥ 20%	OR, 8.38; 95% CI, 2.67-26.33	= 0.001



Procedimiento de Valve-in-Valve

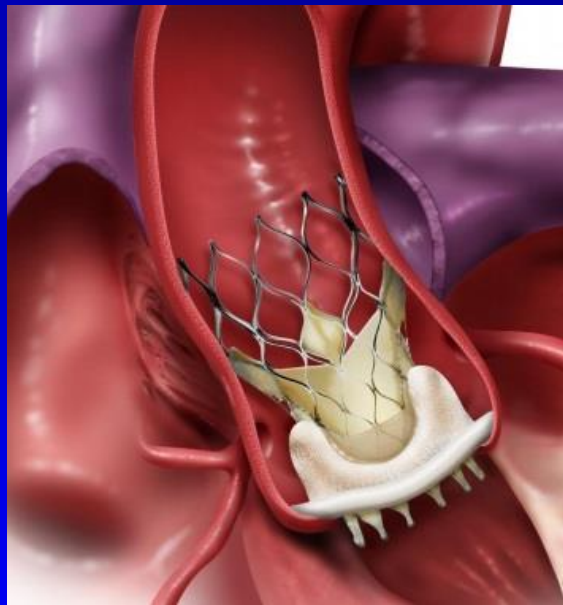
Diámetro Interno de la Prótesis

Inner diameter of aortic bioprostheses with corresponding suggested Sapien™ XT valve size			
	Labelled size (mm)	Measured inner diameter (mm)	Suggested Sapien™ size (mm)
Sorin Biomedica MITROFLOW™	21	17	23*
	23	19	23
	25	20	23
	27	22	23
Edwards PERIMOUNT™ Magna Ease	21	18	23*
	23	21	23
	25	22	23
	27	24	26
St. Jude medical TRIFECTA™	21	18	23*
	23	20	23
	25	22	23
	27	24	26
Medtronic HANCOCK II™	21	18	23*
	23	20	23
	25	22	23
	27	24	26

*The suggested Sapien™ size will create high trans-valvular gradients. This option should be used only in inoperable patients

Procedimiento de Valve-in-Valve

CoreValve vs Edwards Sapien

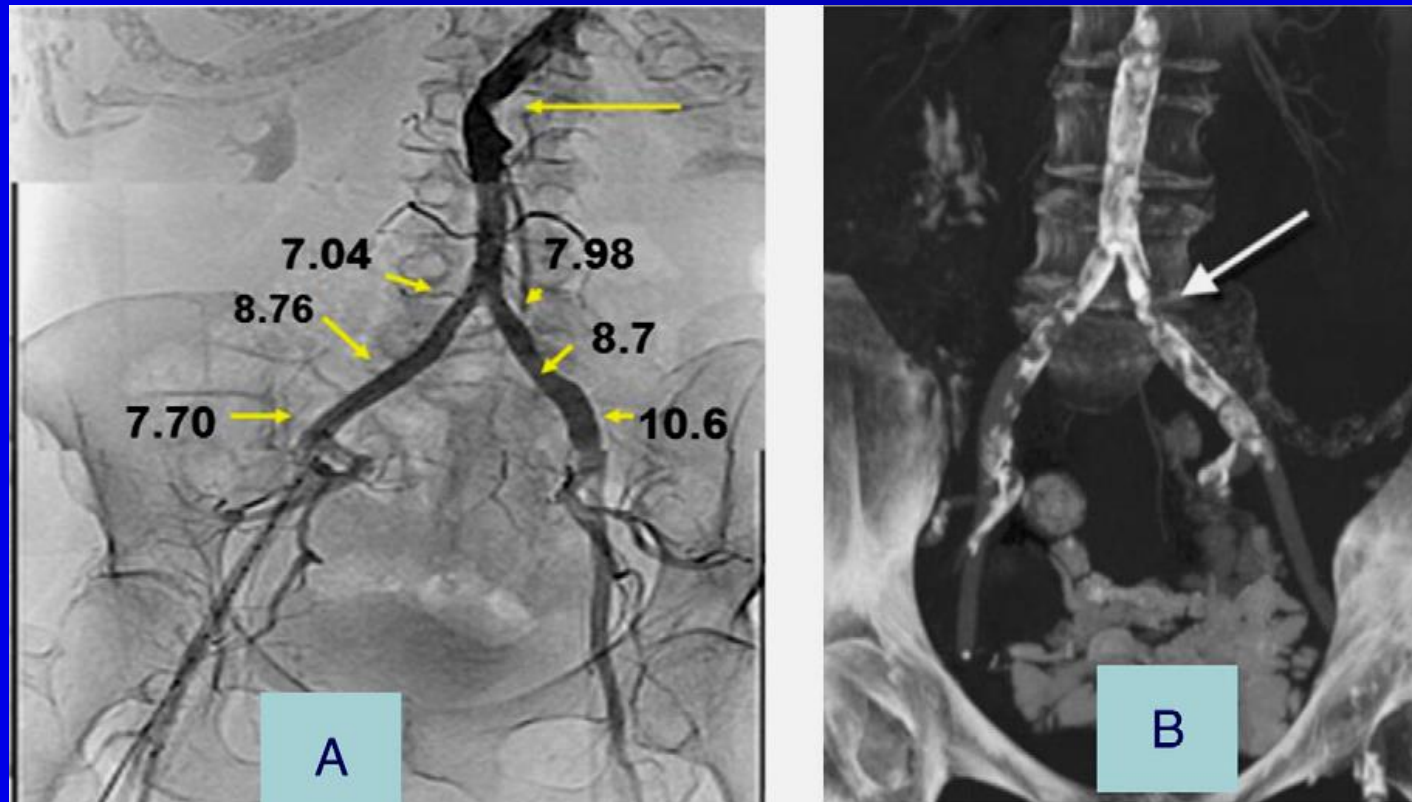


CoreValve - Las valvas se posicionan por encima de la prótesis quirúrgica; disminuyendo la dependencia de la dimensión interna de la bioprótesis

Evaluación Vascular Periférica

Grado de Calcificación e Calibre

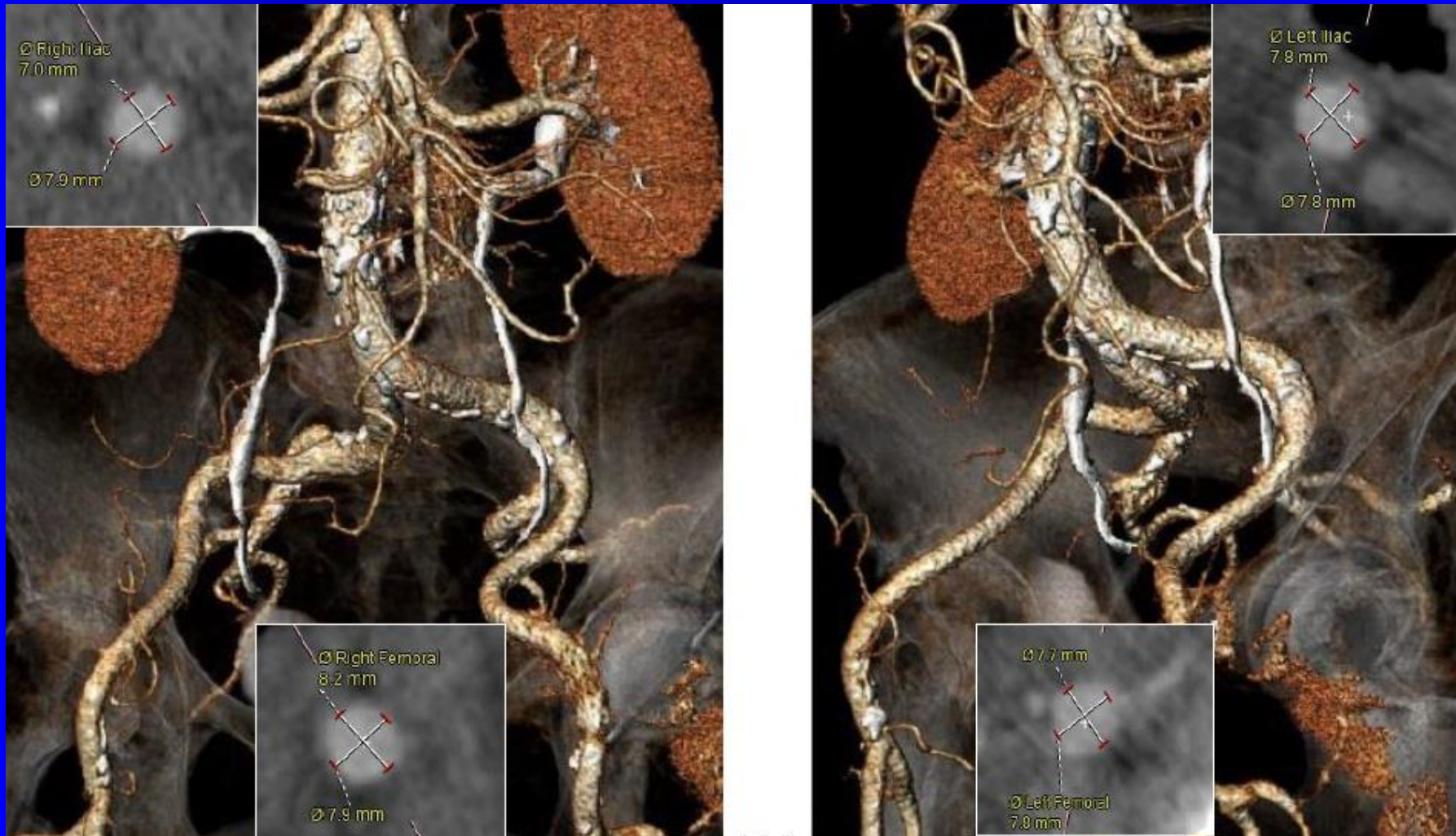
Calcificación puede no ser correctamente evaluada por angiografía



Diámetro Mínimo da Artéria Femoral
CoreValve - 6 mm
Edwards Sapien – 6 mm (23) e 6,5 mm (26-29)

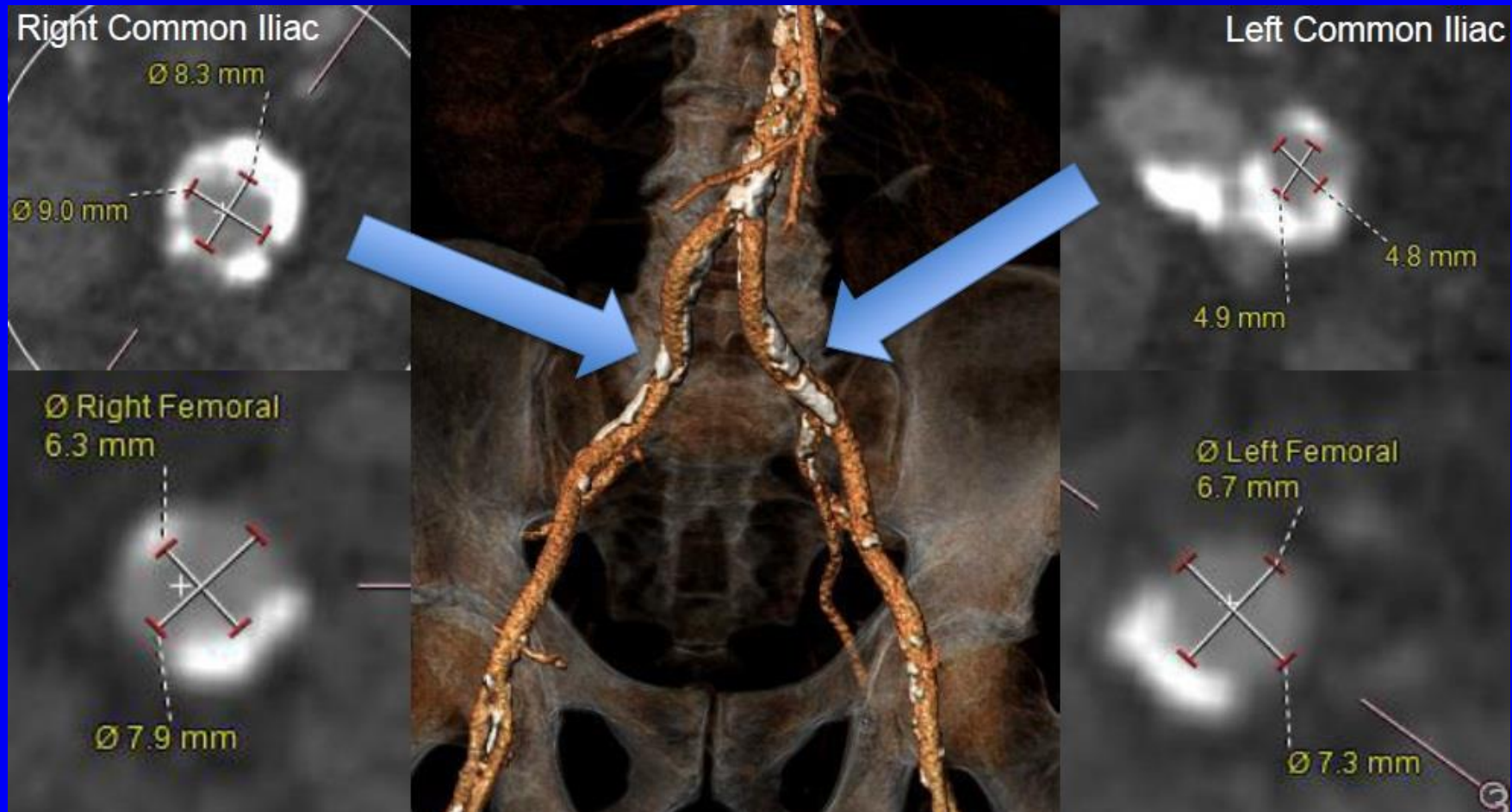
Evaluación Vascular Periférica

Grado de Tortuosidad y Calibre



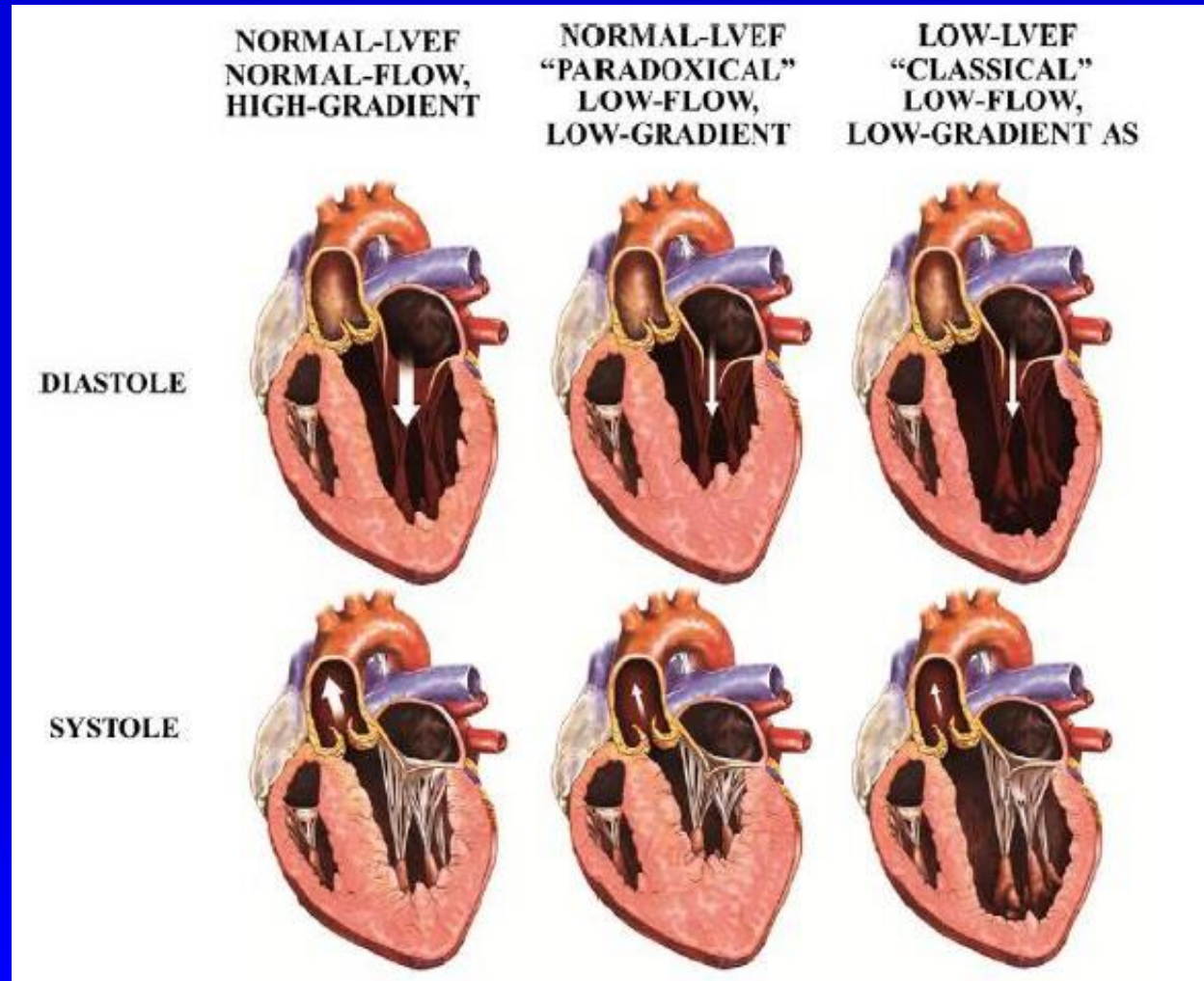
Evaluación Vascular periférica

Grado de Calcificación y Calibre



Estenosis Aórtica Grave

Bajo Flujo – Bajo Gradiente



Estenosis Aórtica Grave

Clasificación

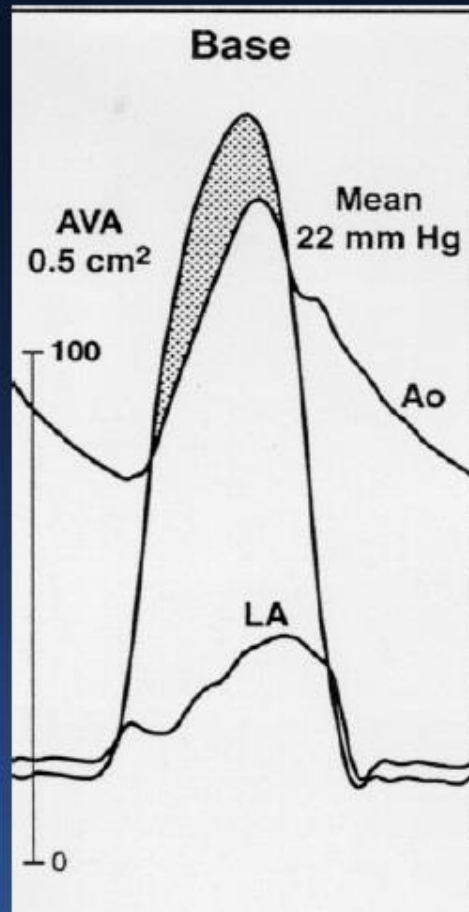
D: Symptomatic severe AS

D1	Symptomatic severe high-gradient AS	<ul style="list-style-type: none"> Severe leaflet calcification or congenital stenosis with severely reduced leaflet opening 	<ul style="list-style-type: none"> Aortic $V_{max} \geq 4$ m/s or mean $\Delta P \geq 40$ mm Hg AVA typically ≤ 1.0 cm² (or AVAi ≤ 0.6 cm²/m²) but may be larger with mixed AS/AR 	<ul style="list-style-type: none"> LV diastolic dysfunction LV hypertrophy Pulmonary hypertension may be present 	<ul style="list-style-type: none"> Exertional dyspnea or decreased exercise tolerance Exertional angina Exertional syncope or presyncope
D2	Symptomatic severe low-flow/low-gradient AS with reduced LVEF	<ul style="list-style-type: none"> Severe leaflet calcification with severely reduced leaflet motion 	<ul style="list-style-type: none"> AVA ≤ 1.0 cm² with resting aortic $V_{max} < 4$ m/s or mean $\Delta P < 40$ mm Hg <u>Dobutamine stress echocardiography shows AVA ≤ 1.0 cm² with $V_{max} \geq 4$ m/s at any flow rate</u> 	<ul style="list-style-type: none"> LV diastolic dysfunction LV hypertrophy LVEF $< 50\%$ 	<ul style="list-style-type: none"> HF Angina Syncope or presyncope
D3	Symptomatic severe low-gradient AS with normal LVEF or paradoxical low-flow severe AS	<ul style="list-style-type: none"> Severe leaflet calcification with severely reduced leaflet motion 	<ul style="list-style-type: none"> AVA ≤ 1.0 cm² with aortic $V_{max} < 4$ m/s or mean $\Delta P < 40$ mm Hg Indexed AVA ≤ 0.6 cm²/m² and Stroke volume index < 35 mL/m² Measured when patient is normotensive (systolic BP < 140 mm Hg) 	<ul style="list-style-type: none"> Increased LV relative wall thickness Small LV chamber with low stroke volume Restrictive diastolic filling LVEF $\geq 50\%$ 	<ul style="list-style-type: none"> HF Angina Syncope or presyncope

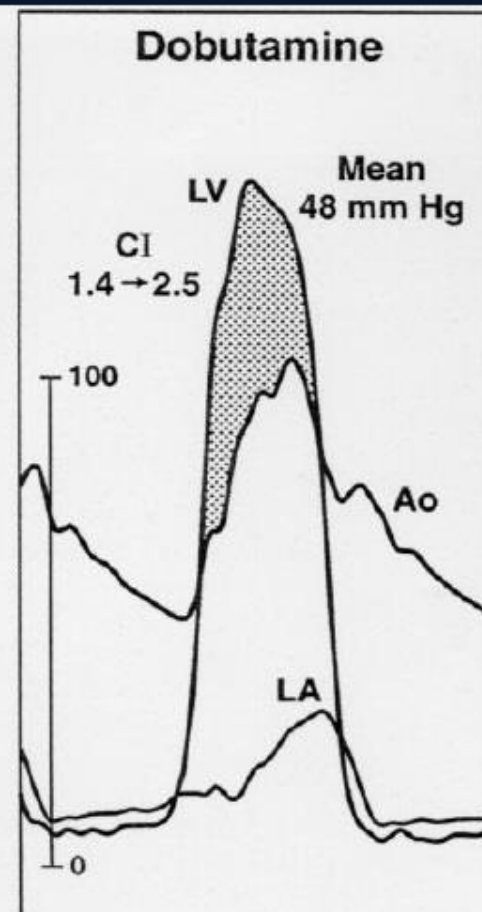
Low Flow - Low Gradient

$FE \leq 50\%$

**Gradient
22 mmHg**



**Gradient
48 mmHg**



Implante Transcateter Valvar Aórtico

Conclusiones

- 1) Lo ITVA es una alternativa atractiva de tratamiento para los pacientes portadores de EAo calcificada grave con alto riesgo quirúrgico o inoperables**
- 2) La selección criteriosa previa es de fundamental importancia para el éxito del procedimiento y reducción de las complicaciones**
- 3) La MSCT demuestra ser un examen extrema importancia permitiendo una evaluación global y con alta precision de estos pacientes**