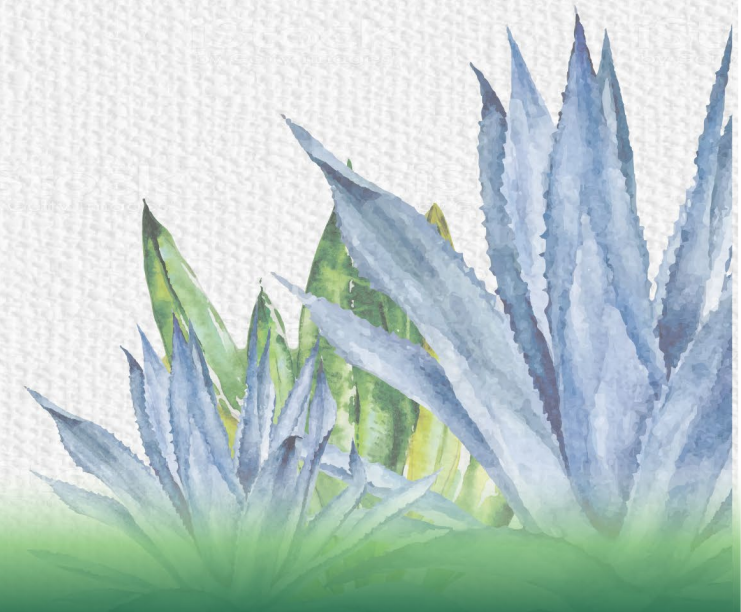


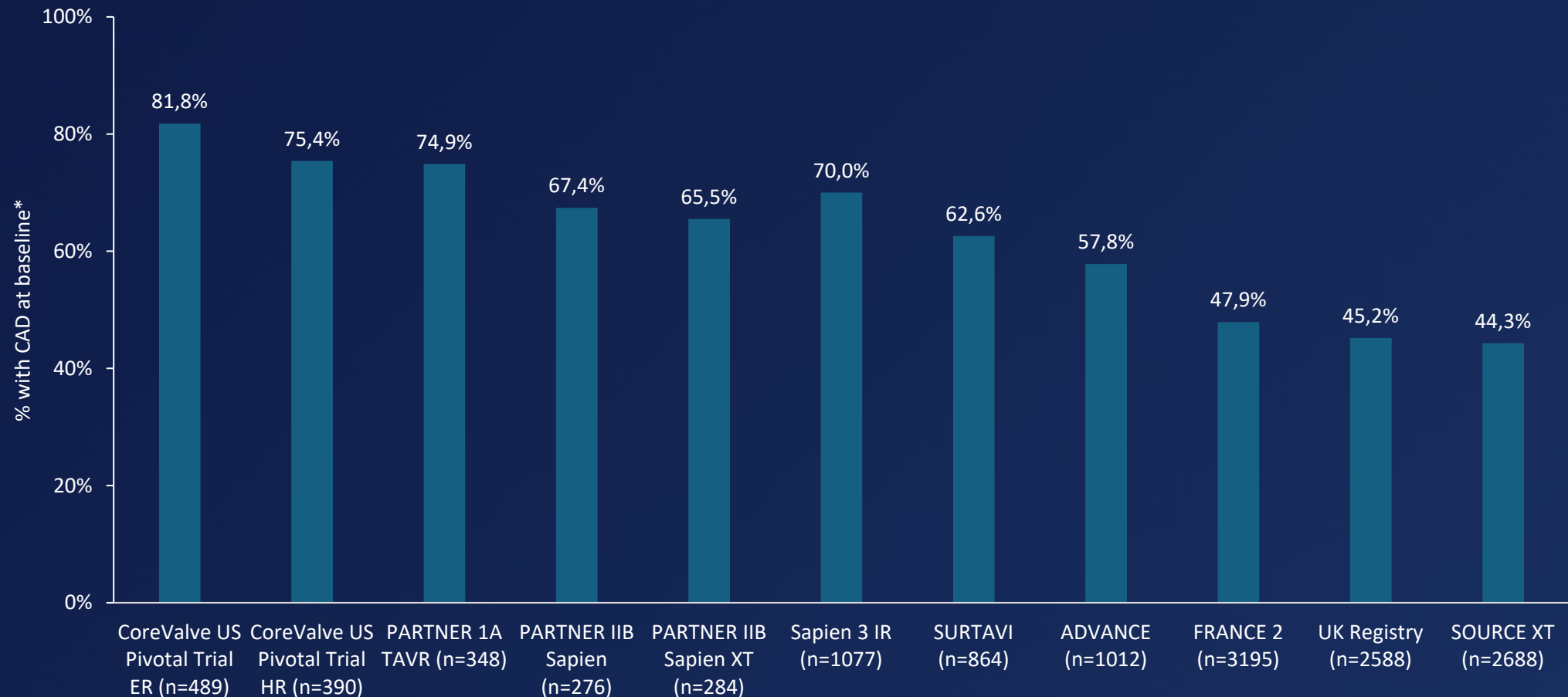
My best strategy in TAVI with CAD: When to treat

Flavio L. Ribichini
Universidad de Verona
Italia



Coronary Artery Disease: Prevalence in TAVI Patients

- Coronary artery disease is highly prevalent in the TAVR population, possibly affecting up to 80% of the cohort



References from left to right: 1. Popma JJ, et al. J Am Coll Cardiol 2014;63:1972-1981. 2. Adams, et al., N Engl J Med 2014; 370: 1790-8; 3. Smith GR, et al. NEJM 2011;364:2187-98. 4&5. Webb, et. al. JACC Cardiovasc Interv 2015;8:1797-806; 6. Thourani VH, et al. Lancet 2016; 387:2218-25. 7. Reardon MJ, et al. NEJM 2017;376:1321-31. 8. Bosmans J, et al. JACC 2015;66:209-17. 9. Gilard M, et al. NEJM 2012;366:1705-15. 10. . Snow TM, et al. Int J Cardiol 2015;199:253-60; 11. Schymik G, et al. European experience with the second-generation Edwards SAPIEN XT transcatheter heart valve in patients with severe aortic stenosis: 1-year outcomes from the SOURCE XT Registry JACC Cardiovasc Interv 2015;8:657-69. CoreValve is a trademark of Medtronic. Third party brands are trademarks of their respective owners.

2017 ESC/EACTS Guidelines for the management of valvular heart disease

2021 ESC/EACTS Guidelines for the management of valvular heart disease

These are all «C»
LOE

Indications for myocardial revascularization

CABG is recommended in patients with a primary indication for aortic/mitral valve surgery and coronary artery diameter stenosis $\geq 70\%$.^e

I

C

CABG should be considered in patients with a primary indication for aortic/mitral valve surgery and coronary artery diameter stenosis $\geq 50\text{--}70\%$.

IIa

C

PCI should be considered in patients with a primary indication to undergo TAVI and coronary artery diameter stenosis $>70\%$ in proximal segments.

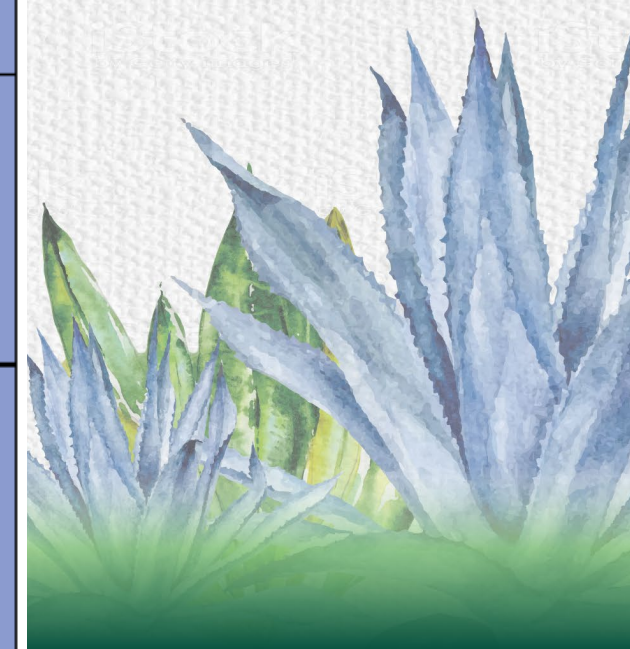
IIa

C

PCI should be considered in patients with a primary indication to undergo transcatheter mitral valve interventions and coronary artery diameter stenosis $>70\%$ in proximal segments.

IIa

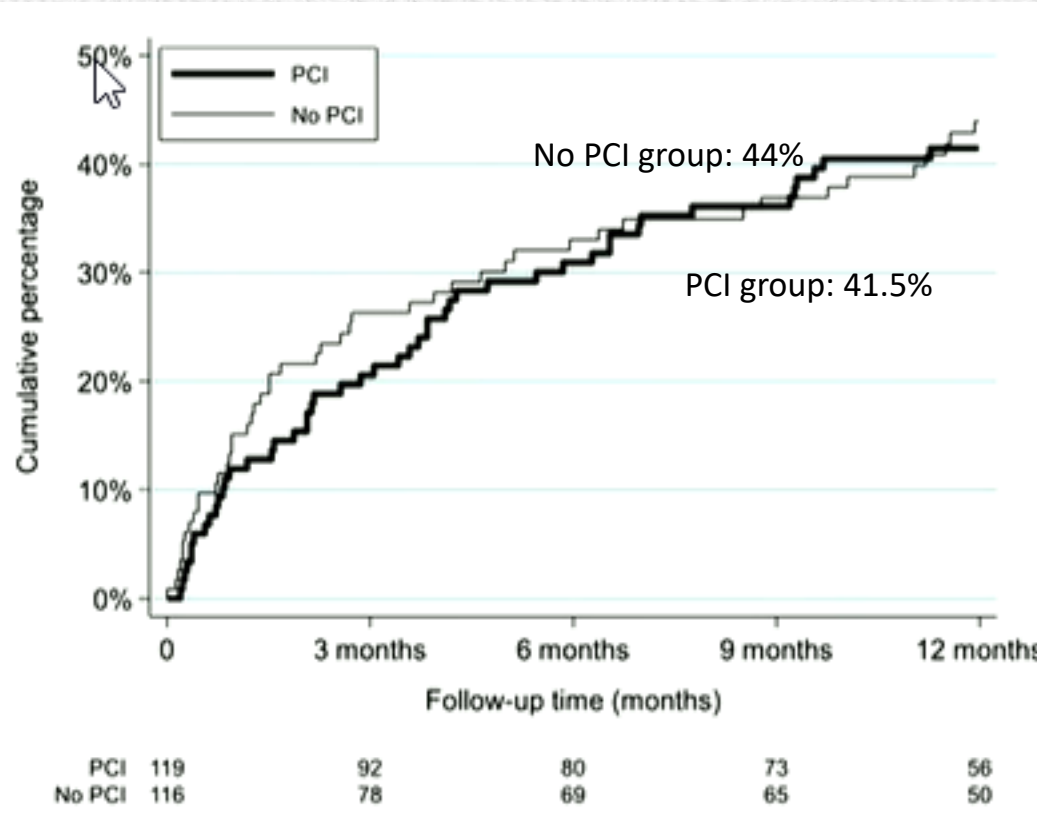
C



ACTIVATION: PCI versus no PCI in patients with AS & CAD undergoing TAVI

Primary Endpoint:
Comparable rates of death & cardiac rehospitalization at 1 year

Limitation:
RCT prematurely stopped due to slow recruitment



	PCI	No PCI	Total
N of patients	119	116	235
Age (years)	83.6±5.0	84.3±5.0	83.9±5.0
Males (%)	58	65	61
STS-PROM	6.7±6.0	6.8±6.0	6.8±7.7
Number of vessels treated			
1	85 (71.4%)		
2	29 (24.4)		
≥3	3 (2.5%)		

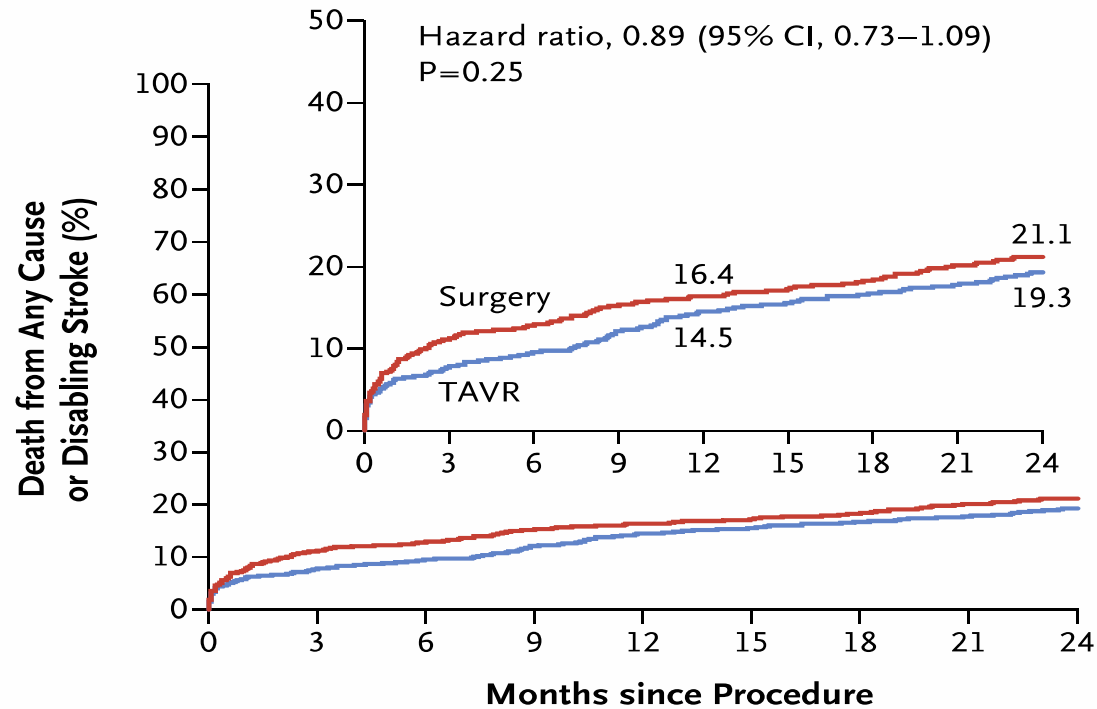
**Higher bleeding in PCI vs no PCI group
(44.5% vs 28.4%, p=0.02)**

Redwood S. Presented at: PCR Valves 2020. November 22, 2020

PARTNER 2 Trial: intermediate risk patients

TAVI 30 days death+ stroke= 4%

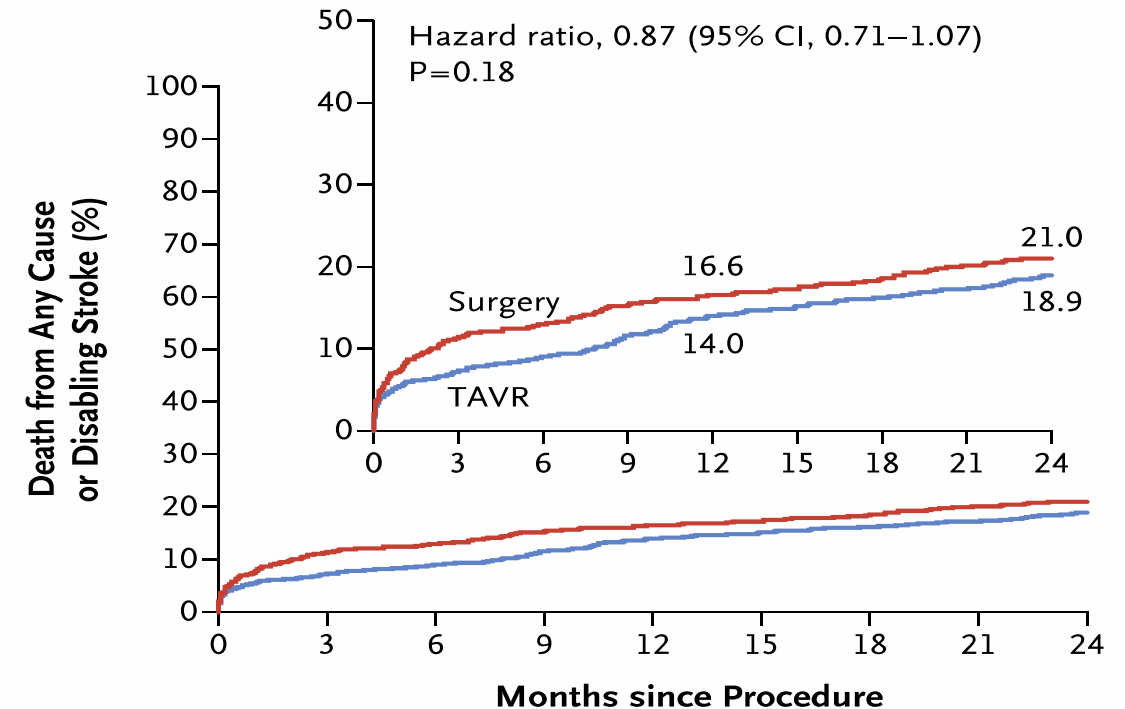
A Intention-to-Treat Population



No. at Risk

	1011	918	901	870	842	825	811	801	774
TAVR									
Surgery	1021	838	812	783	770	747	735	717	695

B As-Treated Population



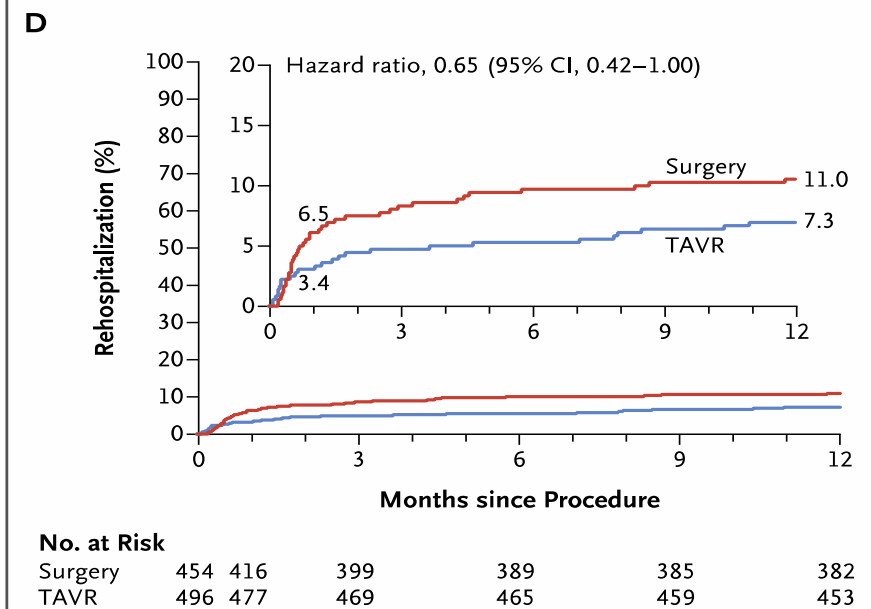
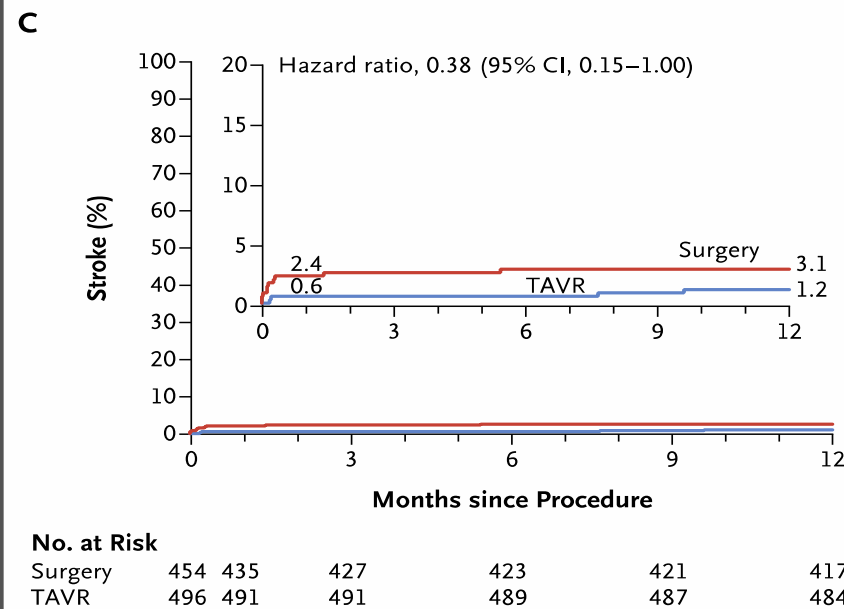
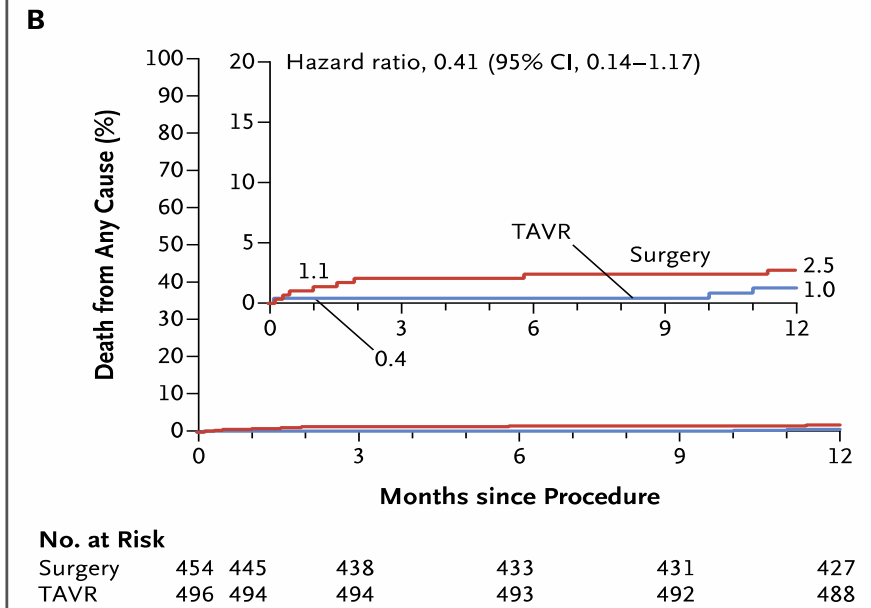
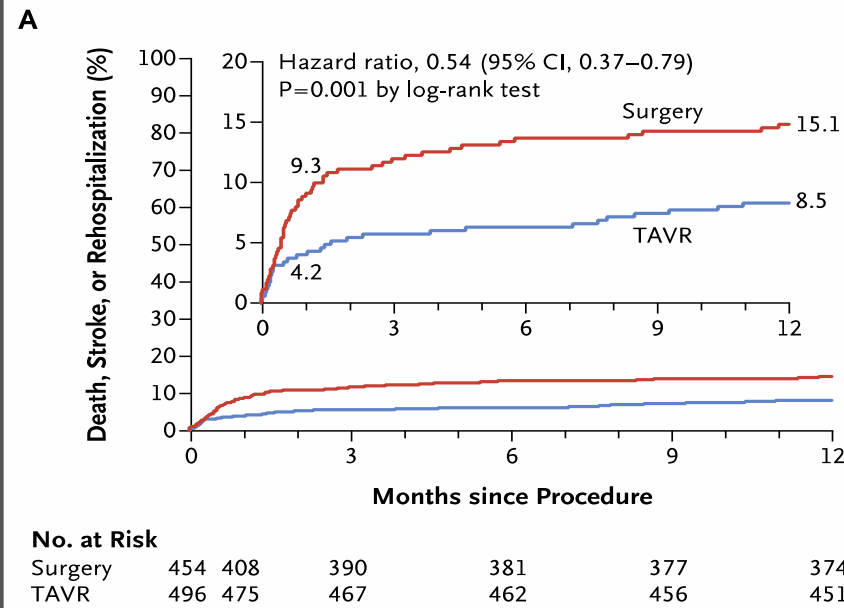
No. at Risk

	994	917	900	870	842	825	811	801	774
TAVR									
Surgery	944	826	807	779	766	743	731	715	694

PARTNER 3 Trial: **low risk** patients (NEJM 2017)

TAVI
One month

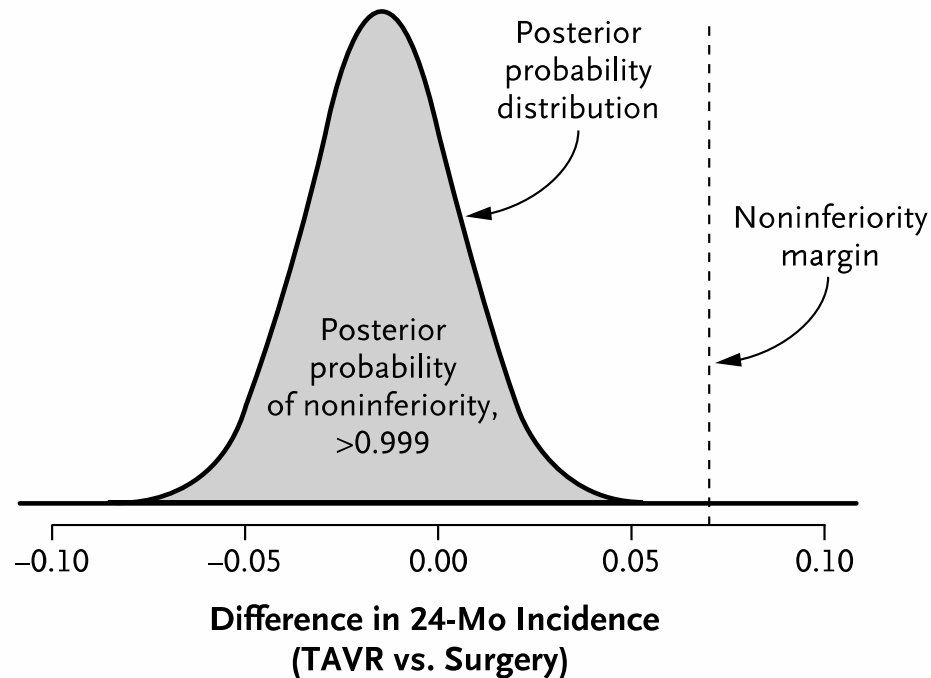
Death 0.4%
Stroke 0.6%



SURTA VI: intermediate risk patients

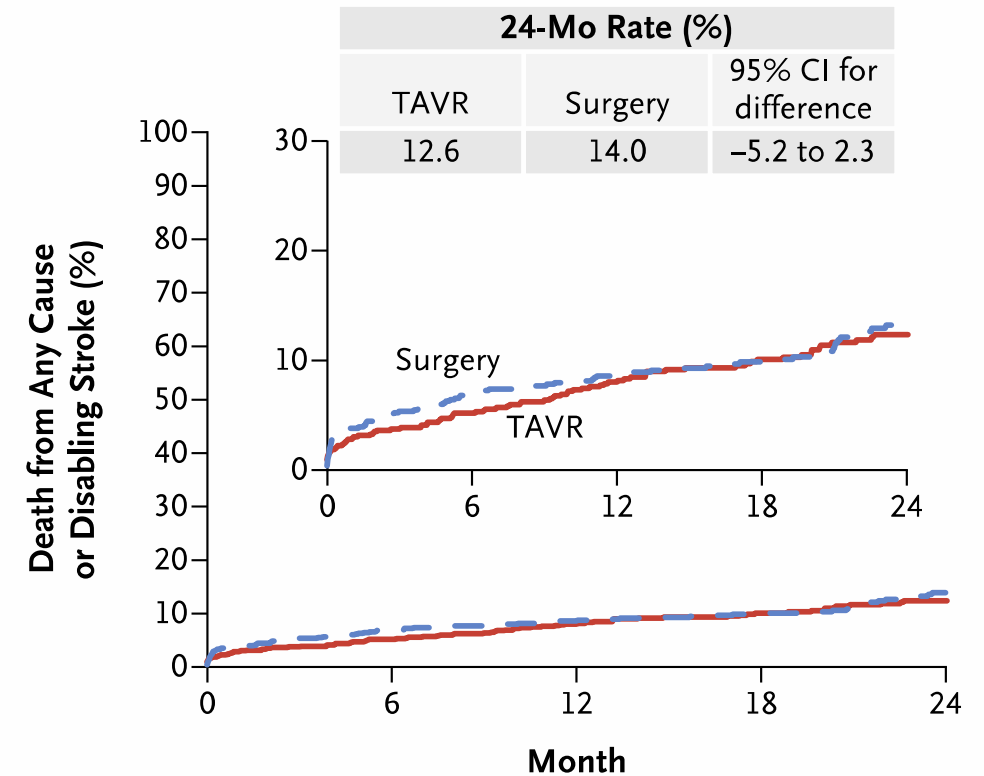
TAVI 30 days death+ stroke= 3%

A Noninferiority Margin of TAVR



TAVR Posterior Median	Surgery Posterior Median	Difference Posterior Median
% (95% CI)		
12.6 (10.2 to 15.3)	14.0 (11.4 to 17.0)	-1.4 (-5.2 to 2.3)

B Primary Outcome



No. at Risk

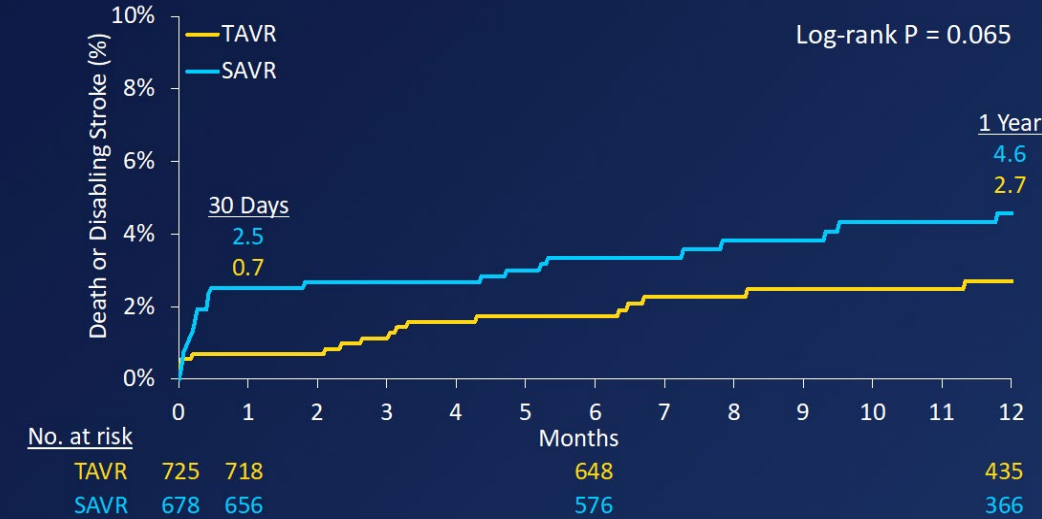
	0	6	12	18	24
TAVR	864	755	612	456	272
Surgery	796	674	555	407	241

Evolut **low risk** trial (NEJM 2017)

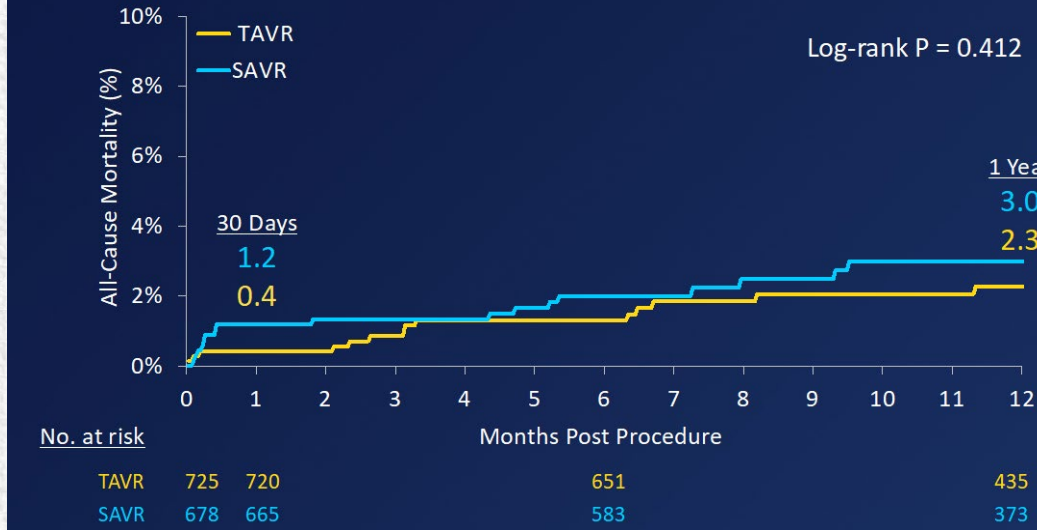
TAVI
One month

Death 0.4%
Stroke 0.3%

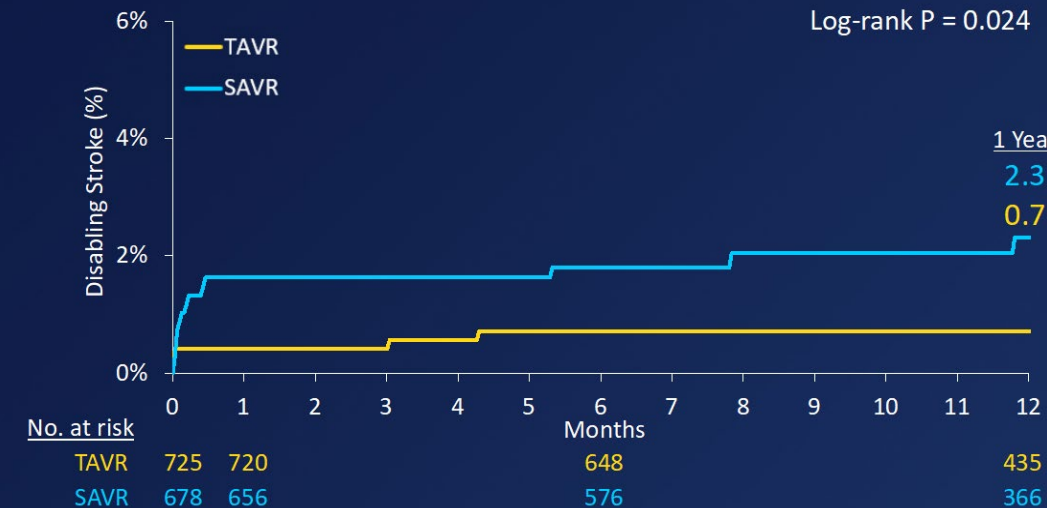
K-M All-Cause Mortality or Disabling Stroke at 1 Year



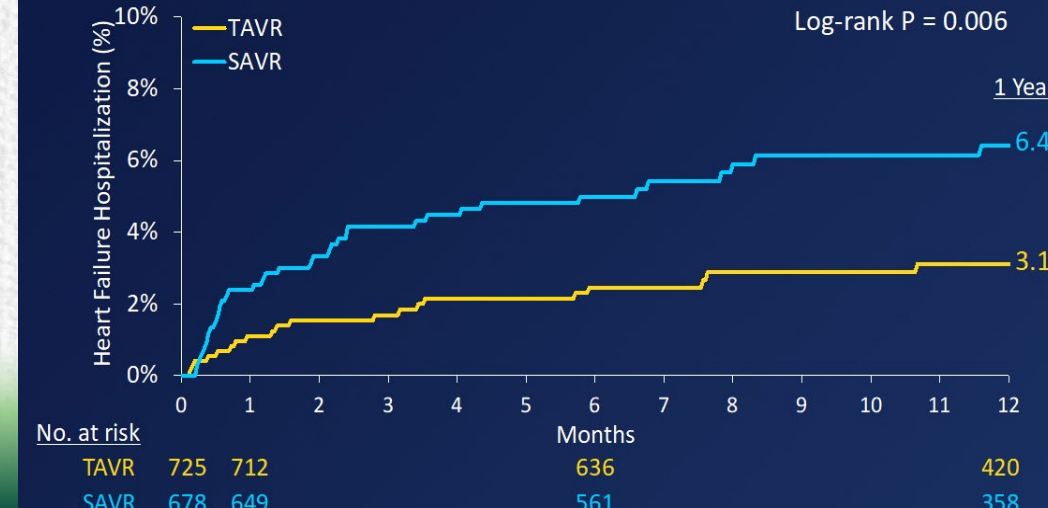
K-M Rates of All-Cause Mortality at 1 Year



K-M Disabling Stroke at 1 Year



K-M Heart Failure Hospitalization at 1 Year



CAD Treatment Timing

TAVR vs SAVR in Intermediate-risk Patients

- The CoreValve™ SURTAVI trial was a randomized trial of TAVR vs. SAVR in intermediate surgical risk patients.¹
- A sub-analysis on TAVR+PCI (concurrent and staged) vs. SAVR+CABG found comparable results between the two groups. However, patients with TAVR + PCI had shorter procedure time, hospital stay, ICU stay, and less AKI.

As-Treated Procedural and 30-Day Outcomes

% or mean ± SD	Revascularization		P
	TAVR + PCI	SAVR + CABG	
	N=128	N=176	
Index procedure time, min	53 ± 35	249 ± 71	<0.01
Length of index hospital stay, days	7 ± 5	11 ± 8	<0.01
Length of index ICU stay, hours	52 ± 46	78 ± 84	<0.01
All-cause mortality or disabling stroke*	6.3	TAVI alone 3%	0.37
All-cause mortality*	4.7	2.3	0.25
Disabling stroke*	2.3	2.3	0.96
Acute kidney injury*	7.8	19.9	<0.01

¹Sondergaard et al., presented at TCT 2017. CoreValve is a trademark of Medtronic.

Procedural Characteristics and Late Outcomes of Percutaneous Coronary Intervention in the Workup Pre-TAVR

Laurent Faroux, MD,^a Francis Quentin Fischer, MD,^e Pierre Victoria Vilalta, MD,^j Alberto Anthony Matta, MD,^b Antonio Eduard Rodenas-Alesina, MD, Isaac Pascual, MD,^k Pedro Per David del Val, MD,^a Thomas

**30-d death
& stroke
na%**

unoz-Garcia, MD,^c Luis Nombela-Franco, MD,^d Gabriela Veiga, MD,^h Enrique Gutiérrez, MD,ⁱ Lluís Asmarats, MD,^m Henrique B. Ribeiro, MD,ⁿ Chijo, MD,^g Marina Urena, MD,^c Damien Metz, MD,^f ez, MD,^h Eduard Fernandez-Nofrerias, MD,^j i, MD,^m Diego Carter Campanha-Borges, MD,ⁿ MD^a



ORIGINAL ARTICLE

Surgical or Transcatheter Aortic-Valve Replacement in Intermediate-Risk Patients

M.J. Reardon, N.M. V M. Mumtaz, D.H. Adam J. Heiser, R. Lange, V S. Windecker, S.J. Y H. Nguyen, Y. Chang

**30-d death
& stroke 3%**

L. Søndergaard, netcuti, T. Gleason, a, M. Williams, Conte, E. Vang, A.P. Kappetein,

VALVULAR AND STRUCTURAL HEART DISEASES

Impact of coronary artery disease and percutaneous coronary intervention in women undergoing transcatheter aortic valve replacement: From the WIN-TAVI registry

Paul Guedeney MD, Didier Tchétché MD, Anna Sonia Petronio MD, Julia Sartori PhD, Thierry Lefèvre MD, Patrizia Presbitero MD, Piera Capranzano MD

**30-d death
& stroke
8%**

Circulation

ORIGINAL RESEARCH ARTICLE

Comparison of a Complete Percutaneous Versus Surgical Approach to Aortic Valve Replacement and Revascularization in Patients at Intermediate Surgical Risk Results From the Randomized SURTAVI Trial

**30-d death
& stroke
6.3%**

SYSTEMATIC REVIEW AND META-ANALYSIS

Transcatheter Aortic Valve Implantation With or Without Percutaneous Coronary Artery Revascularization: A Systematic Review and Meta-Analysis

Rafail A. Kotronias, MBChB, MSc; Chun Shing Kwok, MBBS, MSc; Sudhakar Ge Ludman, MD, FRCP, FESC; Jonathan N. Townend, MD, FRCP; Sagar N. Doshi, MB Philippe Gagnéux, MD; Howard C. Herrmann, MD, FACC, MSCAI; Mamas A. Ma

**30-d death
& stroke
9%**



Impact of coronary artery disease and percutaneous coronary intervention on outcomes in patients with severe aortic stenosis undergoing transcatheter aortic valve implantation

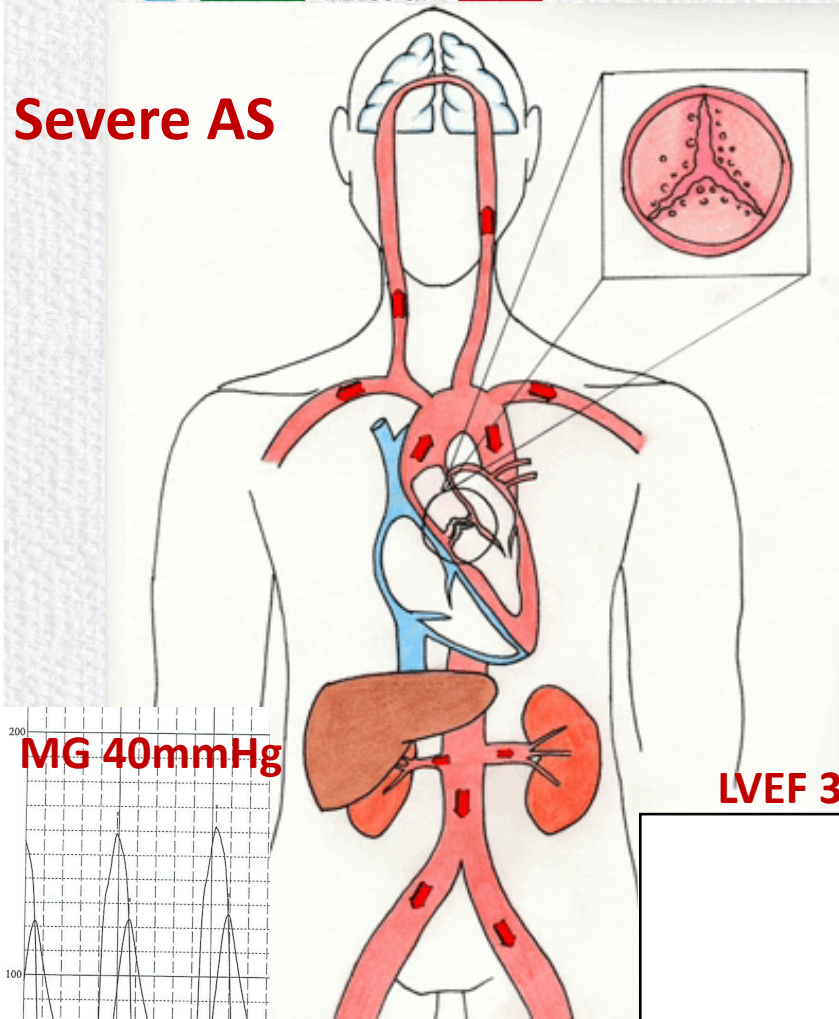
Peter Wenaweser^{1*}, MD; Thomas Pilgrim², MD; Stefan Stortecky¹, MD; Christoph Huber², MD; Ahmed A. Khatib¹, MD; Lutz Buellesfeld¹, MD; Steffen Gloekler¹, MD; Bernhard Meier¹, MD; Stephan Windecker¹, MD

**30-d death
& stroke
13%**

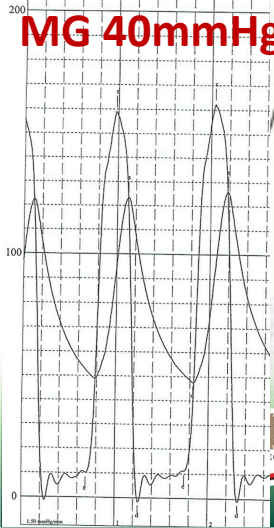
CI= 3.0 l/min/m²

CI= 4.0 l/min/m²

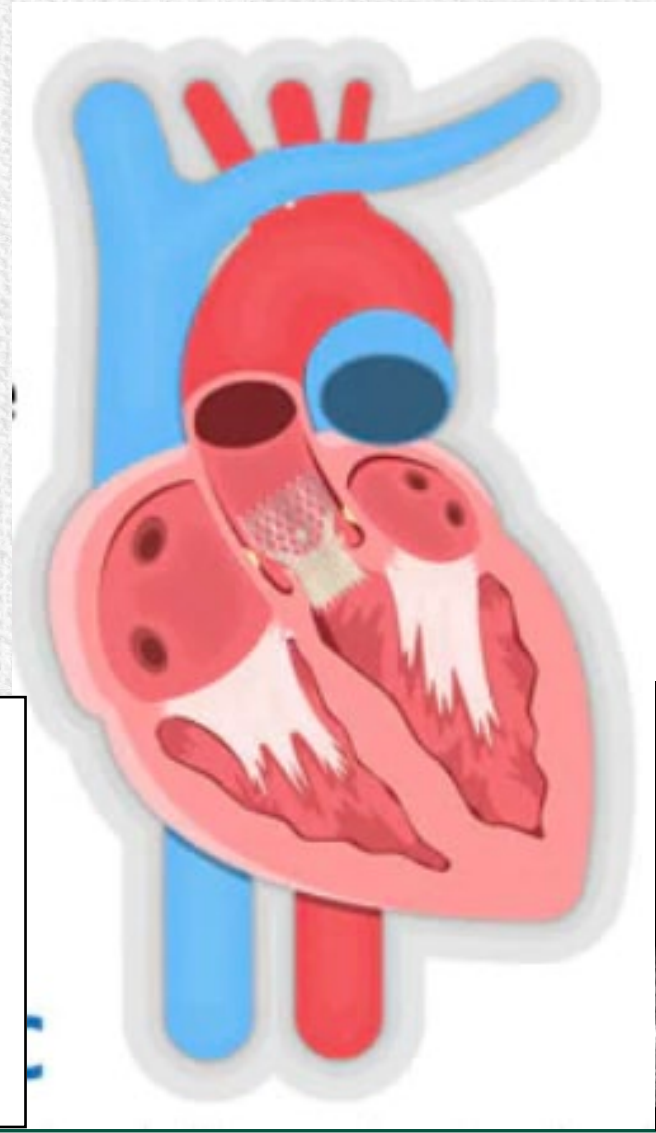
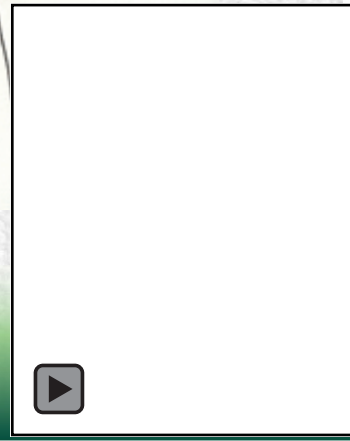
Severe AS



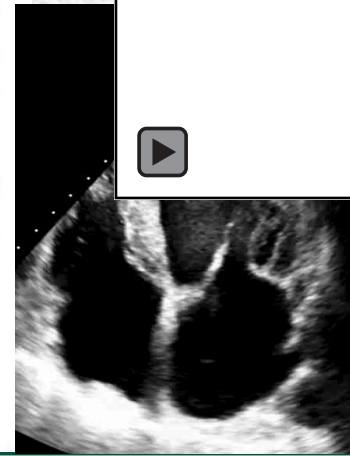
MG 40mmHg



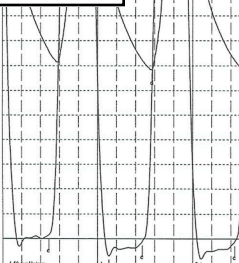
LVEF 35%



LV



mmHg



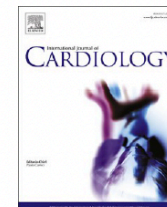


ELSEVIER

Contents lists available at [ScienceDirect](http://www.sciencedirect.com)

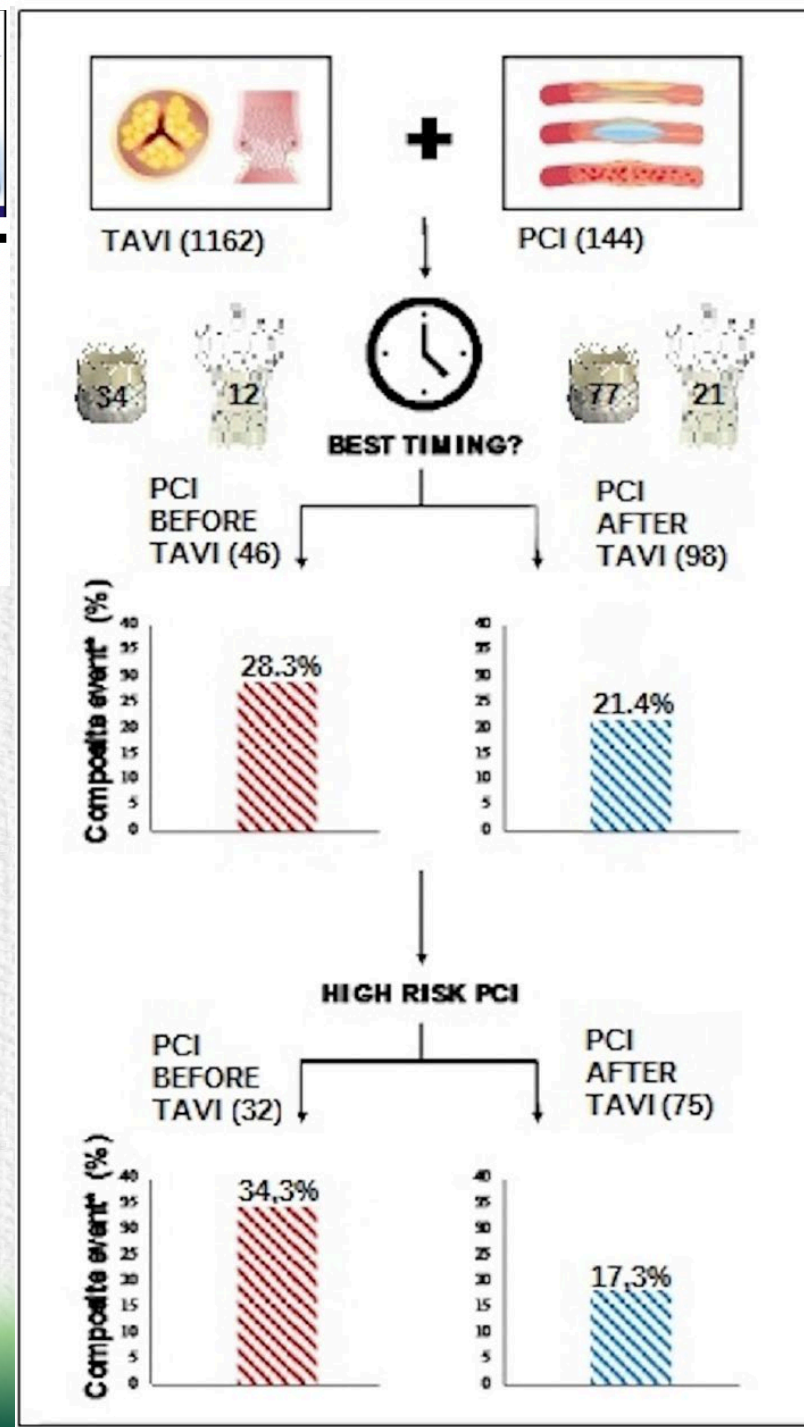
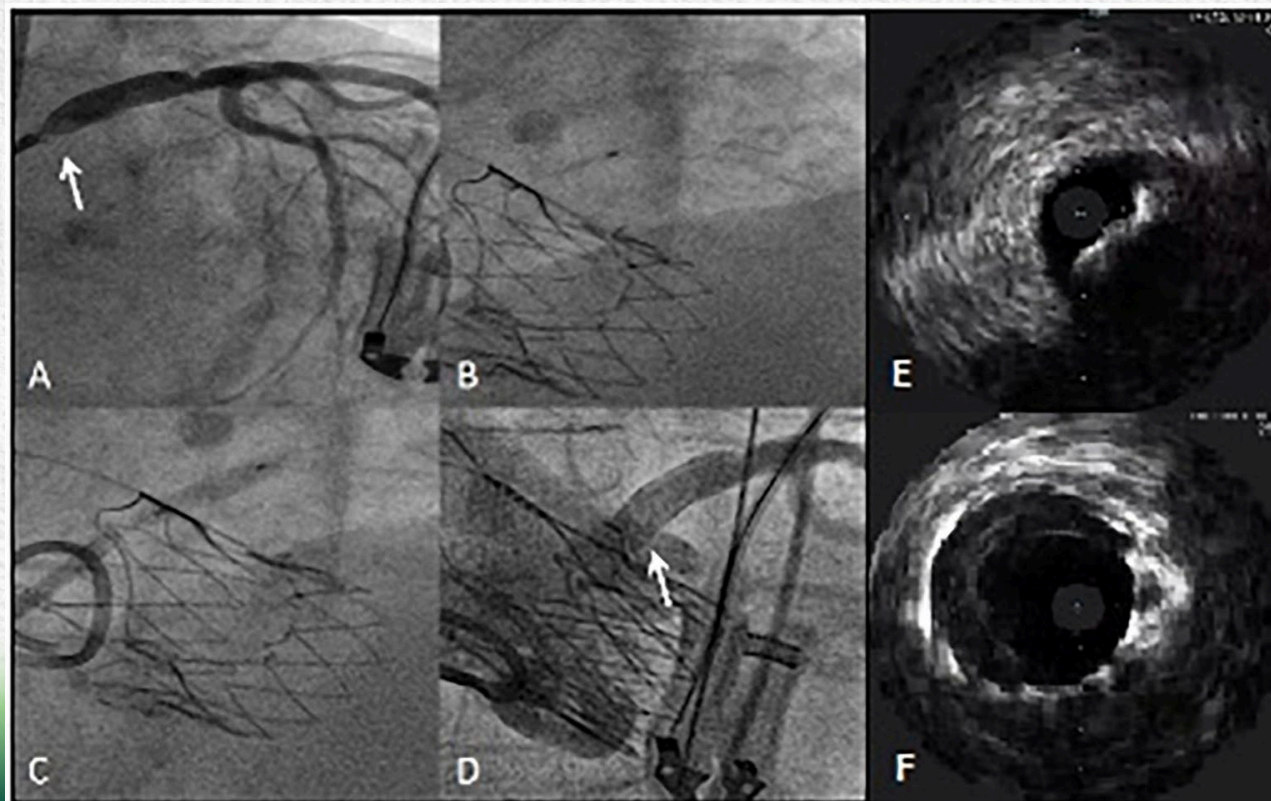
International Journal of Cardiology

journal homepage: www.elsevier.com/locate/ijcard



Optimal timing for percutaneous coronary intervention in patients undergoing transcatheter aortic valve implantation

Mattia Lunardi^{a,1}, Gabriele Venturi^{a,1}, Paolo Alberto Del Sole^a, Alessandro Ruzzarin^a, Andrea Mainardi^a, Michele Pighi^a, Gabriele Pesarini^a, Roberto Scarsini^a, Domenico Tavella^a, Leonardo Götting^b, Flavio Luciano Ribichini^{a,*}

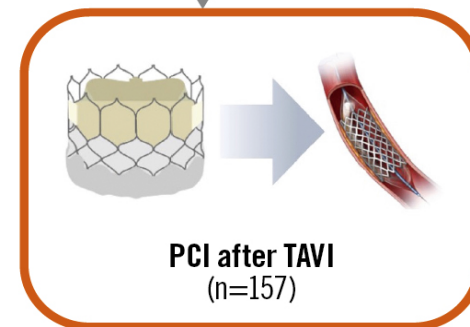
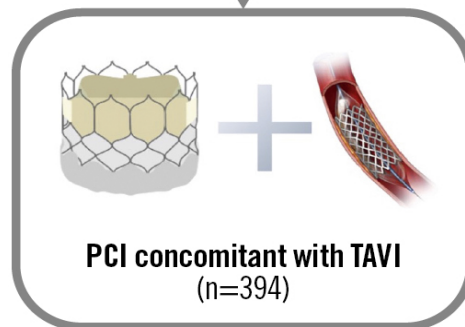
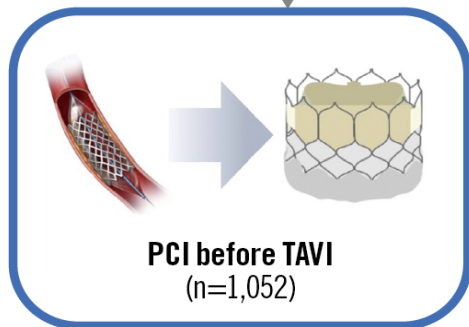


TAVI one month: death 2.1% stroke 0.9% (3%)

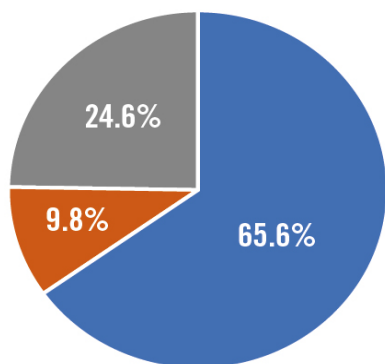
	PCI pre-TAVI (46)	PCI post-TAVI (98)	p.	High-risk PCI pre TAVI (32)	High-risk PCI post-TAVI (75)	p.
<u>Procedural complications</u>						
Procedural death	0 (0.0%)	0 (0.0%)	-	0 (0.0%)	0 (0.0%)	-
Arrhythmias	0 (0.0%)	2 (2.0%)	1.000	0 (0.0%)	2 (2.6%)	0.987
Periprocedural MI	0 (0.0%)	4 (4.1%)	0.126	0 (0.0%)	0 (0.0%)	-
Cardiogenic shock	1 (2.2%)	4 (4.1%)	0.889	1 (3.1%)	2 (2.6%)	0.977
<u>In hospital events</u>						
Stroke	3 (6.5%)	0 (0.0%)	0.031	3 (9.3%)	0 (0.0%)	0.025
AKI	7 (15.2%)	11 (11.2%)	0.590	7 (21.9%)	7 (9.3%)	0.115
Major VC	4 (8.6%)	5 (5.1%)	0.467	4 (12.5%)	3 (4.0%)	0.193
Major Bleeding	4 (8.6%)	4 (4.1%)	0.267	3 (9.3%)	3 (4.0%)	0.361
TLR or TVR	0 (0.0%)	0 (0.0%)	-	0 (0.0%)	0 (0.0%)	-
Cardiac death	2 (4.4%)	0 (0.0%)	0.098	2 (6.2%)	0 (0.0%)	0.081
In hospital complications	11 (23.9%)	13 (13.3%)	0.149	9 (28.1%)	9 (12.0)	0.051
Composite procedural and in hospital event	13 (28.3%)	21 (21.4%)	0.403	11 (34.4%)	13 (17.3%)	0.075

TAVI patients undergoing PCI for stable CAD in the REVASC-TAVI registry
(n=1,617)

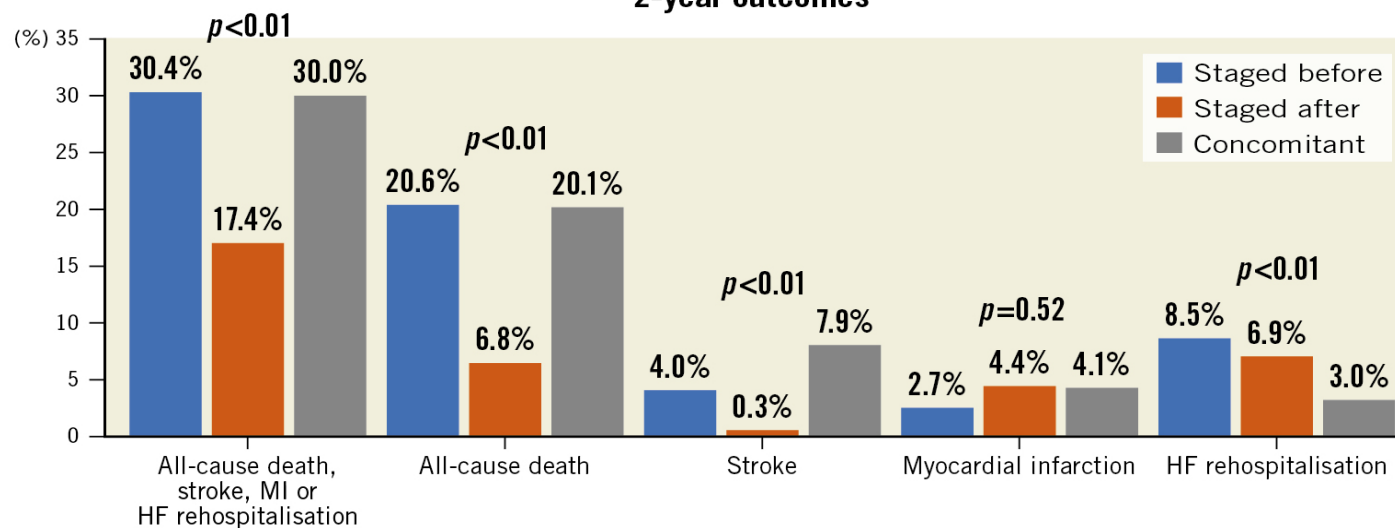
Patients excluded
Data of timing not available (n=7)
Unplanned PCI (n=7)



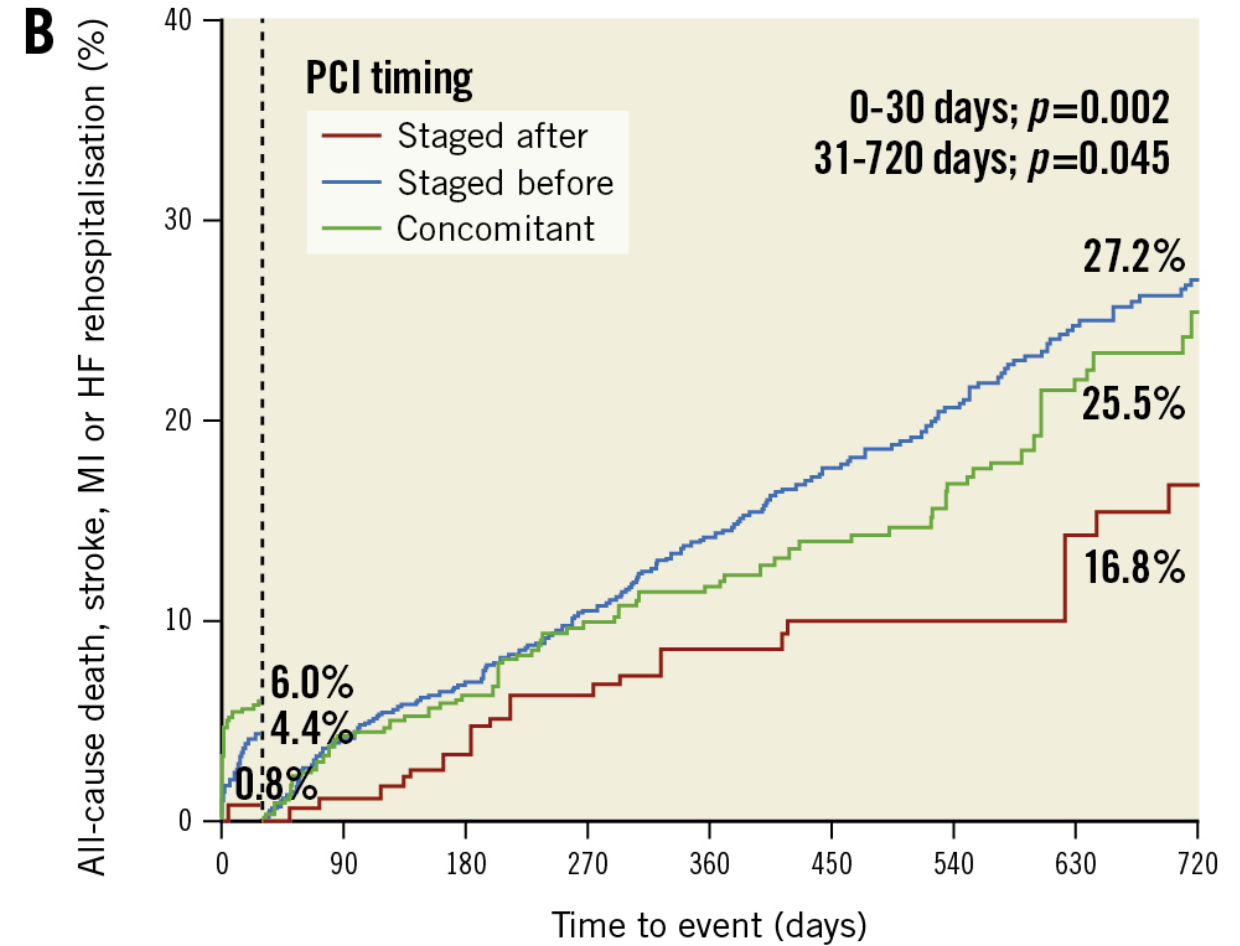
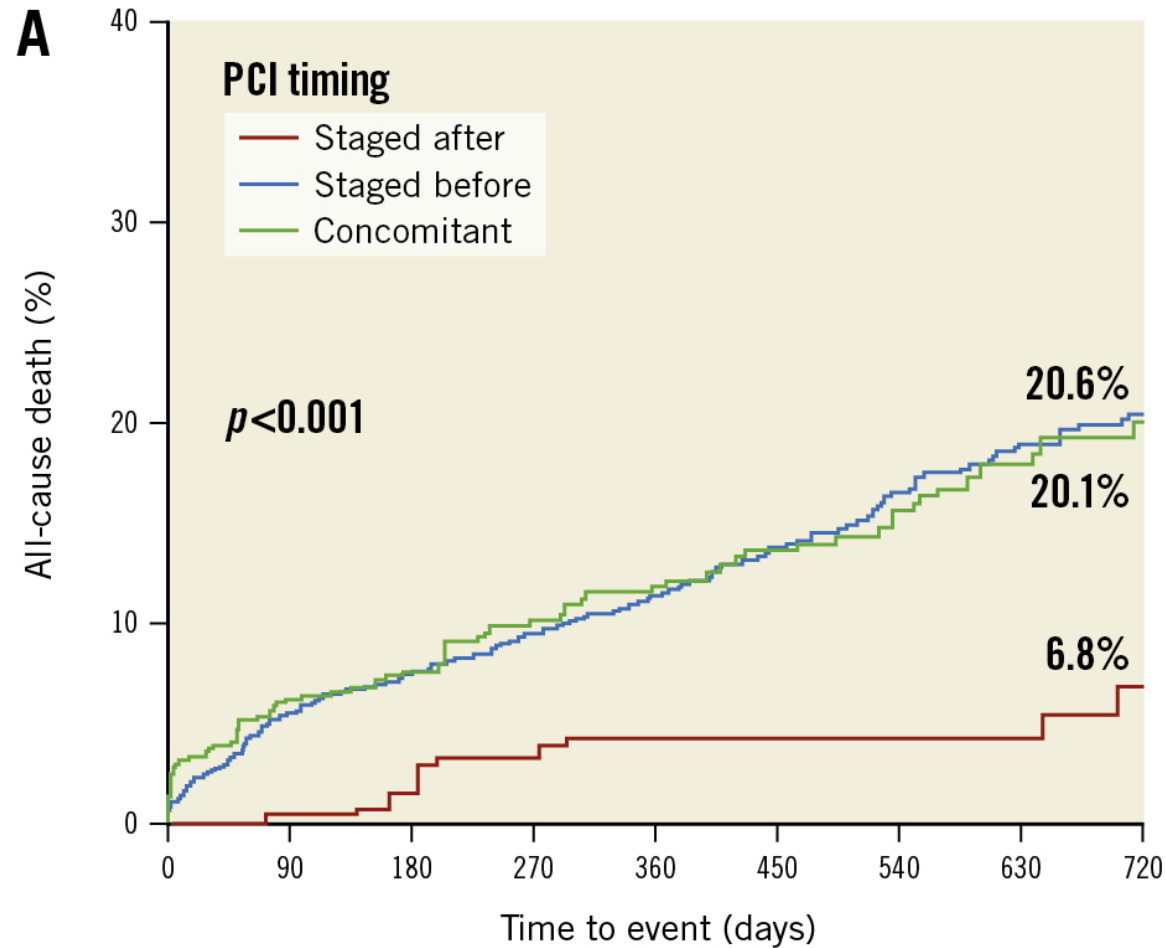
PCI timing distribution



2-year outcomes



All-cause death and MACEs according to PCI timing strategy in patients undergoing TAVI



TAVI PCI
NCT04310046
=986

- Angiography-guided PCI Before TAVI
- Angiography -guided PCI After TAVI (1-45/d)

Ongoing

Primary Endpoint:
Composite of all-cause **death**, myocardial **infarction**, , major **bleeding** and need for target vessel **revascularization** or re-hospitalization at 12 months

FAITAVI
NCT03360591
n=320

- Angiography-guided PCI diameter stenosis of $\geq 50\%$
- Physiology-guided PCI if FFR ≤ 0.80

Presented PCR 25

Primary Endpoint:
Composite of all-cause **death**, myocardial **infarction**, **stroke**, major **bleeding** and need for target vessel **revascularization** at 12 months

NOTION-3
NCT03058627
N=452

- Complete revasc with PCI
- Conservative management

Published

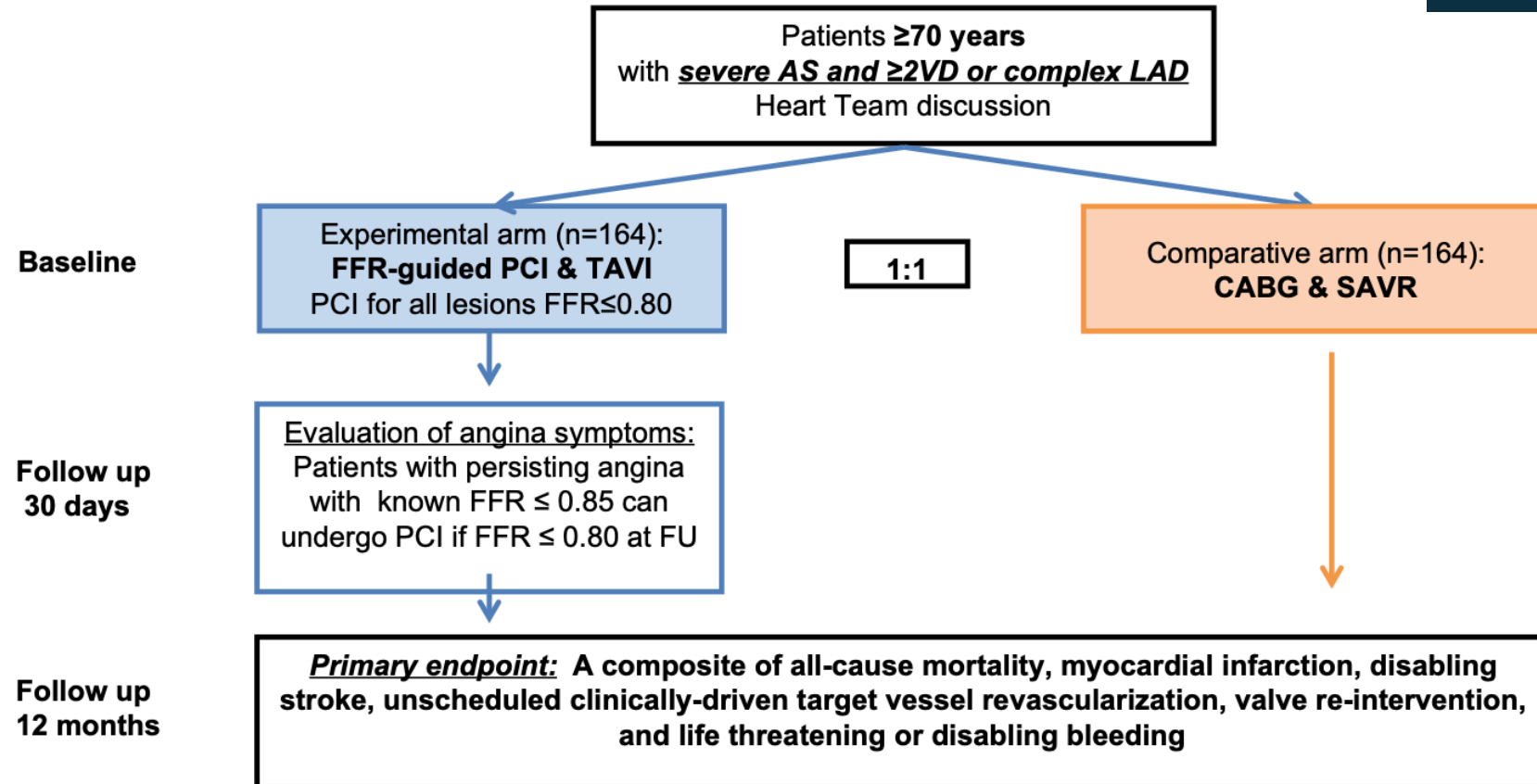
Primary Endpoint:
Composite of all-cause **mortality**, myocardial **infarction**, or urgent **revascularization** at 12 months

TCW
NCT03424941
N=328

- FFR-guided PCI and TAVI
- CABG and SAVR

Published

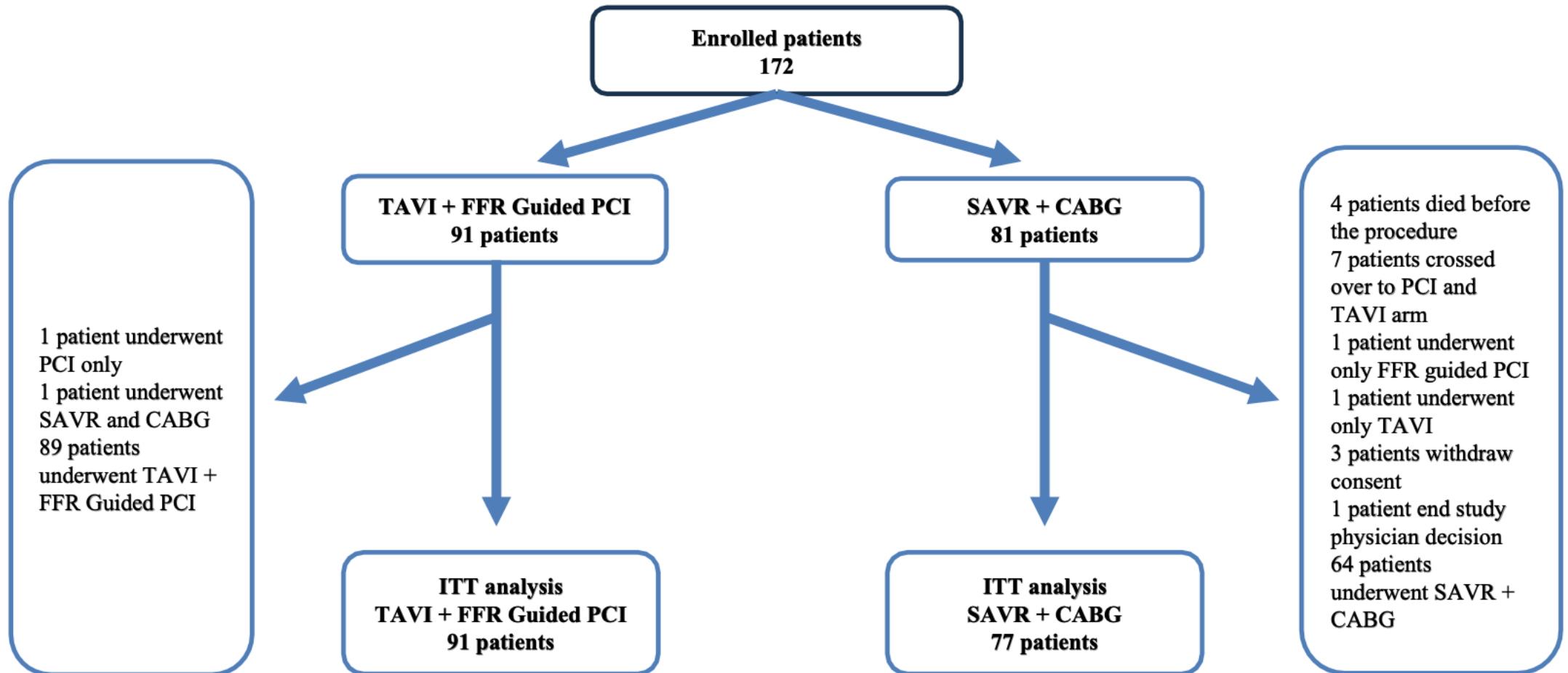
Primary Endpoint:
Composite of all-cause **mortality**, myocardial **infarction**, disabling **stroke**, unscheduled clinically-driven target vessel **revascularization**, valve **re-intervention**, and life threatening or disabling **bleeding** at one year



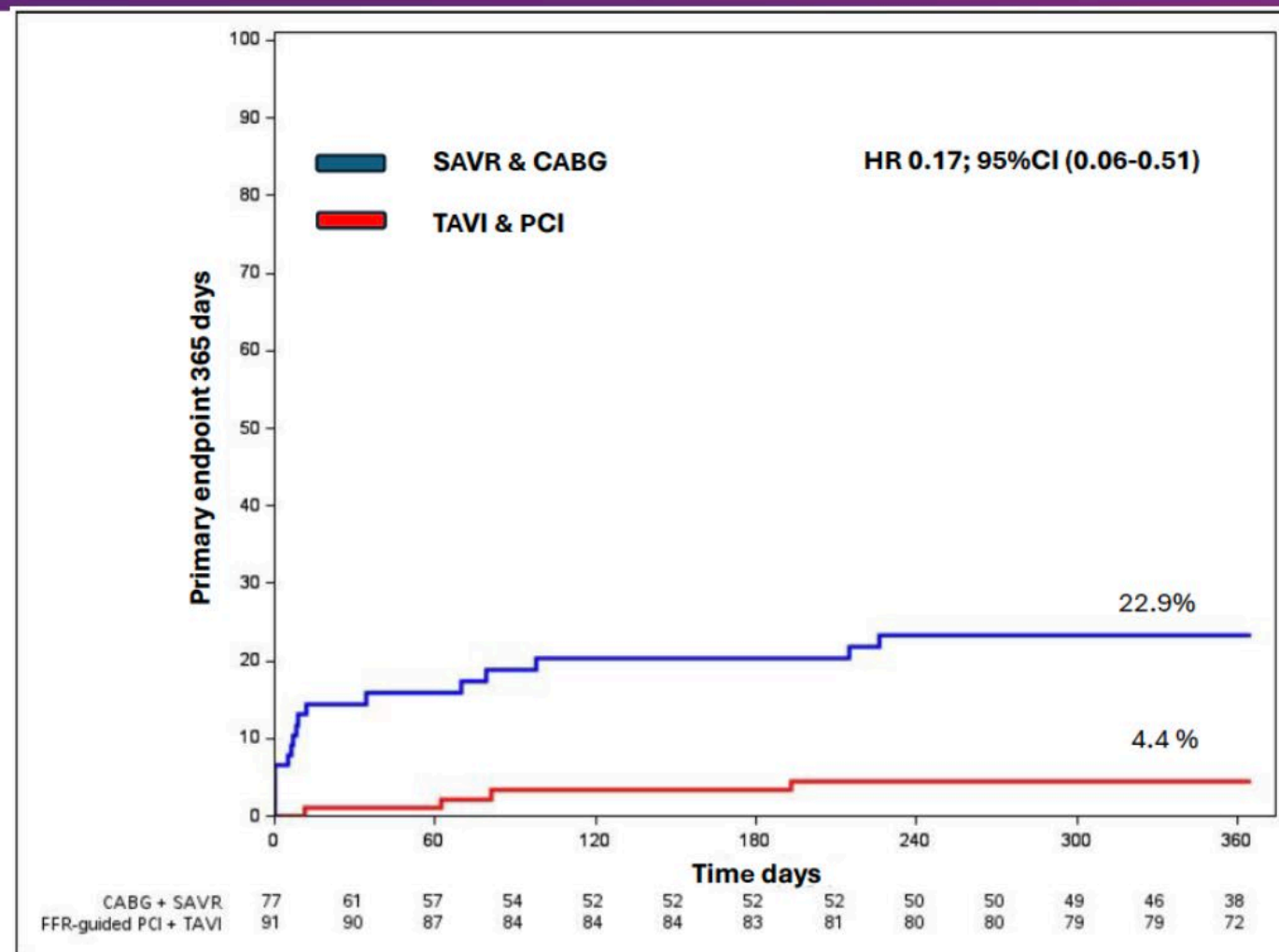
Study outcomes and definitions

- **Primary study endpoint:** a patient-oriented composite of ***all-cause mortality, myocardial infarction, disabling stroke, unscheduled clinically-driven target vessel revascularization, valve reintervention, and life threatening or disabling bleeding*** at 1 year
- **Major secondary endpoints:**
 - **MACE** a composite of ***cardiovascular mortality, all stroke, myocardial infarction, unscheduled coronary or valve re-intervention*** at 1 year
 - **All-cause mortality and all stroke** at 30 days and at 1 year

Patient Flow Chart



Primary Endpoint outcomes IT analysis



ORIGINAL ARTICLE

NOTION-3
NCT03058627
N=452

PCI in Patients Undergoing Transcatheter Aortic-Valve Implantation

J. Lønborg, R. Jabbari, M. Sabbah, K.T. Veien, M. Niemelä, P. Freeman, R. Linder, D. Ioanes, C.J. Terkelsen, O.A. Kajander, S. Koul, M. Savontaus, P. Karjalainen, A. Erglis, M. Minkinen, R. Sørensen, H.-H. Tilsted, L. Holmvang, G. Bieliauskas, J. Ellert, J. Piuholo, A. Eftekhari, O. Angerås, A. Rück, E.H. Christiansen, T. Jørgensen, B.T. Özbek, C. Glinge, L. Søndergaard, O. De Backer, and T. Engstrøm, for the NOTION-3 Study Group*

TAVI candidates with at least 1CAD stenosis >90% or FFR <0.80 pre-TAVI

1:1 Randomization

PCI

Conservative Tx

Primary Endpoint:

Composite of all-cause **mortality**, myocardial **infarction**, or urgent **revascularization** at 24 months



Table 1. Characteristics of the Patients at Baseline.*

Characteristic	PCI (N = 227)	Conservative Treatment (N = 228)
Median age (IQR) — yr	82 (78–85)	81 (78–85)
Female sex — no. (%)	73 (32)	75 (33)
Median body-mass index (IQR)†	26 (24–29)	26 (24–29)
Median STS-PROM score (IQR) — %‡	3 (2–4)	3 (2–4)
New York Heart Association class — no. (%)		
I	21 (9)	23 (10)
II	101 (44)	110 (48)
III	103 (45)	89 (39)
IV	2 (1)	6 (3)
Median creatinine level (IQR) — $\mu\text{mol/liter}$ §	89 (74–109)	87 (73–106)
Median aortic-valve area (IQR) — cm^2	1 (1–1)	1 (1–1)
Median peak aortic-valve gradient (IQR) — mm Hg	75 (64–90)	75 (66–90)
Median left ventricular ejection fraction (IQR) — %	60 (50–60)	60 (50–60)
Medical history — no. (%)		
History of PCI	28 (12)	37 (16)
History of myocardial infarction	14 (6)	24 (11)
History of CABG	7 (3)	10 (4)
Treatment of hypercholesterolemia	158 (70)	169 (74)
Diabetes mellitus	59 (26)	61 (27)
Treatment of hypertension	158 (70)	168 (74)
Current or former smoking	113 (50)	133 (58)
History of stroke or transient ischemic attack	48 (21)	47 (21)
Previous malignant disease	45 (20)	45 (20)
Chronic obstructive pulmonary disease	49 (22)	64 (28)
Atrial fibrillation	81 (36)	74 (32)
Peripheral artery disease	19 (8)	26 (11)

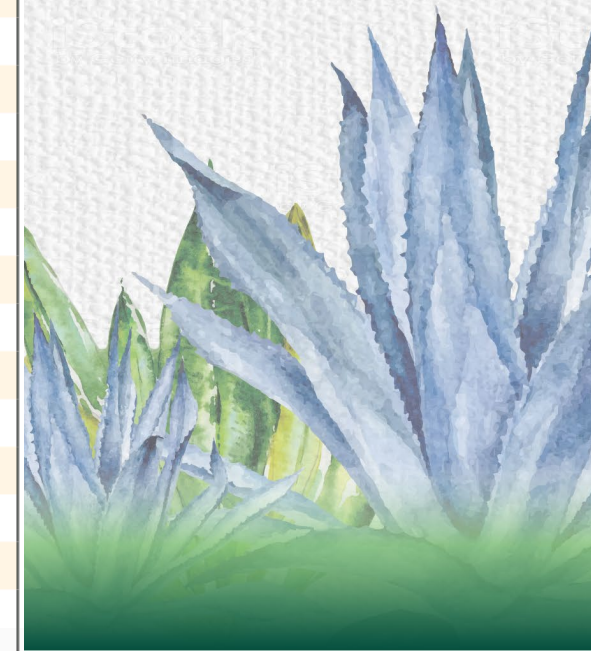
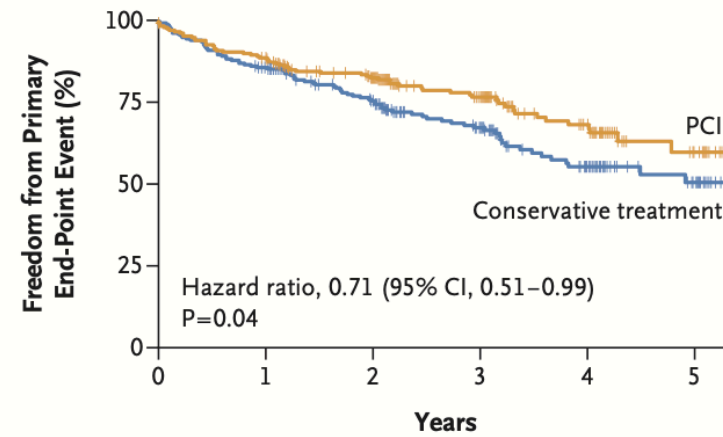


Table 2. Angiographic Findings and Characteristics of the PCI and TAVI Procedures.*

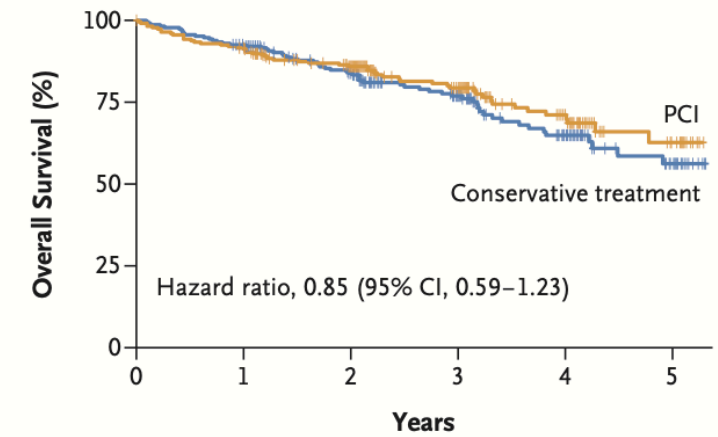
Variable	PCI (N = 227)	Conservative Treatment (N = 228)
Angiographic findings		
Median no. of physiologically significant lesions per patient (IQR) [†]	1 (1–2)	1 (1–2)
No. of lesions with fractional flow reserve ≤ 0.80	167	155
No. of lesions with diameter stenosis $\geq 90\%$	184	162
Median largest diameter stenosis (IQR) — %	90 (80–90)	90 (71–90)
Median SYNTAX score (IQR) [‡]	9 (6–14)	9 (5–14)
PCI procedure[†]		
Median no. of days from randomization to PCI (IQR)	9 (1–26)	—
Timing of PCI — no./total no. (%)		
Before TAVI	163/219 (74)	—
Concomitant with TAVI	37/219 (17)	—
After TAVI	19/219 (9)	—
Complete revascularization achieved — no./total no. (%) [§]	194/219 (89)	—
TAVI procedure		
Median no. of days from randomization to TAVI (IQR)	34 (7–62)	25 (2–54)
Balloon-expandable heart valve — no. (%)	90 (40)	95 (42)

A Death from Any Cause, Myocardial Infarction, or Urgent Revascularization (primary end point)



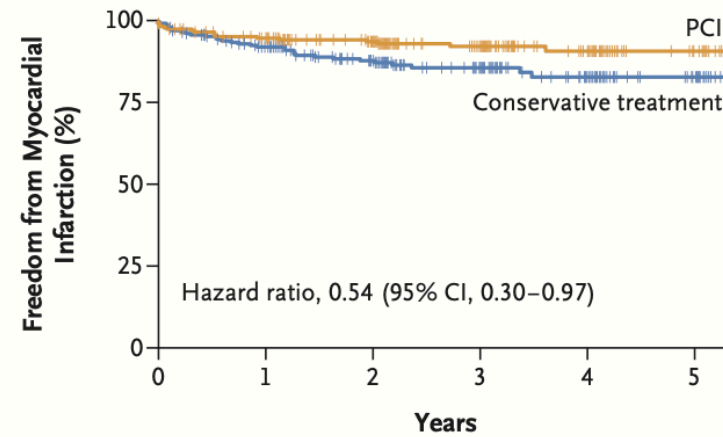
No. at Risk						
PCI	227	199	161	109	58	14
Conservative treatment	228	191	148	91	46	18

B Death from Any Cause



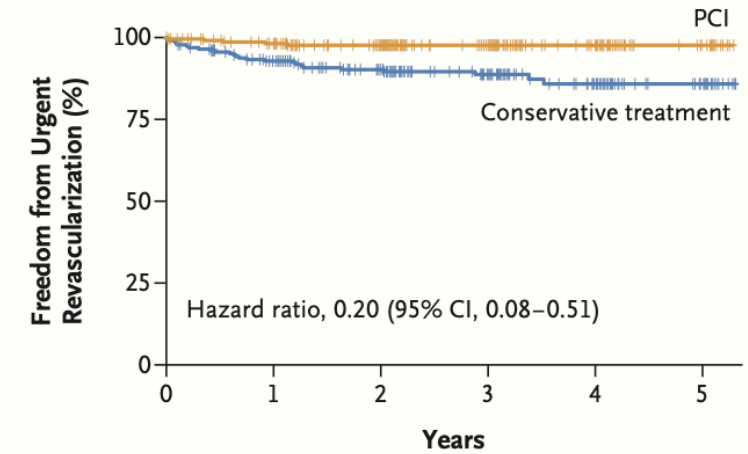
No. at Risk						
PCI	227	206	168	115	60	15
Conservative treatment	228	207	166	103	54	20

C Myocardial Infarction



No. at Risk						
PCI	227	199	161	109	58	14
Conservative treatment	228	193	150	92	47	18

D Urgent Revascularization



No. at Risk						
PCI	227	202	164	112	59	15
Conservative treatment	228	195	153	94	48	19

Figure 1. Primary End Point: Major Adverse Cardiac Events.



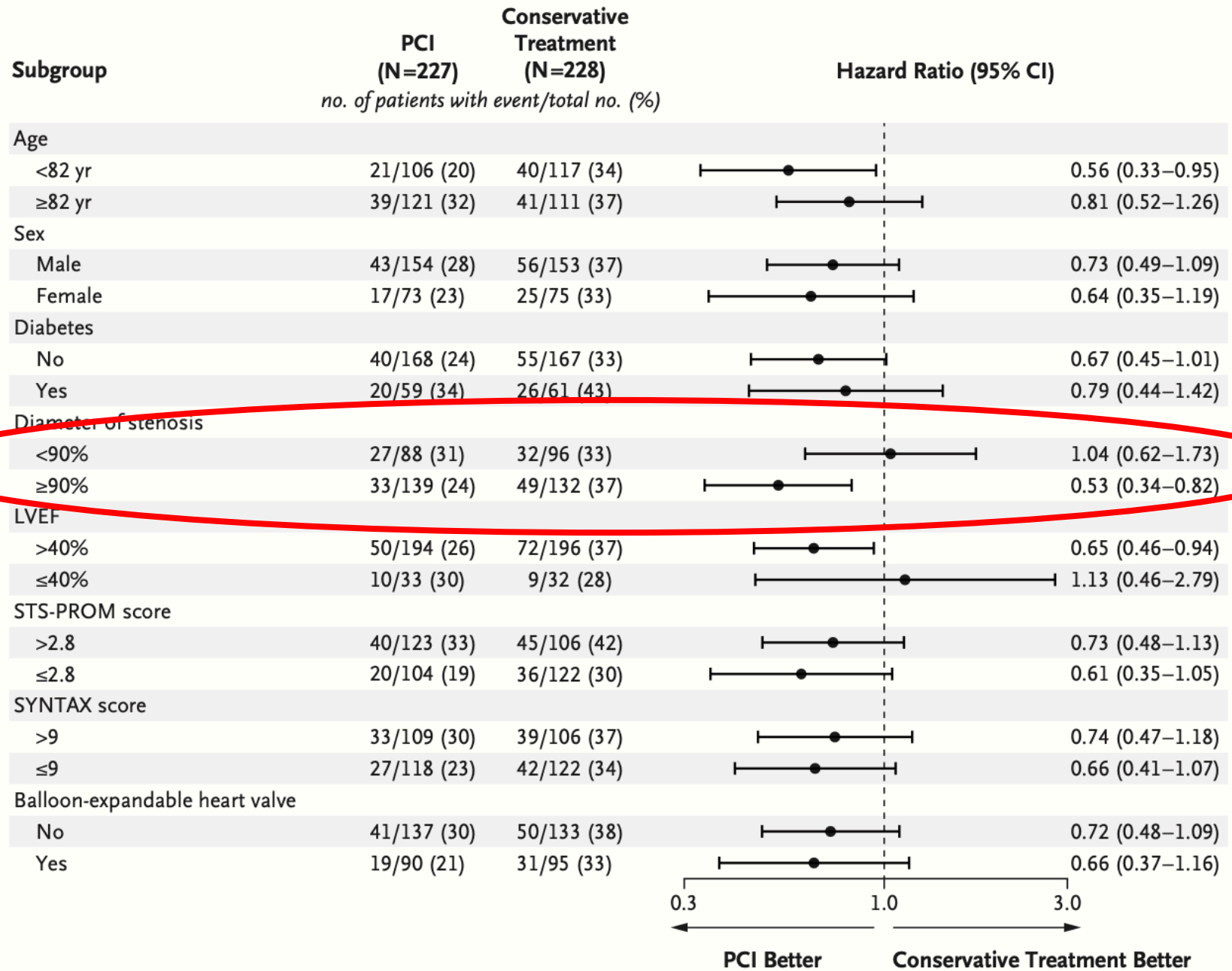


Figure 2. Subgroup Analysis of the Primary End Point.

FAITAVI: Functional Assessment In TAVI

Eurointervention 2024;20:504-10



Study design: multicentre, prospective, randomized, open-label, superiority trial with blinded adjudication of clinical events



Study population: 320 patients



Randomization: 1:1 to FFR-guided versus angio-guided PCI before or after TAVI

Left Main lesions (>50-<90%) included. ACS and EF<35% excluded



Primary endpoint: 12-month MACCE, a composite of all-cause death, MI, ischemia-driven TVR, stroke, or major bleeding (BARC ≥ 3)



Interventional strategy: In the FFR-guided arm, lesions with $\text{FFR} \leq 0.80$ were treated, those >0.85 were deferred, and borderline lesions ($\text{FFR} 0.81\text{--}0.85$) were re-assessed after TAVI

In the angio-guided arm, all lesions $\geq 50\%$ in vessels >2.5 mm were treated

FAITAVI: Functional Assessment In TAVI

Eurointervention 2024;20:504-10

TAVI candidates as per Heart Team indication (inclusion/exclusion criteria satisfied)
Signed informed consent
Enrolment period Nov 2017-June 2023

320 patients enrolled and randomized 1:1

Angio-guided PCI (156)

FFR-guided PCI (164)

TAVI

Angio-guided PCI if
 $\%DS \geq 50$ (before or after TAVI)

FFR-guided PCI if
 $FFR \leq 0.80$ (before or after TAVI)

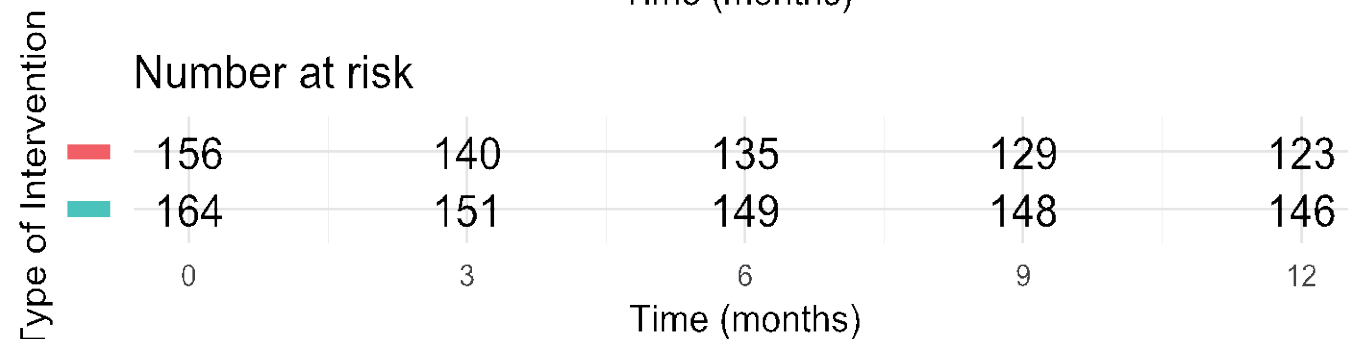
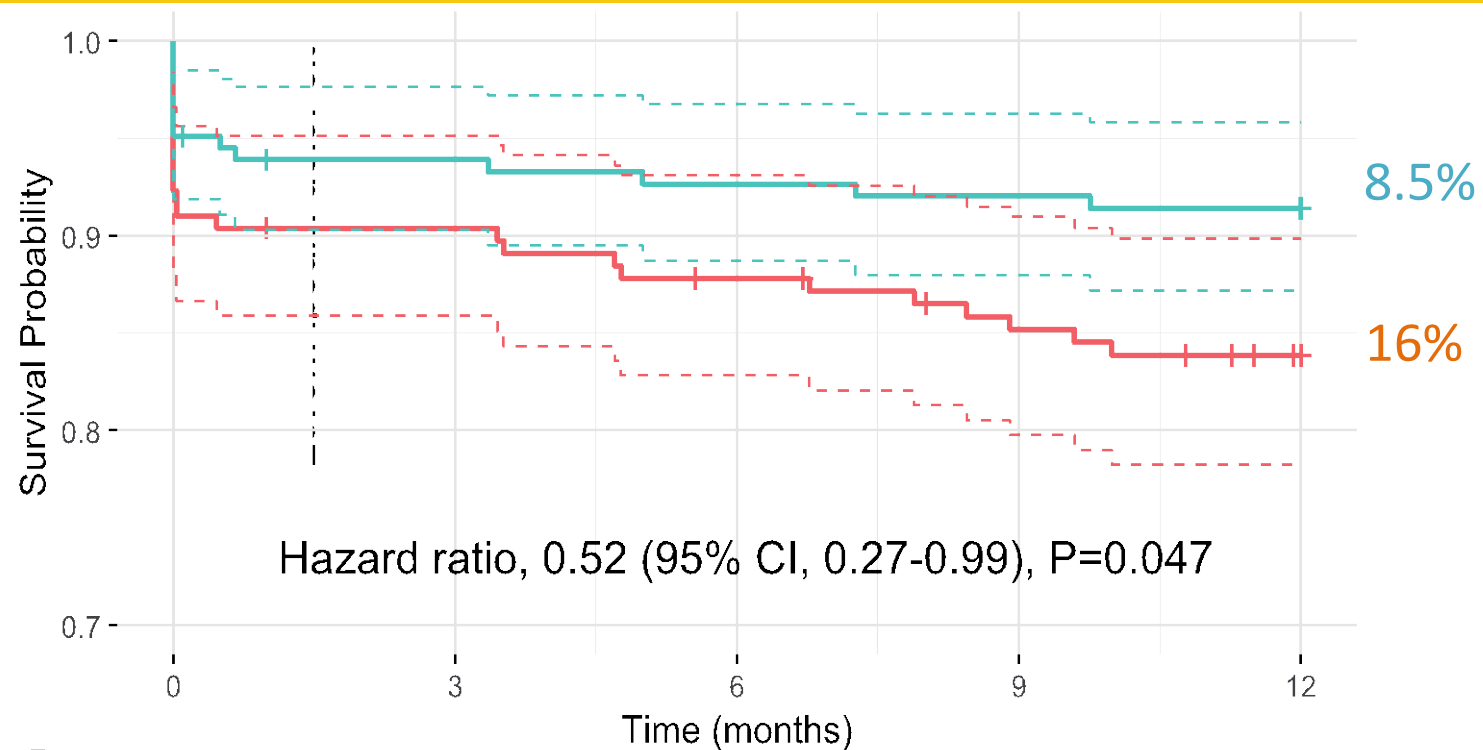
Immediate outcome in the cathlab

Primary endpoint at 12 months (97%)

Study cohort

Characteristics of the study cohort at baseline		
Characteristic	FFR-guided PCI (n=164)	Angiography-guided PCI (n=156)
Age - year	87 (83-90)	86 (82-89)
Female sex - %	41	44
STS-PROM score - %	3 (2-4)	4 (2-5)
Left ventricular ejection fraction (IQR) - %	58 (53-64)	59 (55-64)
Median aortic valve area (IQR) cm ²	0.7 (0.6-0.9)	0.7 (0.6-0.9)
Median mean aortic valve gradient (IQR) - mmHg	46 (40-53)	43 (38-52)
History of PCI - %	14	17
History of myocardial infarction - %	13	11
Diabetes mellitus - %	34	38
Chronic kidney disease	26	29
Atrial fibrillation	29	28
Peripheral artery disease	25	26
Median SYNTAX score (IQR) [†]	7 (5-11)	7 (5-11)
Median diameter stenosis (IQR) - %	55 (46-63)	56 (48-63)
Median FFR (IQR) before TAVI	0.80 (0.74-0.87)	-

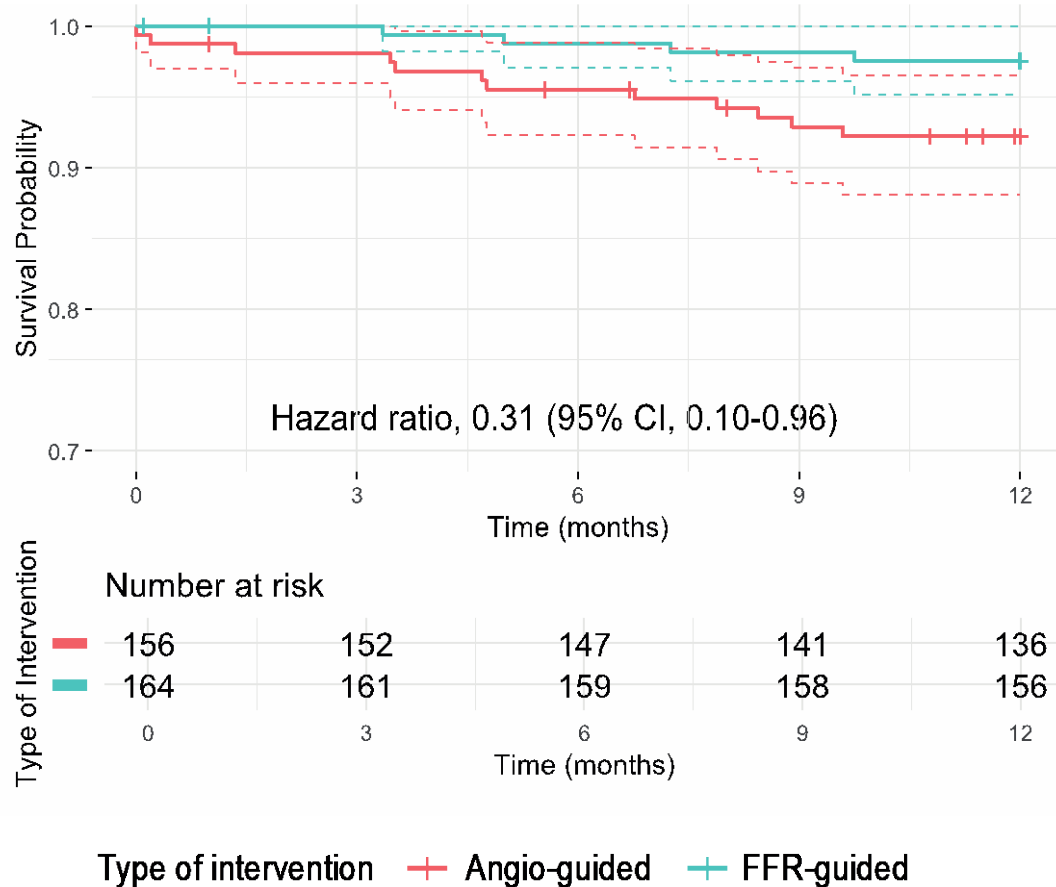
Primary endpoint



Type of intervention + Angio-guided + FFR-guided

Secondary and safety endpoints

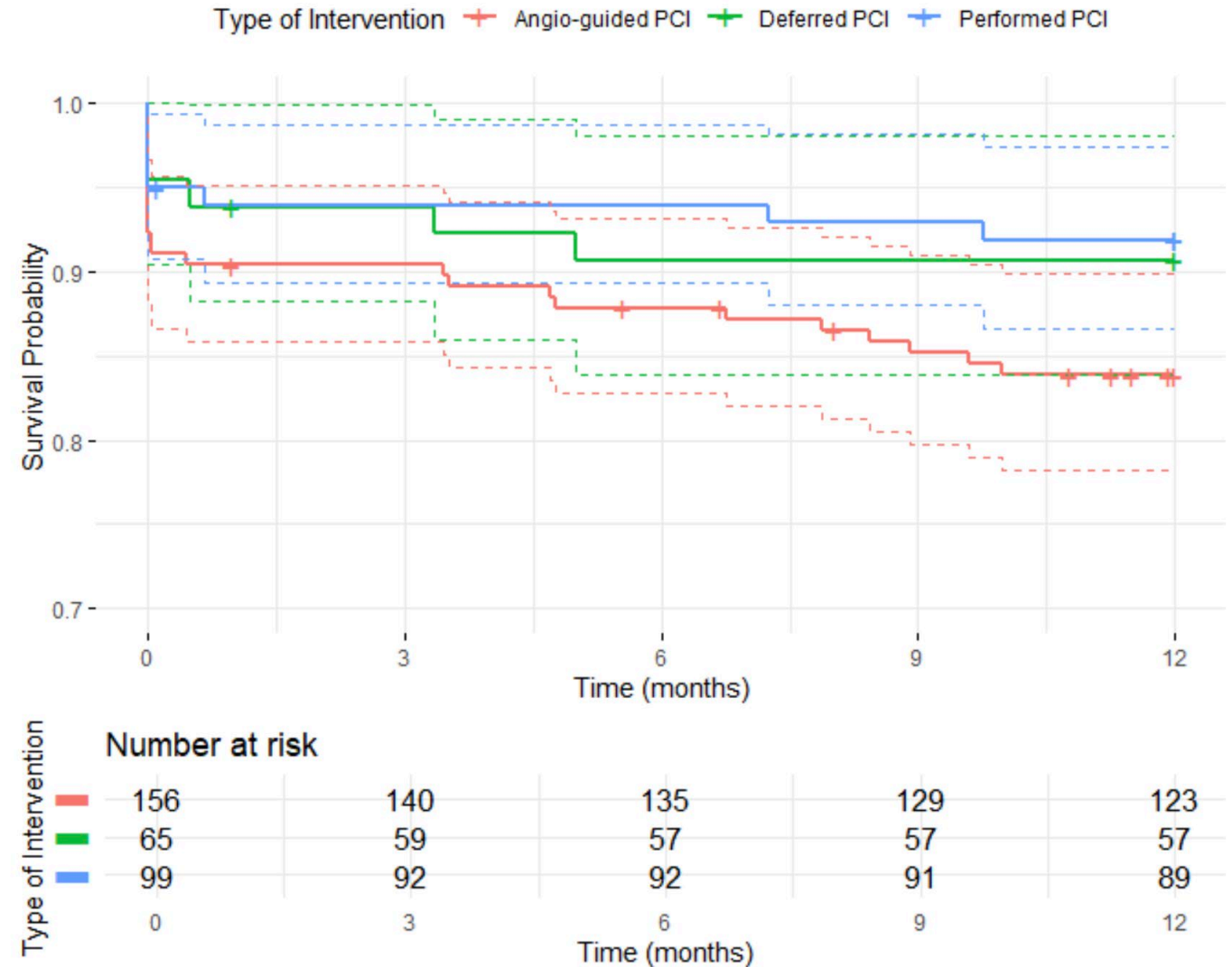
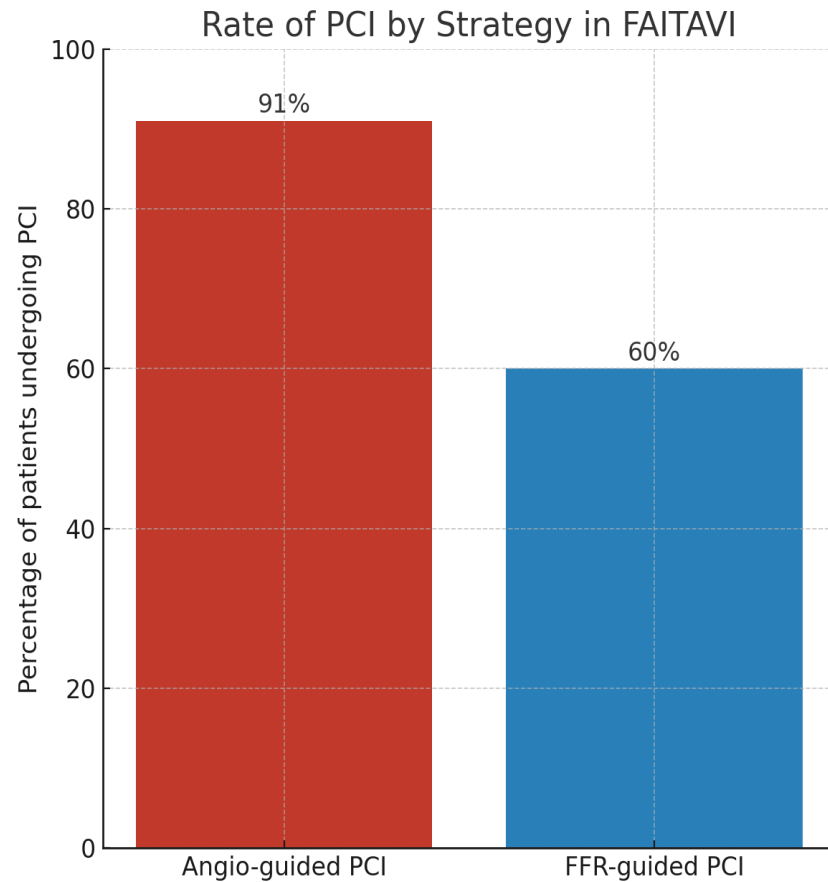
All cause death



Secondary endpoints	FFR-guided	Angio-guided	HR (95% CI)
Death from any cause	4 (2.4)	12 (7.7)	0.31 (0.10-0.96)
Death from cardiovascular causes	0 (0)	3 (1.9)	
Myocardial infarction	1 (0.6)	2 (1.2)	0.47 (0.04-5.23)
Ischemia driven revascularization	0 (0)	3 (1.9)	
Stroke	1 (0.6)	4 (2.5)	0.24 (0.03-2.11)

Safety endpoints	FFR-guided	Angio-guided	HR (95% CI)
Major bleeding	8 (4.9)	10 (6.4)	0.75 (0.30-1.91)
Acute kidney failure	5 (3.0)	5 (3.2)	0.95 (0.26-3.48)
Periprocedural myocardial infarction	1 (0.6)	2 (1.2)	0.47 (0.02-4.98)
Vascular complications	13 (7.9)	9 (5.7)	1.41 (0.59-3.50)

Deferred vs treated coronary lesions



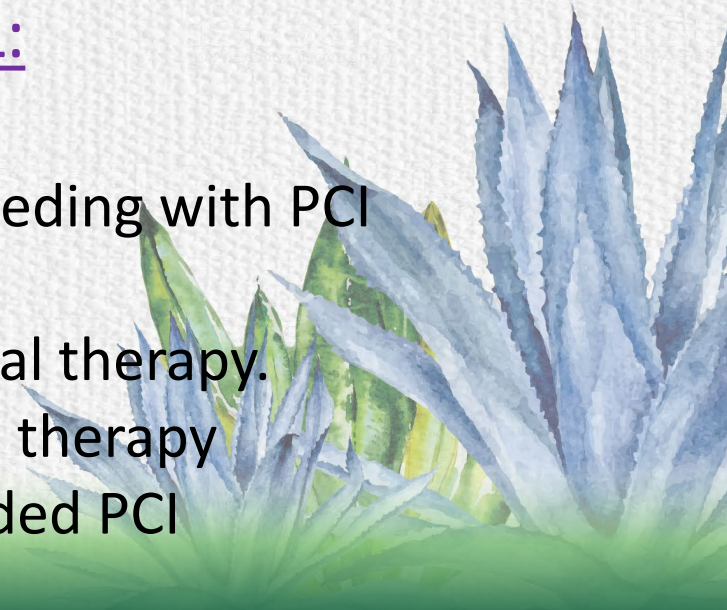
PCI before TAVI? or TAVI before PCI?

Driving considerations:

1. Most patients came to our observation because of AS, not CAD
2. Most AS patients with CAD have no angina...
3. Detection of inducible ischemia in AS is not standardized
4. Any complication of PCI could be better tolerated without AS
5. Organ perfusion is better after removal of AS

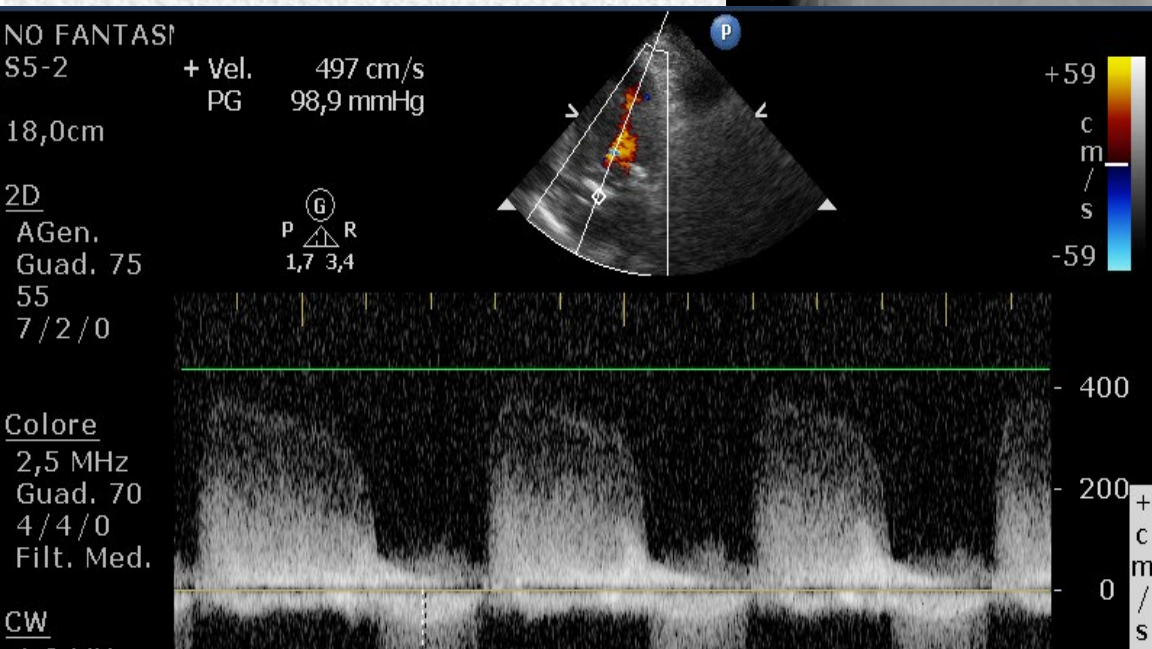
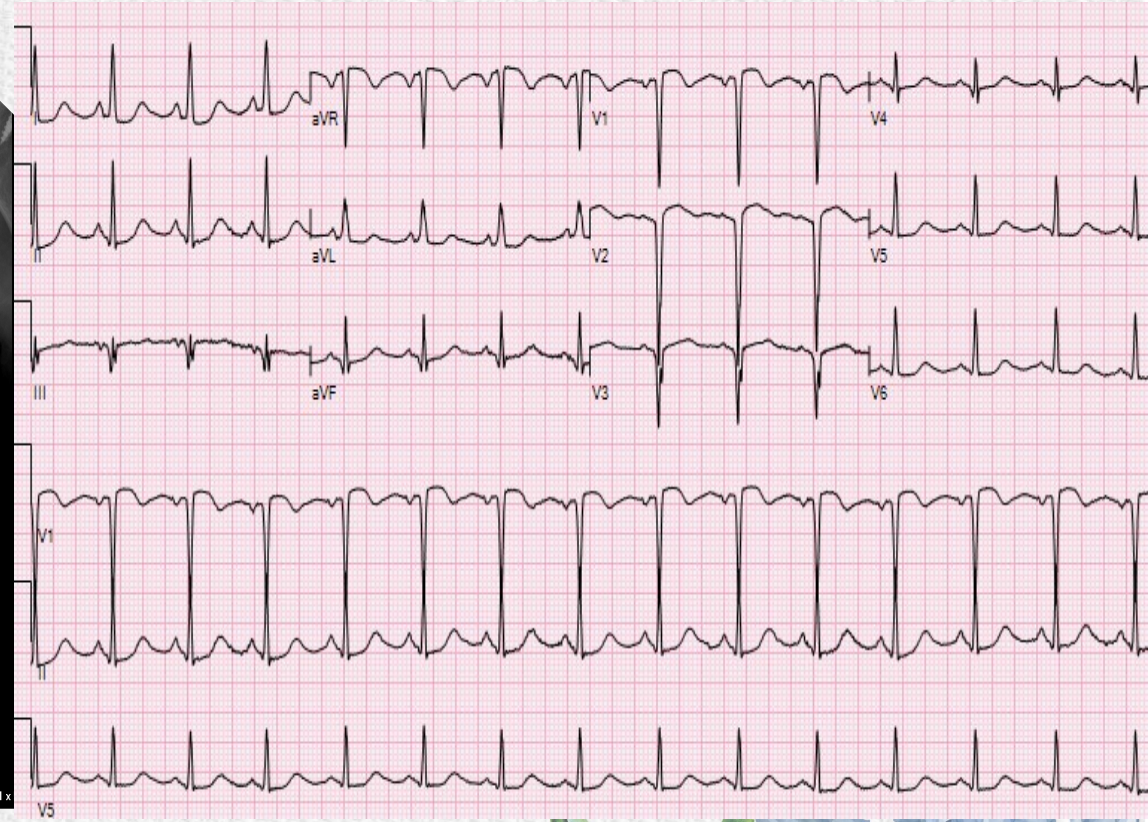
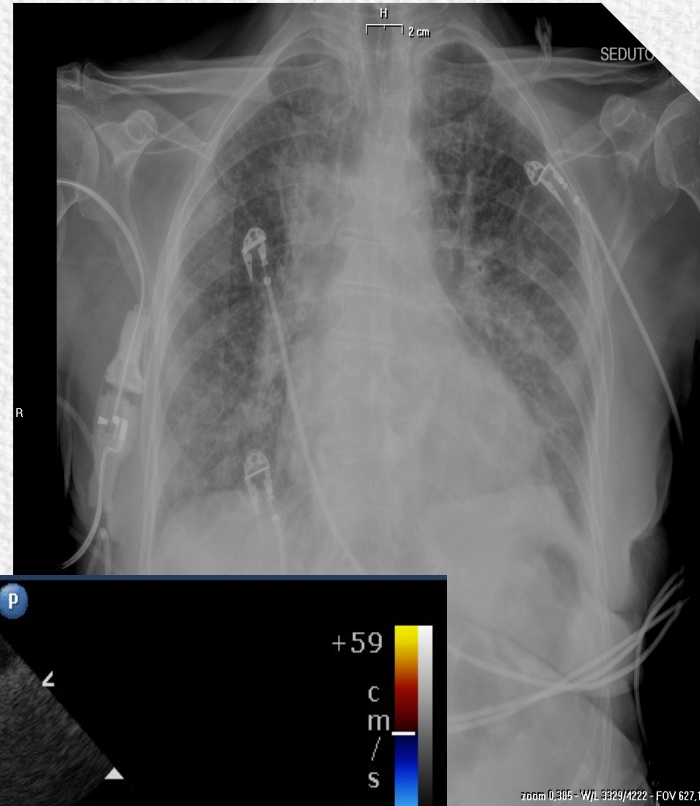
Available evidence released after the latest GL:

- **ACTIVATION:** PCI vs medical therapy: only higher bleeding with PCI
- **TCW:** TAVI and PCI better than SAVR and CABG
- **NOTION-3:** PCI of lesions >90% better than medical therapy.
PCI of lesions <90% same as medical therapy
- **FAITAVI:** FFR-guided PCI better than angio-guided PCI



Case 1: ACS and AVS

- 83 yo female
- Known severe AVS symptomatic for angina, refused any intervention
- Diagnosis of anterior STEMI with heart failure

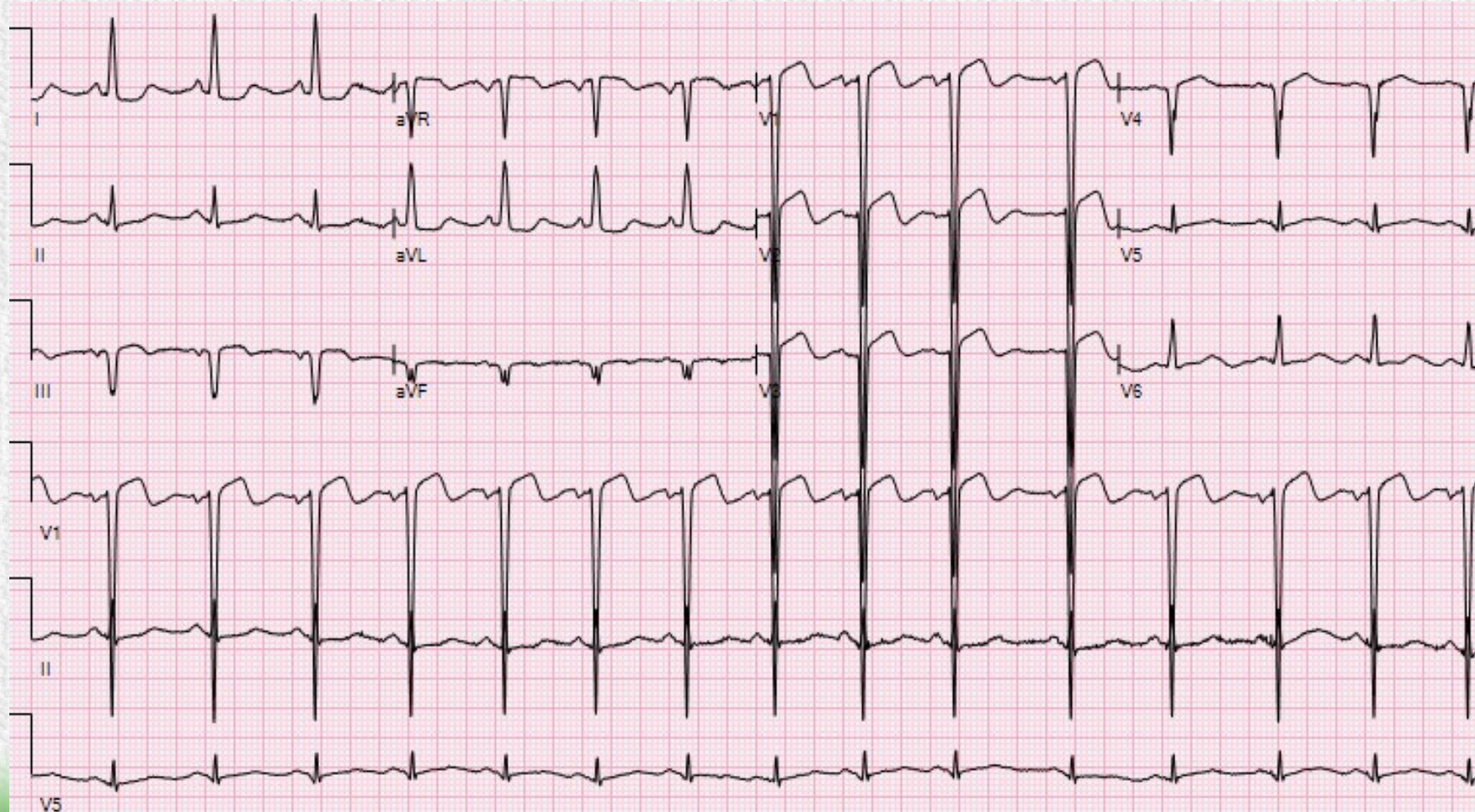
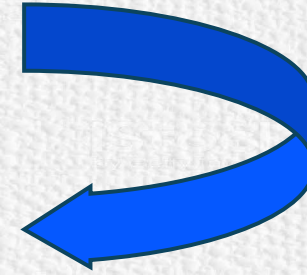


Due to the lack of angina and the acute renal insufficiency she was managed medically with diuretics, inotropes, DAPT and non invasive ventilation with c-pap



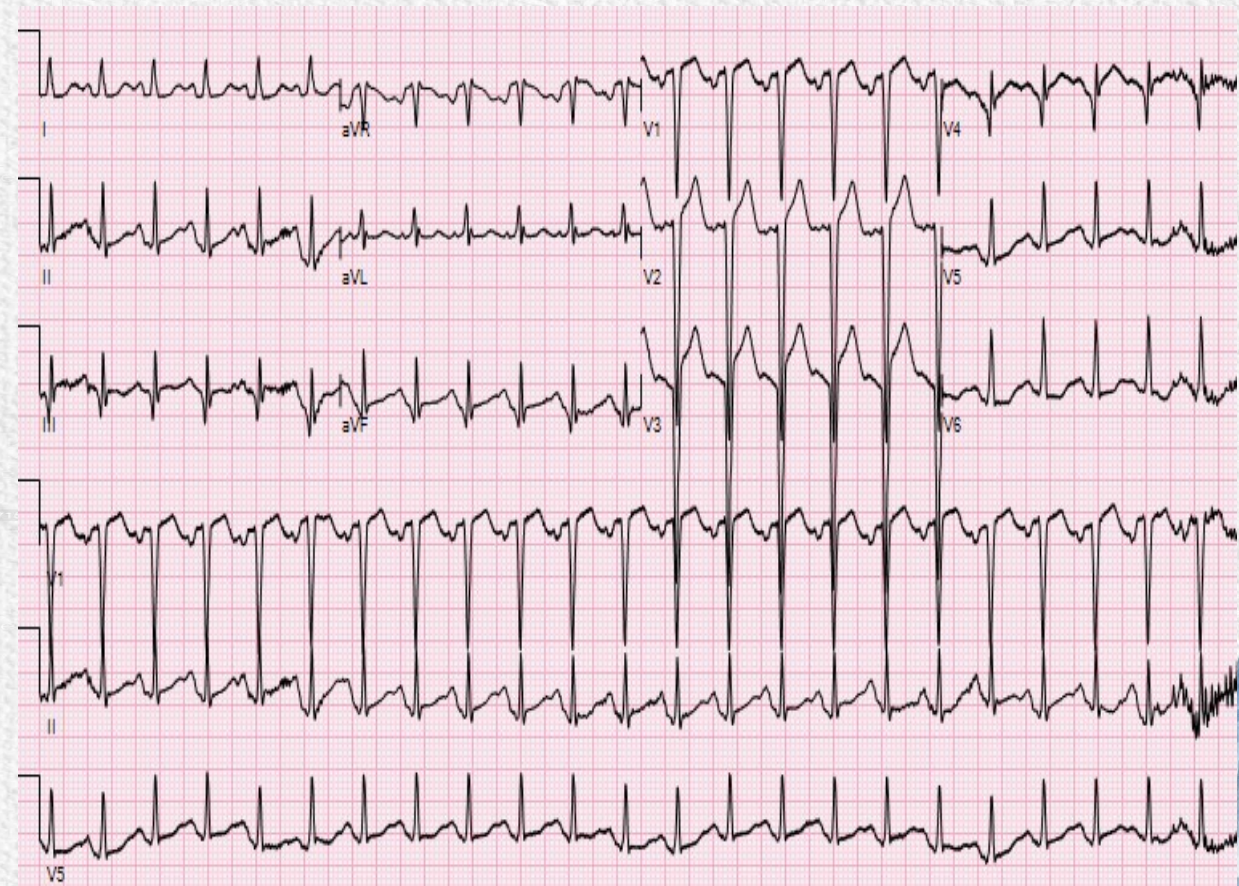
48h after admission recurrence of chest
pain and hemodynamic shock

Emergency angiography



Ostial LM stenting

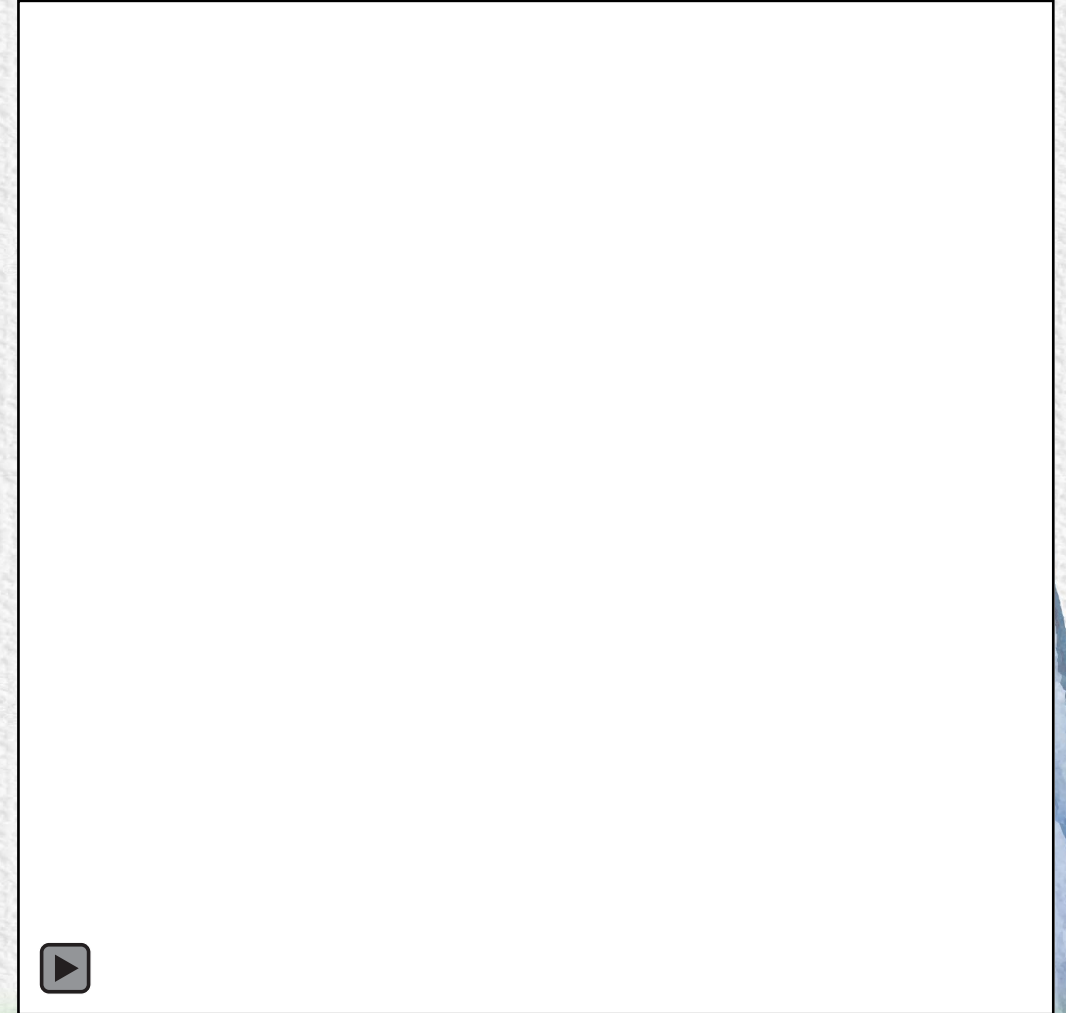
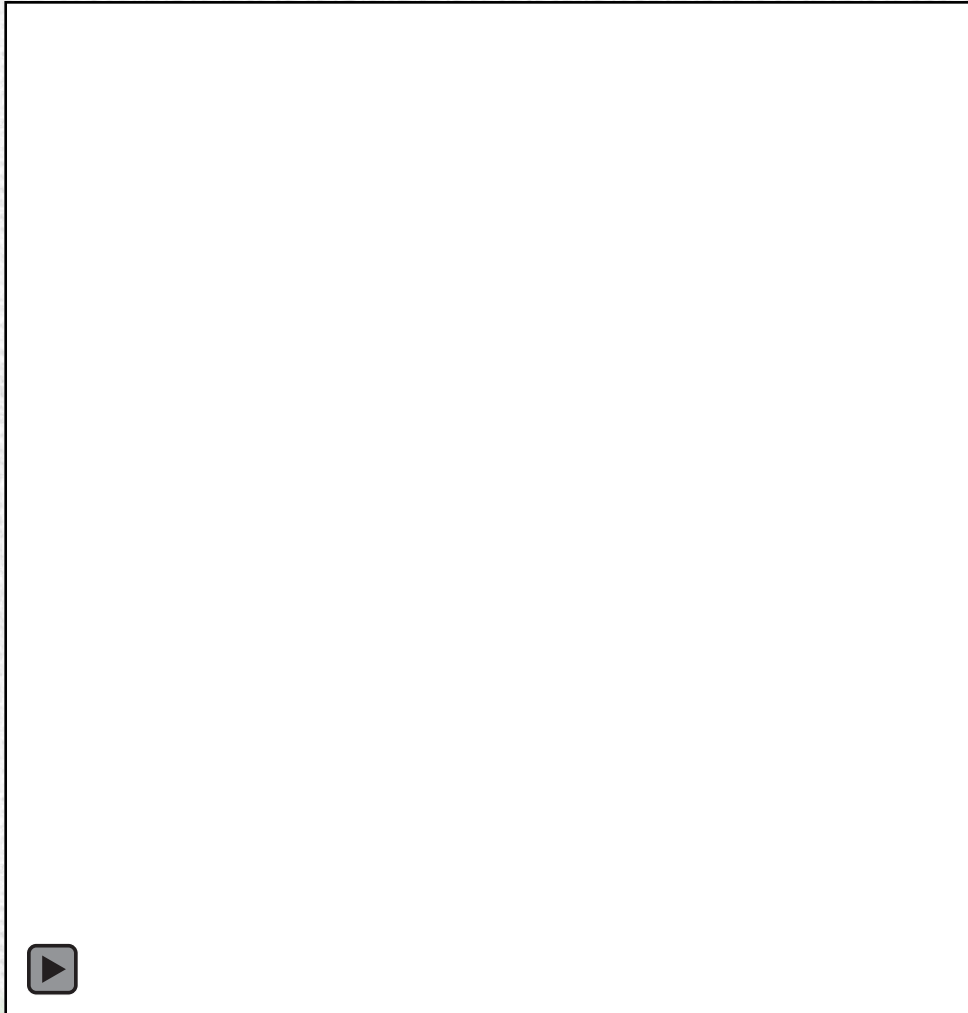




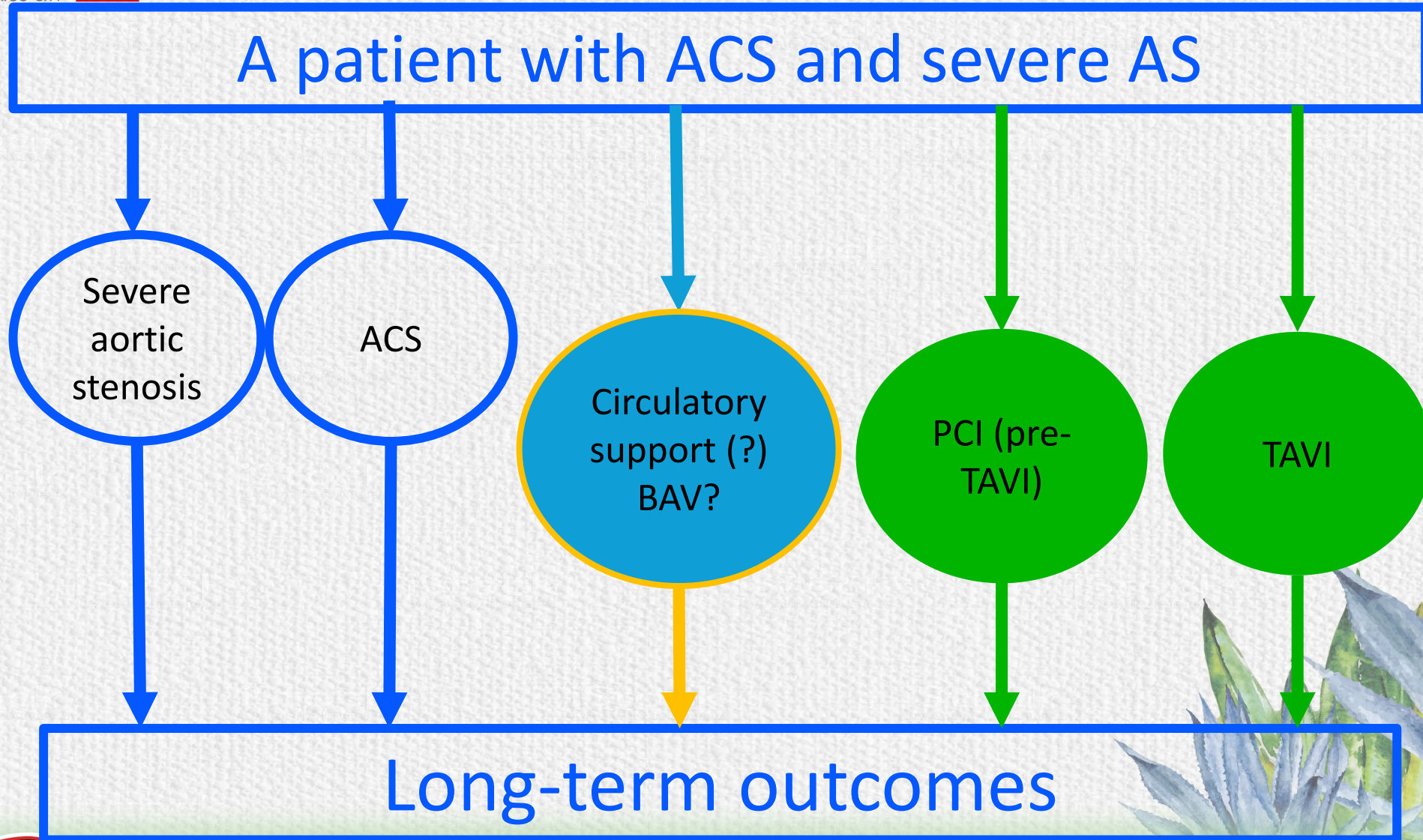
ECG post-PCI
31/05/2020



Staged TAVI procedure: 23mm Sapien Ultra valve



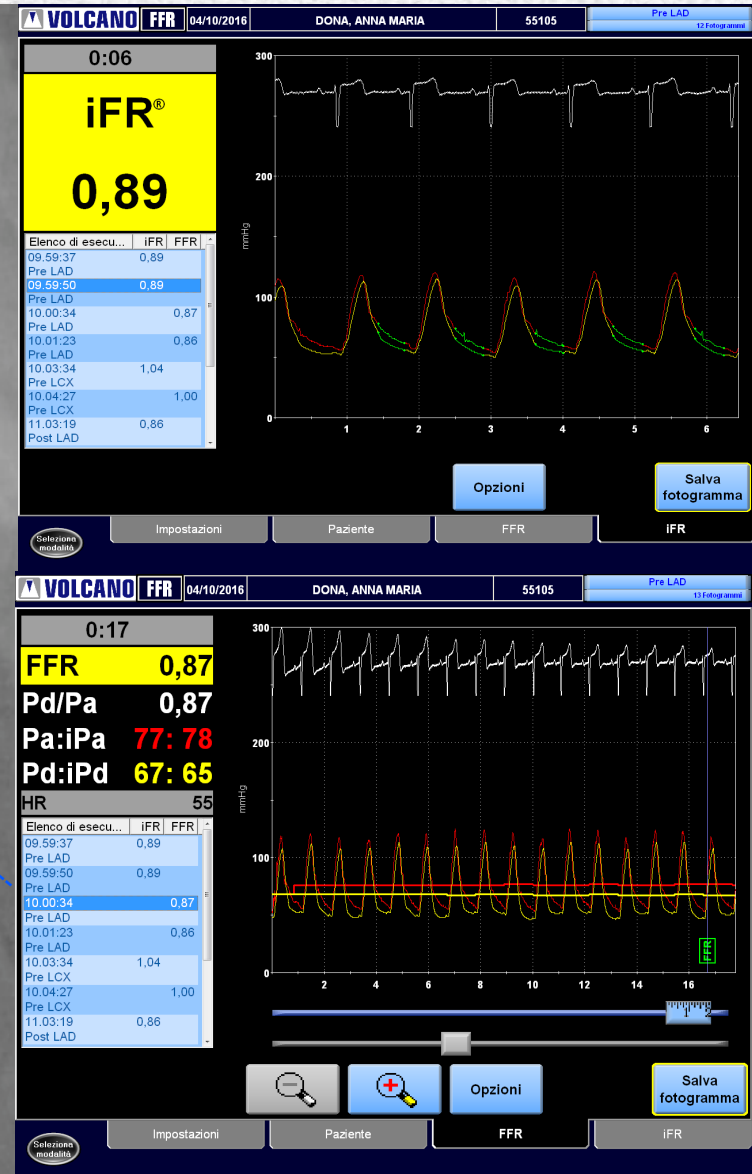
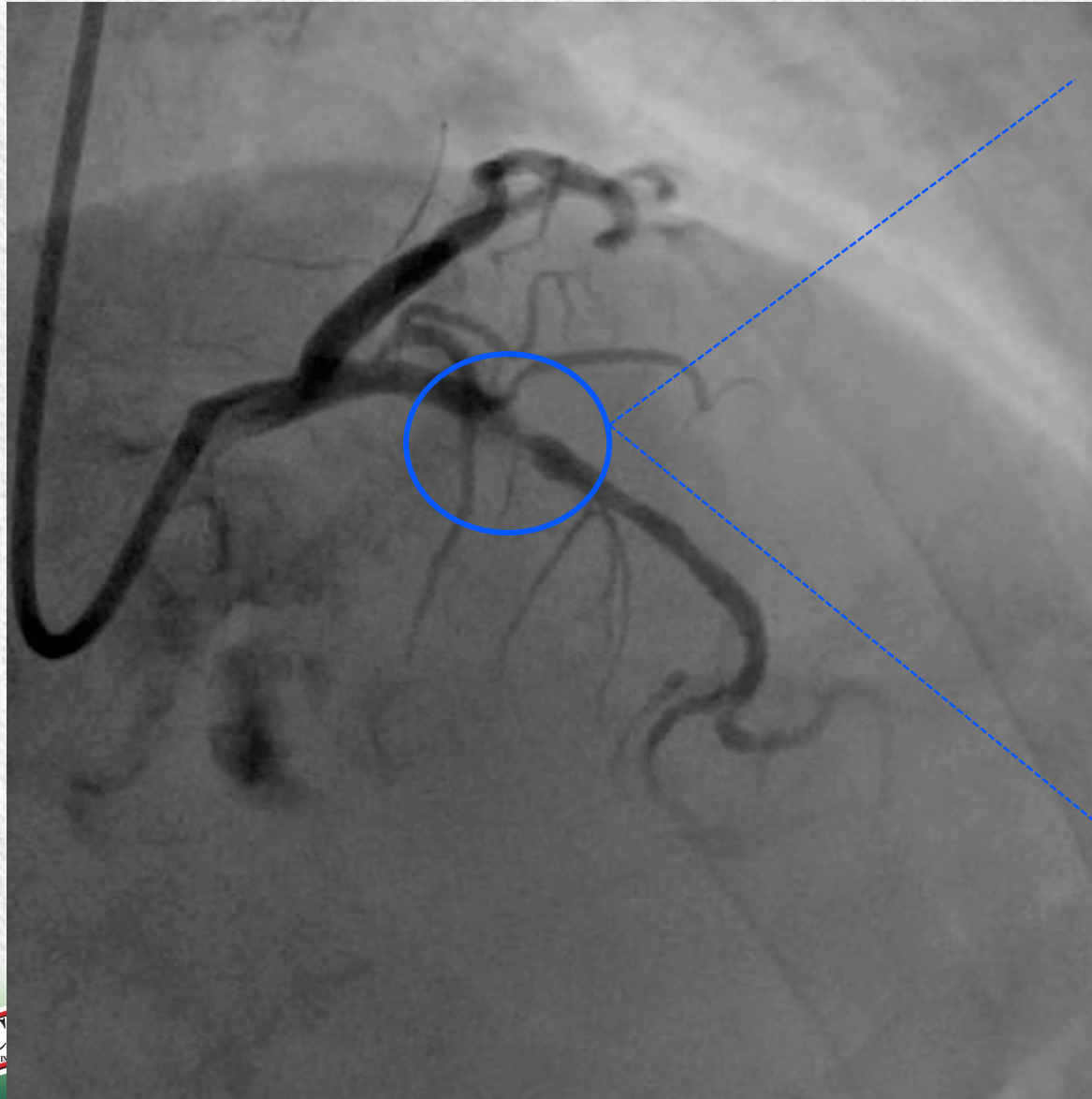
Timing of PCI: clinical & procedural factors



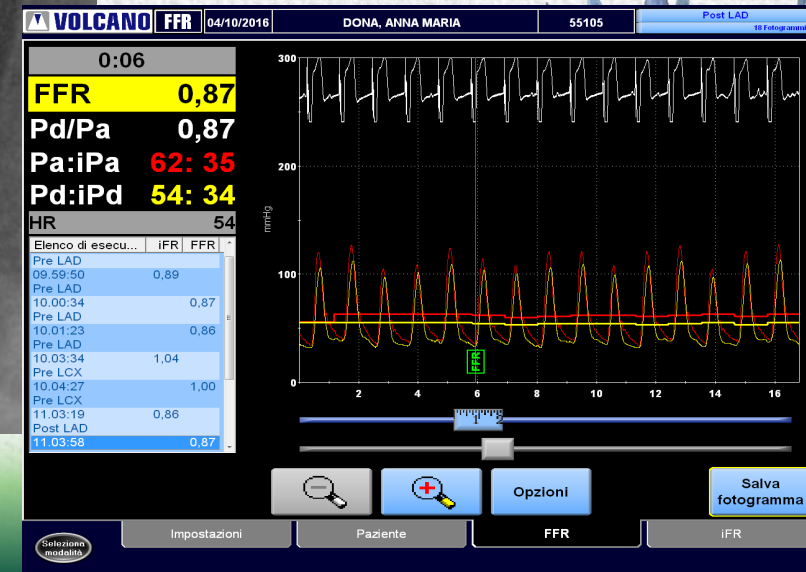
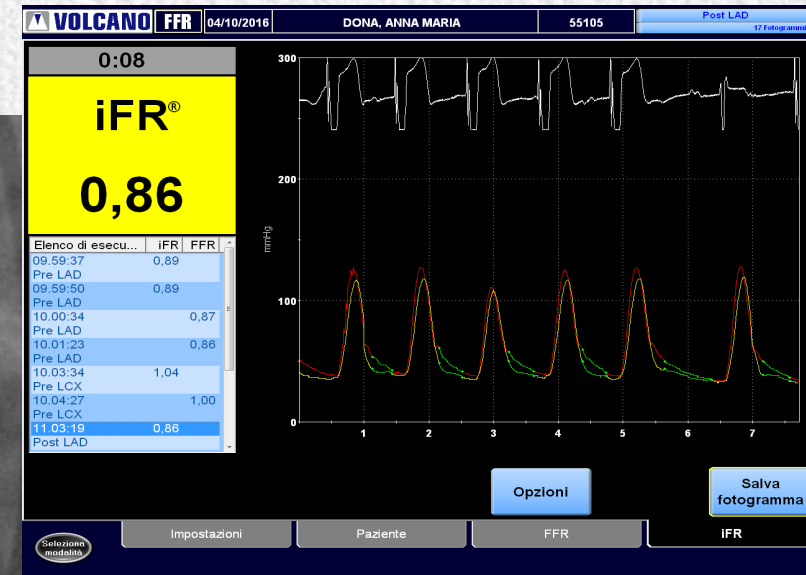
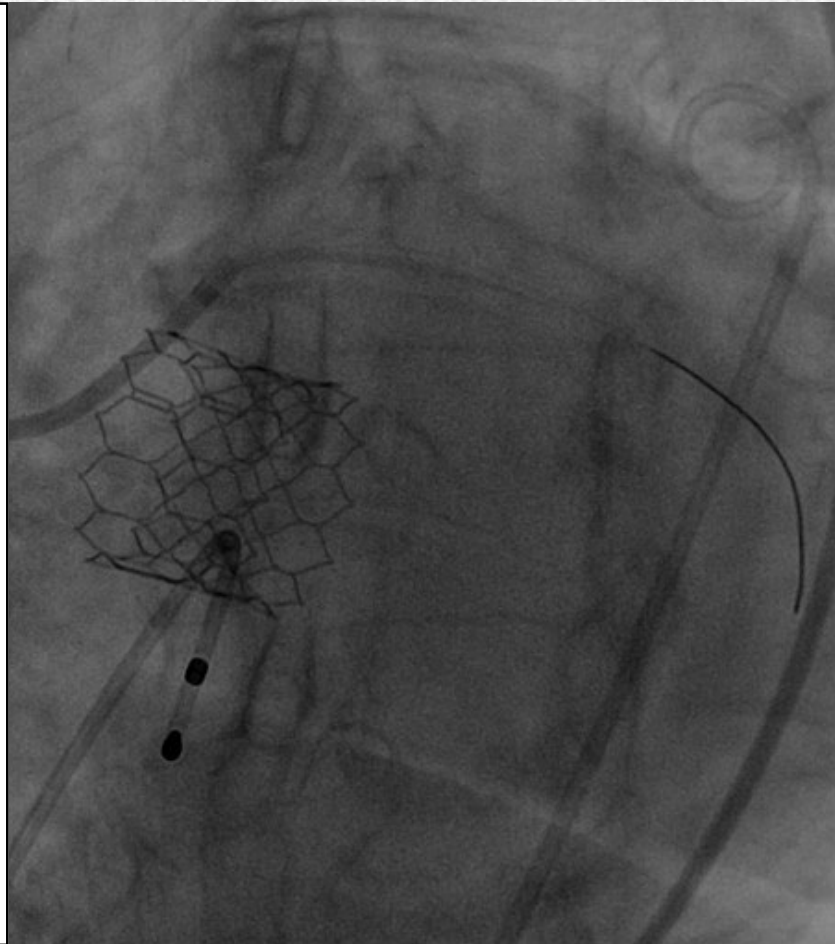
Case 2: AVS with asymptomatic and low-risk CAD

- Female, 80 years old
- Hypertension, Dyslipidemia
- Severe aortic stenosis (MG= 55mmHg)
- Normal LV EF (60%)
- GFR 35ml/min
- Symptomatic for NYHA class II, never had chest pain

Coronary physiology pre-TAVI



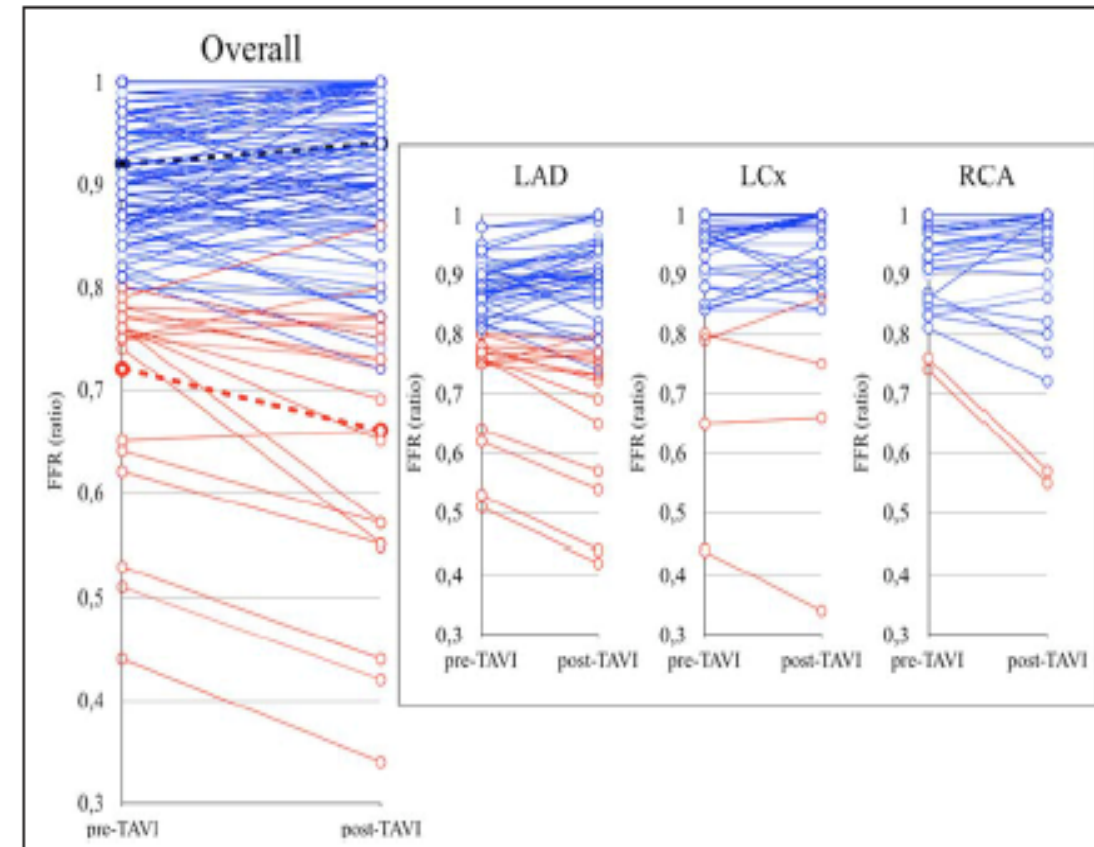
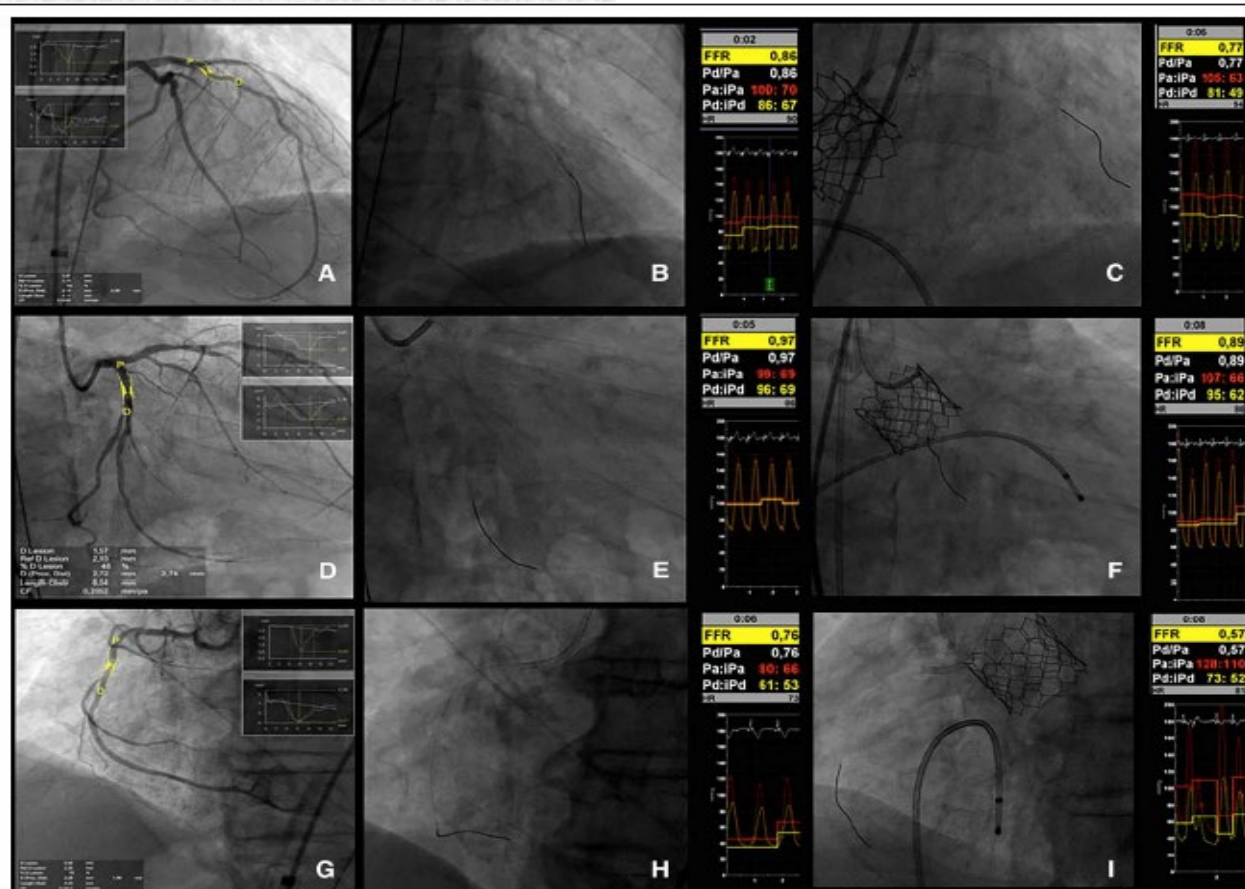
Coronary physiology post-TAVI





Functional Assessment of Coronary Artery Disease in Patients Undergoing Transcatheter Aortic Valve Implantation Influence of Pressure Overload on the Evaluation of Lesions Severity

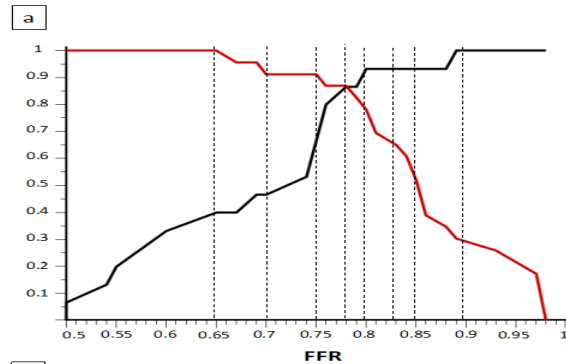
Gabriele Pesarini, MD; Roberto Scarsini, MD; Carlo Zivelonghi, MD; Anna Piccoli, MD;
Alessia Gambaro, MD; Leonardo Gottin, MD; Andrea Rossi, MD; Valeria Ferrero, MD;
Corrado Vassanelli, MD; Flavio Ribichini, MD



Short communication

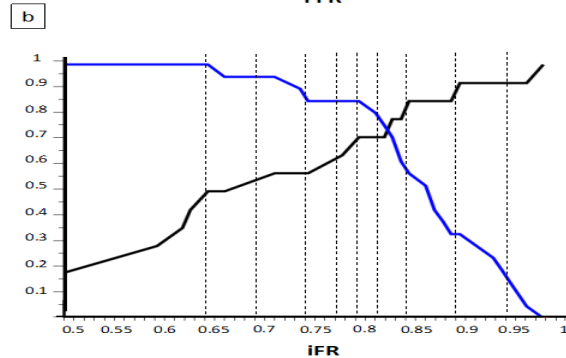
Correlation between intracoronary physiology and myocardial perfusion imaging in patients with severe aortic stenosis

Roberto Scarsini ^{a,b}, Rosaria Cantone ^a, Gabriele Venturi ^a, Giovanni Luigi De Maria ^b, Andrea Variola ^{a,c}, Paolo Braggio ^c, Mattia Lunardi ^a, Gabriele Pesarini ^a, Marco Ferdeghini ^c, Anna Piccoli ^{b,e}, Mauro Feola ^d, Rajesh K. Kharbada ^b, Adrian P. Banning ^b, Flavio Ribichini ^{a,*}



— Sensitivity
— Specificity

FFR cut-off	Accuracy	Sensitivity	Specificity
0.65	78%	40%	100%
0.70	78%	53%	92%
0.75	83%	67%	92%
0.78*	88%	87%	88%
0.80	85%	93%	81%
0.83**	78%	93%	69%
0.85	68%	93%	54%
0.90	54%	100%	27%



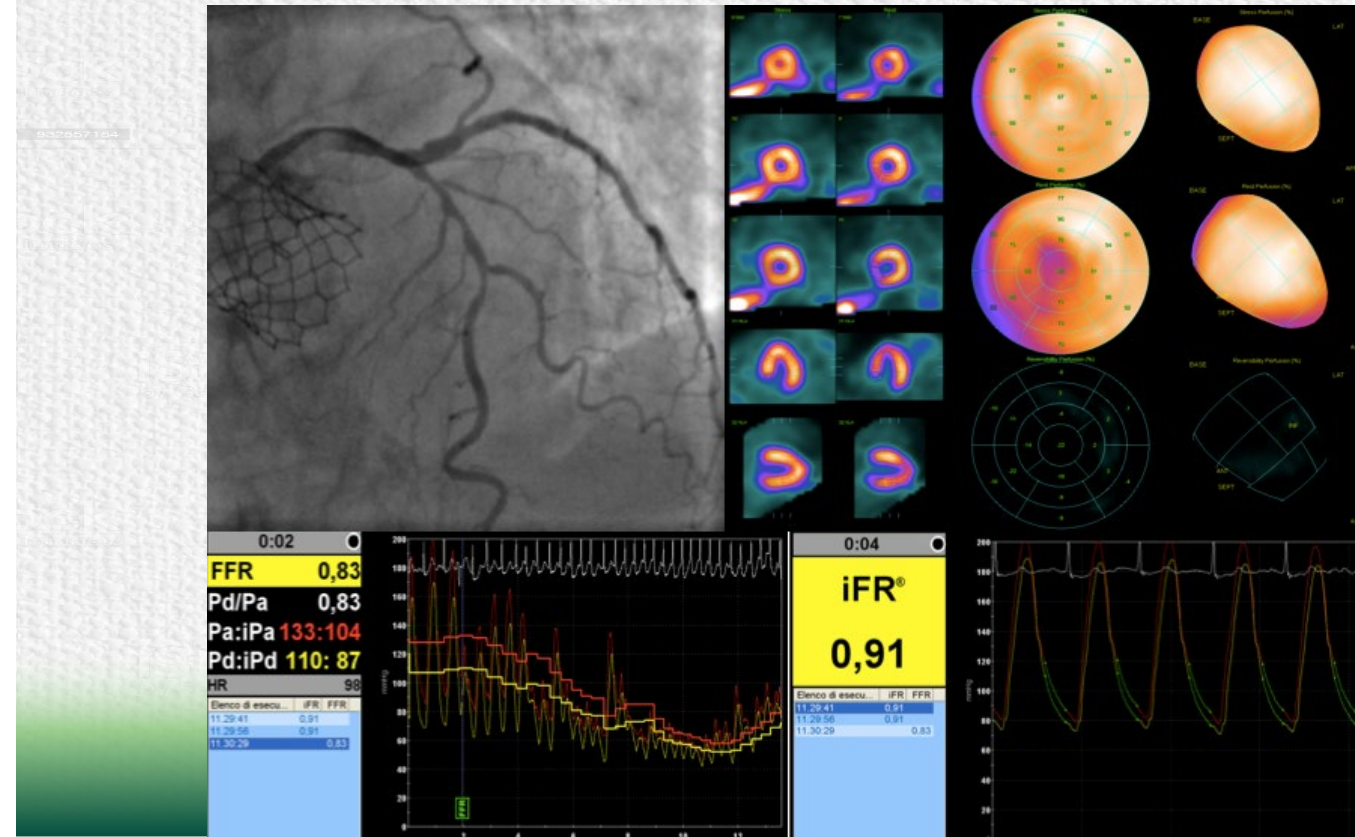
— Sensitivity
— Specificity

iFR cut-off	Accuracy	Sensitivity	Specificity
0.65	78%	47%	96%
0.70	80%	60%	92%
0.75	78%	60%	88%
0.78*	80%	73%	85%
0.80	78%	73%	81%
0.82***	73%	80%	69%
0.85	68%	87%	58%
0.89	59%	93%	38%
0.95	49%	100%	19%

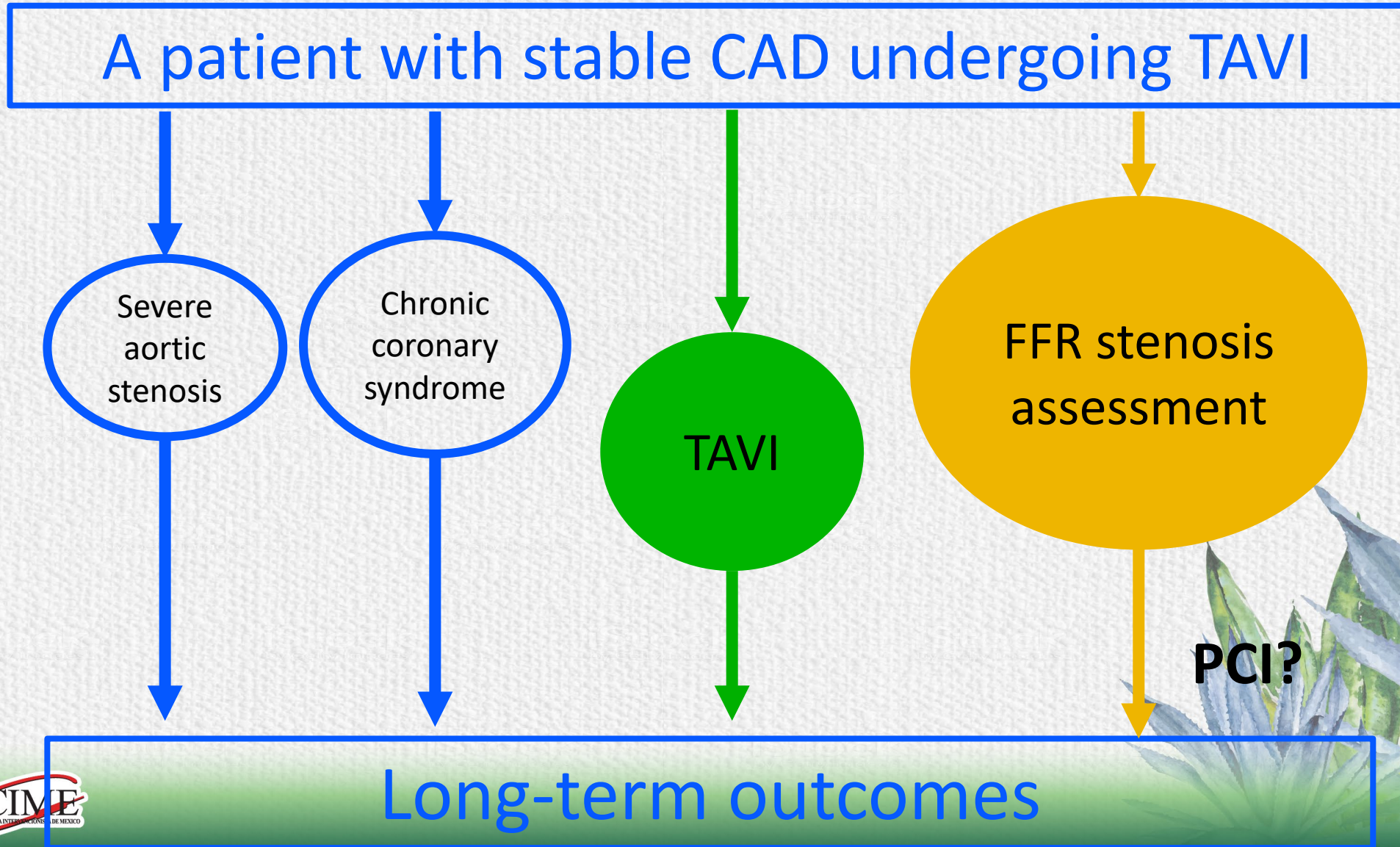
*Cut-off identified by ROC derived Youden index

** prespecified FFR cut-off<0.83

***prespecified iFR cut-off<0.82



Timing of PCI: clinical & procedural factors

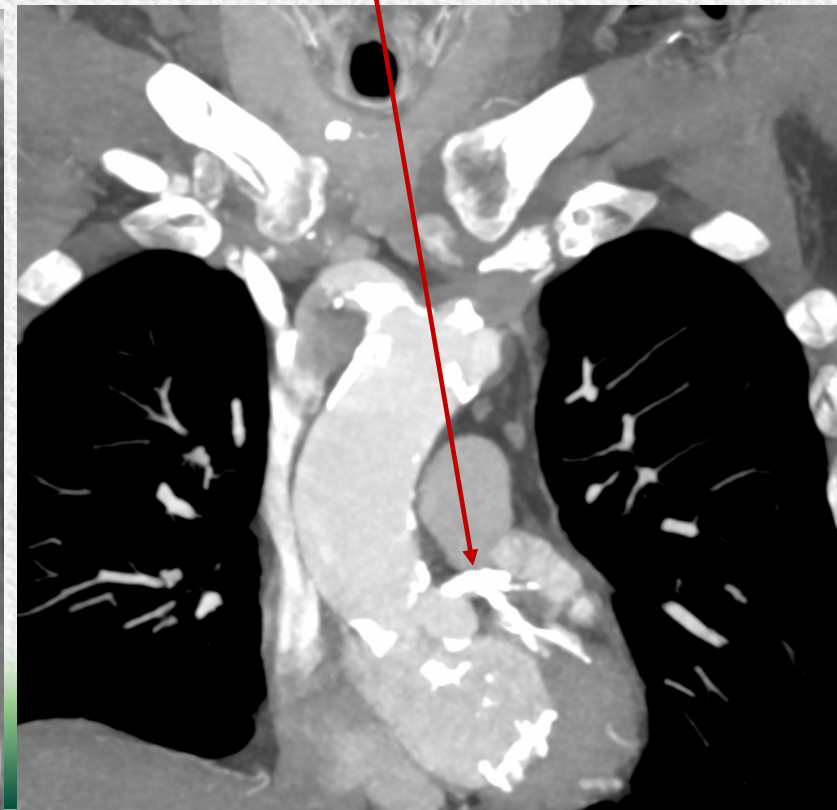
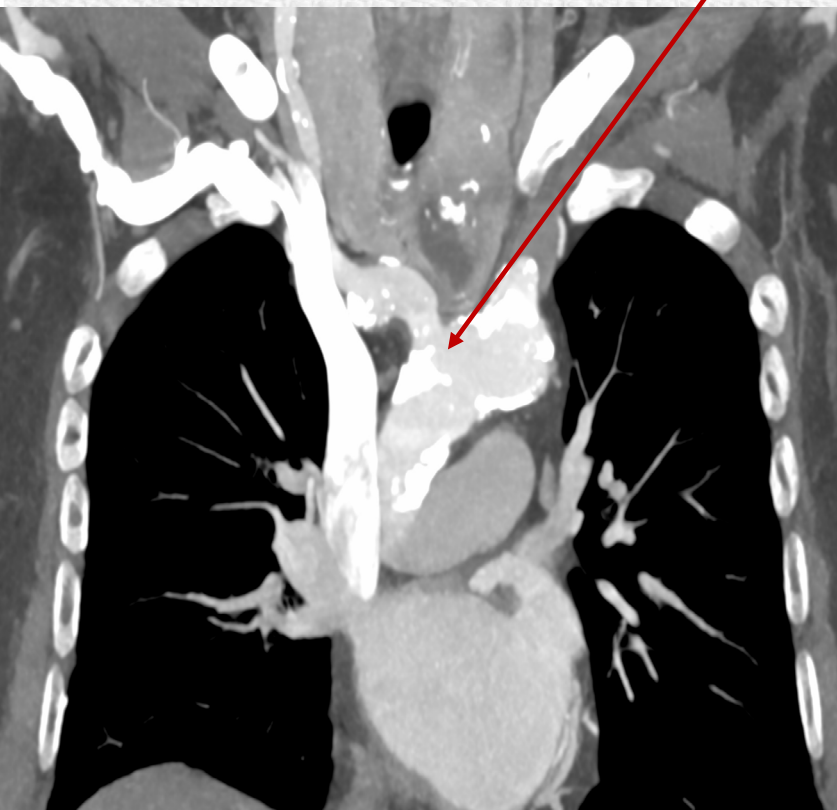


Case 3: AVS with symptomatic and high-risk CAD

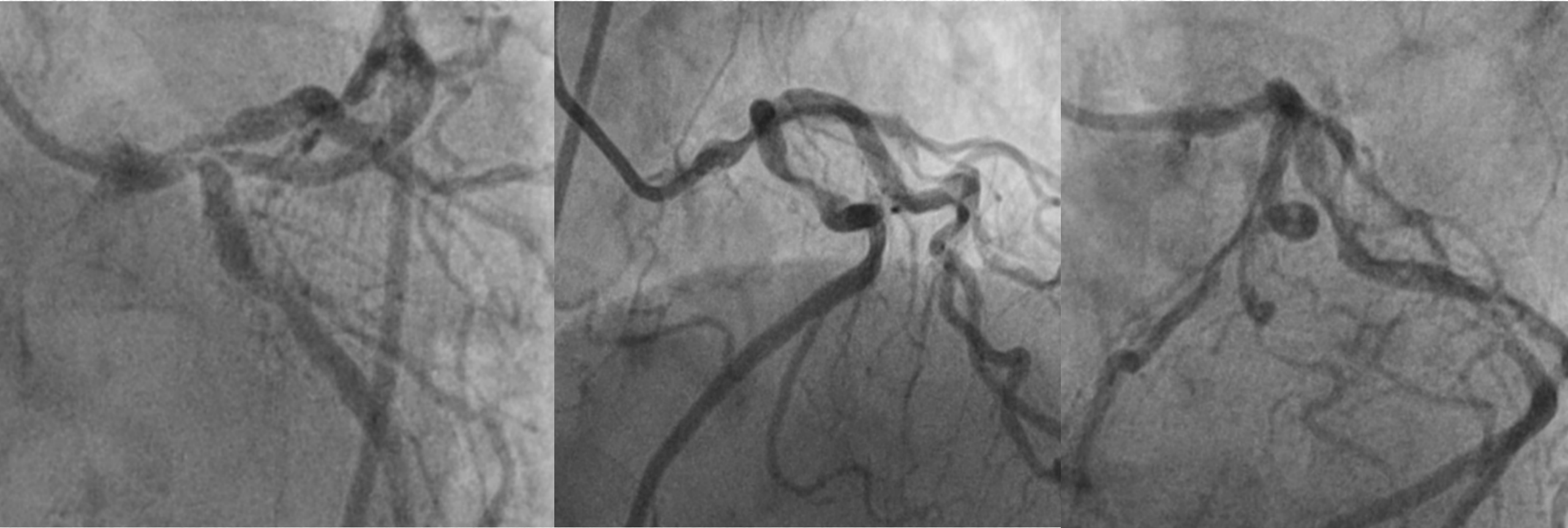
Porcelain aorta and total occlusion of the left CCA and subclavian / left vertebral artery and massive coronary calcification

- A 76 years old male
- Impaired LV function (EF 40%) and renal GFR (40ml/min)
- Previous bilateral CEA with total occlusion of the left CEA

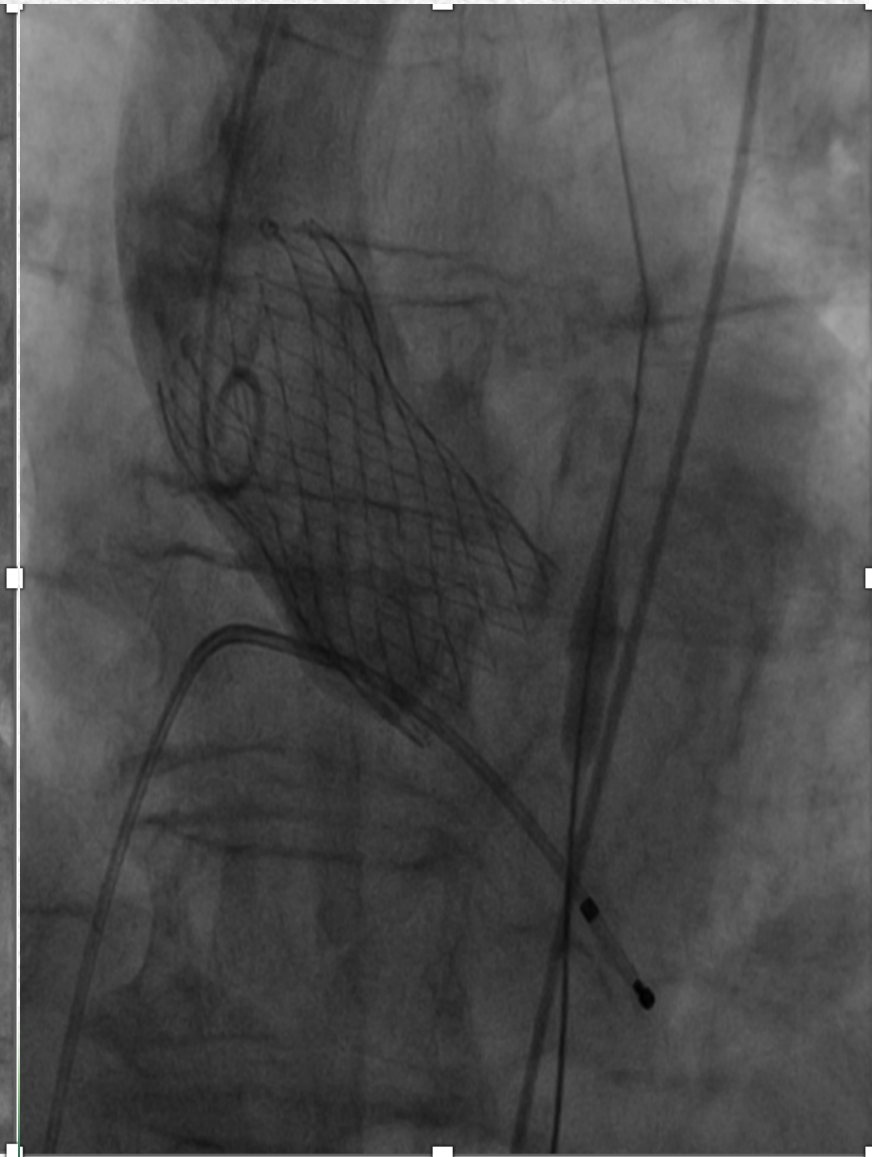
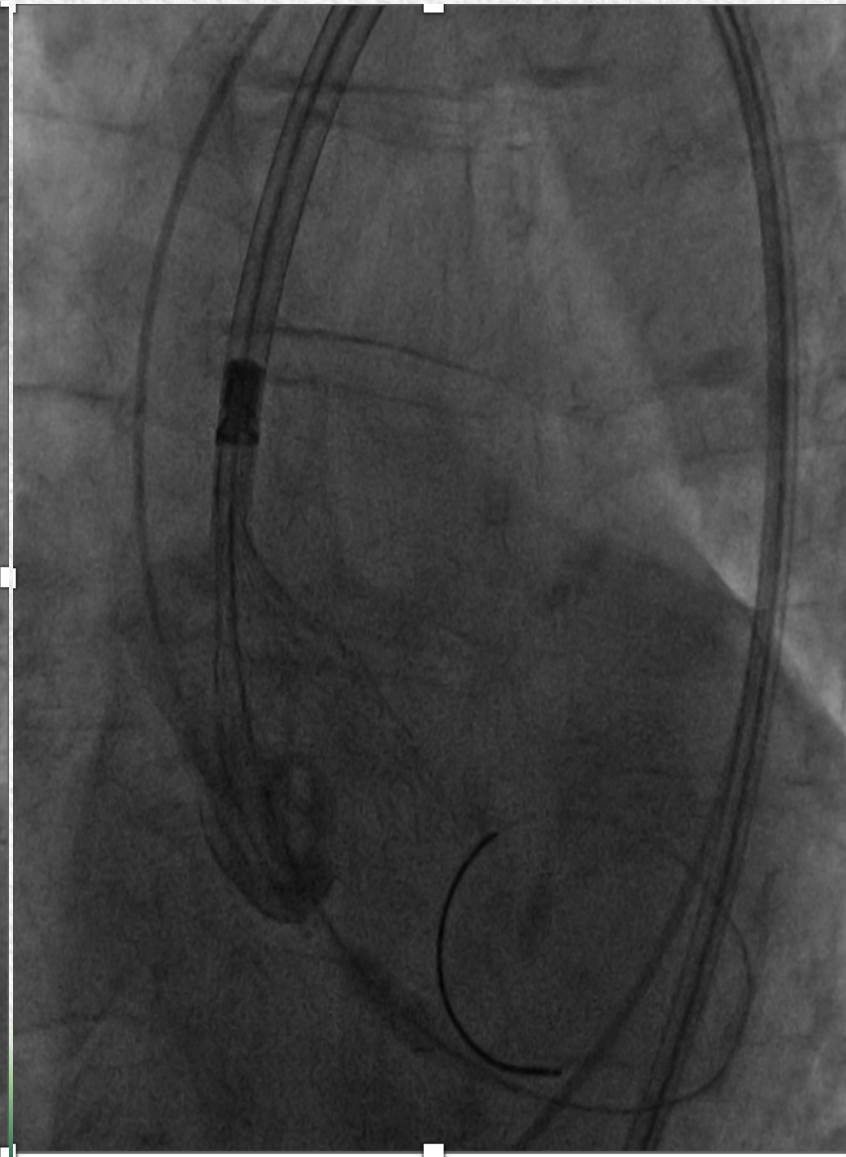
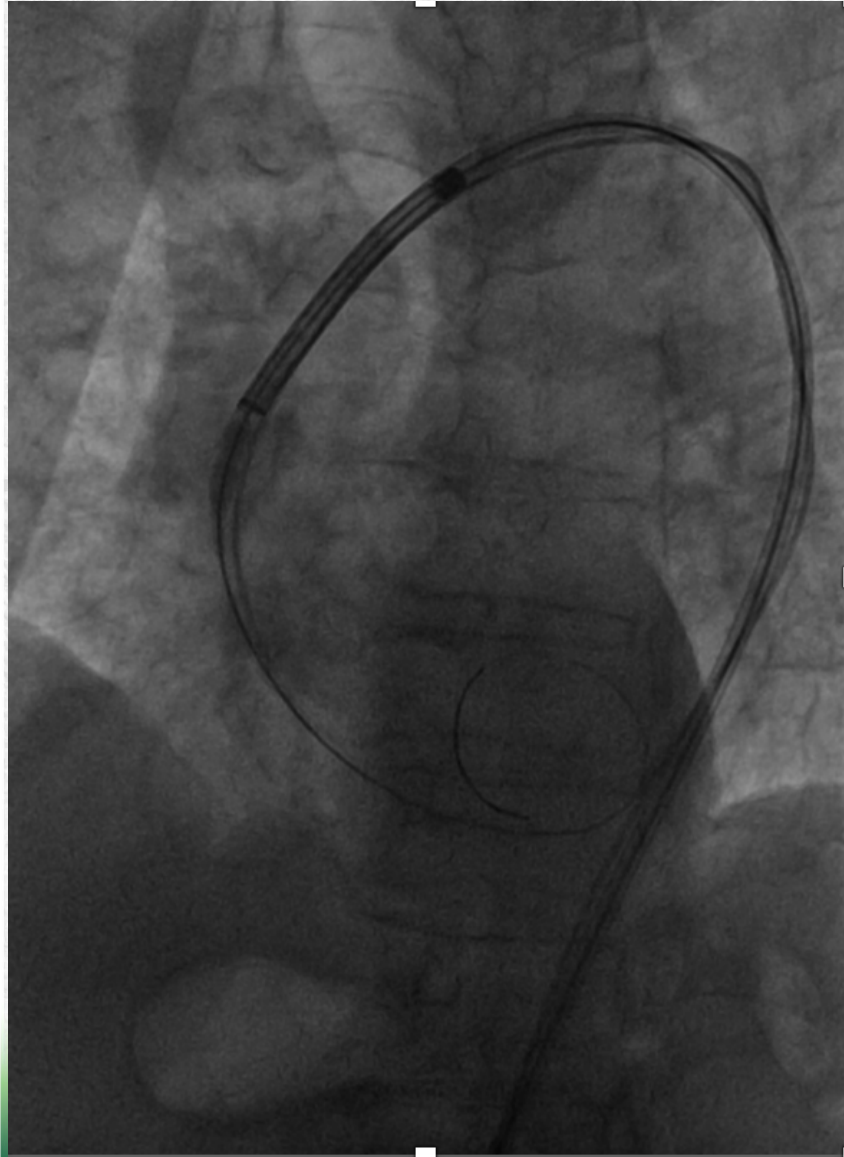
Severe AVS symptomatic for angina (CCS class II and worsening to class III + NYHA class II)



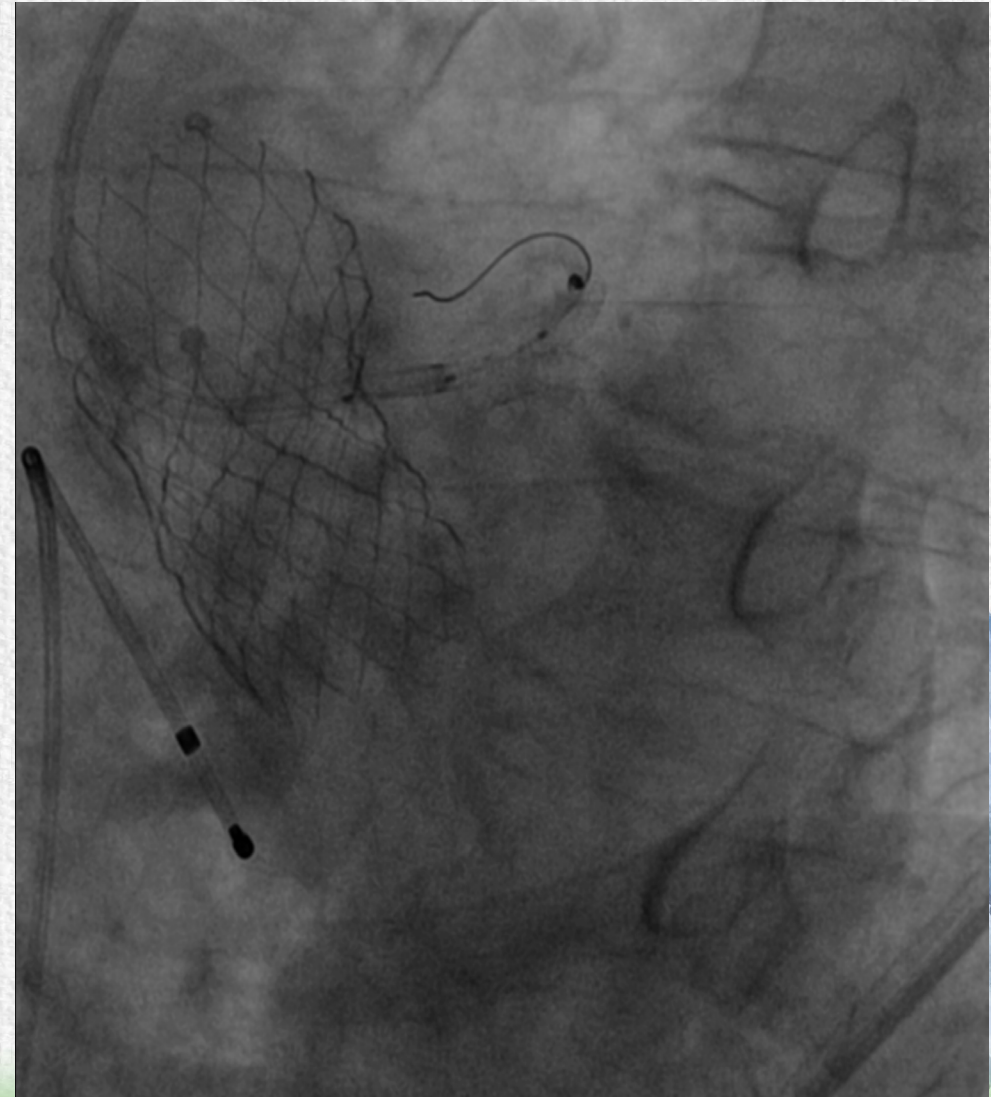
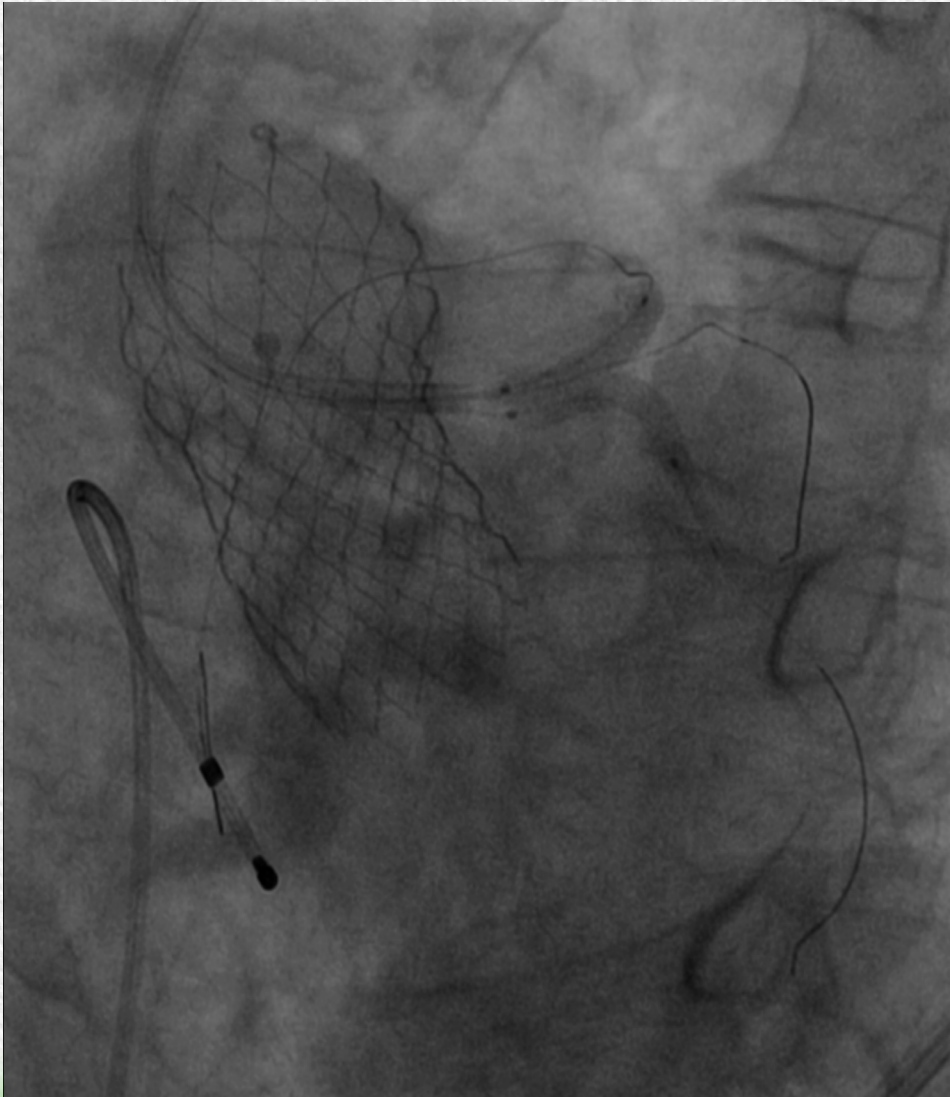
Coronary angiogram



Trans-femoral catheter valve Implantation

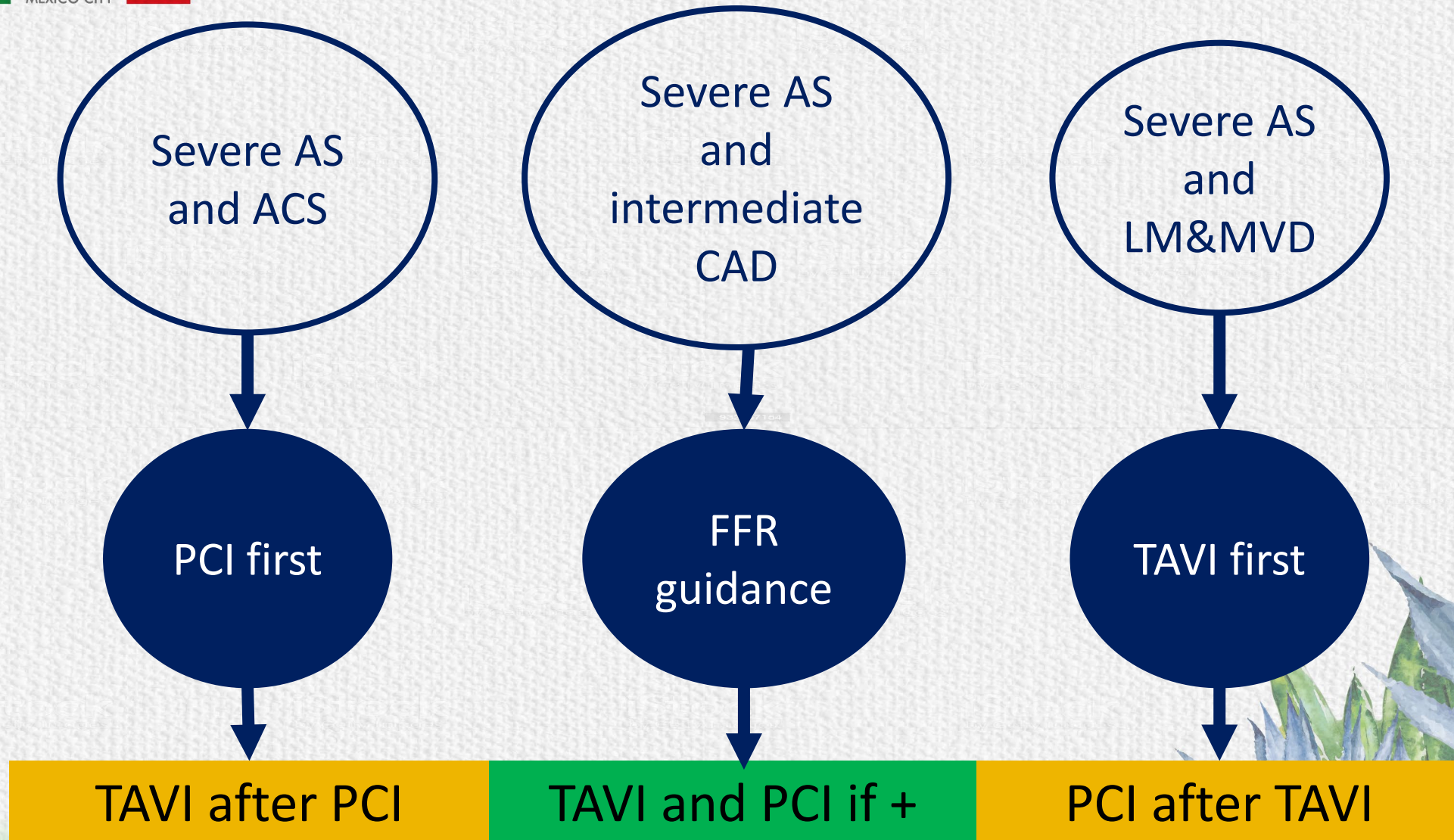


and then, the left main





Conclusions

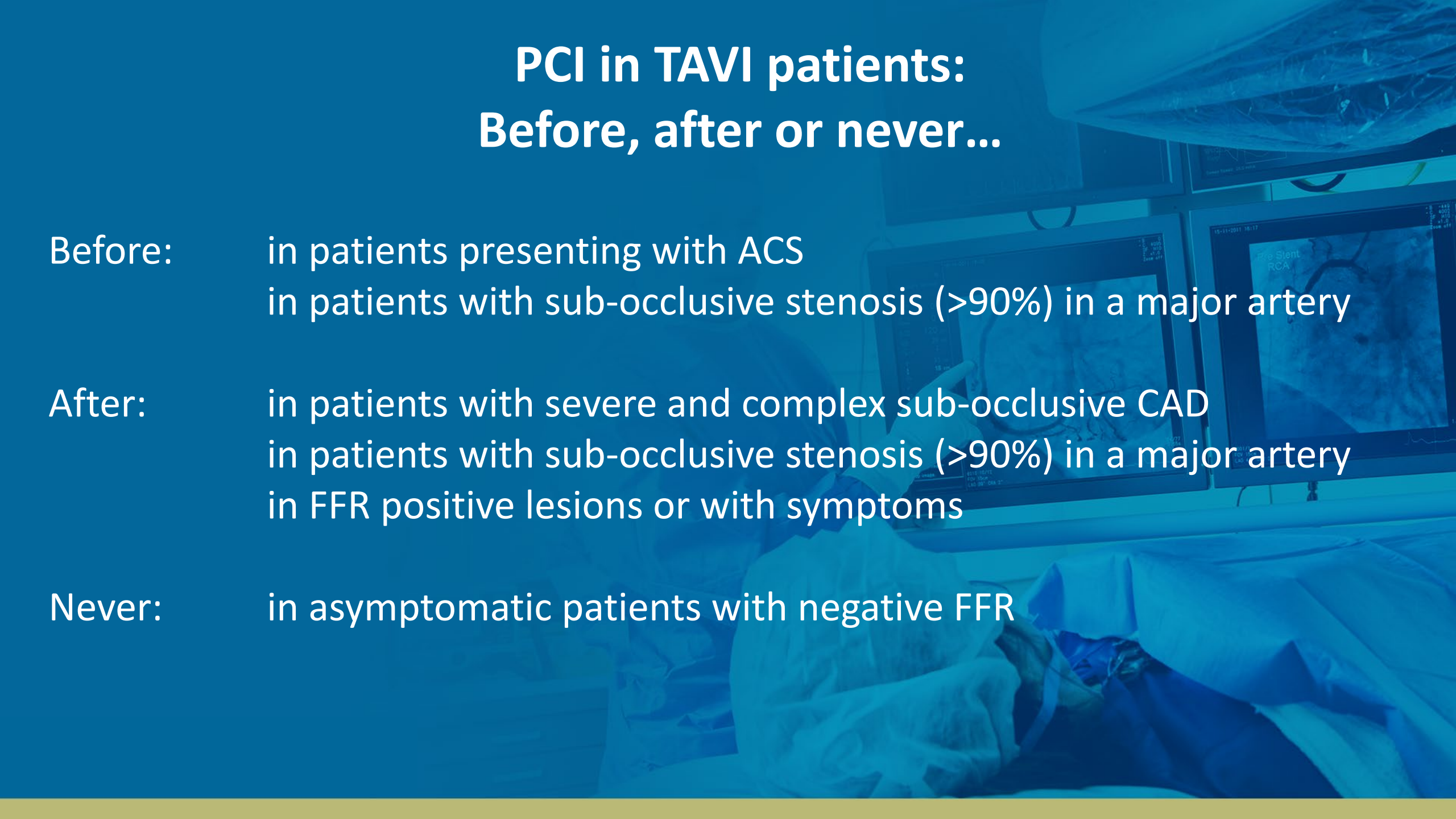


PCI in TAVI patients: Before, after or never...

Stable CAD in AS should follow the same recommendations as stable CAD based on ISCHEMIA and FAME... after removing the stenotic valve:

- 1 : Improve cardiac output first by removing the aortic stenosis
- 2 : Define significant coronary lesions with physiology after valve replacement
- 3: Limit contrast, use IVI
- 4: In calcified CAD use appropriate devices and IVI
- 5: Apply HBR concepts

PCI in TAVI patients: Before, after or never...

- 
- Before:**
- in patients presenting with ACS
 - in patients with sub-occlusive stenosis (>90%) in a major artery
- After:**
- in patients with severe and complex sub-occlusive CAD
 - in patients with sub-occlusive stenosis (>90%) in a major artery
 - in FFR positive lesions or with symptoms
- Never:**
- in asymptomatic patients with negative FFR

Gracias

