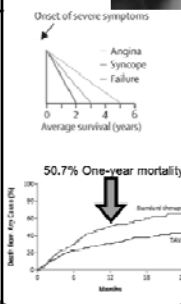


Transcatheter Aortic Valve Replacement

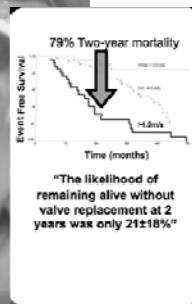
Juan Crestanello, MD
Interim Director, Division of Cardiac Surgery
Associate Professor
Division of Cardiac Surgery
The Ohio State University Wexner Medical Center

Aortic Stenosis

Symptomatic



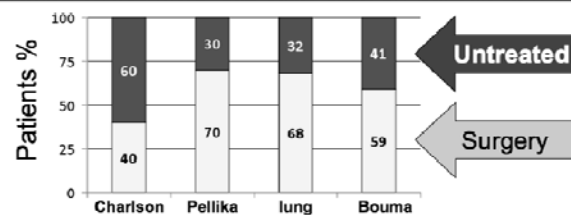
Asymptomatic



Leon ME et al. NEJM 2010, Otto CM et al. Circulation 1997

Transcatheter AVR

Patients with Severe AS Treated with Surgery



- Advanced age
- Frailty
- Left ventricular dysfunction
- Increased "surgical risk"
- Multiple comorbidities

Transcatheter AVR

Edwards Sapien



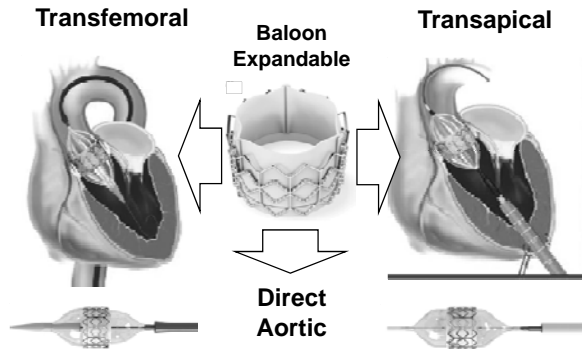
FDA approved

Medtronic CoreValve

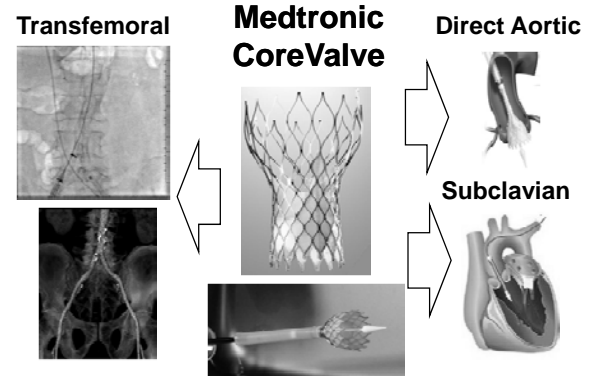


Inoperable FDA approved
High risk: in trials

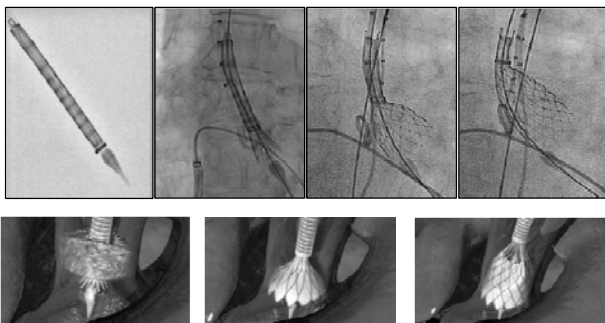
Edwards Sapien TAVR



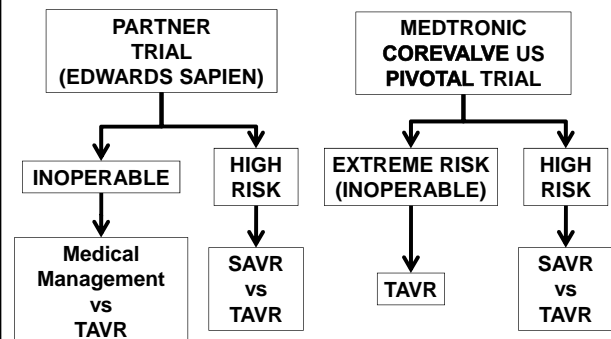
Transcatheter AVR



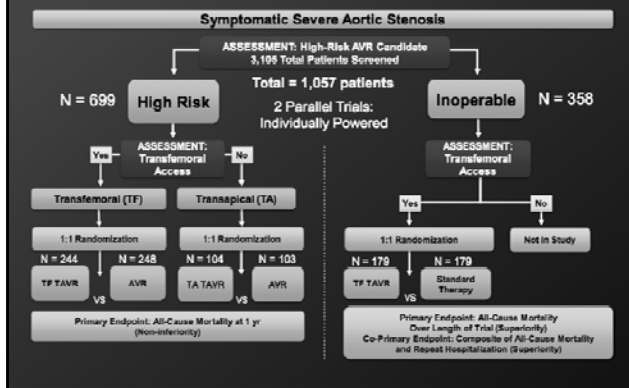
CoreValve Deployment Self Expandable



The Trials

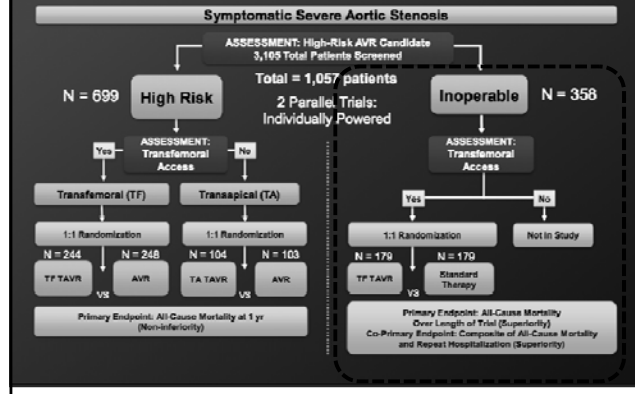


PARTNER TRIAL



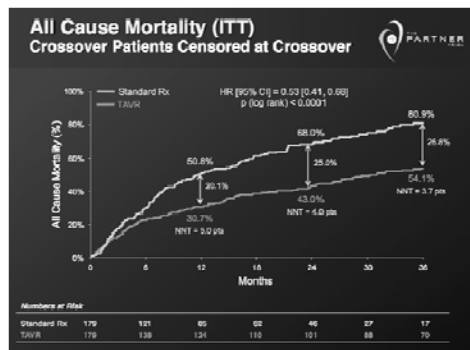
Permission to use Partner Trial information and images provided courtesy of Edwards Lifesciences.

PARTNER TRIAL



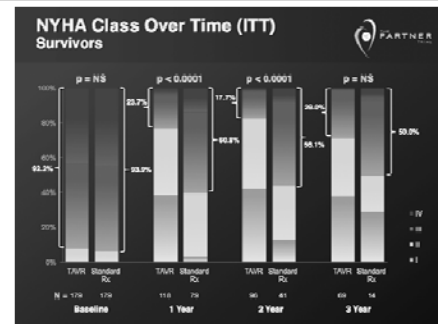
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Inoperable Aortic Stenosis TAVR vs. Medical Management



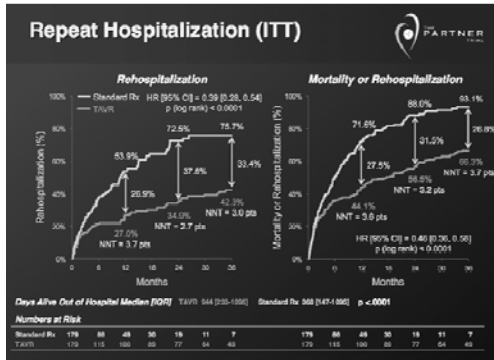
Permission to use Partner Trial information and images provided courtesy of Edwards Lifesciences.

Inoperable Aortic Stenosis TAVR vs. Medical Management



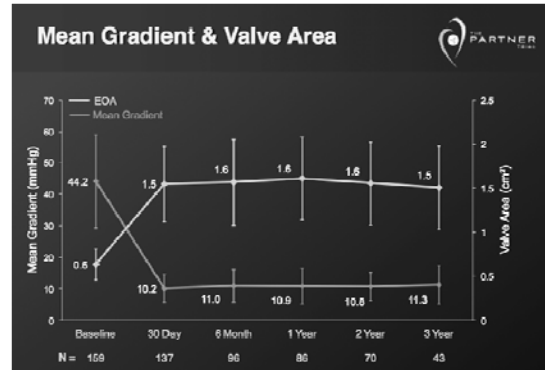
Permission to use Partner Trial information and images provided courtesy of Edwards Lifesciences.

Inoperable Aortic Stenosis TAVR vs. Medical Management



Permission to use Partner Trial information and images provided courtesy of Edwards Lifesciences.

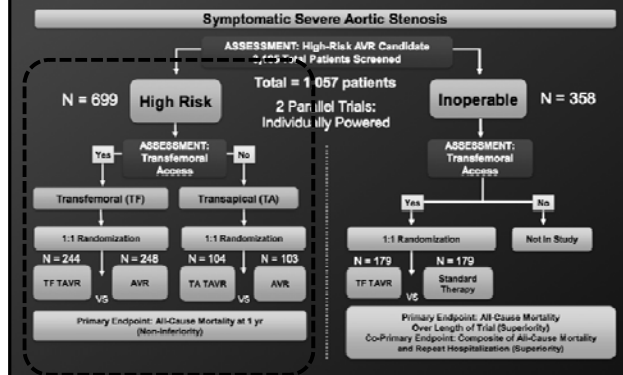
Inoperable Aortic Stenosis TAVR vs. Medical Management



Inoperable Aortic Stenosis TAVR vs. Medical Management

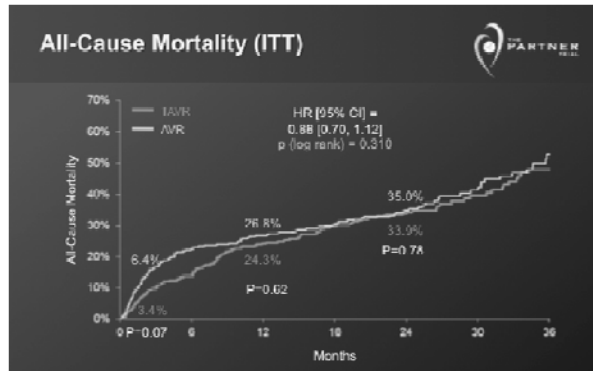
- Medical management did not change the natural history of aortic stenosis
- TAVR relieved aortic stenosis
- TAVR was superior to medical therapy:
 - Decreased all cause mortality
 - Decreased cardiovascular mortality
 - Decreased rate of rehospitalization
 - Improved NYHA functional class

PARTNER TRIAL



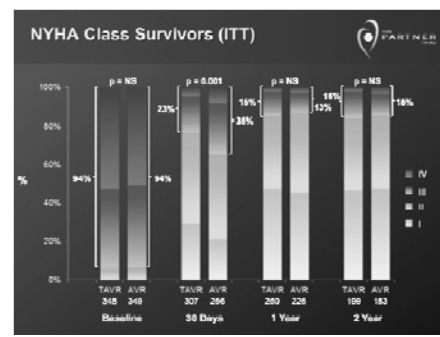
Permission to use Partner Trial information and images provided courtesy of Edwards Lifesciences.

High Risk Aortic Stenosis TAVR vs. AVR



Permission to use Partner Trial information and images provided courtesy of Edwards Lifesciences.

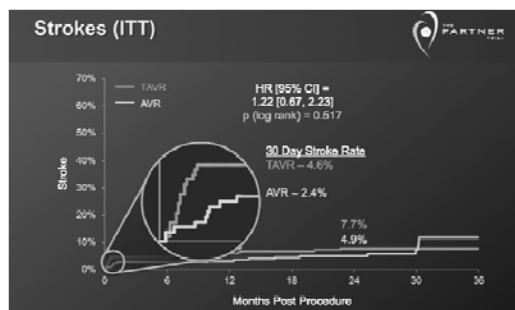
High Risk Aortic Stenosis TAVR vs. AVR



80% on NYHA class I or II

Permission to use Partner Trial information and images provided courtesy of Edwards Lifesciences.

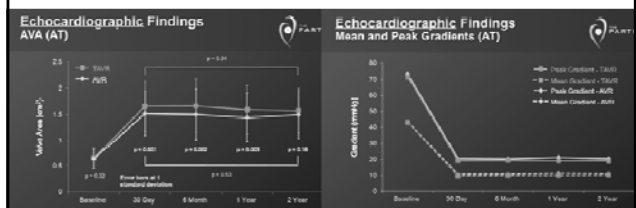
High Risk Aortic Stenosis TAVR vs. AVR



TAVR increased risk of early stroke

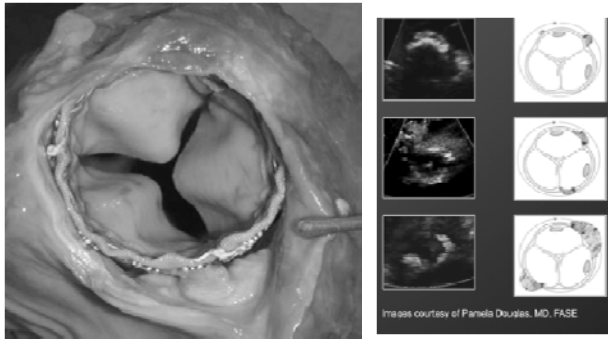
Permission to use Partner Trial information and images provided courtesy of Edwards Lifesciences.

High Risk Aortic Stenosis TAVR vs. AVR

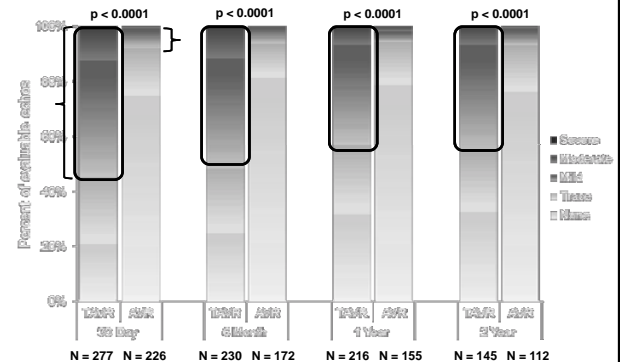


Permission to use Partner Trial information and images provided courtesy of Edwards Lifesciences.

High Risk Aortic Stenosis Paravalvular Aortic Regurgitation

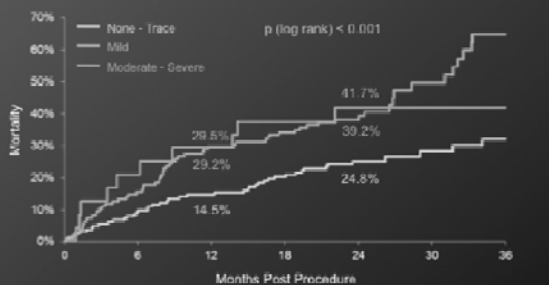


Paravalvular Aortic Regurgitation



High Risk Aortic Stenosis Paravalvular Aortic Regurgitation

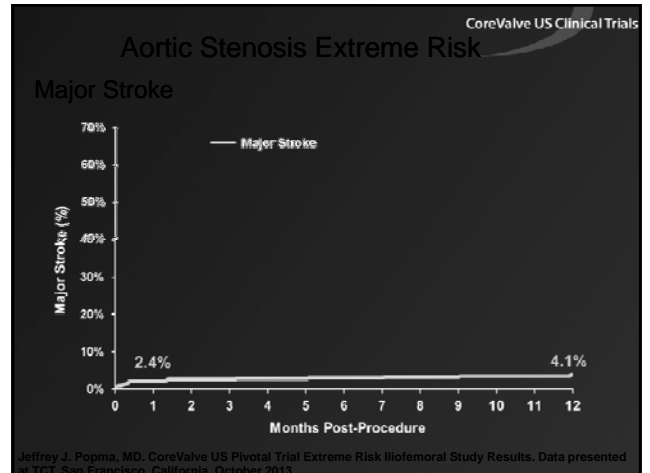
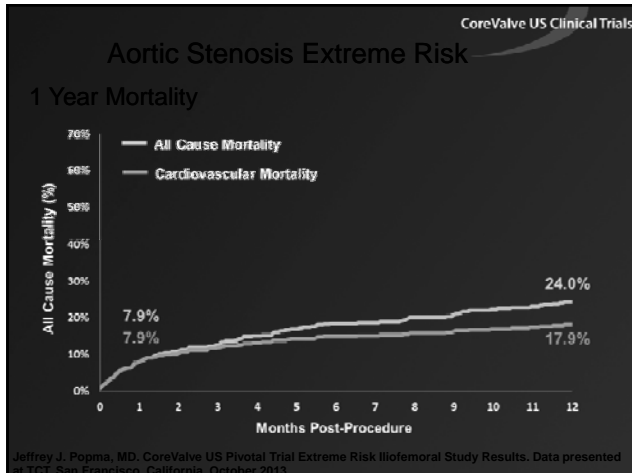
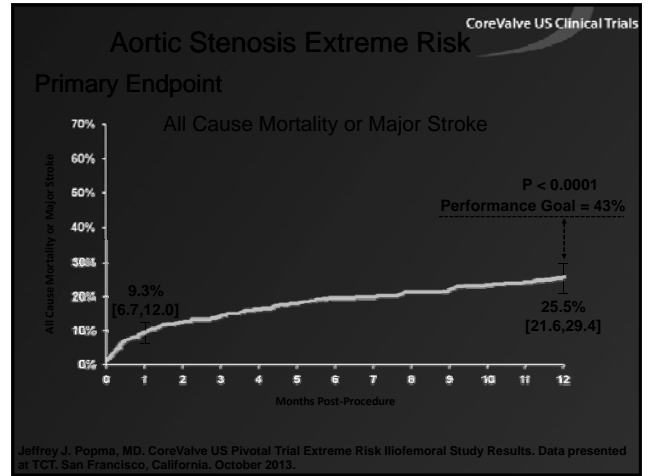
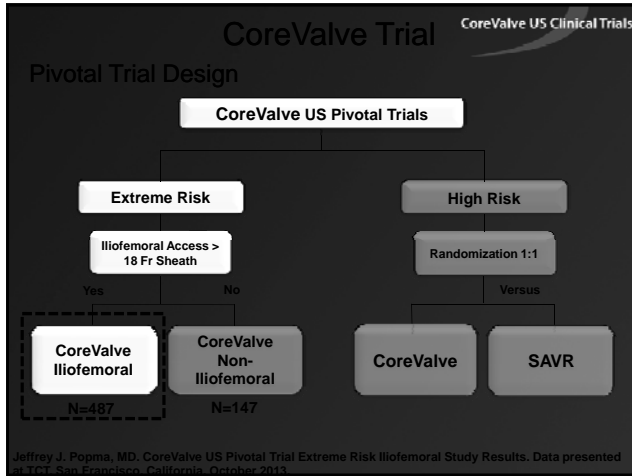
Mild Paravalvular AR and Mortality TAVR Patients (AT)



Permission to use Partner Trial information and images provided courtesy of Edwards Lifesciences.

High Risk Aortic Stenosis TAVR vs. SAVR

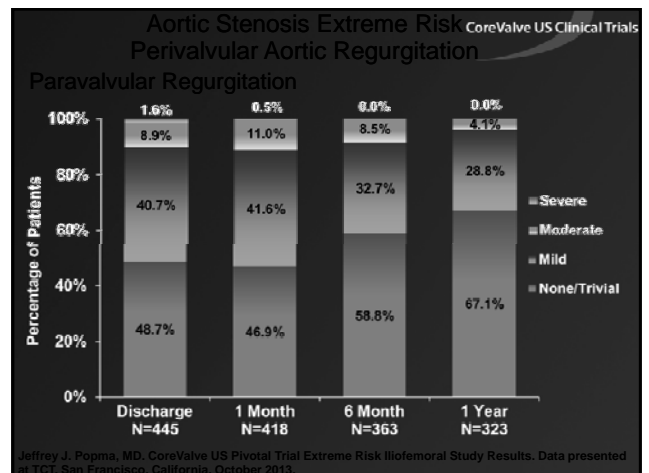
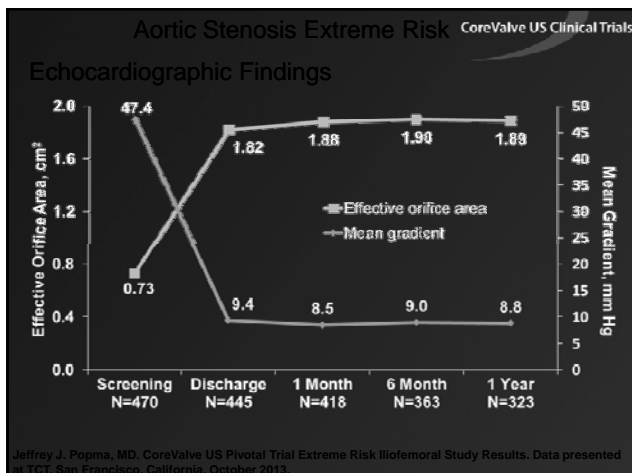
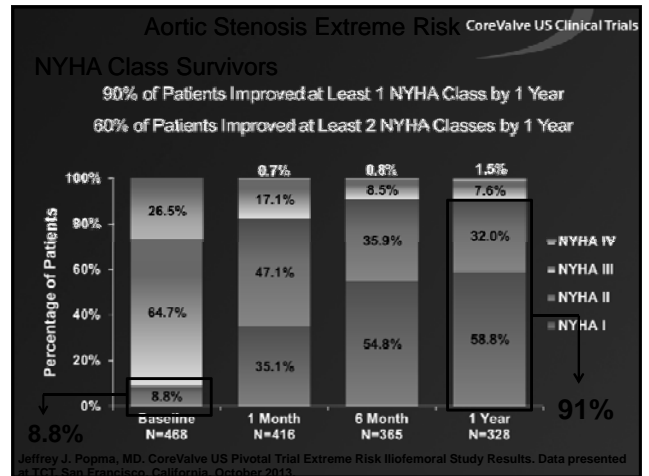
- TAVR and SAVR effectively relieved AS
- Mortality was similar up to 2 years
- NYHA class was similar
- 30 day stroke rate was higher in TAVR
- TAVR was associated with PVL
- Mild moderate and severe PVL resulted in increased mortality

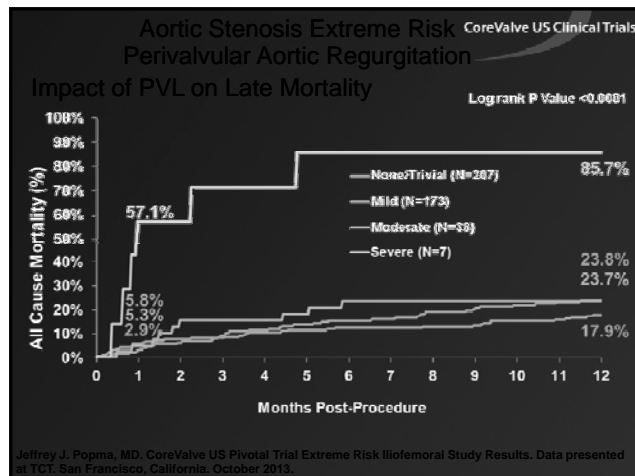
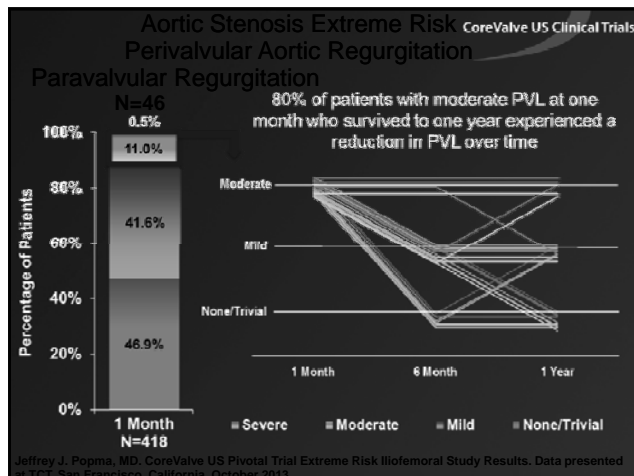


Aortic Stenosis Extreme Risk CoreValve US Clinical Trials		
Secondary Endpoints		
Events*	1 Month	1 Year
Any Stroke, %	3.9	6.7
Major, %	2.4	4.1
Minor, %	1.7	3.1
Myocardial Infarction, %	1.3	2.0
Reintervention, %	1.3	2.0
VARC Bleeding, %	35.1	41.4
Life Threatening or Disabling, %	11.7	16.6
Major, %	24.1	27.6
Major Vascular Complications, %	8.3	8.5
Permanent Pacemaker Implant, %	22.2	27.1
Per ACC Guidelines, %	17.4	19.9

* Percentages obtained from Kaplan Meier estimates

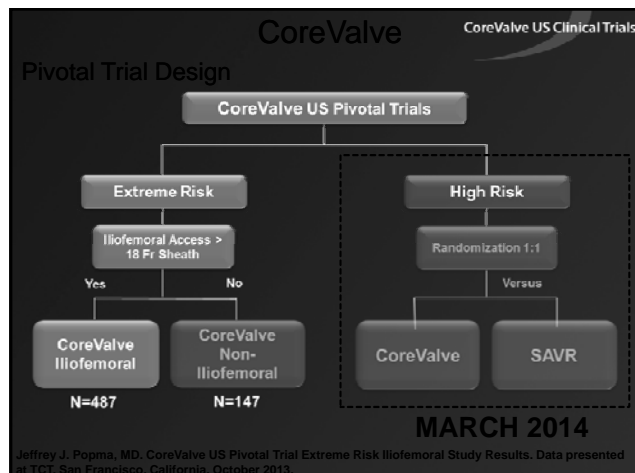
Jeffrey J. Popma, MD. CoreValve US Pivotal Trial Extreme Risk Iliofemoral Study Results. Data presented at TCT, San Francisco, California, October 2013.

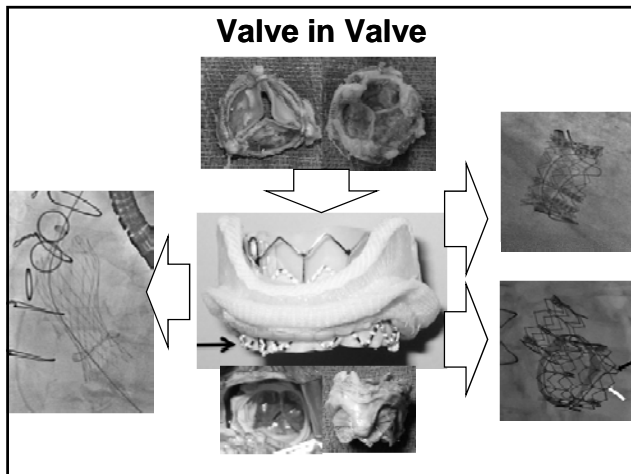




CoreValve Extreme Risk

- TAVR relieved aortic stenosis
- TAVR reduced mortality and stroke rate at one year
- Low rate of stroke
- Mild and moderate PVL was not associated with increased mortality





Ongoing and Upcoming Trials

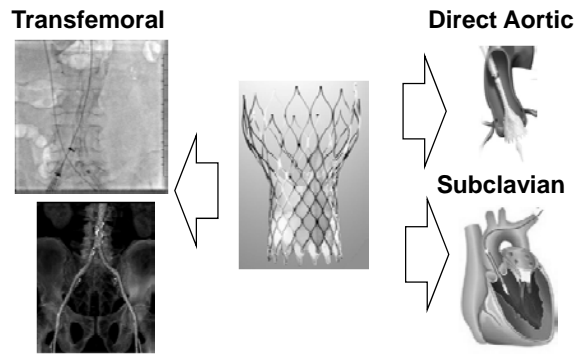
Intermediate Risk	High Risk	Extreme Risk
SURTAVAL Trial	CoreValve Pivotal Trial	CoreValve Pivotal Trial
	Expanded Use	Expanded Use
PARTNER II	PARTNER A	PARTNER B



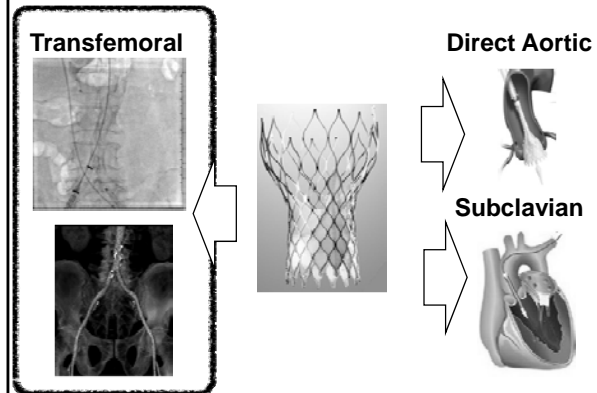
TAVR with Medtronic CoreValve: Femoral Approach

Barry George, MD
 Director
 Advanced Catheter-Based Therapeutics
 and Structural Heart Disease
 Associate Professor – Clinical
 Department of Cardiovascular Medicine
 The Ohio State University Wexner Medical Center

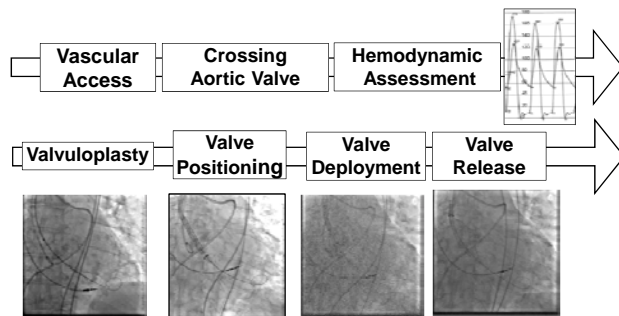
CoreValve Delivery Options



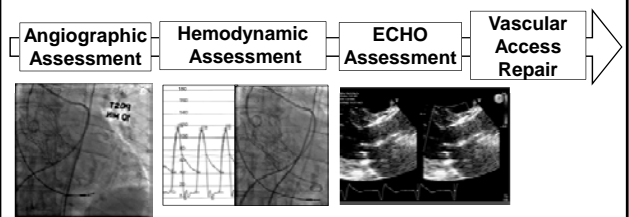
CoreValve Delivery Options



Transfemoral Step by Step



Transfemoral Step by Step



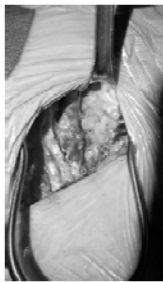
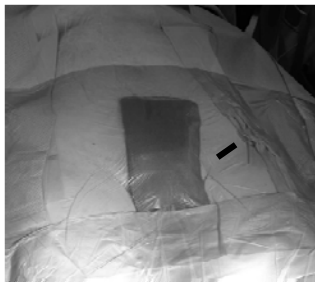
Vascular Access

Vascular Access

- Femoral artery cut down (percutaneous) for delivery sheath
- Contralateral femoral artery placement of 6F sheath.

Vascular Access

- Femoral artery cut down (percutaneous) for delivery sheath



Vascular Access

- Femoral artery cut down (percutaneous) for delivery sheath



5-0 Prolene purse string Needle and guidewire

Administer Heparin to achieve ACT>300 s

Vascular Access

- Femoral artery cut down (percutaneous) for delivery sheath



6 French Sheath placement

Pigtail placement into descending thoracic aorta

Vascular Access

- Femoral artery cut down (percutaneous) for delivery sheath

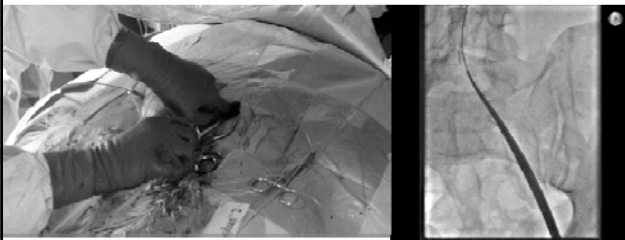


Insertion Amplatz Super Stiff guidewire over pigtail catheter



Vascular Access

- Femoral artery cut down (percutaneous) for delivery sheath



Arterial dilatation with 12,14 and 18 F dilators over Amplatz Super Stiff guidewire

Vascular Access

- Femoral artery cut down (percutaneous) for delivery sheath



Placement of 18 F sheath



Vascular Access

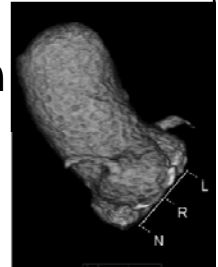
- Contralateral femoral artery placement of 6F sheath.



Percutaneous
arterial access

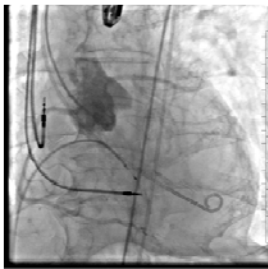
6F Sheath and pigtail

Implant projection

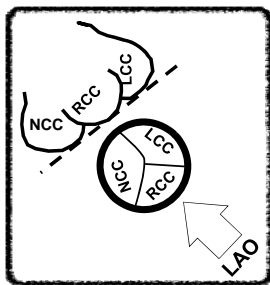


Determination of implantation projection

- Alignment of all 3 cusps of aortic valve in a single plane.



Pigtail advance to non-
coronary sinus

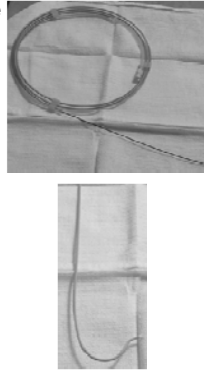
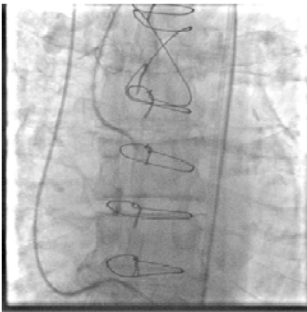


Crossing Aortic Valve

Initial Hemodynamic Assessment

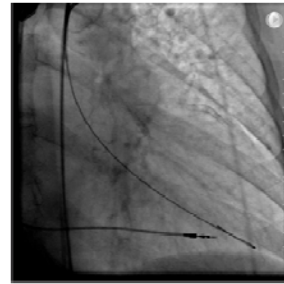
Crossing the Aortic Valve

- AL 2 catheter with straight guidewire
- AP projection



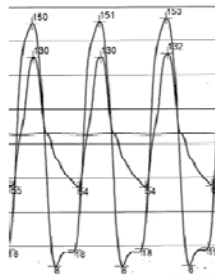
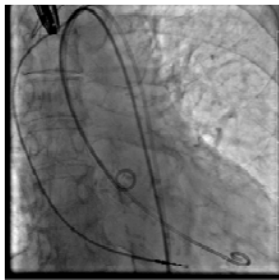
Crossing the Aortic Valve

- Exchange guidewire
- ROA projection
- Pigtail placement in LV apex



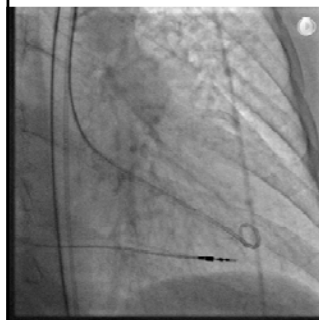
Initial Hemodynamic Assessment

- LV and Aortic pressure



Placement of stiff guidewire in the LV apex

- RAO projection
- Amplatz Super Stiff ST1 (pigtail configuration)



Aortic Valvuloplasty

Aortic balloon valvuloplasty

- Balloons:
 - Z MED (NuMED Inc)
 - NuCLEUS Ballon (NuMED Inc)
 - True Balloon (Loma Vista Medical)
- Balloon size: smaller diameter of aortic annulus
- Pacing rate: 160-180 bpm
- Screw-in temporary pacing lead
- If patient has a PPM,transvenous pacer in the OR.
- No valvuloplasty in:
 - Low EF patients <30-35%
 - Large plaque-calcifications in the aorta or sinus of Valsalva
 - Valve in valve

Aortic balloon valvuloplasty

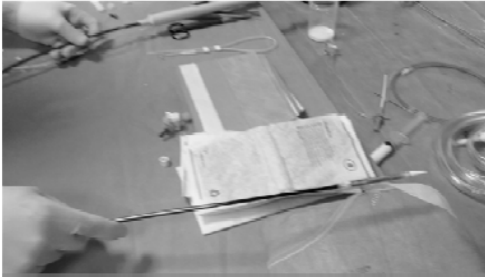
- Pacer on
- Balloon up
- Ballon down
- Pacer off



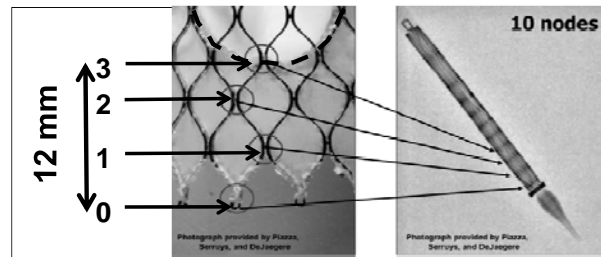
CoreValve Implantation

CoreValve Implantation

- CoreValve is loaded into the delivery system before the valvuloplasty

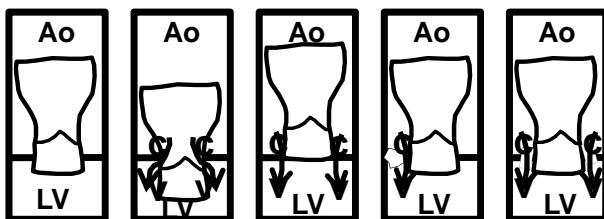


CoreValve in the delivery capsule



Optimal implantation depth 2 to 6 mm

Depth of implantation



Correct
No AI

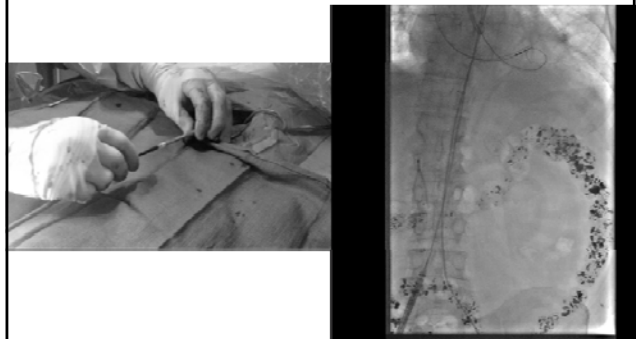
Too deep
AI
AV block

Too shallow
AI
Pop out

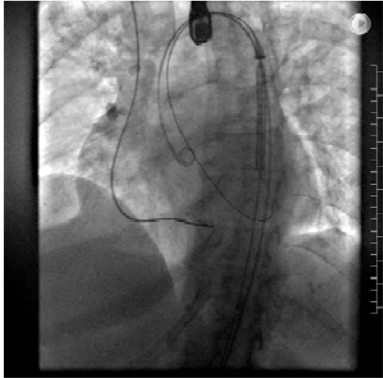
Irregular
annulus
AI

Large
annulus
AI

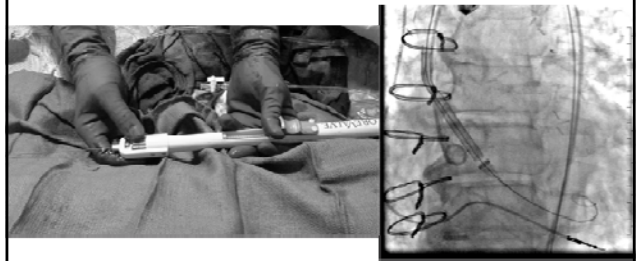
CoreValve insertion and advancement into the aortic root



CoreValve insertion and advancement into the aortic root

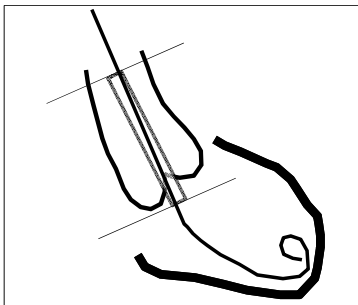


CoreValve Deployment

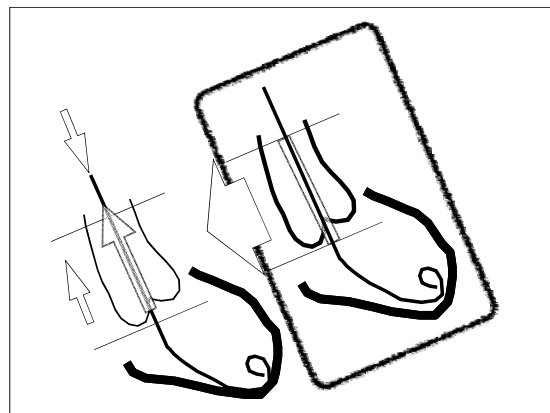


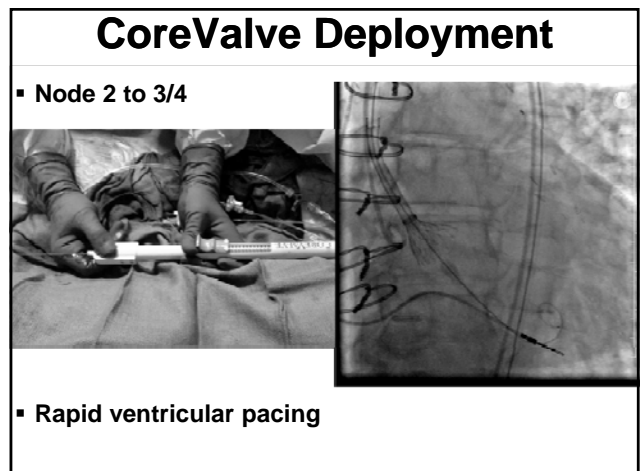
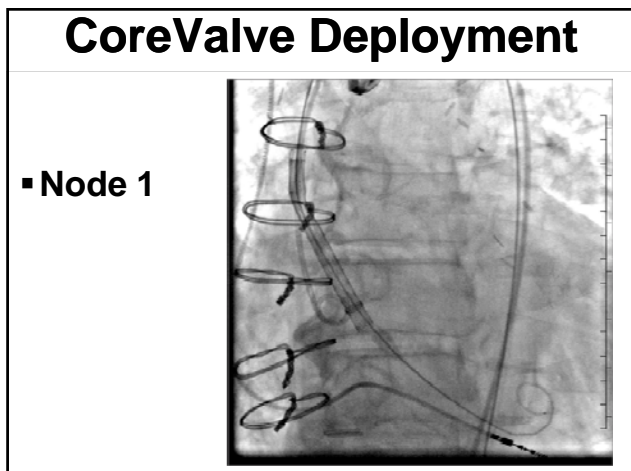
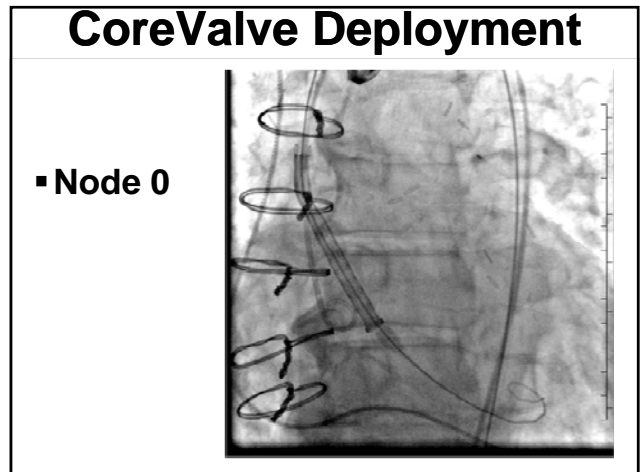
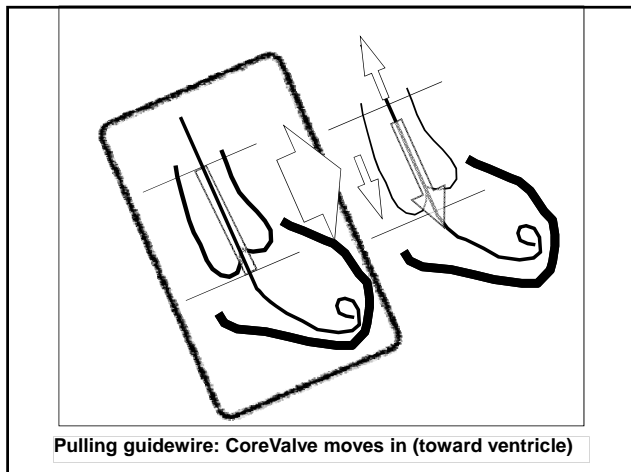
CoreValve Deployment

- Position adjustments
 - Operator #1: Pulling or pushing delivery catheter
 - Operator #2: Pulling or pushing guidewire



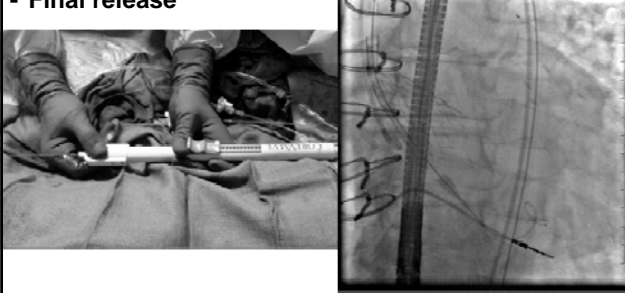
Pushing guidewire: CoreValve moves out (toward the aorta)





CoreValve Deployment

- Final release



Nosecone

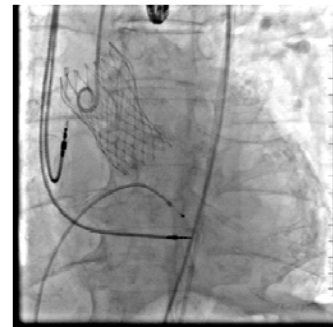
- Recapture of the nosecone in the descending aorta and removal of the delivery catheter



Valve Function Assessment

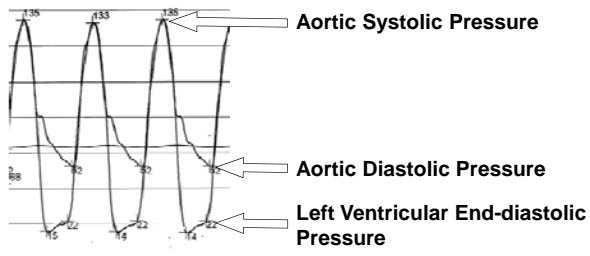
Valve Function Assessment

- Angiography



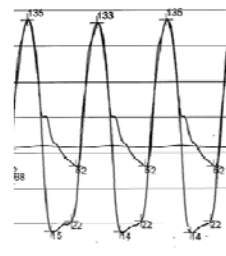
Valve Function Assessment

▪ Hemodynamics



Valve Function Assessment

▪ Hemodynamics

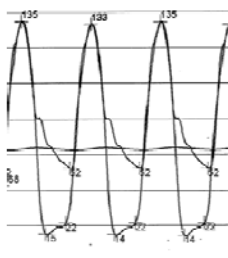


$$\text{Aortic Regurgitation Index} = \frac{\text{Ao Diastolic} - \text{LVEDP}}{\text{Ao Systolic}} > 0.25$$

Aortic Diastolic Pressure
>40 mmHg

Valve Function Assessment

▪ Hemodynamics

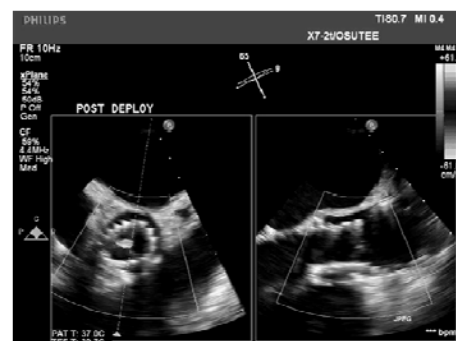


$$\text{Aortic Regurgitation Index} = \frac{\text{Ao Diastolic} - \text{LVEDP}}{\text{Ao Systolic}} > 0.25$$

$$\frac{(82-22)}{135} = 0.29$$

Valve Function Assessment

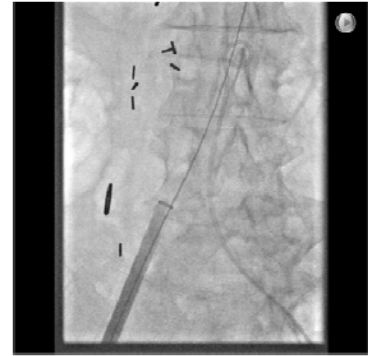
▪ Echocardiography



Sheath Removal Access Repair

Sheath removal

- Over stiff guidewire
- Inject contrast through sheath
- Contralateral pigtail in the aortic bifurcation



Sheath removal

- If iliac artery rupture:
 - Advance sheath and dilator over stiff guidewire
 - Place Coda aortic occlusion balloon through contralateral pigtail (arterial sheath may need to be exchange)
 - Place coverstent through ipsilateral sheath

Arteriotomy repair

