

# The Upstream Management of Aortic Stenosis

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# Disclosure Statement of Financial Interest

Within the past 12 months, I or my spouse/partner have had a financial interest/arrangement or affiliation with the organization(s) listed below.

## Affiliation/Financial Relationship

- Grant/Research Support
- Scientific Advisory Board
- Executive Physician Council

## Company

- Edwards Lifesciences, Abbott
- Medtronic
- Boston Scientific Corp



# Upstream Management of Aortic Stenosis

?



# Upstream Management of CAD

- Pathophysiology
- Risk Factor Modification (i.e. lipids, smoking etc)
- Detection (CAC, CCTA, stress testing, Invasive angio)
- Criterion for Treatment (symptoms, anatomy, FFR)
- Treatment Options (Medical, PCI, CABG)
- Effectiveness and Durability of Treatment Options

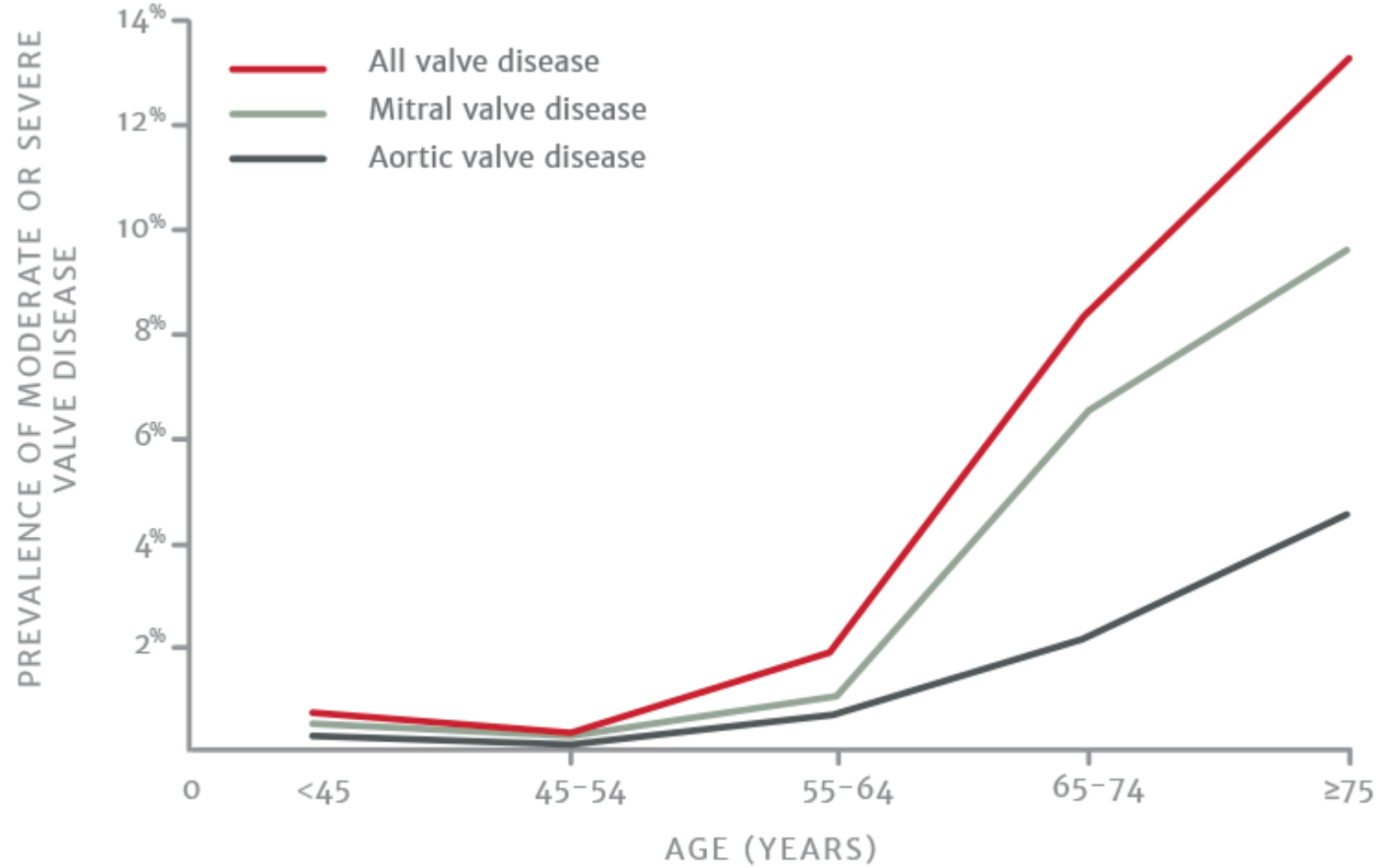


# Upstream Management of Aortic Stenosis

- Pathophysiology
- Risk Factors

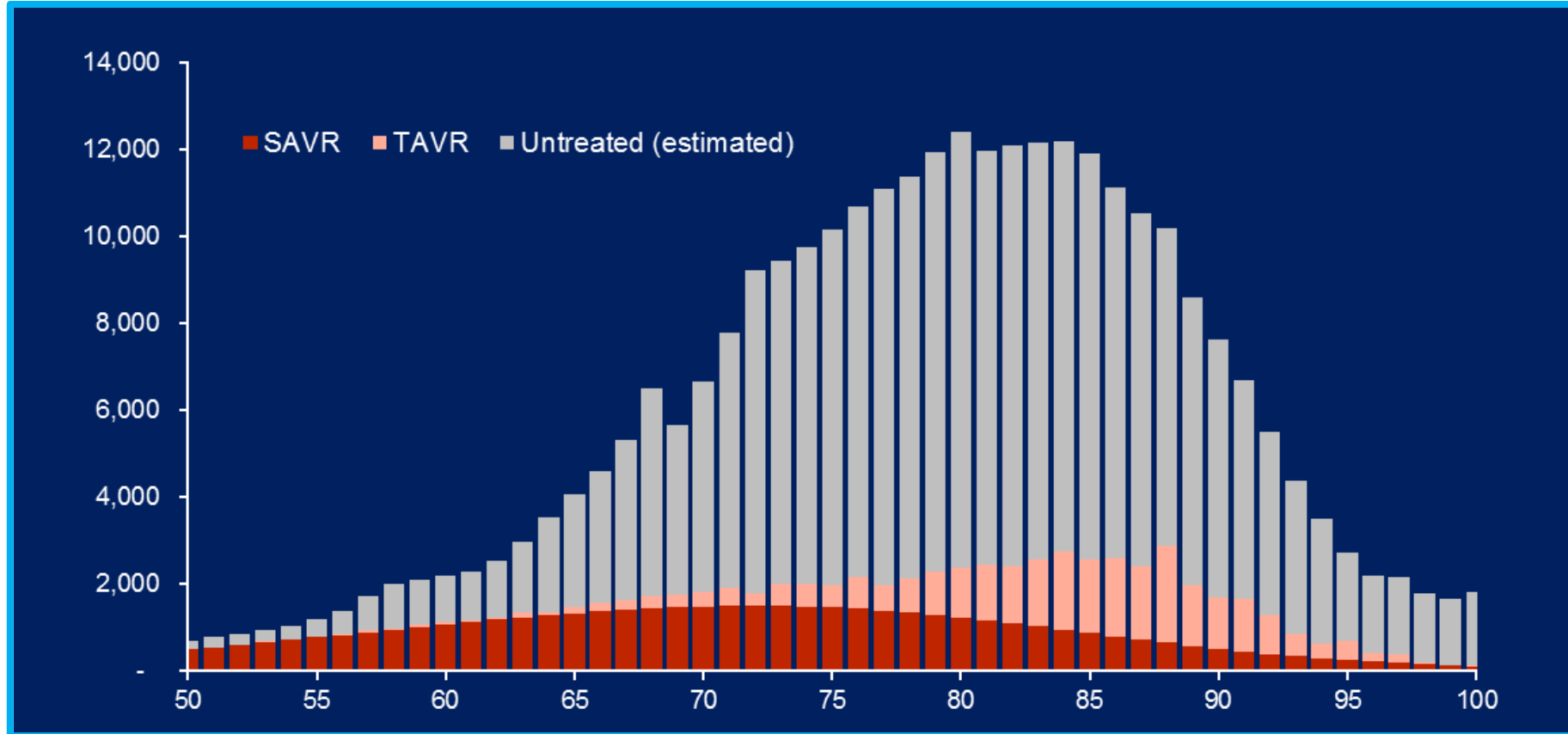


# Incidence of AS according to Age

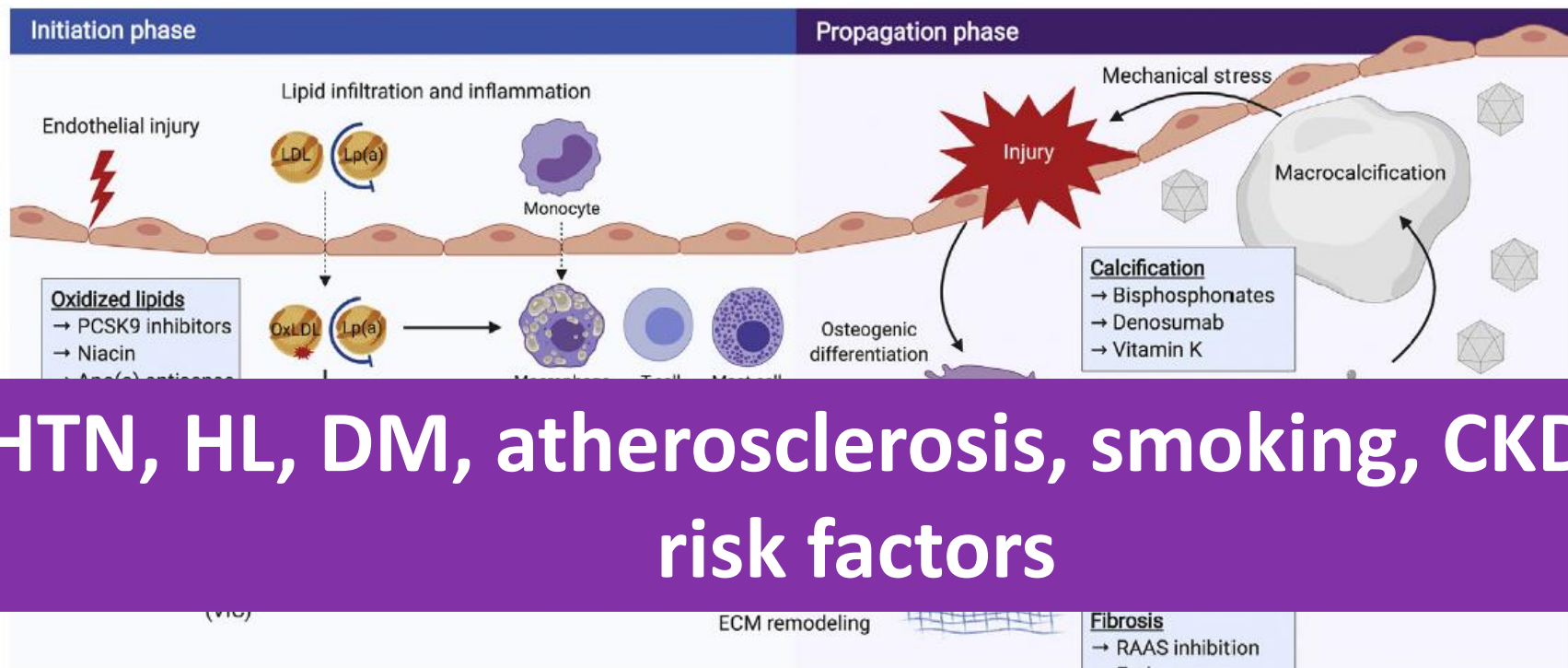


# Incidence of AS according to Age (6% above age 75)

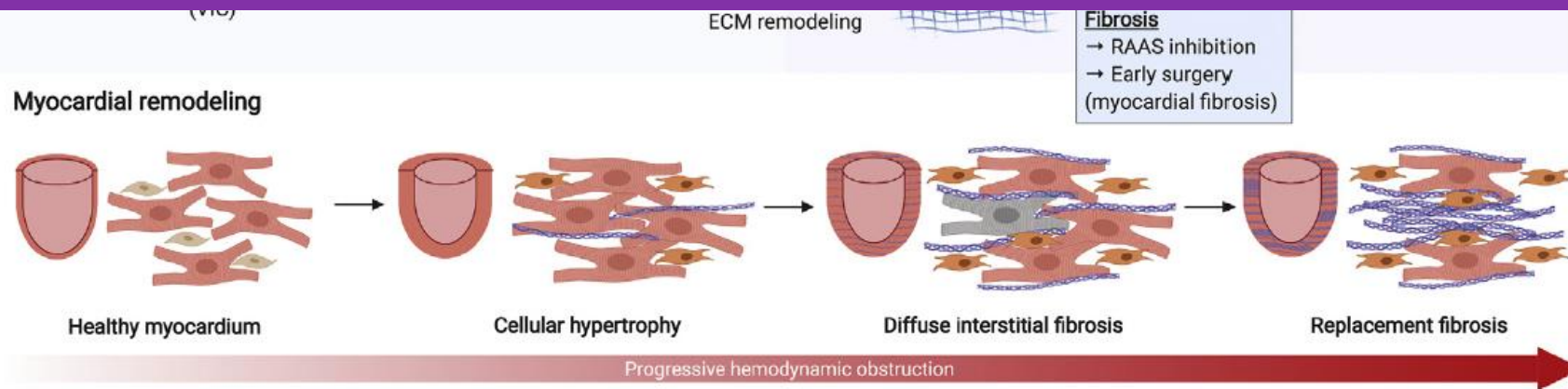
## 2015 Severe Symptomatic AS Patients in the U.S.



# Pathophysiology of Aortic Stenosis



**HTN, HL, DM, atherosclerosis, smoking, CKD are risk factors**





# Statin Trials to Treat CAVD

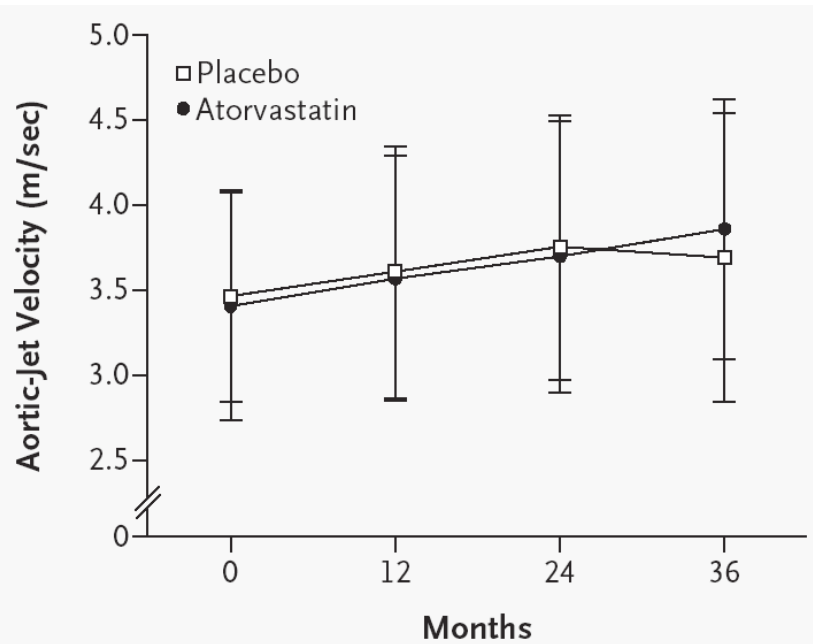
Study (year)	Participants	Outcomes
Retrospective study (2001) <sup>73</sup>	180 participants aged >60 years	Significant decrease in peak systolic pressure gradient (a marker of aortic valve function) in patients taking statins
Retrospective study (2001) <sup>75</sup>	174 patients with mild-to-moderate CAVD (57 statin-treated, 117 not taking statins)	Patients taking statins had a slight improvement in aortic valve remodelling
Prospective analysis (2002) <sup>82</sup>	156 patients (38 statin-treated, 118 not receiving any lipid-lowering treatment)	Patients taking statins had a slight improvement in aortic valve area
SALTIRE trial (2005) <sup>83</sup>	Randomized, double-blind trial of 155 patients given atorvastatin or placebo (mean follow-up 25 months)	No significant difference in aortic jet velocity or valve calcification between atorvastatin and placebo
RAAVE trial (2007) <sup>74</sup>	Prospective study of 121 patients with moderate-to-severe CAVD (61 patients received rosuvastatin, 60 received no treatment)	Statin treatment seemed to slow haemodynamic progression of CAVD, as indicated by changes in aortic jet velocity
SEAS trial (2008) <sup>84</sup>	Randomized, double-blind trial of 1,873 patients with mild-to-moderate asymptomatic aortic stenosis receiving either simvastatin or placebo daily	Statin treatment did not reduce cardiovascular events associated with CAVD
SEAS follow-up (2010) <sup>85</sup>	1,763 patients from the SEAS trial divided into tertiles according to CAVD severity on the basis of peak aortic jet velocity	Statins did not improve CAVD outcomes regardless of initial severity of disease
ASTRONOMER trial (2010) <sup>86</sup>	Randomized, double-blind trial of 269 patients given rosuvastatin or placebo	Statin treatment did not reduce deterioration in peak aortic pressure gradient

Abbreviation: CAVD, calcific aortic valve disease.

# Failure of Statin Rx to Treat CAVD

## SALTIRE (2005)

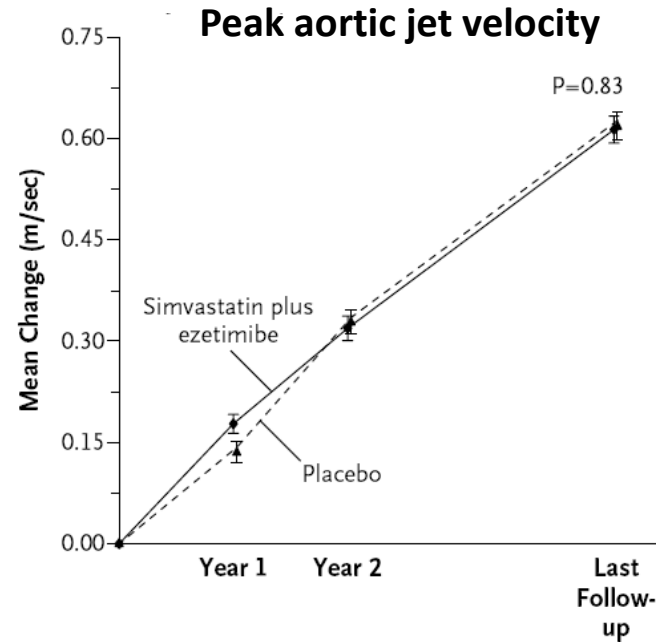
N = 155 pts



Cowell et al, NEJM,  
352:2389-97,2005

## SEAS (2008)

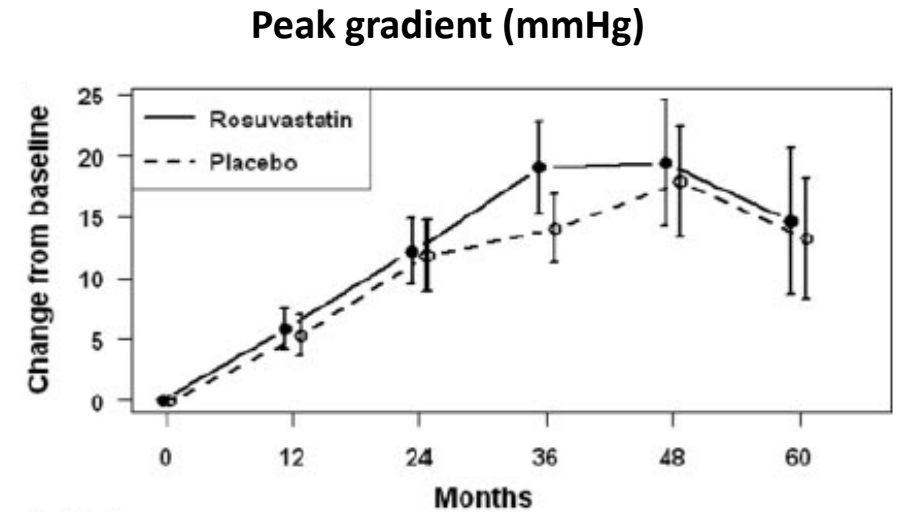
N = 1,873 pts



Rossebo et al, NEJM,  
359:1343-56, 2008

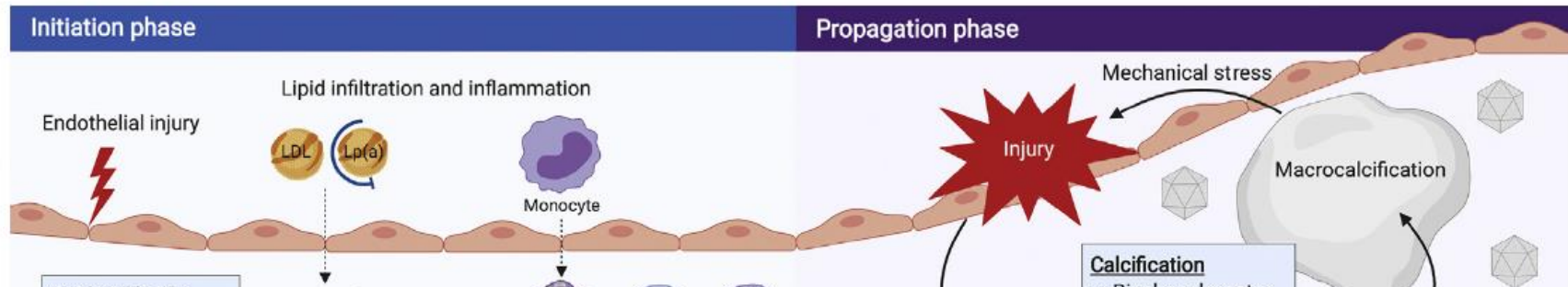
## ASTRONOMER (2010)

N = 269 pts

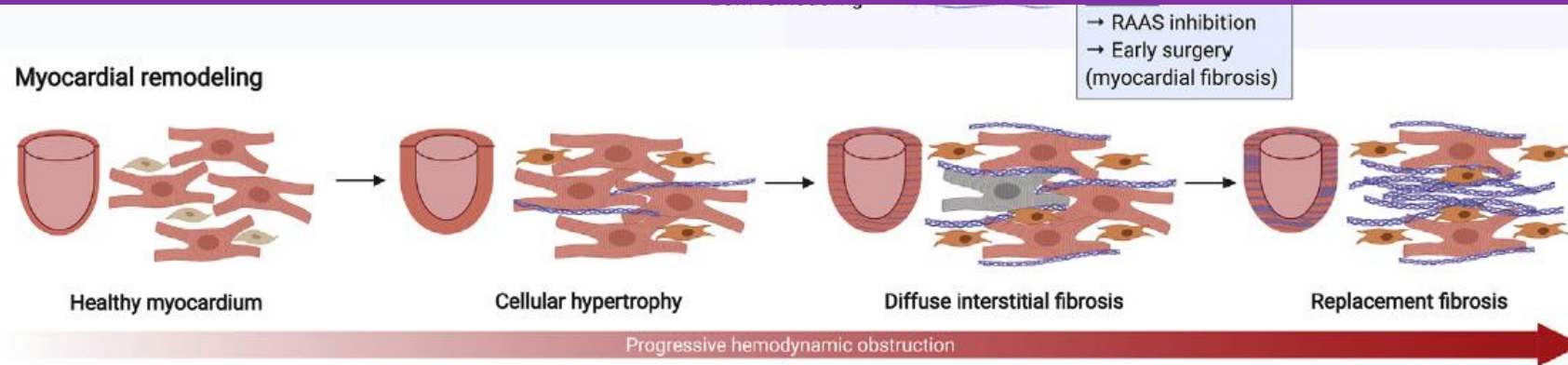


Chan et al, Circulation  
121:306-314, 2010

# Pathophysiology of Aortic Stenosis



**As of today, there are NO known proven medical therapies (e.g. statins and ACEI) to slow or prevent the progression of AS.**



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ESTABLISHED IN 1812

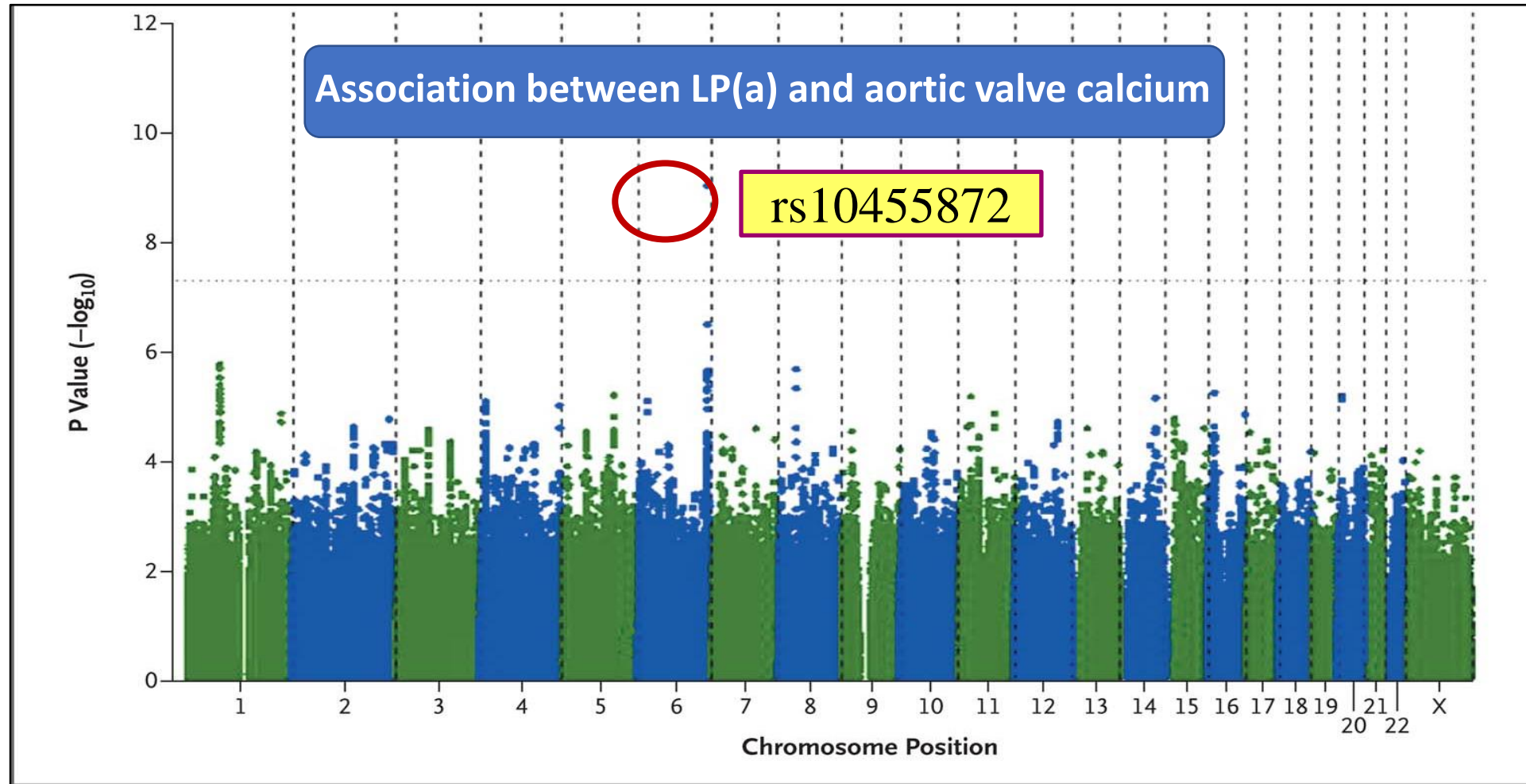
FEBRUARY 7, 2013

VOL. 368 NO. 6

## Genetic Associations with Valvular Calcification and Aortic Stenosis

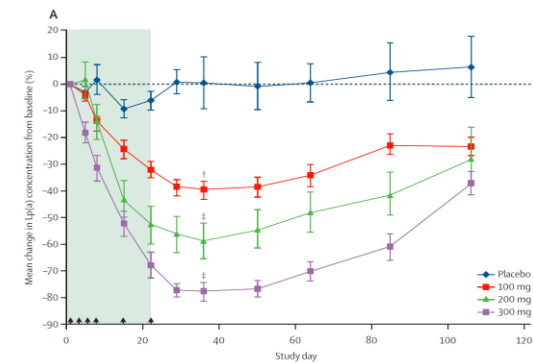
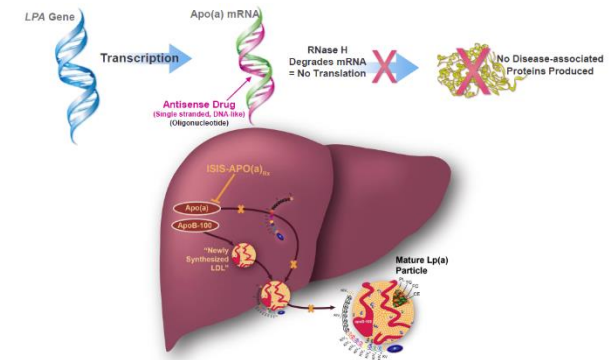
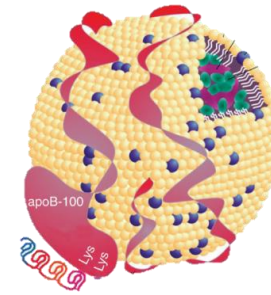
George Thanassoulis, M.D., Catherine Y. Campbell, M.D., David S. Owens, M.D., J. Gustav Smith, M.D., Ph.D., Albert V. Smith, Ph.D., Gina M. Peloso, Ph.D., Kathleen F. Kerr, Ph.D., Sonali Pechlivanis, Ph.D., Matthew J. Budoff, M.D., Tamara B. Harris, M.D., Rajeev Malhotra, M.D., Kevin D. O'Brien, M.D., Pia R. Kamstrup, M.D., Ph.D., Børge G. Nordestgaard, M.D., D.M.Sc., Anne Tybjaerg-Hansen, M.D., D.M.Sc., Matthew A. Allison, M.D., M.P.H., Thor Aspelund, Ph.D., Michael H. Criqui, M.D., M.P.H., Susan R. Heckbert, M.D., Ph.D., Shih-Jen Hwang, Ph.D., Yongmei Liu, Ph.D., Marketa Sjogren, Ph.D., Jesper van der Pals, M.D., Ph.D., Hagen Kälsch, M.D., Thomas W. Mühleisen, Ph.D., Markus M. Nöthen, M.D., L. Adrienne Cupples, Ph.D., Muriel Caslake, Ph.D., Emanuele Di Angelantonio, M.D., Ph.D., John Danesh, F.R.C.P., Jerome I. Rotter, M.D., Sigurdur Sigurdsson, M.Sc., Quenna Wong, M.S., Raimund Erbel, M.D., Sekar Kathiresan, M.D., Olle Melander, M.D., Ph.D., Vilmundur Gudnason, M.D., Ph.D., Christopher J. O'Donnell, M.D., M.P.H., and Wendy S. Post, M.D.,  
for the CHARGE Extracoronary Calcium Working Group

# Lp (a) Reigns Supreme in GWAS!

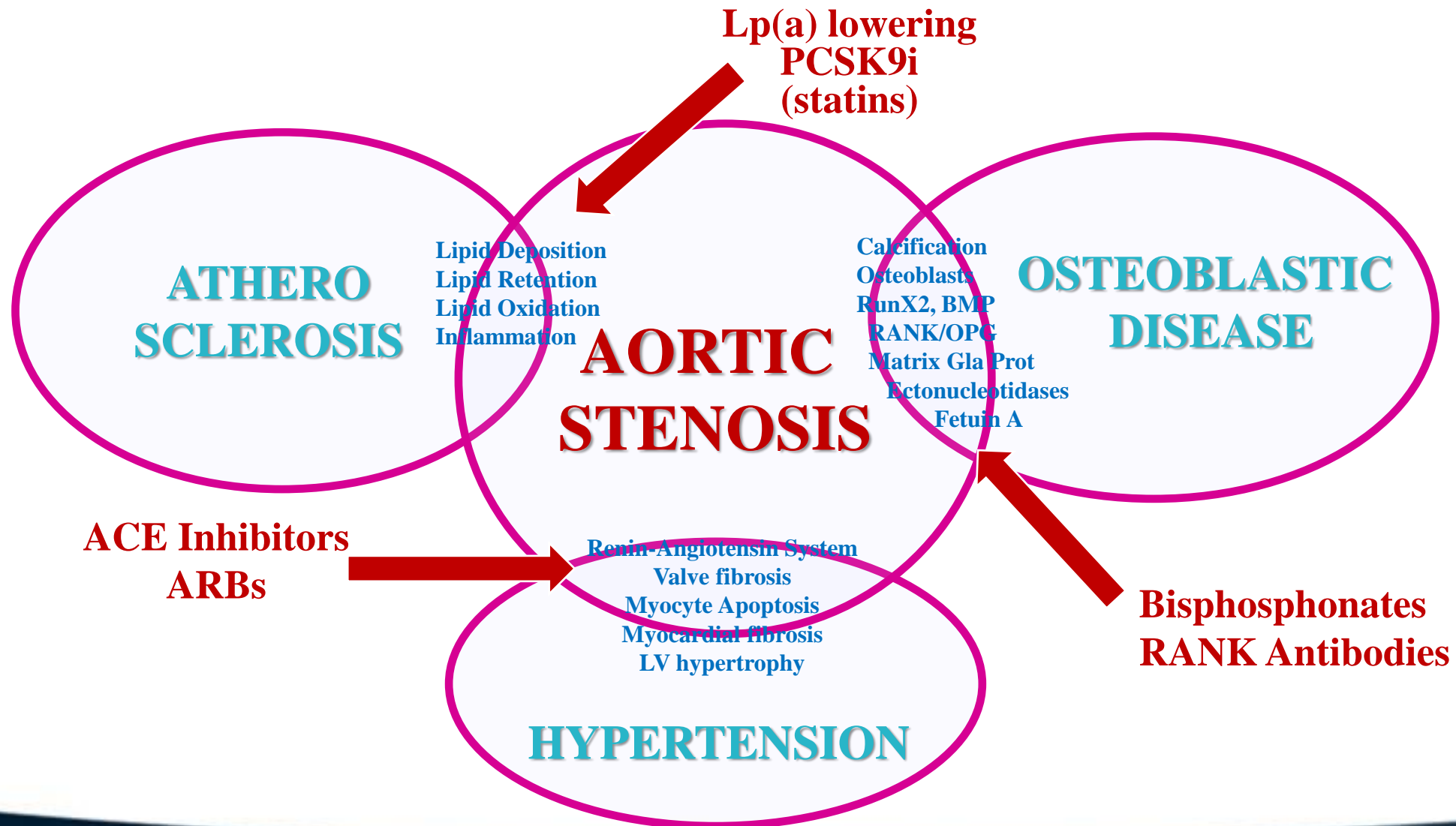


# Lp (a) Reigns Supreme!

- About 20% of the general population (65 Million people in North America) have elevated Lp(a)
- Lp(a) circulating levels are determined genetically and currently available drugs (Niacin) only achieve modest reduction in Lp(a)
- Phase I and II trials report that oligonucleotide antisense directed to Apo(a), reduces Lp(a) levels by >80% with minimal side effects



# Modern Thinking – Medical Rx for AS



# Future Perspectives on Medical Rx for CAVD

Study	Target	Treatment	Main Inclusion Criteria	Follow-up	Primary Efficacy Endpoints
PCSK9 inhibitors in the progression of aortic stenosis (NCT03051360)	ApoB-containing lipoproteins; PCSK9	Biweekly injection of PCSK9 inhibitor vs placebo	Mild-moderate aortic stenosis (n = 140)	2 years	Change in aortic valve CT calcium score and <sup>18</sup> F-NaF uptake
EAVaLL—Early Aortic Valve Lipoprotein (a) Lowering (NCT02109614)	Lipoprotein(a)	Daily extended-release niacin 1500–2000 mg vs Placebo	Aortic sclerosis or mild aortic stenosis + elevated Lp(a) levels (>50 mg/dL) (n = 150)	2 years	Change in aortic valve CT calcium score
SALTIRE II—Study Investigating the Effect of Drugs Used to Treat Osteoporosis on the Progression of Calcific Aortic Stenosis (NCT02132026)	Mineral metabolism	<ul style="list-style-type: none"> <li>Alendronic acid (n = 50) vs placebo tablets (n = 25)</li> <li>Denosumab (n = 50) vs placebo injections (n = 25)</li> </ul>	Aortic stenosis ( $V_{max} > 2.5$ m/s)	2 years	Change in aortic valve calcium score, aortic valve <sup>18</sup> F-NaF uptake
BASIK2—Bicuspid Aortic Valve Stenosis and the Effect of Vitamin K2 on calcium metabolism on <sup>18</sup> F-NaF PET/MRI (NCT02917525)	Vitamin K2-Matrix Gla protein	Daily vitamin K2 360 $\mu$ g (n = 22) vs placebo (n = 22)	Bicuspid aortic valve and calcified mild to moderate aortic stenosis	18 months	Change in aortic valve <sup>18</sup> F-NaF uptake at 6 mo; change in aortic valve calcium score (secondary endpoint at 6 + 18 mo)
EvoLveD—Early Valve Replacement Guided by Biomarkers of LV Decompensation in Asymptomatic Patients With Severe AS (NCT03094143)	Midwall fibrosis and timing of intervention	Early aortic valve replacement vs routine care	Asymptomatic severe aortic stenosis ( $V_{max} > 4.0$ m/s; or $V_{max} > 3.5$ with AVA $< 0.6$ cm <sup>2</sup> /m <sup>2</sup> )	$\pm$ 3 y (until 88 events accrue)	Composite of all-cause mortality or unplanned aortic stenosis-related hospitalisation

Abbreviations: ApoB, apolipoprotein B; AVA, aortic valve area;  $V_{max}$ , peak aortic jet velocity.



# Incomplete Understanding of Pathogenesis

- Are there octogenarian with pristine aortic valves?
- Are risk factor modifications occur early enough in trials?
- Should we be studying aortic stenosis progression or prevention?
- Are all tricuspid valves born equal?
- LV reserve/response to aortic stenosis variations

# Upstream Management of Aortic Stenosis

- Pathophysiology
- Risk Factors
- Detection



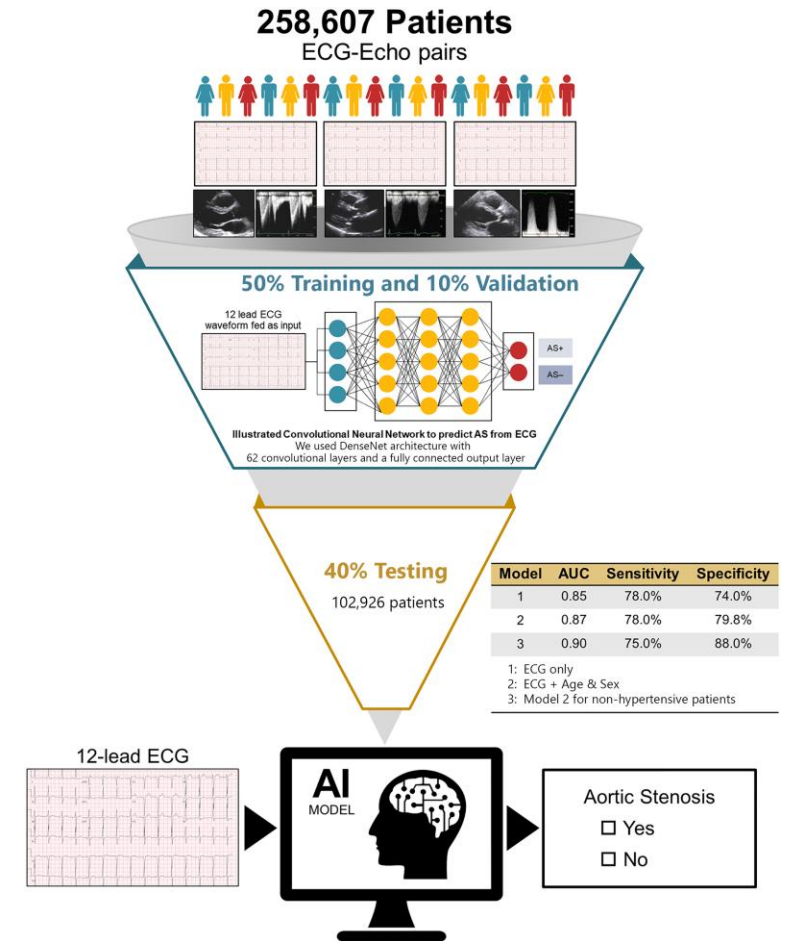
# Future Screening Tools for Valvular Heart Disease

## Artificial Intelligence/Machine Learning

### Electrocardiogram screening for aortic valve stenosis using artificial intelligence

Michal Cohen-Shelly <sup>1</sup>, Zachi I. Attia <sup>1</sup>, Paul A. Friedman <sup>1</sup>, Saki Ito <sup>1</sup>, Benjamin A. Essayagh <sup>1</sup>, Wei-Yin Ko <sup>1</sup>, Dennis H. Murphree <sup>1</sup>, Hector I. Michelena <sup>1</sup>, Maurice Enriquez-Sarano <sup>1</sup>, Rickey E. Carter <sup>2</sup>, Patrick W. Johnson <sup>2</sup>, Peter A. Noseworthy <sup>1</sup>, Francisco Lopez-Jimenez <sup>1</sup>, and Jae K. Oh <sup>1\*</sup>

In the test group, the AI-ECG labelled 3833 (3.7%) patients as positive with the area under the curve (AUC) of 0.85. The sensitivity, specificity, and accuracy were 78%, 74%, and 74%, respectively.



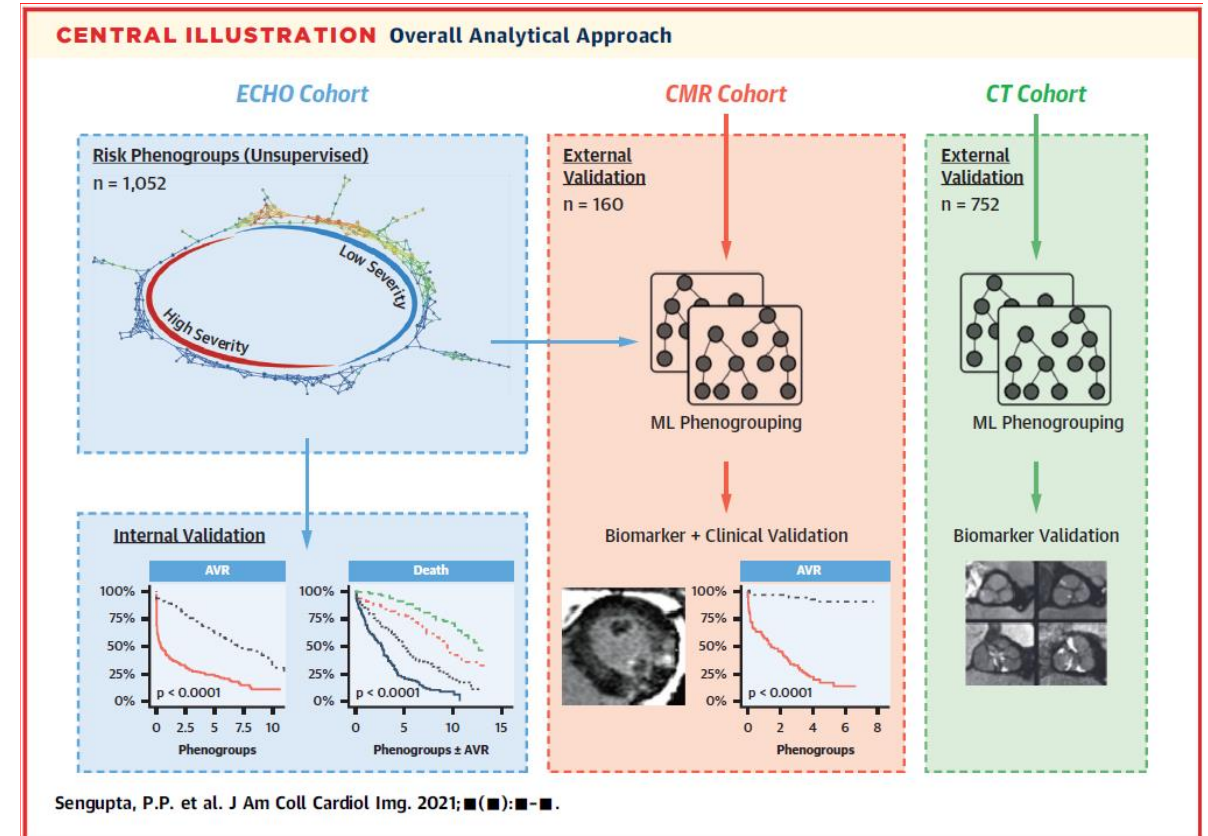
# Future Screening Tools for Valvular Heart Disease

## *Artificial Intelligence/Machine Learning*

### A Machine-Learning Framework to Identify Distinct Phenotypes of Aortic Stenosis Severity

Partho P. Sengupta, MD, DM,<sup>a</sup> Sirish Shrestha, MS,<sup>a</sup> Nobuyuki Kagiyama, MD, PhD,<sup>a</sup> Yasmin Hamirani, MD,<sup>a</sup> Hemant Kulkarni, MD,<sup>a,b</sup> Naveena Yanamala, PhD,<sup>a</sup> Rong Bing, MBBS,<sup>c</sup> Calvin W.L. Chin, MD, PhD,<sup>d</sup> Tania A. Pawade, MD, PhD,<sup>c</sup> David Messika-Zeitoun, MD,<sup>e</sup> Lionel Tastet, MSc,<sup>f</sup> Mylène Shen, PhD,<sup>f</sup> David E. Newby, MD, PhD,<sup>c</sup> Marie-Annick Clavel, DVM, PhD,<sup>f</sup> Philippe Pibarot, DVM, PhD,<sup>f</sup> Marc R. Dweck, MD, PhD,<sup>c</sup> for the Artificial Intelligence for Aortic Stenosis at Risk International Consortium

**Conclusions:**  
Machine learning can integrate ECHO measurements to augment the classification of disease severity in most patients with AS, with major potential to optimize the timing of AVR. (JACC Imaging 2021)



# Upstream Management of Aortic Stenosis

- Pathophysiology
- Risk Factor Modification
- Detection
  - Is there a CAC score equivalent for early aortic stenosis?
    - Score of 1648 is associated with severe AS



# Upstream Management of Aortic Stenosis

- Pathophysiology
- Risk Factor Modification
- Detection
- **Criteria for Treatment**



# Traditional Thinking – Aortic Stenosis

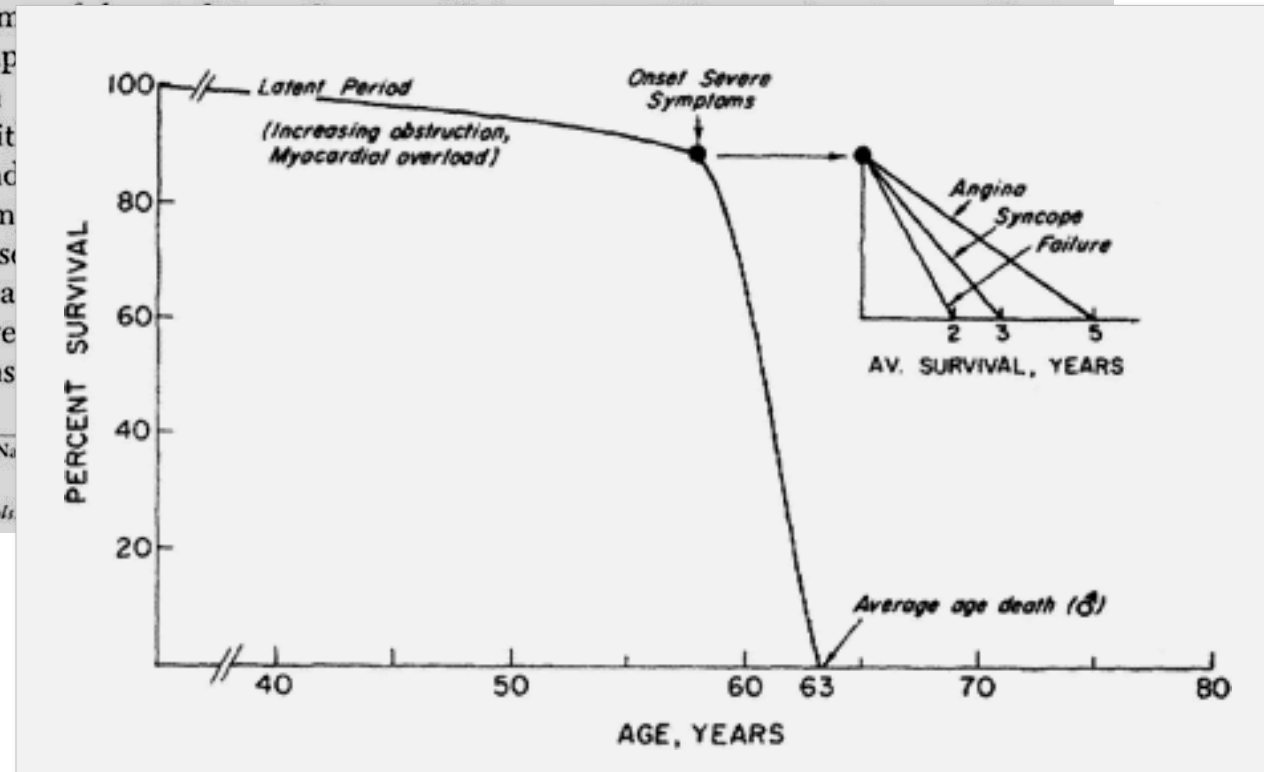
## Aortic Stenosis

By JOHN ROSS, JR., M.D. AND EUGENE BRAUNWALD, M.D.

THE ADVENT of corrective operations for various forms of aortic stenosis has placed increasing emphasis on the need for accurate information regarding the natural history of patients with this disease. An understanding of the course of aortic stenosis because of its potential for sudden death and the grave consequences that may accompany the onset of severe symptoms.

From the Cardiology Branch, National Institutes of Health, Bethesda, Maryland.

Supplement V to *Circulation*, Vol. 38



# Traditional Thinking – Aortic Stenosis

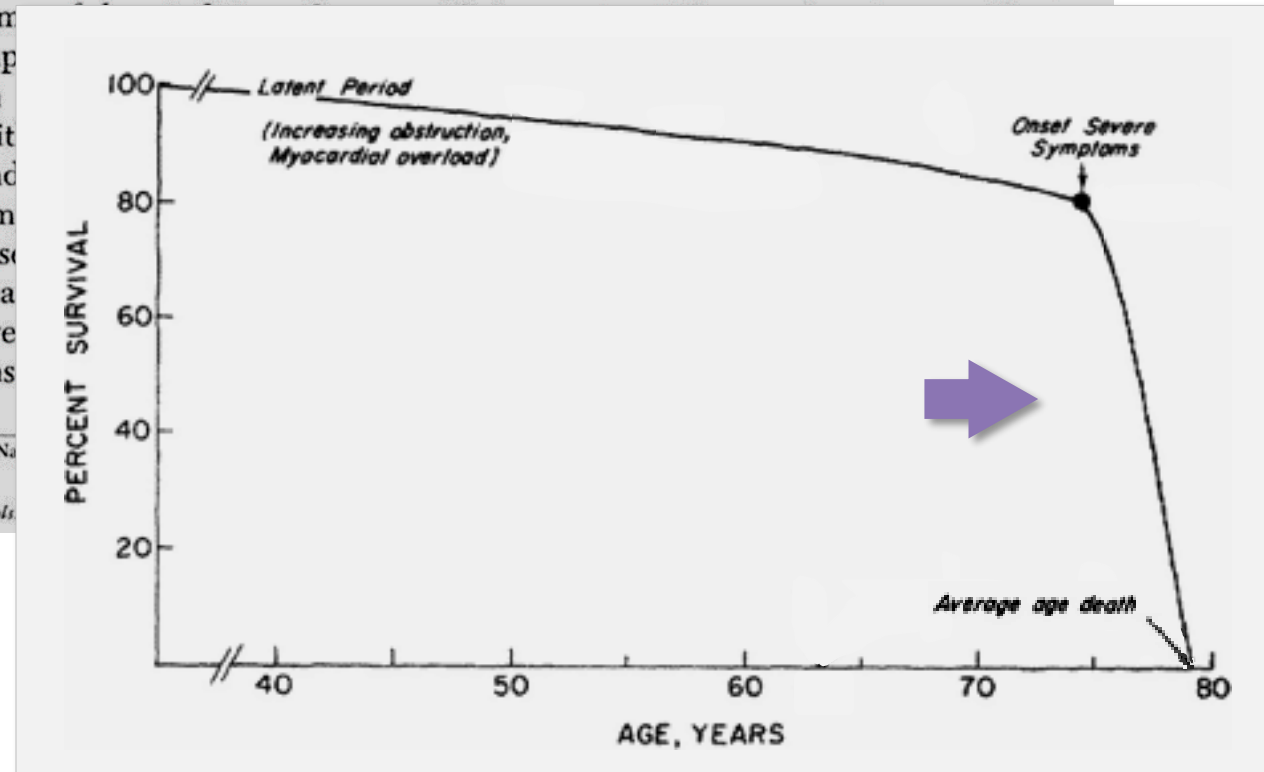
## Aortic Stenosis

By JOHN ROSS, JR., M.D. AND EUGENE BRAUNWALD, M.D.

THE ADVENT of corrective operations for various forms of aortic stenosis has placed increasing emphasis on the need for accurate information regarding the natural history of patients with isolated valvular aortic stenosis. An understanding of the disease process assumes particular importance in aortic stenosis because of the prevalence of sudden death and the grave consequences that may accompany the onset of severe symptoms.

From the Cardiology Branch, National Institutes of Health, Bethesda, Maryland.

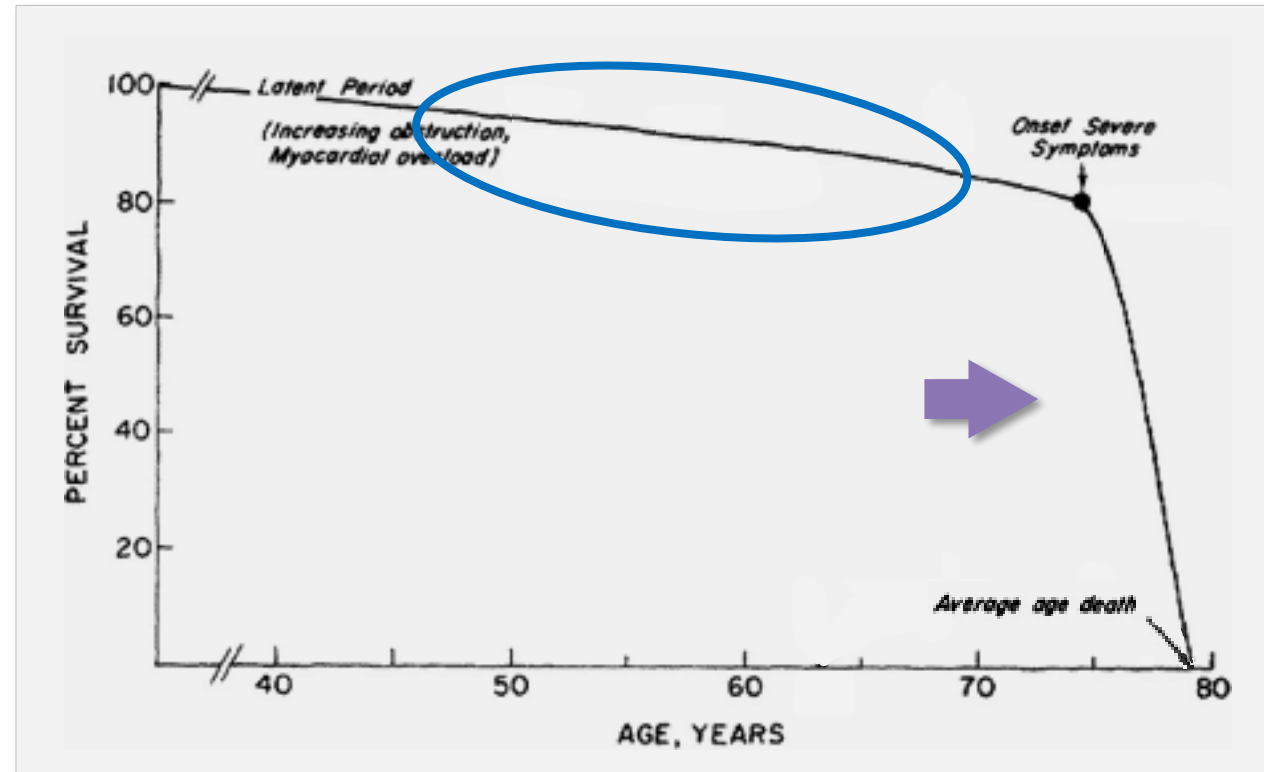
Supplement V to *Circulation*, Vol. 38





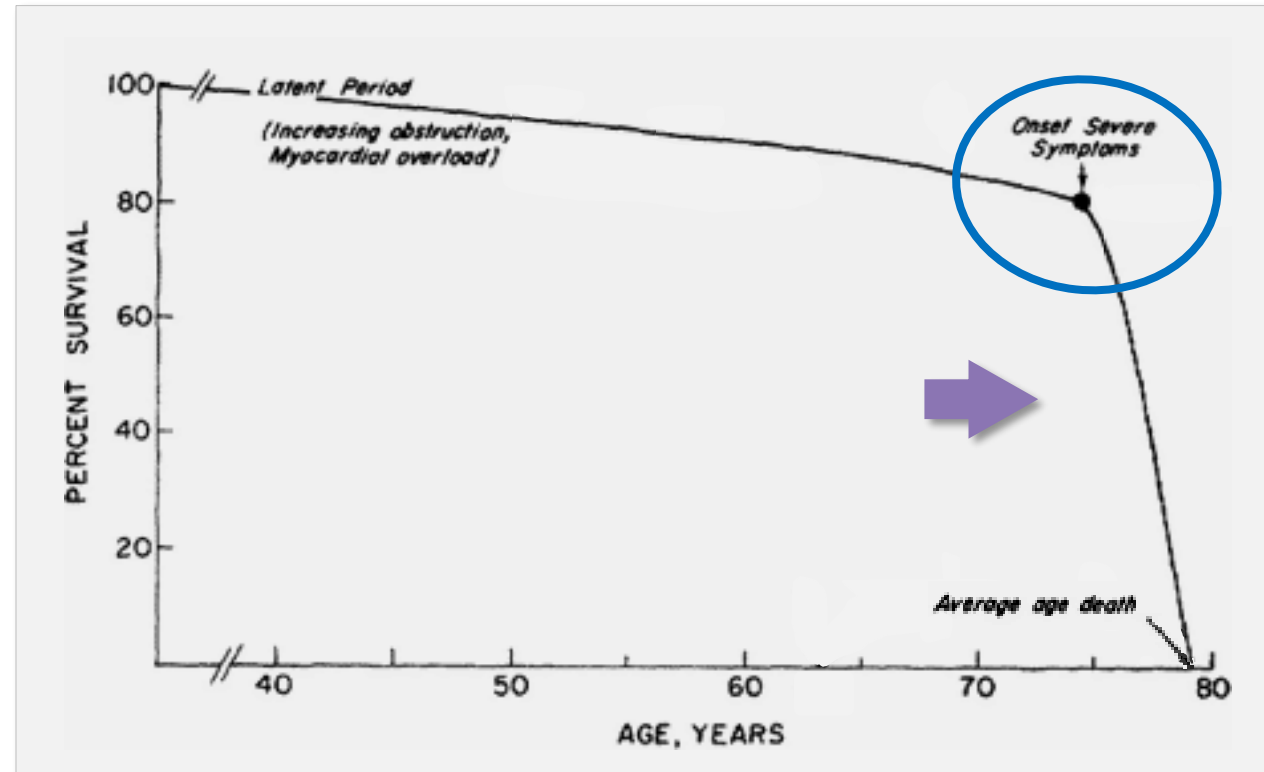
# Traditional Thinking – Aortic Stenosis

*Fundamental fallacies: 1. there are no important reversible and irreversible structural changes during the so-called latent period which negatively impact subsequent clinical outcomes*

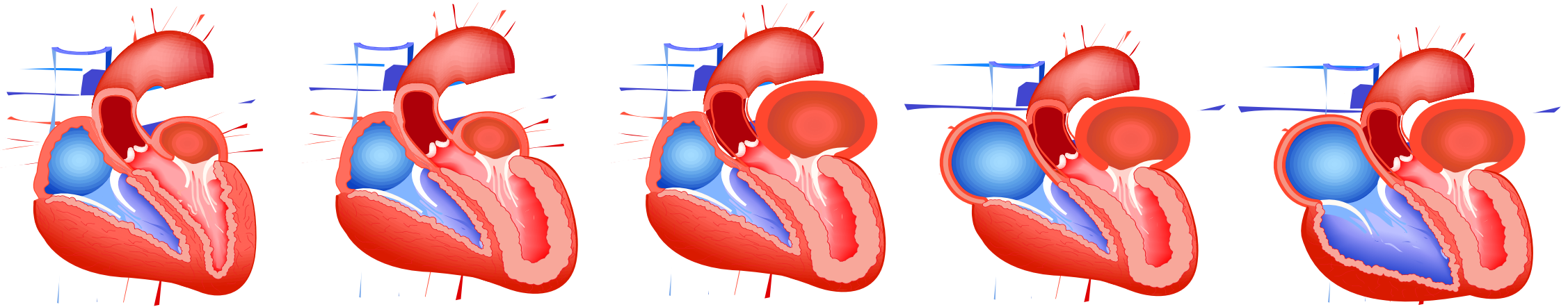


# Traditional Thinking – Aortic Stenosis

*Fundamental fallacies: 2. the onset of symptoms is discrete, easily identifiable (even in the elderly), and is inexorably linked to aortic stenosis severity*



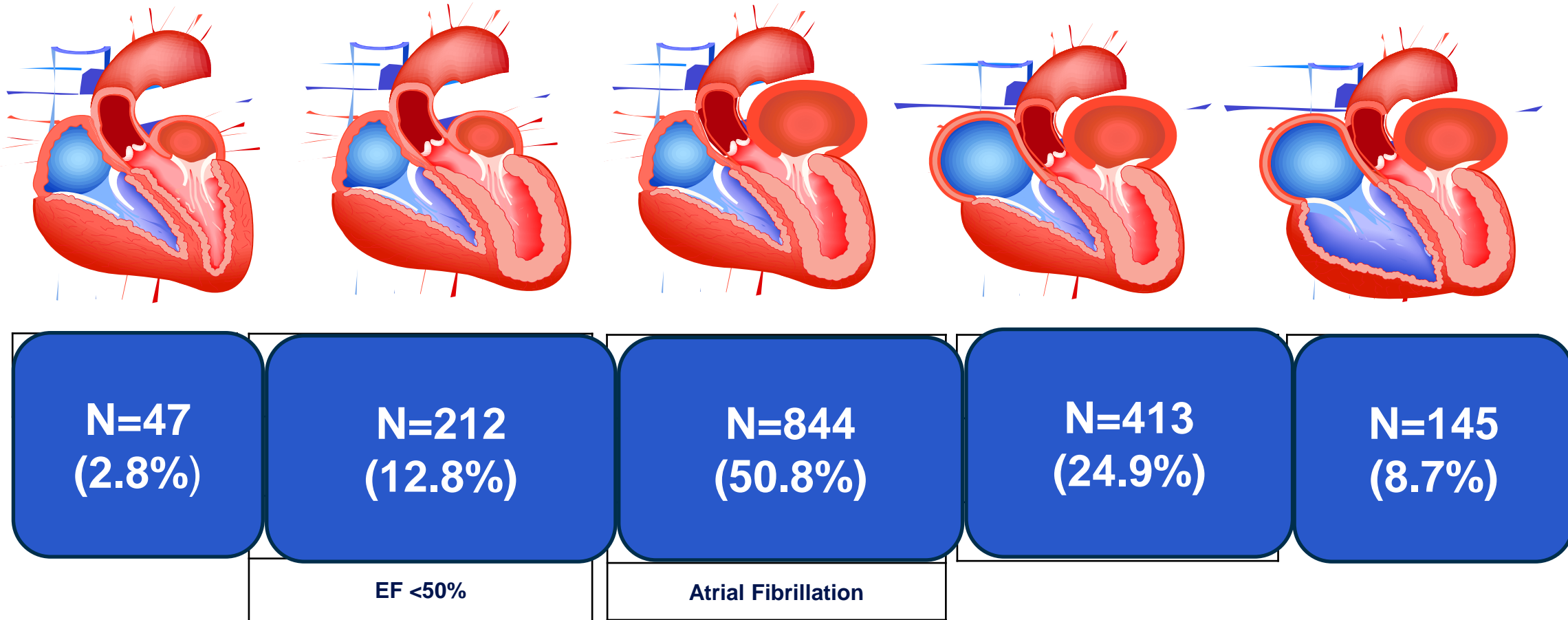
# Staging Classification in Severe AS (n=1,661 pts)



<b>Stage 0</b> No damage	<b>Stage 1</b> LV damage	<b>Stage 2</b> LA/Mitral damage	<b>Stage 3</b> PA/Tricuspid damage	<b>Stage 4</b> RV damage
	Increased LV Mass Index >115 g/m <sup>2</sup> Male >95 g/m <sup>2</sup> Female	Indexed left atrial volume >34mL/m <sup>2</sup>	PAS ≥60mmhg	Moderate-Severe RV dysfunction
	E/e' >14	Moderate-Severe MR	Moderate-Severe TR	
	EF <50%	Atrial Fibrillation		

*Patients hierarchically classified based on the presence of at least one variable in the highest stage (independent, not additive)*

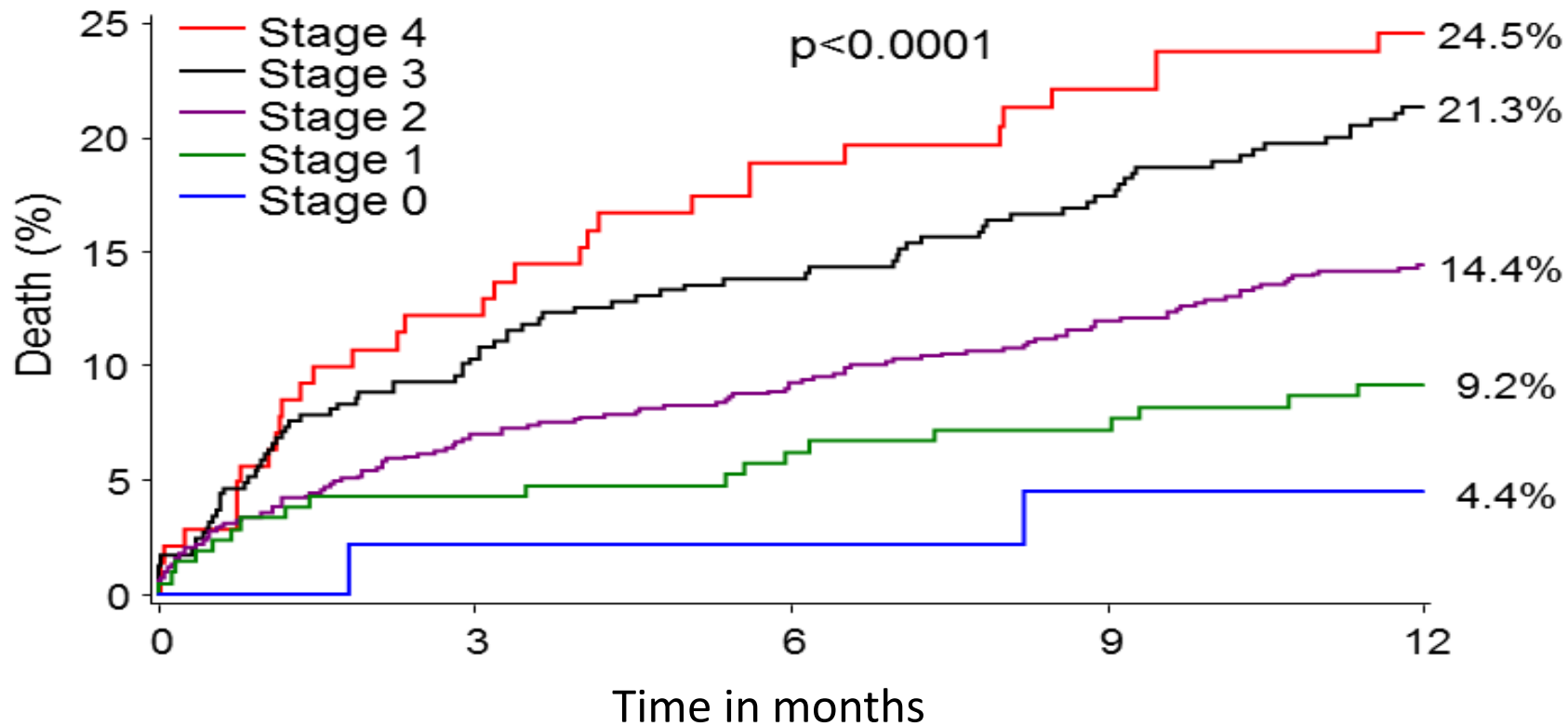
# Staging Classification in Severe AS (n=1,661 pts)



*Patients hierarchically classified based on the presence of at least one variable in the highest stage (independent, not additive) ; Severe AS: AVA 1.0 cm<sup>2</sup>; mean gradient 40 mmHg*

# Staging Classification in Severe AS (n=1,661 pts)

## One-year Mortality after AVR

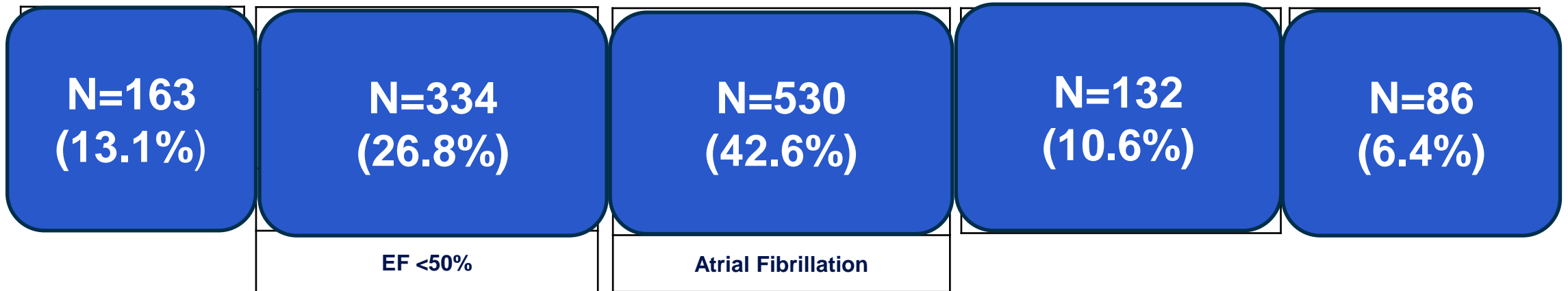
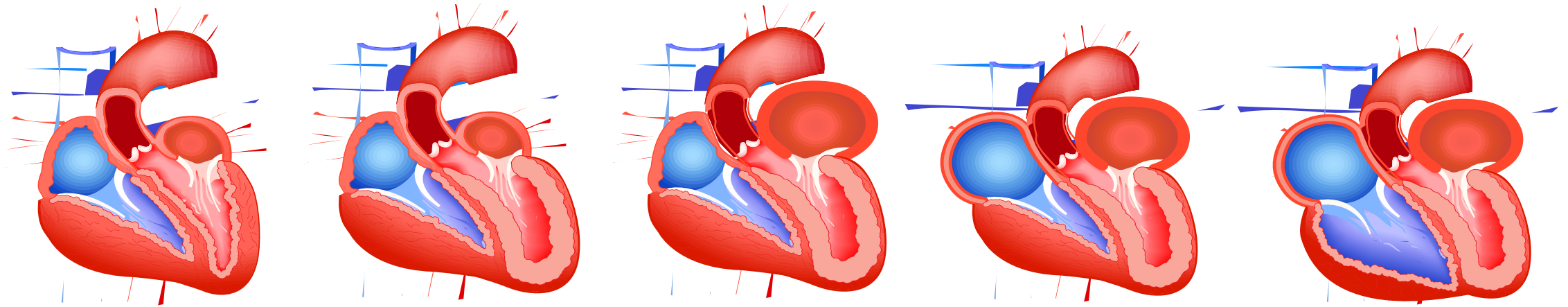


# Prognostic Implications of Associated Cardiac Abnormalities Detected on Echocardiography in Patients With Moderate Aortic Stenosis

Mohammed Rizwan Amanullah, MBBS,<sup>a,\*</sup> Stephan Milhorini Pio, MD,<sup>b,\*</sup> Arnold C.T. Ng, MBBS, PhD,<sup>c</sup> Kenny Y.K. Sin, MBBS,<sup>d</sup> Nina Ajmone Marsan, MD, PhD,<sup>b</sup> Zee Pin Ding, MBBS,<sup>a</sup> Martin B. Leon, MD,<sup>e</sup> Philippe Généreux, MD,<sup>f</sup> Victoria Delgado, MD, PhD,<sup>b</sup> See Hooi Ewe, MBBS, PhD,<sup>a</sup> Jeroen J. Bax, MD, PhD<sup>b</sup>

- 1245 patients with moderate AS followed in a longitudinal database
- Patients grouped according to index echocardiograms into 5 categories of severity of cardiac damage
- Significant higher mortality rates with increasing extent of extra-aortic valvular cardiac abnormalities (log-rank  $p < 0.001$ )

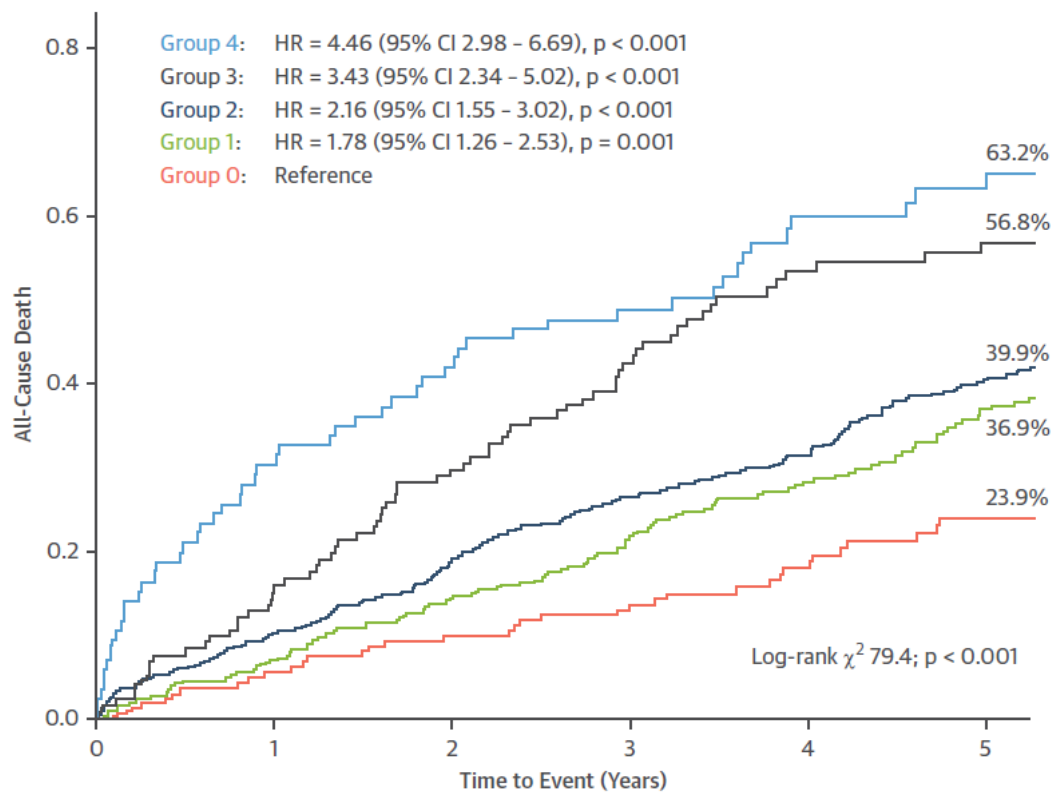
# Staging Classification in Moderate AS (n=1,245 pts)



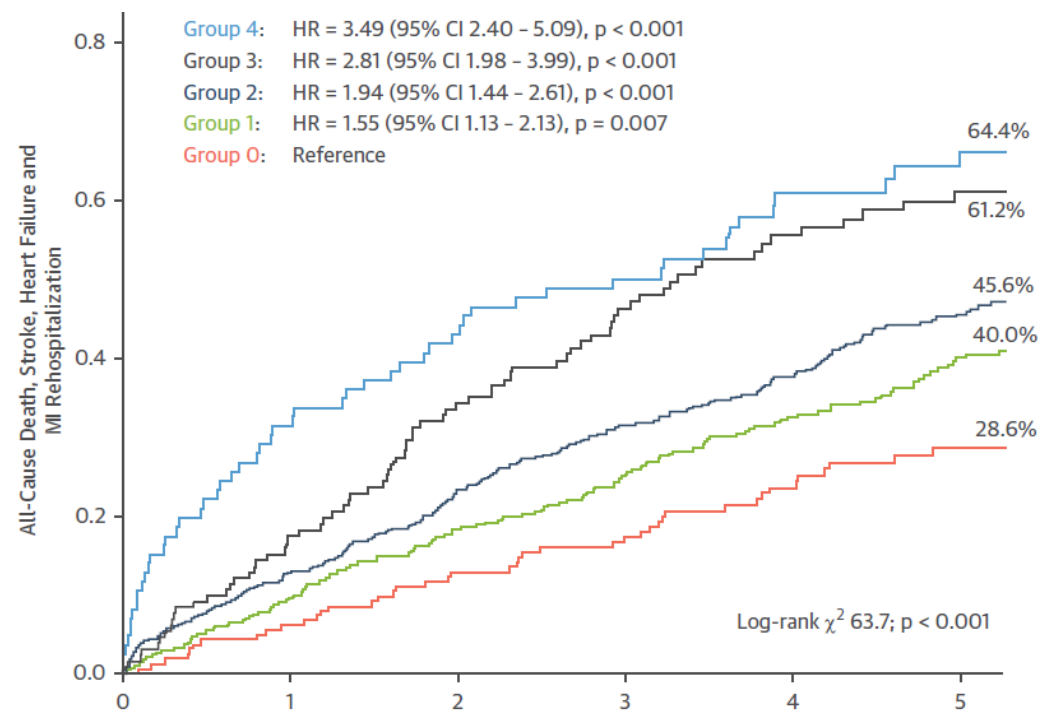
*Patients hierarchically classified based on the presence of at least one variable in the highest stage (independent, not additive) ; Moderate AS: AVA 1.2 cm<sup>2</sup>; mean gradient 24.4mmHg*

# Staging Classification in Moderate AS (n=1,245 pts)

## 5-yr Mortality



## 5-yr Mortality, Stroke or Rehosp

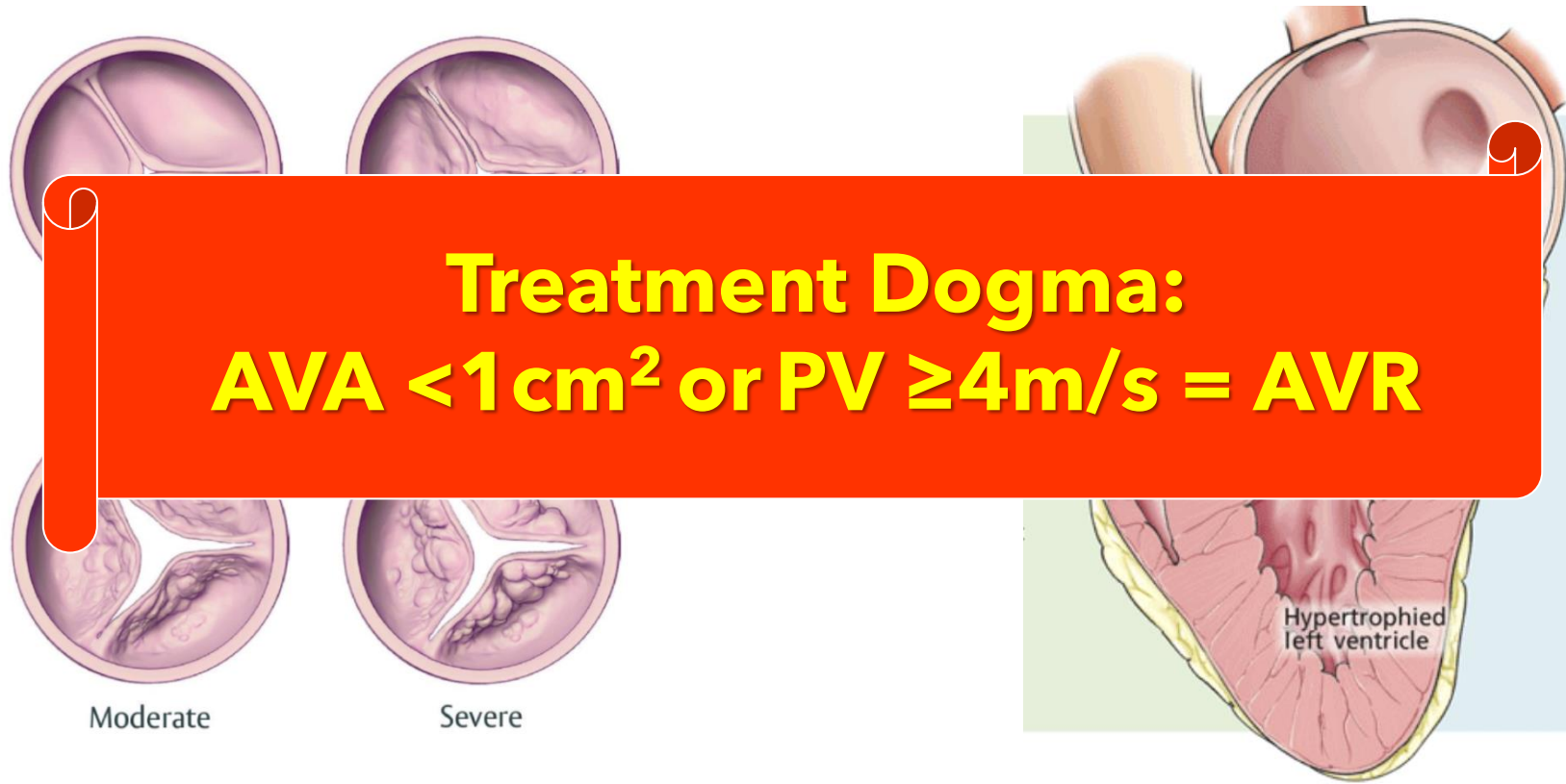




# Aortic Valve Therapies: The Future?

## *UPSTREAM AS Treatment*

*Two parallel processes with 'variable' linkage*



**Treatment Dogma:**  
 **$AVA < 1 \text{ cm}^2$  or  $PV \geq 4 \text{ m/s} = \text{AVR}$**

# The RECOVERY Surgical AVR Trial

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ESTABLISHED IN 1812

JANUARY 9, 2020

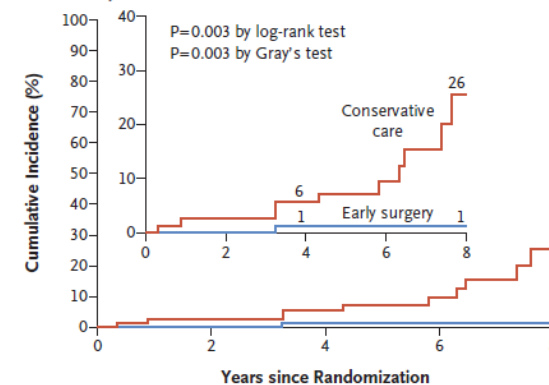
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## Early Surgery or Conservative Care for Asymptomatic Aortic Stenosis

Duk-Hyun Kang, M.D., Ph.D., Sung-Ji Park, M.D., Ph.D., Seung-Ah Lee, M.D., Sahmin Lee, M.D., Ph.D., Dae-Hee Kim, M.D., Ph.D., Hyung-Kwan Kim, M.D., Ph.D., Sung-Cheol Yun, Ph.D., Geu-Ru Hong, M.D., Ph.D., Jong-Min Song, M.D., Ph.D., Cheol-Hyun Chung, M.D., Ph.D., Jae-Kwan Song, M.D., Ph.D., Jae-Won Lee, M.D., Ph.D., and Seung-Woo Park, M.D., Ph.D.

- 145 asymptomatic patients w very severe AS randomized to early surgery or conservative care
- 1<sup>ry</sup> endpoint (operative and FU death) was 1% vs. 15% in early surgery vs. conservative care (P=0.003)

A Operative Mortality or Death from Cardiovascular Causes

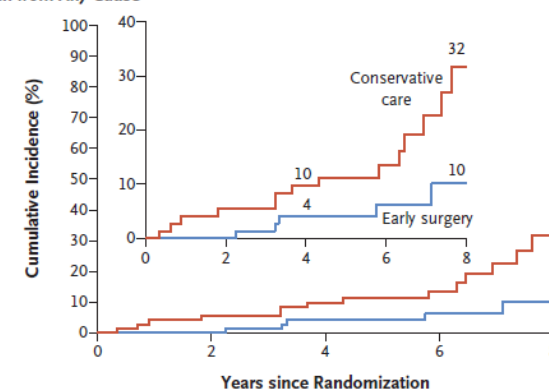


No. at Risk

Conservative care 72 68 65 36 12

Early surgery 73 73 70 38 13

B Death from Any Cause



No. at Risk

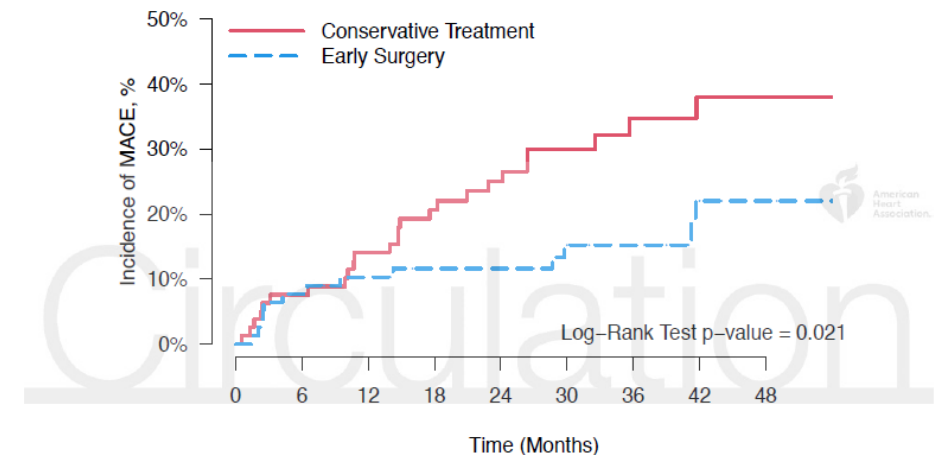
Conservative care 72 68 65 36 12

Early surgery 73 73 70 38 13

# The AVATAR Surgical AVR Trial

## Aortic Valve Replacement versus Conservative Treatment In Asymptomatic Severe Aortic Stenosis: The AVATAR Trial

- 157 asymptomatic patients (ETT confirmed) w severe AS, randomized to early surgery or conservative care at 9 centers from 7 EU countries; median FU 32 months
- Early surgery operative mortality 1.4%
- 1<sup>ry</sup> endpoint (MACE = death, MI, stroke and HF re hosp) was lower with early surgery vs. conservative care (HR 0.46, 95% CI 0.23-0.90; p=0.02)

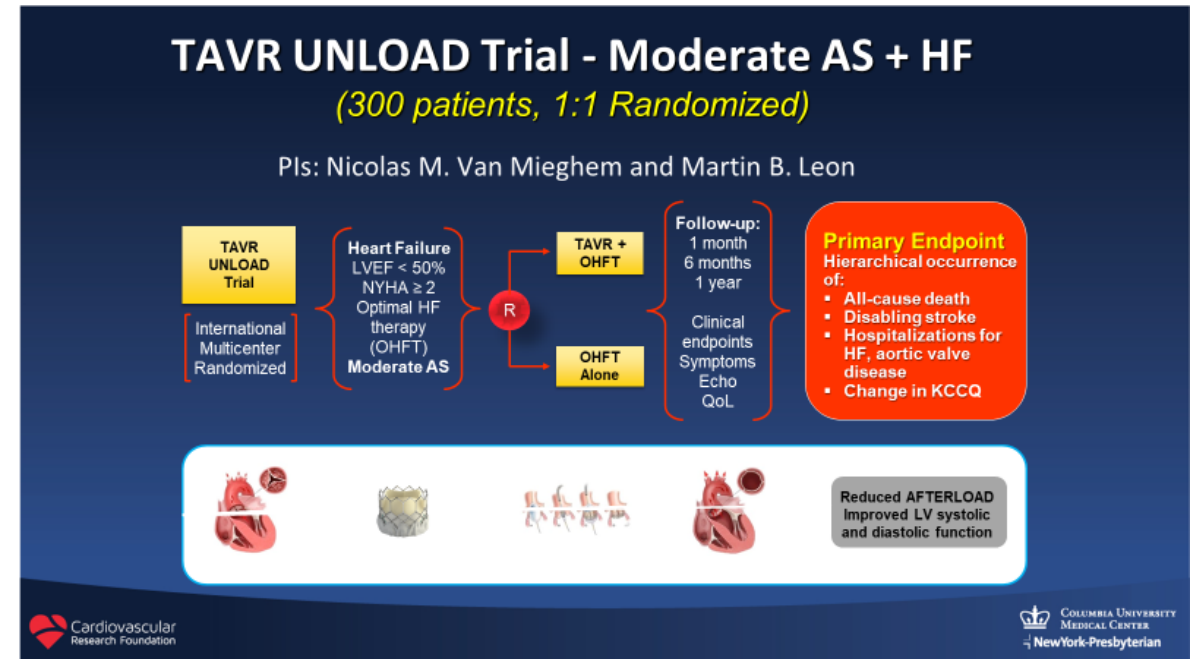
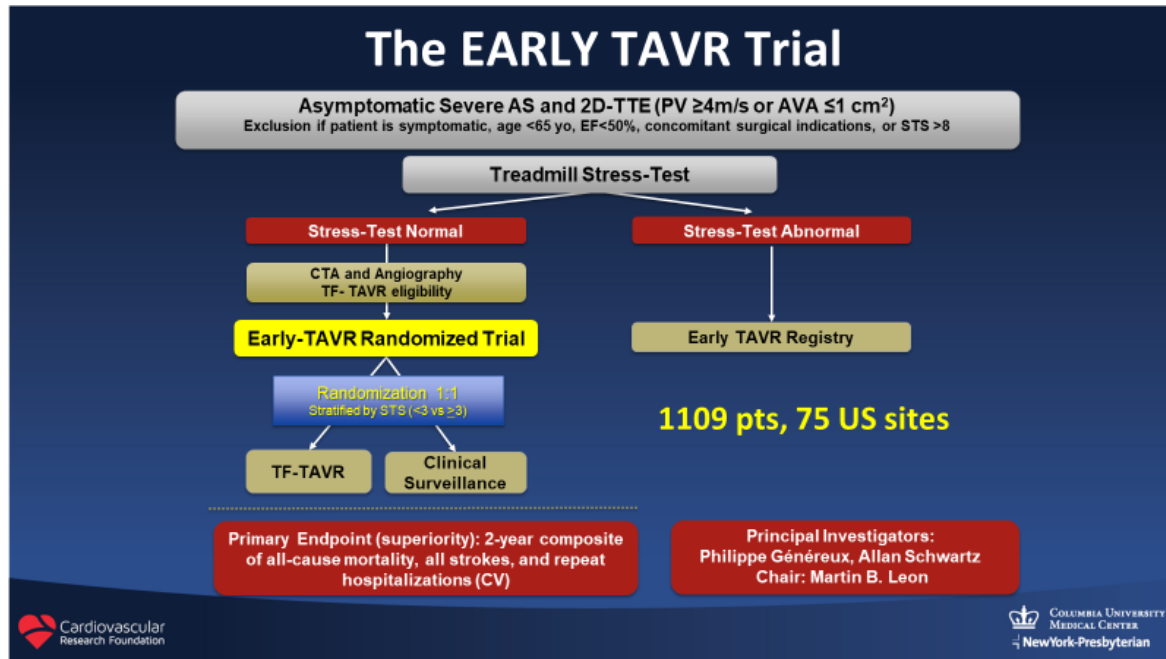


	Patients, n								
Conservative Treat.	79	73	66	59	49	36	25	19	12
Early Surgery	78	72	68	63	56	46	38	23	13

# Pre-emptive (earlier) TAVR

## *EARLY TAVR and UNLOAD Trials*

### Expanding TAVR Clinical Indications to 'Earlier' Treatment Scenarios



# Upstream Mod AS Treatment

## *“At Risk” Predictors*

- Cardiac symptoms (esp. heart failure NYHA 3 or 4)
- Low ejection fraction (< 60% LVEF)
- Atrial fibrillation (persistent or recent paroxysmal)
- Low stroke volume (SVI < 35 cc/m<sup>2</sup>)
- Severe diastolic dysfunction (by echo criteria)
- Rapid AS progression (increase PV > 0.3 m/sec/year)
- Elevated cardiac biomarkers (BNP)
- Elevated AV calcium score by CT

# AS Severity Grading and Cardiac Staging

Grade or Stage	Stage 0 None	Stage 1 LV	Stage 2 LA-mitral	Stage 3 PA-tricuspid	Stage 4 RV
<b>Grade 0</b> $V_{\max} < 2\text{m/s}$					
<b>Grade 1</b> $V_{\max} 2\text{-}2.9\text{m/s}$					
<b>Grade 2</b> $V_{\max} 3\text{-}3.9\text{m/s}$					
<b>Grade 3</b> $V_{\max} \geq 4\text{m/s}$					

# AS Severity Grading and Cardiac Staging

Grade or Stage	Stage 0 None	Stage 1 LV	Stage 2 LA-mitral	Stage 3 PA-tricuspid	Stage 4 RV
<b>Grade 0</b> $V_{\max} < 2\text{m/s}$					
<b>Grade 1</b> $V_{\max} 2\text{-}2.9\text{m/s}$					
<b>Grade 2</b> $V_{\max} 3\text{-}3.9\text{m/s}$					
<b>Grade 3</b> $V_{\max} \geq 4\text{m/s}$		<b>AVR</b>	<b>AVR</b>	<b>AVR</b>	<b>AVR</b>

# AS Severity Grading and Cardiac Staging

Grade or Stage	Stage 0 None	Stage 1 LV	Stage 2 LA-mitral	Stage 3 PA-tricuspid	Stage 4 RV
Grade 0 $V_{\max} < 2\text{m/s}$					
Grade 1 $V_{\max} 2-2.9\text{m/s}$					
Grade 2 $V_{\max} 3-3.9\text{m/s}$	<b>PROGRESS</b>	<b>PROGRESS</b>	<b>PROGRESS</b>	<b>PROGRESS</b>	<b>PROGRESS</b>
Grade 3 $V_{\max} \geq 4\text{m/s}$	<b>EARLY TAVR</b>				



# AS Severity Grading and Cardiac Staging

Grade or Stage	Stage 0 None	Stage 1 LV	Stage 2 LA-mitral	Stage 3 PA-tricuspid	Stage 4 RV
Grade 0 $V_{max} < 2\text{m/s}$					
Grade 1 $V_{max} 2-2.9\text{m/s}$					
Grade 2 $V_{max} 3-3.9\text{m/s}$	PROGRESS	PROGRESS	PROGRESS	PROGRESS	PROGRESS
Grade 3 $V_{max} \geq 4\text{m/s}$	EARLY TAVR	AVR	AVR	AVR	AVR

?Multi-drug 'precision' medical Rx